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Disruption of *CK2β* in Embryonic Neural Stem Cells Compromises Proliferation and Oligodendrogenesis in the Mouse Telencephalon^{∇†}

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Genetic programs that govern neural stem/progenitor cell (NSC) proliferation and differentiation are dependent on extracellular cues and a network of transcription factors, which can be regulated posttranslationally by phosphorylation. However, little is known about the kinase-dependent pathways regulating NSC maintenance and oligodendrocyte development. We used a conditional knockout approach to target the murine regulatory subunit (beta) of protein kinase casein kinase 2 (CK2β) in embryonic neural progenitors. Loss of CK2β leads to defects in proliferation and differentiation of embryonic NSCs. We establish CK2β as a key positive regulator for the development of oligodendrocyte precursor cells (OPCs), both *in vivo* and *in vitro*. We show that CK2β directly interacts with the basic helix-loop-helix (bHLH) transcription factor Olig2, a critical modulator of OPC development, and activates the CK2-dependent phosphorylation of its serine-threonine-rich (STR) domain. Finally, we reveal that the CK2-targeted STR domain is required for the oligodendroglial function of Olig2. These findings suggest that CK2 may control oligodendrogenesis, in part, by regulating the activity of the lineage-specific transcription factor Olig2. Thus, CK2β appears to play an essential and uncompensated role in central nervous system development.

Casein kinase 2 (CK2) is a conserved serine/threonine kinase with more than 300 substrates, mostly proteins related to transcription-directed signaling (27). CK2 is a heterotetrameric holoenzyme formed by two catalytic subunits, α and α' , that associate with a dimeric building block of regulatory β subunits ($\alpha_2\beta_2$). CK2 β modulates the substrate specificity of the CK2 enzymatic activity, and its architecture is consistent with its role as a docking partner for other interacting proteins (4). We previously demonstrated that disruption of CK2 β function in mice results in postimplantation lethality. Moreover, many attempts to generate CK2 $\beta^{-/-}$ embryonic stem (ES) cells failed, suggesting an essential role of mammalian CK2 β for ES cell viability (2).

The function of CK2 β for cell cycle progression has been investigated, but its precise role is largely unknown. The cell proliferation function of CK2 β was first characterized in human fibroblasts, by use of antisense oligodeoxynucleotides and microinjection of specific antibodies (19, 26). Recently, a genome-wide survey of protein kinases required for cell progression into cultured *Drosophila melanogaster* S2 cells by double-stranded RNA demonstrated that CK2 β is required for centrosomal normality (1). In the same vein, downregulation of CK2 β by small interfering RNA (siRNA) results in delayed

cell cycle progression in cultured mammalian cells (42). Finally, in a genetic screen for mutations affecting the central brain of *Drosophila*, a hypomorphic allele of *D. melanogaster* CK2 β has been isolated, and that study suggested a role for CK2 β in cell proliferation or cell survival during brain development (15).

In the developing mouse brain, neural stem/progenitor cells (NSCs), which reside in the ventricular zone (VZ), give rise to neurons and glial cells (astrocytes and oligodendrocytes) (23). Chronologically, neurons are generated first, and glial cells are generated subsequent to neurogenesis. Oligodendrocytes originate from ventral neural progenitors in the VZ, and once committed, they migrate laterally and dorsally as parenchymal precursors (oligodendrocyte precursor cells [OPCs]) to populate the entire embryonic telencephalon, where they terminally differentiate (reviewed in reference 29). Distinct genetic programs using combinations of transcription factors govern multiple spatiotemporal origins of oligodendroglial specification (reviewed in reference 28). The basic helix-loop-helix (bHLH) transcription factor Olig2 plays an essential role in the embryonic specification of neural progenitors into the oligodendrocyte lineage, and *Olig2* gene function is absolutely required for OPC development (17, 21, 36, 43). Accordingly, Olig2 expression is detected first in ventral neural progenitors, precedes OPC-specific expression markers, and is maintained at all stages of oligodendrocyte development (29). It has been reported that the oligodendroglial function of Olig2 can be modulated by protein kinase AKT-mediated phosphorylation of its N-terminal domain (31). Interestingly, neural bHLH transcrip-

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tion factors have been described as being phosphorylated and regulated by the CK2 holoenzyme (9, 39). In the mouse brain, CK2 subunits (α , α' , and β) are expressed in all neural cells (A. Kolding and B. Boldyreff, unpublished data) in which the CK2 β -to-CK2 α -plus-CK2 α' ratio is largely elevated compared to that in other organs (41), suggesting a higher-threshold requirement in the amount of the holoenzyme in the central nervous system (CNS). All together, these observations suggest that CK2 β may modulate neural homeostasis.

In this study, we have used Cre/loxP-mediated recombination to generate mice with a CK2 β null allele in NSCs of the developing brain. Disruption of CK2 β results in inhibition of NSC proliferation and in a severe deficiency for progenitors to specify OPCs. In addition, we identified, *in vitro*, the serine-threonine-rich (STR) domain of Olig2 as a strict CK2 β -dependent, CK2-targeted substrate. We also show that the STR domain of Olig2 is a modulator of its oligodendroglial activity when tested in neurosphere assays, suggesting that CK2 β could mediate its effect via Olig2 oligodendroglial activity. Hence, these findings provide insight into the genetic basis underlying neural progenitor proliferation and oligodendrocyte development.

MATERIALS AND METHODS

Mouse breeding. Mice were of a C57BL/6 background. *Nestin-cre* transgenic mice (37) kept as heterozygotes were bred with CK2 $\beta^{loxP/loxP}$ mice (2). Double-heterozygous mice (CK2 $\beta^{loxP/wt}$; *Nestin-cre*) were bred back to CK2 $\beta^{loxP/loxP}$ mice to produce the genotypes used in this study, (i) CK2 $\beta^{loxP/wt}$, (ii) CK2 $\beta^{loxP/loxP}$, (iii) CK2 $\beta^{loxP/wt}$; *Nestin-cre*, and (iv) mutant CK2 $\beta^{loxP/loxP}$; *Nestin-cre* (referred to here as CK2 β^{-}). CK2 $\beta^{loxP/wt}$, CK2 $\beta^{loxP/loxP}$, and CK2 $\beta^{loxP/wt}$; *Nestin-cre* embryonic day 18.5 (E18.5) embryo phenotypes were identical and are referred to here as wild type (WT). Live-born mice, postnatal day 0 (P0) pups, and E18.5 embryos were genotyped by tail DNA PCR (2). Animal treatment was performed in accordance with the ethics committee (ComEth) of Grenoble, France.

Western blot analysis. Olfactory bulb, cerebellum, and meningeal cells were removed from dissected brains. Forebrain protein extracts were obtained after homogenization and sonication in 25 mM Tris (pH 8.5)–1 mM dithiothreitol (DTT)–200 mM NaCl–5 mM EDTA. Cells were lysed in 50 mM Tris (pH 7.5)–5 mM EDTA–500 mM NaCl–1% Triton X-100. Extractions were done with protease and phosphatase inhibitor cocktails (Sigma). Samples (40 μ g) were analyzed by 12% SDS-PAGE and processed for Western blot analysis. Antibodies used were rabbit anti-CK2 β (1/500, directed against the C terminus of CK2 β), rabbit anti-CK2 α (1/1,000; Calbiochem), mouse anti- β -tubulin class III isoform (1/1,000; Chemicon), rabbit anti-poly(ADP-ribose) polymerase (anti-PARP) antibody (1/1,000; Cell Signaling), and mouse antihemagglutinin (anti-HA) antibody (clone 12CA5, 1/1,000; Roche).

Histological, immunohistochemical, and *in situ* hybridization analyses. Embryos were perfused intracardially with 4% paraformaldehyde (PFA). Dissected brains were postfixed overnight at 4°C and processed for paraffin or OCT embedding. Paraffin samples were cut at 5 μ m and used for hematoxylin and eosin (H&E) or for immunohistochemical analysis. In order to detect proliferating cells in S phase (bromodeoxyuridine [BrdU] labeling), a 1-h short pulse was performed as described previously (2). Paraffin-embedded sections were incubated with an anti-BrdU rat monoclonal antibody (1/75; Harlan, Indianapolis, IN). Mitotic cell activities were detected with a rabbit antibody, anti-Ser10 phospho-H3 (PH3, 1/2,000; Upstate). Frozen sections (16 μ m) were used for immunostaining. Cryosections were incubated in a blocking solution (phosphate-buffered saline [PBS]–0.2% Triton X-100–5% normal goat serum) for 20 min and with a mouse IgG blocking reagent (Vector) for 1 h. The primary antibodies used were mouse anti-RC2 (1/10; Hybridoma Bank, University of Iowa), rat anti-GFAP (1/500; U.S. Biological), rabbit anti-NG2 antibody (1/100; Chemicon), rat anti-PECAM antibody (1/2; gift from A. Vecchi, Milan, Italy), rabbit anti-Olig2 antibody (1/20,000; gift from C. Stiles, Boston, MA), and mouse anti-Ki67 (clone MM1, 1/100; Novocastra). Primary antibodies were diluted in PBS–1% normal goat serum, followed by appropriate cyanin-3 dye-conjugated, cyanin-2 dye-conjugated (1/500; Jackson ImmunoResearch Laboratories), Alexa488 dye-conjugated (1/500; Molecular Probe), or horseradish peroxidase (HRP) 3',3'-diamino-

benzidine staining system secondary antibodies. Cells were stained with Hoechst dye 33342 (2 μ g/ml) to visualize nuclei. Images were acquired using a Zeiss fluorescent microscope (Axiovert 200 M) with 16 \times and 40 \times objectives. Images were combined for figures by using Adobe Photoshop 8.0. *In situ* hybridization using antisense, digoxigenin-labeled *Pdgfra* riboprobe was performed as described previously (20).

Neurosphere culture. E18.5 forebrains from diverse CK2 β genetic backgrounds were subjected to trypsin digestion for 15 min at 37°C. Tissues were mechanically dissociated into single-cell suspensions, filtered through a 70- μ m cell strainer, and further treated with 10 μ g DNase I (Roche) for 10 min at room temperature. Cells from individual embryos, 2 \times 10⁶ cells/10-cm-diameter uncoated dish, were cultured for primary neurosphere assays in Dulbecco's modified Eagle's medium (DMEM)-F12 medium with B27 complement (D/F-B27) containing 10 ng/ml epidermal growth factor (EGF; PeproTech) and 10 ng/ml fibroblast growth factor-2 (FGF; gift from G. Bouche, Toulouse, France). Numbers of generated primary spheres were counted after 4 days of *in vitro* culture (4DIV). To assess for proliferation, 7DIV primary spheres were mechanically dissociated and individual viable cells were counted. These cells (6 \times 10⁵ cells/10-cm-diameter uncoated dish) were also used to assess for self-renewal in secondary neurosphere assays. Olig neurospheres were generated from neural stem/progenitor cells (NSCs) isolated from ganglionic eminences of *Olig2*^{+/-} or *Olig2*^{-/-} E14.5 embryos (18). For rescue experiments, *Olig2*^{-/-} NSCs were subsequently infected with HA-tagged Olig2-encoding viruses (see below) and selected with blasticidin (2 μ g/ml) 48 h after infection. Expression of HA-Olig2 proteins was confirmed by immunocytochemistry. For NSC differentiation assays, 4DIV neurosphere cultures were plated on poly-L-lysine-coated glass coverslips (at a density of ~50 neurospheres/cm²) and shifted into D/F-B27 medium in the presence of 3.5% fetal bovine serum (FBS; Biowest) (differentiating medium) (11). 3DIV after plating, differentiated cultures were fixed with the PFA solution and processed for immunocytochemistry. Additional primary antibodies were mouse anti- β -tubulin III (1/500; Babco), mouse anti-O4 (1/10; gift from C. Soula, Toulouse, France), rat anti-Pdgf receptor α chain (1/100; BD PharMingen), and rat anti-HA (clone 3F10, 1/500; Roche).

Embryonic stem cell culture. CK2 $\beta^{loxP/wt}$ and CK2 $\beta^{loxP/-}$ ES cell lines cultured in the presence of 10³ units/ml leukemia inhibitory factor (LIF; Chemicon) and Cre-pMSCV-puro viral supernatant have been described previously (2). The PCR product corresponding to HA-tagged CK2 β^{wt} coding regions was amplified using oligonucleotide DNA primers (5' primer A [5'-CGGAATTCGCCCA CCATGATCCTTATGATGTTCTGATTATGCTATGAGCAGTCCGAG GAGGTG-3'] and 3' primer B [5'-GAAGATCTTCAGCGGATGGTCTTCAC G-3']) and cloned into EcoRI/BglII pMSCV-neo viral vector sites (Clontech). CK2 β viral supernatants were generated by transfection of BOSC23 cells. Viral HA-CK2 β -pMSCV-neo supernatants were first used to infect 2.5 \times 10⁴/cm² CK2 $\beta^{loxP/-}$ ES cells. To avoid chimera colonies, one day after infection, ES cells were subjected to trypsin treatment and individual cells were plated. Neomycin (Neo) selection (350 μ g/ml) was started 48 h after infection, and Neo^r clones were expanded and screened for expression of exogenous the HA-CK2 β^{wt} protein by Western blot analysis using mouse anti-HA antibody. Selected Neo^r clones were further infected with Cre-pMSCV-puro supernatant. (CK2 $\beta^{loxP/wt}$ ES cells were used as positive controls.) One day later, individualized ES cells were plated, and puromycin (puro) selection (1.5 μ g/ml) was started 48 h postinfection for 6DIV or 9DIV. Individual clones were expanded and analyzed by PCR analysis and Southern blot analysis for the absence of the CK2 β^{loxP} allele (2) and by Western blot analysis for the absence of the endogenous wild-type CK2 β protein. Neo^r puro-resistant CK2 $\beta^{-/-}$ ES cells expressing only the exogenous HA-CK2 β^{wt} protein were seeded in the absence of LIF (3 \times 10⁶ cells/10-cm-diameter uncoated plate/10 ml) to form floating embryoid bodies (EBs). In order to stimulate neural differentiation, 5 μ M retinoic acid was added after 4DIV of EB cultures and left for additional 4 days. EBs were further dissociated with trypsin, and individual cells (3 \times 10⁶ cells/10-cm-diameter uncoated plate/10 ml) were used in neurosphere assays.

CK2 activity. Crude embryonic E18.5 forebrain extracts (adjusted to 5 mg/ml of protein concentration) were assayed for CK2 β -dependent CK2 kinase activity at 22°C for 10 min using a 3 mM concentration of the CK2 β -specific synthetic peptide RRREETEEEE (13) and [γ -³²P]ATP. One unit of CK2 activity is defined as the amount of activity necessary to transfer 1 fmol of phosphate per minute into the synthetic peptide. Chromatographic fractions (5 μ l) were assayed for 3 min with a 150 μ M concentration of the CK2 consensus synthetic peptide RRREEDSDDEE (32) or with a purified fusion protein fragment consisting of glutathione S-transferase (GST) and Olig2 amino acids 1 to 177 [Olig2(1-177)]. The PCR product corresponding to the mouse Olig2(1-177) coding region was amplified from the Olig2 CMV2 template (34) by using the Expand high-fidelity PCR system (Roche) with oligonucleotide DNA primers (5' primer C [5'-TTG

TABLE 1. Genotypes of live-born mice and embryos from *CK2β^{loxP/wt}, Nestin-cre × CK2β^{loxP/loxP}* crossings^a

Offspring or embryo	No. with genotype:			
	<i>CK2β^{loxP/wt}</i> (WT)	<i>CK2β^{loxP/loxP}</i> (WT)	<i>CK2β^{loxP/wt}, Nestin-cre</i> (WT)	<i>CK2β^{loxP/loxP}, Nestin-cre</i> (<i>CK2β^{-/-}</i>)
Live-born mice	22	18	15	0
P0	3	5	4	5
E14.5/E16.5/E18.5	142	146	148	154

^a For details, see Materials and Methods.

AATTCATATGGACTCGGACGCCAGCCT-3'] and 3' primer D [5'-AACTC GAGTCAGTAGATCTCGCTCACCAGTC-3']) and subcloned into EcoRI/XhoI pGEX4T2 vector sites (Amersham Biosciences) in order to generate the GST-Olig2(1-177) fusion protein. Purified GST-Olig2 proteins (20 μg) were incubated at 22°C for 0 to 30 min with 25 μM [γ -³²P]ATP and 12 mM MgCl₂ in the presence of 60 to 240 ng of CK2α subunit with increasing amounts of

equivalent CK2β subunit (0 to 230 ng). Phosphorylated proteins were analyzed by SDS electrophoresis and autoradiography.

Phosphopeptide mapping and automated Edman degradation. The phosphorylated GST-Olig2(1-177) fusion protein was excised from the nitrocellulose membrane and digested *in situ* with trypsin (modified sequencing grade; Promega). Two-dimensional phosphopeptide mapping was done using thin-layer electrophoresis (HTLE-7000; CBS Scientific). First-dimension electrophoresis was performed in pH 1.9 buffer (formic acid, glacial acetic acid, and water at 50:156:1,794, vol/vol/vol) for 30 min at 2,000 V, and chromatography in the second dimension was performed in isobutyric acid-*n*-butyl alcohol-pyridine-glacial acetic acid-water (1,250:38:96:58:558, vol/vol/vol/vol). After exposure, phosphopeptides were eluted from the plates in pH 1.9 buffer and lyophilized; aliquots of the samples were coupled to Sequelon-AA membrane (Millipore Corp.) by use of carbodiimide coupling and subjected to automated Edman degradation by using an Applied Biosystems sequencer (model 477A). Released phenylthiohydantoin derivatives from each cycle were spotted onto thin-layer chromatography plates. The radioactivity in each spot was revealed by exposure on a FujiX Bio-imager.

GST pull-down assay. Additional PCR products corresponding to coding regions for mouse Olig2 N-terminal amino acids 1 to 108 [N-ter(1-108)] and Olig2 bHLH amino acids 109 to 177 [bHLH(109-177)] were additionally ampli-

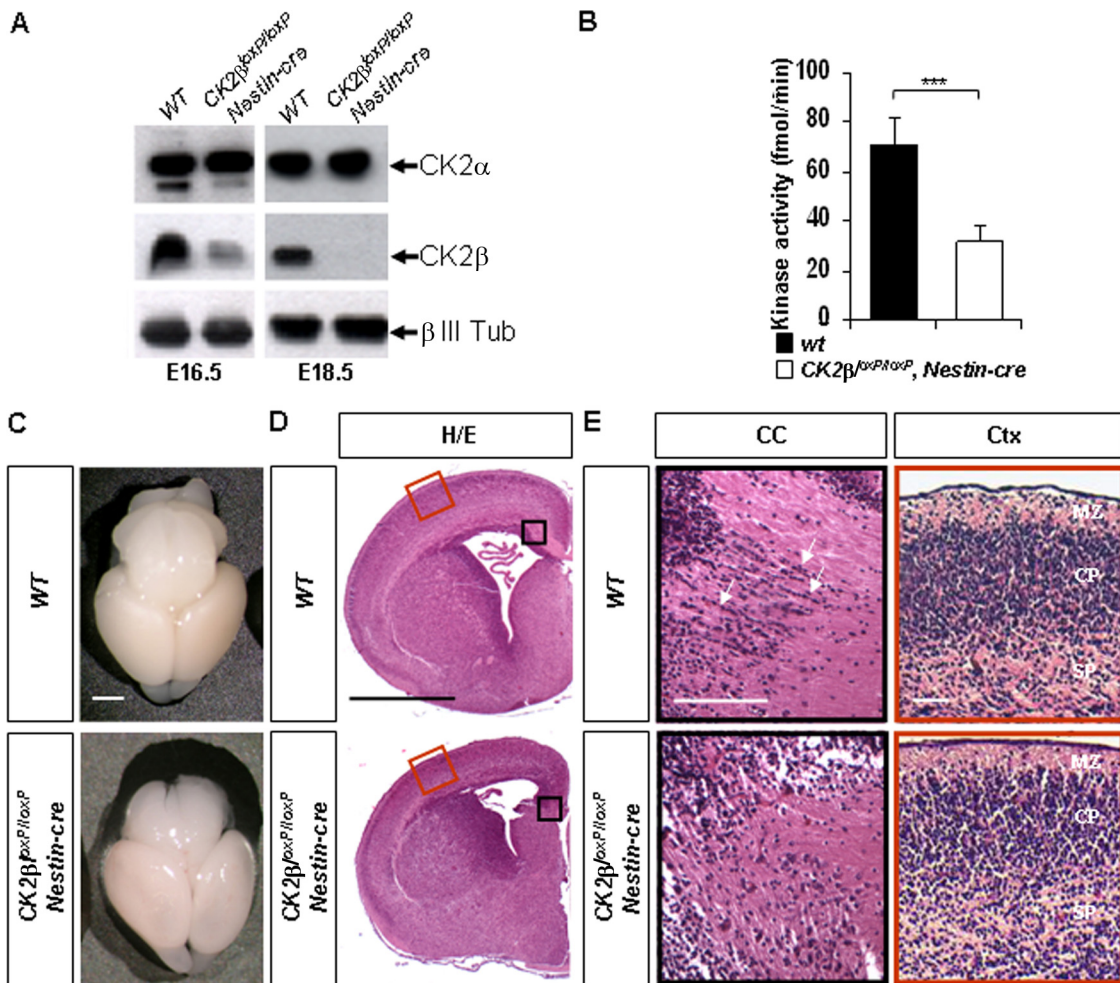


FIG. 1. *CK2β* loss in the central nervous system impairs telencephalon development. (A) Western blot analysis of *CK2β* and *CK2α* expressions in E16.5 and E18.5 forebrain extracts. A specific neuronal (class III β -tubulin [β III Tub]) antibody was used to control protein normalization. (B) *CK2β*-dependent CK2 activity in E18.5 *CK2β^{loxP/loxP}, Nestin-cre* forebrain extracts was significantly reduced (***, $P = 2.7 \times 10^{-9}$ [3 extracts in quadruplicate per genotype]) compared to that in wild-type controls. (C) Brain morphology of E18.5 *CK2β^{loxP/loxP}, Nestin-cre* embryos compared to that of the wild type. (D) Representative H&E staining of E18.5 anterior forebrains. Black and red boxed areas are shown at higher magnification in panel E. (E) At the level of the corpus callosum (CC; black boxes), linear arrays of cells (arrows) detected in wild-type embryos were absent in the mutants. In contrast, laminar patterns appeared similar in the cortex (Ctx; red boxes). MZ, mantle zone; CP, cortical plate zone; SP, subplate zone. Scale bars, 1 mm (C and D) and 100 μm (E).

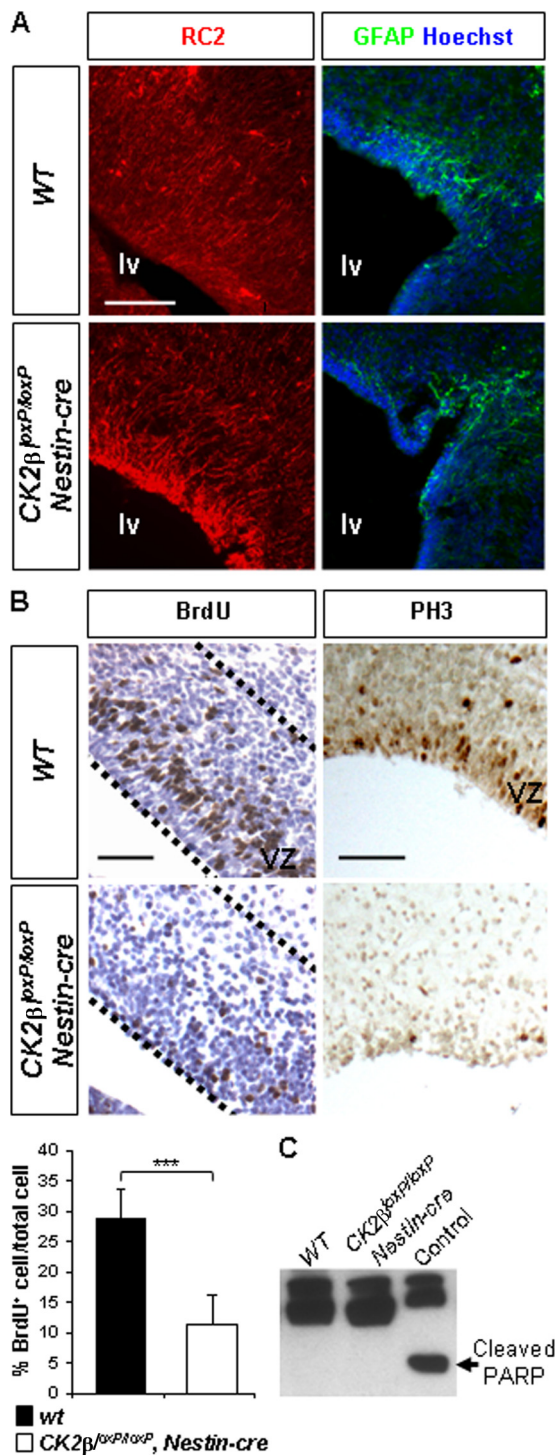


FIG. 2. *CK2β* positively controls forebrain NSC proliferation but did not interfere with NSC specification. (A) Analysis of RC2 (red) and GFAP (green) expressions by immunohistochemistry (IHC) in E18.5 wild-type and *CK2β^{loxP/loxP}, Nestin-cre* telencephalons. Cell nuclei were stained with Hoechst dye 33342 (blue). (B) There was a significant reduction (***, $P = 2.7 \times 10^{-12}$ [3 embryos per genotype and 3 nonadjacent sections per ventricular zone {VZ}]) of BrdU⁺ S-phase NSCs in *CK2β^{loxP/loxP}, Nestin-cre* VZs (outlined with a dotted line) of lateral ventricles compared to the number in wild-type controls, as well as a near absence of PH3-positive mitotic cells. (C) Caspase-3-dependent apoptosis analysis in E18.5 forebrains. Western blot analysis of cleaved PARP in E18.5 forebrain extracts. The

control represents the staurosporine-treated (1 μM, 3 h) NIH 3T3 positive extract; cleaved PARP (arrow) serves as a marker of cells undergoing caspase-3-dependent apoptosis. Scale bars, 100 μm (A) and 50 μm (B).

Control of HA-tagged mouse *Olig2* vectors. A PCR product corresponding to the HA-tagged mouse *Olig2*(1-177) coding region was amplified using 5' primer C (5'-AACTCGAGTCACTGCTGACGCTCGGGCTCAG-3') and using 5' primer F (5'-TTGAATTCATATGCTGCGCTGAAGATCAACA GCCGGA-3') with 3' primer D, respectively. Interaction assays were performed using prebound GST or GST-*Olig2* glutathione Sepharose beads (Amersham Biosciences) equilibrated in PBS-0.05% Tween 20-1 mg/ml bovine serum albumin (BSA) and incubated in the presence of [³⁵S]methionine-labeled CK2β (2 × 10⁵ cpm) at 22°C for 90 min. Bound CK2β subunits were analyzed by SDS electrophoresis and autoradiography.

Construction of HA-tagged mouse *Olig2* vectors. A PCR product corresponding to the HA-tagged mouse *Olig2*(1-177) coding region was amplified using 5' primer G (5'-TTGAATTCGGGCCATGTACCCATACGATGTTCCAGATT ACGCTATGGACTCGGACGCCAGCCT-3') and 3' primer H (5'-GAAGAT CTCGCTCACCAGTCGCT-3') and subcloned into EcoRI/BglII sites of the *Olig2* CMV2 vector (34) to generate HA-tagged full-length mouse *Olig2^{wt}*(1-323). The CMV2 vector with HA-*Olig2* STR amino acids 77 to 94 (*STR⁷⁷⁻⁹⁴*) deleted (HA-*Olig2^{ΔSTR}*) was generated by site directed mutagenesis using sense primer I (5'-TGGGCGGCGTGGCTTCAAGAAGAAGACAAGAAGCAG AT-3') and antisense primer J (5'-ATCTGCTTCTTGTCTTCTTCTTGAAG CCACCGCCGCCA-3'). Construction of pWZL-BLAST viral vectors encoding HA-*Olig2^{wt}* and HA-*Olig2^{ΔSTR}* proteins were generated after subcloning EcoRI/XhoI PCR fragments obtained after amplification with respective CMV2 templates by using a GC-rich DNA polymerase (Invitrogen) with 5' primer G and 3' primer K (5'-AACTCGAGTCACTTGGCGCTGGAGGTGAG-3').

Cos7 cell transfection and coimmunoprecipitation assay. For coimmunoprecipitations, a Flag-tagged CK2β construct was produced after PCR amplification using 5' primer L (5'-TTGGATCCGCGCCACCATGGACTACAAGGACG ACGACGACAAGATGAGCAGCTCCGAGG-3') and 3' primer M (5'-GAAG ATCTTCAGCGGATGGTCTTACAG-3') and cloned into BamHI/BglII pSG5 vector sites. The Flag-CK2β pSG5 and the HA-tagged full-length mouse *Olig2^{wt}*(1-323) CMV2 constructs were transiently Lipofectamine-transfected into Cos7 cells cultured in DMEM-10% fetal calf serum (FCS). Transfected cells were lysed in radioimmunoprecipitation assay (RIPA) buffer (48 h posttransfection) in the presence of protease inhibitors, and for each extract, protein expression levels were analyzed by Western blot analysis. Five micrograms of a monoclonal antibody against the Flag epitope (Sigma) were added in precleared lysates for 1 h at 4°C. Next, 50 μl of RIPA-equilibrated "mouse true blot" beads (eBioscience) was added and incubated under constant rotation for 1 h. After the beads were washed with RIPA buffer, the precipitated proteins were analyzed by SDS electrophoresis and Western blotting.

Statistical measures. For kinase activities, three forebrain-derived extracts per genotype were tested in quadruplicate. For telencephalon cell counts, at least three embryos per genotype were used for each day postcoitum (dpc) indicated. Cells were counted on three or more nonadjacent sections per telencephalon. For *in vitro* CK2β neurosphere assays, sphere and cell counts were performed from three or six embryo-derived cultures per genotype. For *in vitro* neurosphere differentiation assays, cell counts were performed from four or more embryo-derived cultures per genotype with six glass coverslips per culture (~50 neurospheres per coverslip). To evaluate the proliferation rate of neural stem cell-derived embryonic stem cells, cell counts were performed from seven differentiation cultures. Data are presented as averages ± standard errors. Statistical comparisons were established using Student's *t* test.

RESULTS

***CK2β* ablation in the central nervous system causes telencephalon defects during late development.** To gain insight into functions of *CK2β* in the developing central nervous system (CNS), we ablated *CK2β* in embryonic neural stem/progenitor cells (NSCs), since loss of *CK2β* is lethal to mouse embryos (2). We crossed *CK2β^{loxP/loxP}* mice with hemizygous *Nestin-cre* transgenic mice (37) to generate conditional mutant *CK2β^{loxP/loxP}*,

control represents the staurosporine-treated (1 μM, 3 h) NIH 3T3 positive extract; cleaved PARP (arrow) serves as a marker of cells undergoing caspase-3-dependent apoptosis. Scale bars, 100 μm (A) and 50 μm (B).

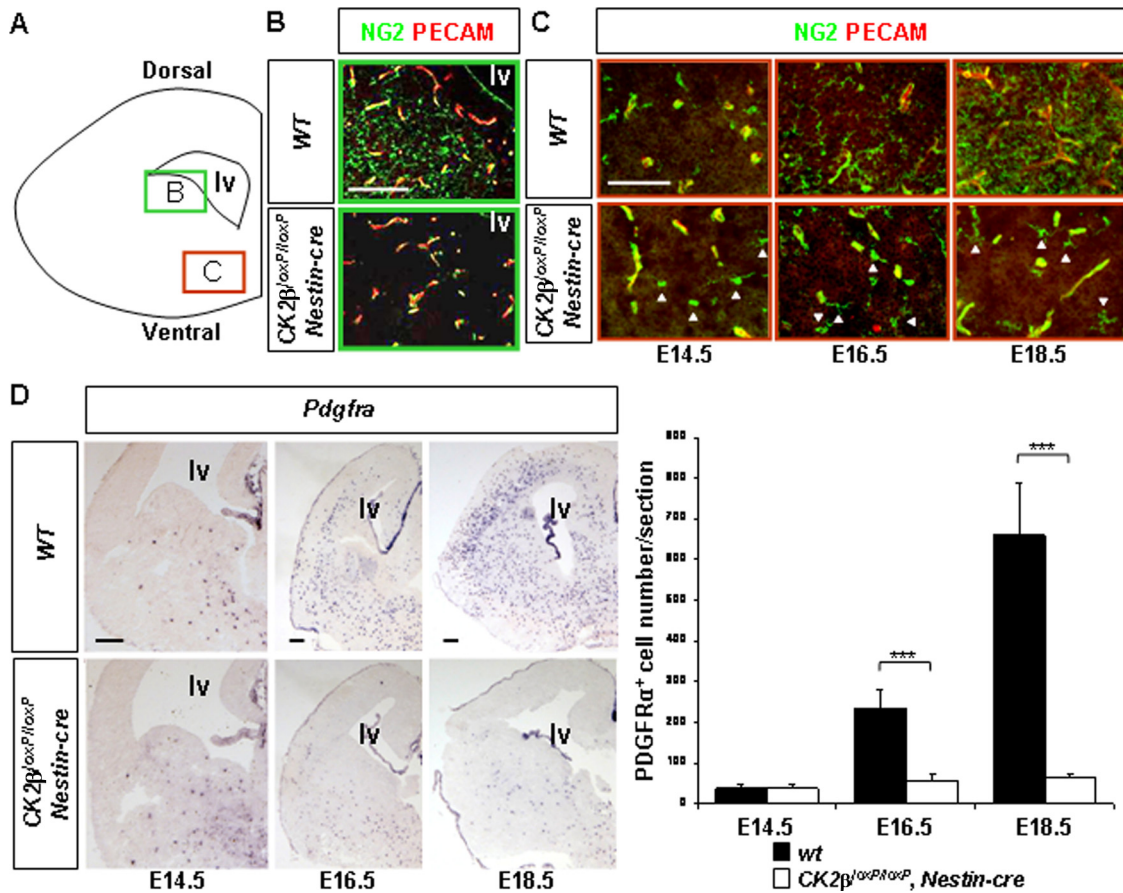


FIG. 3. Defective OPC development in $CK2\beta^{loxP/loxP}$, $Nestin-cre$ telencephalons. (A) Schematic of a coronal telencephalon section. The green and red boxes indicate the regions analyzed for the detection of parenchymal OPCs in panel B and of ventral OPCs in panel C, respectively. (B) Analysis of NG2 (green) and PECAM (red) expressions by immunohistochemistry (IHC) in E18.5 wild-type and $CK2\beta^{loxP/loxP}$, $Nestin-cre$ telencephalons. Note the normal appearance of multiprocessed NG2⁺ parenchymal OPCs in the wild type and their absence in $CK2\beta^{loxP/loxP}$, $Nestin-cre$ embryos. The pattern of NG2 staining in vessel pericytes parallels PECAM endothelial cell labeling (yellow) and was unaffected. (C) NG2 and PECAM expression analysis (IHC) revealed the presence of ventral NG2⁺ OPCs (arrowheads) in $CK2\beta^{loxP/loxP}$, $Nestin-cre$ fetal (E14.5) and late embryonic (E16.5 to E18.5) telencephalons. (D) *In situ* hybridization (ISH) analysis of *Pdgfra* expression in fetal (E14.5) and late embryonic (E16.5 to E18.5) telencephalons. There was a significant reduction (***, P values of 2.5×10^{-10} at E16.5 and 1.0×10^{-6} at E18.5 [4 embryos per dpc and per genotype and 3 nonadjacent sections per telencephalon]) of *Pdgfra*⁺ OPCs in $CK2\beta^{loxP/loxP}$, $Nestin-cre$ telencephalons compared to the number in wild-type controls. lv, lateral ventricle. Scale bars, 100 μ m (B and C) and 200 μ m (D).

Nestin-cre mice (referred to here as $CK2\beta^{-}$). $CK2\beta^{-}$ mutant pups were born with a Mendelian ratio (Table 1) but did not feed and died shortly after birth. Therefore, the phenotype of embryos was analyzed. As expected, the CK2β protein was absent in $CK2\beta^{-}$ mutant E18.5 forebrain extracts (Fig. 1A). However, the CK2β protein was still detected in $CK2\beta^{-}$ mutant E16.5 extracts, although Cre-mediated recombination occurs at E9.5 to E10.5 (10). This observation is in agreement with the stability of the CK2β protein integrated in the structure of tetrameric CK2 holoenzymes (22) and could explain the late $CK2\beta^{-}$ mutant phenotype. Importantly, the expression of the CK2α catalytic subunit was unchanged upon CK2β ablation. We also measured the CK2β-dependent CK2 activity in E18.5 forebrain extracts by using phosphorylation of the CK2β-specific synthetic peptide RRREE ETEE (13). In line with ablation of CK2β in $CK2\beta^{-}$ mutant forebrains, CK2β-dependent CK2 activity was significantly reduced (~60%) in $CK2\beta^{-}$ mutant extracts compared to wild-type controls (Fig. 1B). The remaining activity in mutant extracts could be related to holoenzyme molecules derived from blood vessel

cells present in dissected forebrains, in which Cre recombinase is not active (10). We conclude that expression of CK2β and, thus, the activity of CK2β-dependent phosphorylation are effectively impaired in $CK2\beta^{-}$ E18.5 forebrains.

To investigate the effects of loss of the CK2β regulatory subunit on the development of the central nervous system, we analyzed the morphologies of E18.5 mutant brains. Gross morphologies of wild-type and $CK2\beta^{-}$ E18.5 brains appeared similar (Fig. 1C). However, further histological analyses showed defects in $CK2\beta^{-}$ telencephalons: an absence at the corpus callosum (CC) level of cells that formed linear, pearls-on-a-string arrays characteristic of oligodendroglial cells (Fig. 1E) (33) and an absence of the emerging hippocampal dentate gyrus (data not shown). In contrast, laminar patterns and cellular density were not affected in the cortex (Ctx; Fig. 1D and E), suggesting that cortical neurogenesis occurred normally in $CK2\beta^{-}$ mutants. Thus, CK2β loss in embryonic NSCs leads to morphology defects during late development of the brain.

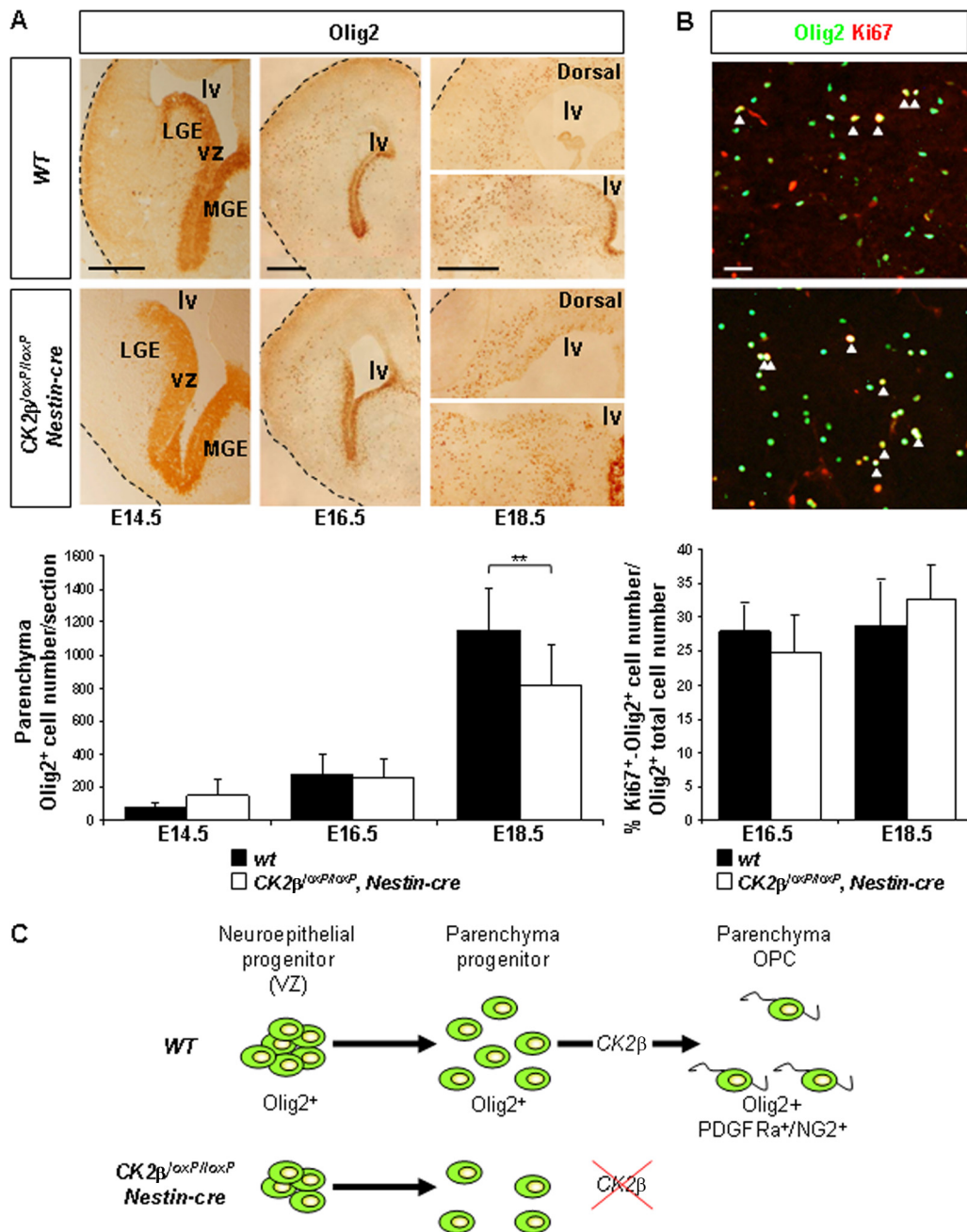


FIG. 4. Olig2⁺ progenitor analysis in fetal and late embryonic telencephalons. (A) IHC analysis of Olig2 expression in embryonic telencephalons. Parenchymal Olig2⁺ progenitors were detected in CK2β^{loxP/loxP}, Nestin-cre E14.5, E16.5, and E18.5 telencephalons. lv, lateral ventricle; vz, ventricular zone; MGE, medial ganglionic eminence; LGE, lateral ganglionic eminence. There was a small decrease (**, $P = 3.4 \times 10^{-3}$ [4 embryos per dpc and per genotype and 3 nonadjacent sections per telencephalon]) in parenchymal Olig2⁺ progenitor numbers in CK2β^{loxP/loxP}, Nestin-cre E18.5 telencephalons compared to the number in wild-type controls. (B) Proliferation analysis of parenchymal Olig2⁺ progenitors/OPCs by double labeling for Ki67 (red) and Olig2 (green). The ratios (percentages) of proliferating cells (Ki67⁺) coexpressing Olig2 in the parenchyma (yellow and arrowheads) to total parenchymal Olig2⁺ cells were not different between CK2β^{loxP/loxP}, Nestin-cre and wild-type E16.5 and E18.5 telencephalons (3 embryos per dpc and per genotype and 4 nonadjacent sections per telencephalon). (C) CK2β regulates OPC specification of parenchymal Olig2⁺ progenitors. Scale bars, 100 μm (A) and 50 μm (B).

CK2β disruption compromises forebrain stem/progenitor cell proliferation. To understand the causes of the defects observed in mutant brains, we studied NSCs. In mice, NSCs in the VZs of developing brains possess radial glia morphology and express RC2. During the late embryonic (E18.5)/neonatal

(P0) transition, radial glia gives rise to adult NSCs and progressively loose RC2 expression in favor of GFAP (23). In CK2β-ablated E18.5 embryos, neither RC2⁺ radial glia nor GFAP⁺ NSCs were substantially altered (Fig. 2A).

Previous results indicated that during mouse embryogenesis,

CK2 β is mainly involved in proliferation rather than in the prevention of apoptosis (2). Therefore, we examined at E18.5 the incorporation of a short pulse of BrdU in the germinal compartment of the telencephalon, the VZ of lateral walls (lw) of lateral ventricles (lv). CK2 β ⁻ VZs displayed a significant reduction (~60%) of NSCs in S phase as determined by the extent of BrdU-labeled nuclei compared to wild-type controls (Fig. 2B). In addition, the BrdU⁺ mutant nuclei showed a rounded morphology instead of an elongated morphology. To further address the effects of CK2 β loss on the cell cycle, sections were stained with the mitosis marker phospho-histone H3. Only few CK2 β ⁻ NSCs were found phospho-histone H3 positive (Fig. 2B). It is possible that the reduced NSC number represents a dysregulation of the cell cycle or an increase in cell death. We did not detect substantial apoptosis by terminal deoxynucleotidyltransferase-mediated dUTP-biotin nick end labeling (TUNEL) assay staining in CK2 β ⁻ E18.5 forebrains (data not shown). Accordingly, cleaved PARP, a hallmark of caspase-3-dependent apoptosis *in vivo*, was undetectable in CK2 β ⁻ E18.5 forebrain protein extracts (Fig. 2C). Taken together, these results suggest that CK2 β is important for the proliferation, but not the genesis, of forebrain NSCs.

CK2 β loss in NSCs impairs oligodendroglial differentiation in developing telencephalons. Next, we examined whether CK2 β regulates oligodendrogenesis, as suggested by disorganization of the corpus callosum of CK2 β ⁻ brains (Fig. 1E). We first analyzed the expression of the oligodendroglial precursor (OPC) marker NG2 (25) by immunohistochemistry (IHC). At E18.5, parenchymal NG2⁺ cells with OPC morphology were absent in CK2 β ⁻ telencephalons, and only NG2⁺ pericytes associated with blood vessel were detected (Fig. 3B). However, in CK2 β ⁻ brains, we were able to detect some NG2⁺ OPCs in a ventral location (Fig. 3C, E18.5 panel). In the mouse telencephalon, early OPCs (E12.5 to E14.5) are of ventral origin (reviewed in references 28 and 29). Thus, it was possible that the remaining E18.5 ventral OPCs in CK2 β ⁻ brains were generated earlier during early embryogenesis, a period when the CK2 β protein is not completely ablated (Fig. 1A). Figure 3C shows that early ventral NG2⁺ OPCs were generated in CK2 β ⁻ mutant E14.5 telencephalons, but in contrast to wild-type controls, they did not expand during the late embryonic period (E16.5 to E18.5). Because NG2 cytoplasmic immunostaining is unsuitable for quantification, we analyzed the expression of another OPC marker, *Pdgfra* (12, 25), by *in situ* hybridization. At E14.5, the numbers of *Pdgfra*⁺ OPCs in wild-type and CK2 β ⁻ brains were similar (Fig. 3D). However, there was a sharp decrease (~75%) in the number of *Pdgfra*⁺ OPCs in E16.5 mutant telencephalons, despite incomplete inactivation of the CK2 β protein. By E18.5, this difference was more pronounced (~90%).

We also investigated the expression of Olig2, a basic helix-loop-helix (bHLH) transcription factor required for OPC development and maintained at all stages of oligodendrocyte development (17, 21, 29, 36, 43). In contrast to the defect observed for NG2⁺ and *Pdgfra*⁺ OPCs, analysis of Olig2 expression revealed identical numbers of parenchymal Olig2⁺ cells in CK2 β ⁻ telencephalons and wild-type controls at E14.5 and E16.5 (Fig. 4A). However, by E18.5, a small decrease (~30%) in the number of parenchymal Olig2⁺ cells was observed in CK2 β ⁻ telencephalons. Because (i) Olig2 identifies

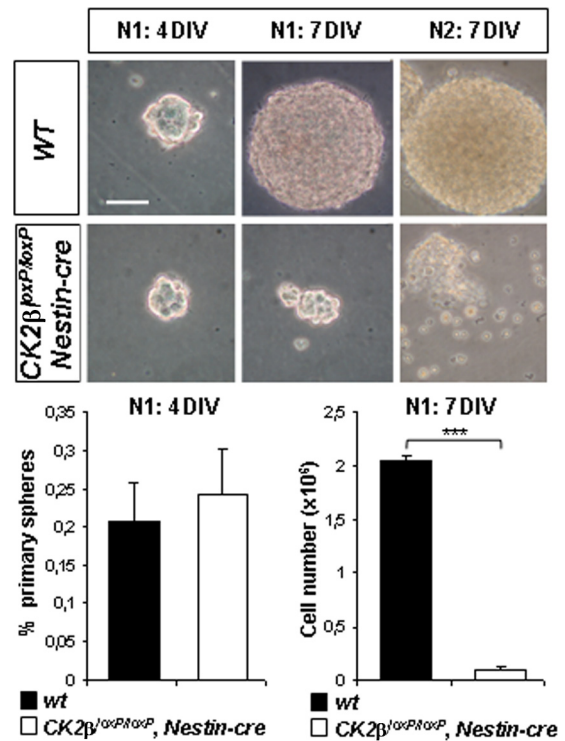


FIG. 5. CK2 β regulates NSC proliferation in neurosphere cultures. Primary neurospheres (N1; 4DIV and 7DIV) derived from E18.5 forebrain NSCs are shown. Compared to wild-type controls, CK2 β ^{loxP/loxP}, Nestin-cre primary neurospheres in 7DIV cultures became smaller and did not generate secondary neurospheres (N2) after dissociation (7DIV). The ratio (percentage) of generated primary neurospheres to E18.5 forebrain-dissociated cells (6 primary neurosphere cultures per genotype) in CK2 β ^{loxP/loxP}, Nestin-cre 4DIV cultures was not different from that of wild-type controls. The number of cells obtained from dissociated CK2 β ^{loxP/loxP}, Nestin-cre primary neurospheres in 7DIV cultures was significantly reduced compared to that from wild-type controls (***, $P = 4.3 \times 10^{-7}$ [3 primary neurosphere cultures per genotype]). Scale bar, 100 μ m.

both neural progenitors and OPCs (27, 28) and (ii) the decrease in the number of parenchymal Olig2⁺ cells is lower than the decrease in the number of *Pdgfra*⁺ OPCs in CK2 β ⁻ brains (0% versus ~75% at E16.5 and ~30% versus ~90% at E18.5), parenchymal Olig2⁺ cells in CK2 β ⁻ brains may represent stalled neural progenitors unable to progress along the oligodendrocyte lineage. Alternatively, these Olig2⁺ cells may represent abnormal glia. To investigate whether the decrease in OPC numbers was due to defective proliferation of parenchymal Olig2⁺ progenitors as described for NSCs (Fig. 2B), we determined the ratio of Ki67⁺ Olig2⁺ cells to total Olig2⁺ cells in the parenchyma of wild-type and CK2 β ⁻ telencephalons at E16.5 and E18.5 (Fig. 4B). Proliferation of parenchymal Olig2⁺ cells was unchanged in CK2 β ⁻ brains and is therefore unlikely to account for OPC deficiency. Because parenchymal Olig2⁺ progenitors originate from Olig2⁺ neuroepithelial cells in the VZ (28, 29), the decrease in the number of Olig2⁺ cells in the parenchyma of CK2 β ⁻ brains may be a consequence of a defective proliferation of Olig2⁺ cells in the VZ (Fig. 4C). The discrepancy observed between the effects of CK2 β disruption on the proliferation of NSCs in the VZ and that of Olig2⁺

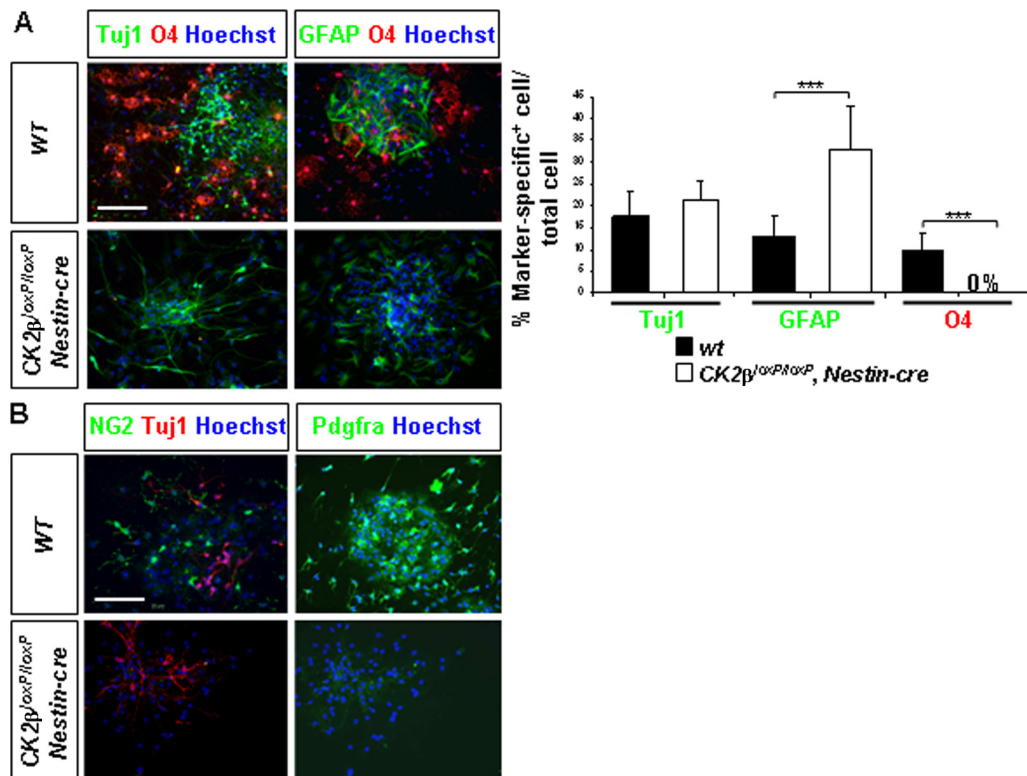


FIG. 6. *CK2β* disruption impairs OPC specification in neurosphere cultures. (A) Primary neurospheres derived from E18.5 forebrains were tested for their multipotency after differentiation. Cells were immunolabeled with the neuronal marker Tuj1 (green), the astrocytic marker GFAP (green), and the oligodendroglial marker O4 (red). The ratio (percentage) of cells expressing lineage-specific markers to the total cell number evaluated by Hoechst-stained nuclei (blue) showed an absence of O4⁺ cells (***, $P = 1.3 \times 10^{-8}$ [12 neurosphere cultures per genotype]) concomitant with a significant increase of GFAP⁺ astrocytes (***, $P = 2.9 \times 10^{-4}$) in *CK2β*^{loxP/loxP}, *Nestin-cre* cultures compared to the number in wild-type controls. The percentages of Tuj1⁺ neurons did not differ between wild-type and mutant cultures. (B) *CK2β* loss altered the specification of NG2⁺ (green) and Pdgfra⁺ (red) OPCs in differentiation cultures. Scale bars, 100 μm.

progenitors in the parenchyma may reflect different mechanisms involved in the control of the proliferation of progenitors downstream of the stem compartment. Collectively, these data suggest that *CK2β* loss leads to a defect in OPC specification of Olig2⁺ progenitors during telencephalon development (Fig. 4C).

***CK2β* is required for neural stem cell proliferation and oligodendroglial differentiation in neurosphere culture.** Embryonic NSCs located in the telencephalic VZ have been identified by their ability *in vitro* to generate neurospheres in response to FGF2 and EGF stimulation (38). In the absence of growth factors, neurosphere cells differentiate into neurons, astrocytes, and oligodendrocytes. To confirm whether *CK2β* loss affects NSC maintenance as shown *in vivo*, we cultured neurospheres derived from *CK2β*^{-/-} E18.5 forebrains and induced their differentiation. There was no significant difference in the numbers of primary neurospheres generated from E18.5 wild-type and *CK2β*^{-/-} forebrains in response to FGF2-EGF after 4 days of *in vitro* culture (4DIV) (Fig. 5). As observed in 7DIV cultures, wild-type cells formed large neurospheres, whereas *CK2β*^{-/-} neurospheres did not expand. This was reflected by a significant reduction (~95%) in the number of cells obtained from dissociated *CK2β*^{-/-} neurospheres. In addition, we were unable to generate *CK2β*^{-/-} secondary neurospheres, suggesting that the self-renewing potency of *CK2β*^{-/-} NSCs was severely compromised. After differentiation, *CK2β*^{-/-} neurosphere cells

formed β-tubulin III-positive neurons and GFAP⁺ astrocytes but failed to generate O4⁺ differentiating OPCs (Fig. 6A). To confirm whether this defect was the result of a failure of OPC specification as observed *in vivo*, we examined NG2 and Pdgfra expression in differentiated neurosphere cultures. In striking contrast to control cultures, *CK2β*^{-/-} neurospheres did not generate any NG2⁺ or Pdgfra⁺ OPCs (Fig. 6B). To further address the requirement of *CK2β* in NSC proliferation and in oligodendrogenesis, we expressed an HA-tagged *CK2β* protein in *CK2β*^{-/-} NSC-derived embryonic stem (ES) cells (Fig. 7) (2). Such *CK2β*^{-/-} NSCs expressing HA-*CK2β* were able to proliferate and to generate O4⁺ differentiating OPCs, excluding a genetic side effect associated with targeting of the *CK2β* locus. Taken together, these results suggest that the cell-autonomous defect(s) caused by *CK2β* disruption results in deficient NSC proliferation and OPC specification.

Olig2 is an interacting partner of *CK2β* subunit and a strict *CK2β*-dependent substrate *in vitro*. Given the essential roles of *Olig2* and *CK2β* for OPC specification in fetal and late embryonic development (17, 21, 36, 43; this study) and the regulation of the oligodendroglial activity of *Olig2* by phosphorylation (31), we investigated whether *Olig2* may be a potential *CK2* substrate. Amino acid sequence analysis of mouse *Olig2* revealed that most putative *CK2* phosphorylation sites (S/T; E/D/S $n - 1, n + 1, n + 2$, and/or $n + 3$) are present within an

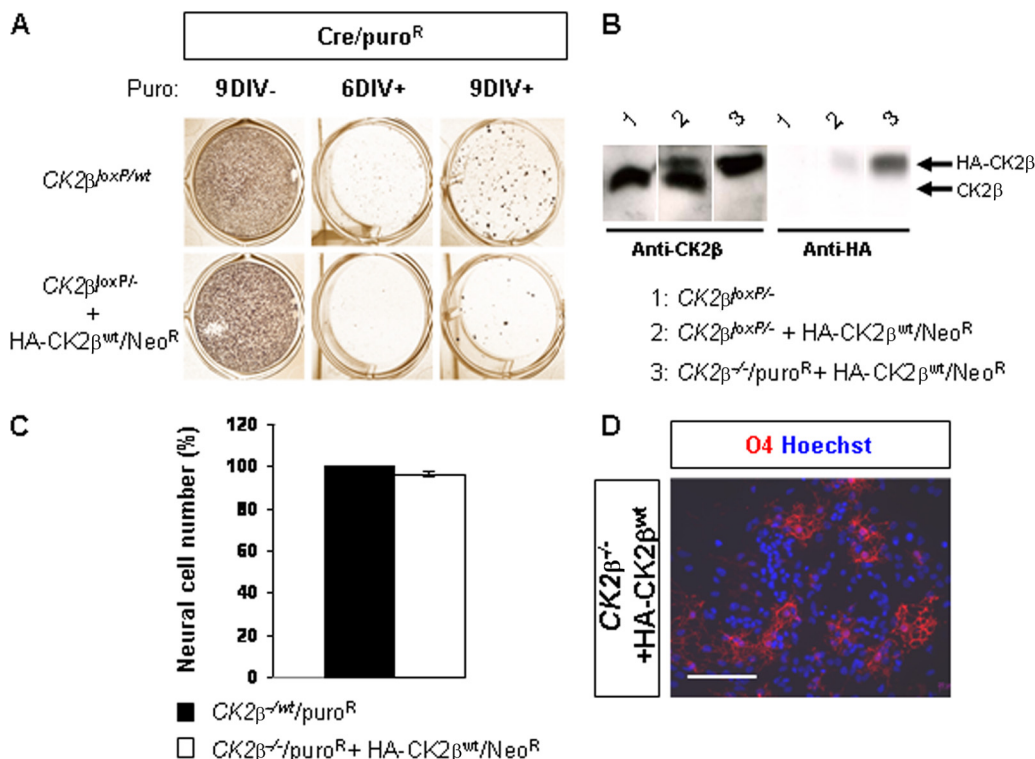


FIG. 7. Exogenous CK2β expression in NSC-derived CK2β^{-/-} embryonic stem (ES) cells promotes proliferation and oligodendroglial differentiation. (A) Clonal selection of CK2β^{-/-}, Cre-pMSCV-puro-cultured (Cre/puro^R) ES cells expressing exogenous HA-CK2β^{wt}/Neof^R protein. CK2β^{-wt}, Cre/puro^R ES cell clones were generated in parallel and served as positive controls. (B) Western blot analysis with anti-CK2β (left panel) and anti-HA (right panel) antibodies of CK2β^{loxP/-} ES cells, CK2β^{loxP/-} and HA-CK2β^{wt}/Neof^R ES cells, and CK2β^{-/-}, Cre/puro^R, and HA-CK2β^{wt}/Neof^R ES cells. Note the complete absence of the endogenous CK2β protein in the CK2β^{-/-} ES cell extract. (C) Proliferation analysis of NSC-derived ES cells. The ratio (percentage) of the number of cells present in neurosphere-derived CK2β^{-/-} ES cells expressing the exogenous HA-CK2β^{wt} protein to the number of cells present in control neurosphere-derived CK2β^{-wt} ES cells was unchanged when normalized to the CK2β^{-wt} ES cell line (7 differentiation cultures). (D) O4⁺ differentiating OPCs were identified from neurosphere-derived CK2β^{-/-} ES cells expressing the exogenous HA-CK2β^{wt} protein and allowed to differentiate. Scale bar, 100 μm.

N-terminal fragment encompassing M¹ to Y¹⁷⁷. We designed a GST-Olig2(1-177) fusion protein that includes mouse Olig2 amino acids 1 to 177 (Fig. 8A) and determined the CK2-dependent incorporation of [³²P]ATP into the purified fragment. As shown in Fig. 8B, the Olig2(1-177) fragment was directly phosphorylated by CK2 only in the presence of the regulatory subunit CK2β. We further investigated CK2β-dependent phosphorylation of Olig2 by analyzing CK2 and Olig2 phosphorylation activities in fractions obtained after ion-exchange chromatography of wild-type E18.5 forebrain extracts (Fig. 8C). Phosphorylation experiments and Western blot analysis revealed coelution of CK2 and Olig2-dependent kinase. This coelution occurred in fractions containing the highest CK2β/CK2α subunits ratio. Moreover, no coelution of CK2 and Olig2-dependent kinase was observed when the chromatography was performed with CK2β⁻ forebrain extracts (data not shown). In addition, Olig2 phosphorylation was abolished in the presence of 10 μM TBB, a specific *in vitro* CK2 inhibitor (data not shown). Taken together, these results suggest that the CK2 tetrameric holoenzyme (α₂β₂) is an Olig2 kinase.

To delineate which particular motif of the mouse Olig2(1-177) fragment was involved in the phosphorylation by the CK2 holoenzyme, we performed two-dimensional tryptic phosphopeptide mapping (Fig. 8D). Analysis of the maps of ³²P-

labeled phosphopeptides revealed multiple spots. A major spot was suitable for Edman degradation. The released detectable radioactivity was observed mainly after cycles 9 and 11. Only one Olig2(1-177) tryptic peptide had serine/threonine residues at positions 9 and 11, allowing the identification of major phosphorylated sites at Ser⁸⁵ and Ser⁸⁷ in a serine-threonine-rich domain (STR⁷⁷⁻⁹⁴, tryptic peptide ⁷⁷SSSSSTSSSTSSAAT SSTK⁹⁵) highly conserved in the human sequence. It should be noted that minor phosphorylation sites were also detected at Ser⁸⁴, Thr⁸⁶, and Ser⁸⁸. Thus, these results identify the Olig2 STR sequence as an unconventional major domain of multiple acceptors for phosphorylation by the CK2 holoenzyme.

To investigate the regulation of Olig2 phosphorylation by CK2β, we tested the capacity of CK2 to interact with Olig2. GST pull-down assays showed that CK2β interacts with the bHLH fragment of Olig2 [Olig2(109-177)] but fails to interact with the fragment containing the STR domain [Olig2(1-108)] (Fig. 8E). Surprisingly, this Olig2(1-108) fragment was not phosphorylated by the CK2 holoenzyme (data not shown). Preincubation of CK2β with CK2α subunits which generate the tetrameric holoenzyme did not prevent its interaction with the bHLH Olig2(109-177) fragment (data not shown). We confirmed that CK2β can interact with Olig2 by coimmunoprecipitation of Flag-tagged CK2β with HA-tagged full-length

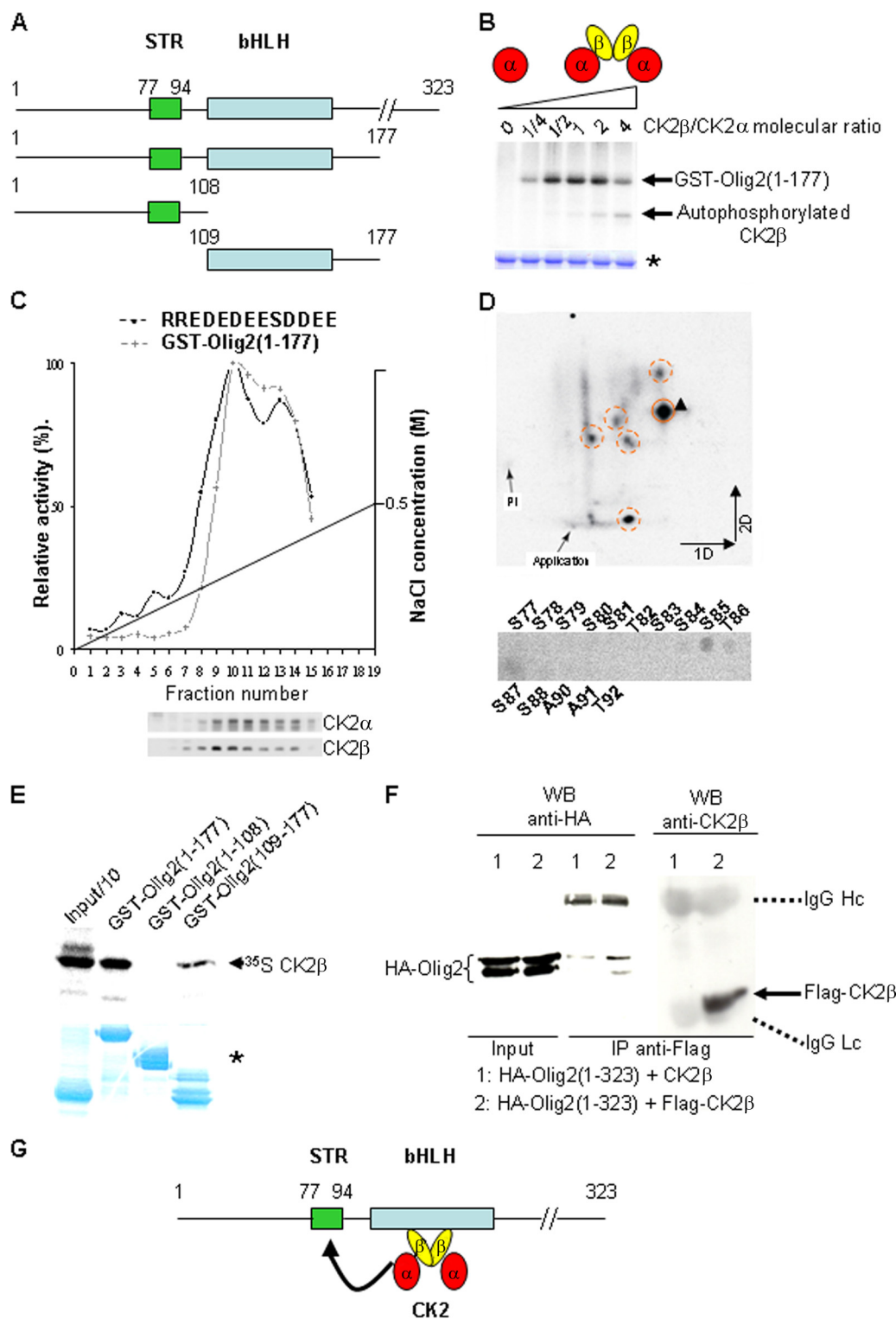


FIG. 8. CK2 β positively modulates CK2-dependent phosphorylation of the Olig2(1-177) fragment *in vitro*. (A) Primary structure of mouse Olig2 and schematic representation of the Olig2 fragments used in this study. Represented are the serine-threonine-rich domain sequence (STR, amino acids 77 to 94), and the basic helix-loop-helix sequence (bHLH, amino acids 109 to 164). (B) Autoradiography of the purified recombinant GST-Olig2(1-177) fusion fragment after *in vitro* phosphorylation by the CK2 catalytic subunit (α) alone or in combination with increasing amounts of the CK2 regulatory subunit (β), which generate increased amounts of the holoenzyme ($\alpha_2\beta_2$). Note that CK2 β , an autosubstrate of the catalytic α subunit, competed in the reaction for the CK2 phosphorylation of Olig2 when present in excess. Protein normalization was controlled by Coomassie blue staining (*). (C) DEAE-Sepharose chromatography of crude wild-type E18.5 forebrain protein extracts. Each fraction was analyzed for consensus relative CK2 activity with the RRREDEESDDEE synthetic substrate/peptide and for relative CK2 β -dependent Olig2(1-177) kinase

mouse Olig2(1-323) after transient expression in Cos7 cells (Fig. 8F). Together, these data suggest that the bHLH domain of Olig2 is a docking site to promote the recruitment of CK2 via physical interaction with its regulatory CK2 β subunit for efficient Olig2 phosphorylation on its STR sequence (Fig. 8G).

The CK2-targeted STR domain is required for the oligodendroglial activity of Olig2 in neurosphere culture. We next asked whether the CK2-phosphorylated STR domain is involved in mediating Olig2 biological function. Because of the many serine and threonine acceptor residues in the Olig2 STR that could potentially compensate for inactivation of the Ser⁸⁵ and Ser⁸⁷ major phosphorylation sites, we generated an Olig2 deletion mutant that lacks the entire STR⁷⁷⁻⁹⁴ domain (HA-Olig2 ^{Δ STR}). Transient transfection into Cos7 cells demonstrated identical levels of expression of HA-Olig2^{wt} (amino acids 1 to 323; full-length) and HA-Olig2 ^{Δ STR} proteins (Fig. 9A). We infected Olig2^{-/-} neurosphere cultures (18) with retroviruses encoding HA-Olig2^{wt} or HA-Olig2 ^{Δ STR} and tested their potential to differentiate into the oligodendroglial lineage (Fig. 9B). As expected, expression of HA-Olig2^{wt} in Olig2^{-/-} NSCs restored the production of O4⁺ differentiating OPCs. In contrast, Olig2^{-/-} NSCs transduced with HA-Olig2 ^{Δ STR} significantly failed to differentiate into O4⁺ cells. These findings demonstrate that, *in vitro*, the CK2-targeted STR domain is required for the oligodendroglial function of Olig2.

DISCUSSION

Kinase-dependent pathways regulating NSC maintenance and fate specification are largely unknown. Here, we inactivated the protein kinase CK2 regulatory subunit (CK2 β) gene in CNS embryonic neural progenitors and showed that CK2 β is an important modulator of telencephalon development. Conditional CK2 β mutant mice display impaired NSC proliferation, as well as defective OPC specification. We show that CK2 β interacts physically with the bHLH transcription factor Olig2, an essential regulator of oligodendroglial differentiation, thereby allowing its serine-threonine-rich (STR) domain to be phosphorylated by CK2. Finally, we show that the Olig2 STR domain mediates the oligodendroglial activity of Olig2. Our results suggest that CK2 β could mediate OPC specification through the modulation of the oligodendroglial activity of Olig2.

We present the first report of an *in vivo* role for CK2 β in NSC proliferation in mammals. Loss of CK2 β results in reduced numbers of dividing cells in the embryonic germinal zone, and our *in vitro* neurosphere assays show that CK2 β is required for NSC proliferation and self-renewal. The altered

nuclear morphology of BrdU⁺ CK2 β ⁻ NSCs in the ventricular zone (Fig. 2B) is reminiscent of a cell cycle arrest event and suggests an interkinetic nuclear migration defect (40). During G₁, the nuclei of NSCs ascend to the basal side of the VZ, where S phase is completed. Nuclei descend back toward the ventricular surface during G₂, and this is where they stay during mitosis. The present study suggests that CK2 β may control this process in the neural stem compartment, leading to a block in the G₁/S phase in CK2 β ⁻ NSCs. Therefore, CK2 β ⁻ NSCs would not proceed through the G₂ phase. Consequently, it would explain the near absence of PH3-positive mitotic CK2 β ⁻ NSCs in the VZ (Fig. 2B) and the normal proliferation of the more downstream population of Olig2⁺ parenchymal progenitors in CK2 β ⁻ embryos (Fig. 4B). Recently, it has been demonstrated that nuclear migration is governed by centrosomal specific microtubule-regulating proteins (40) and reported that CK2 β is required for centrosomal normality (1). However, a strict CK2 β -dependent CK2 substrate(s) involved in NSC proliferation has not been described.

We also show that CK2 β ⁻ NSCs are specifically devoid of oligodendroglial potency. Our *in vivo* and *in vitro* studies demonstrate that depletion of the CK2 β gene in embryonic neural progenitors blocks the production of NG2⁺ and Pdgfra⁺ OPCs. Several observations are in favor of a cell-intrinsic regulation by CK2 β in OPC specification *per se*. (i) In E16.5 CK2 β ⁻ telencephalons, the number of parenchymal Olig2⁺ progenitors was normal, despite a strong defect in NG2⁺ and Pdgfra⁺ OPCs, and by E18.5, whereas the number of Olig2⁺ progenitors increased, the number of OPCs in CK2 β ⁻ telencephalons remained low compared to that in wild-type controls (Fig. 3 and 4). (ii) Olig2⁺ cell proliferation was unchanged in CK2 β ⁻ brains (Fig. 4B). (iii) CK2 β ⁻ neurosphere cultures recapitulated the deficiency in OPC specification observed *in vivo* (Fig. 6). (iv) CK2 β ⁻ neurospheres cultured in the presence of other factors (PDGF-AA and insulin-like growth factor I [IGF-I]) were unable to differentiate into oligodendrocytes (data not shown), suggesting that a defect in a particular extracellular instruction for NSC specification to oligodendrogenesis cannot account for OPC deficiency (3, 5, 8, 14, 35). (v) CK2 β is required to promote oligodendrogenesis in NSC-derived embryonic stem (ES) cells (Fig. 7). Thus, CK2 β -dependent specification of OPCs may correspond to a molecular mechanism(s) shared by the multiple independent molecular, regional, and temporal origins of progenitors in the forebrain (reviewed in reference 28) and controlled by CK2 β .

Genetic studies in mice demonstrated that during fetal and late embryogenesis, the bHLH transcription factor Olig2 is necessary for specification of progenitors into OPCs/oligoden-

activity. Activity profiles were normalized to the activity detected in the peak fractions. In parallel, a sample of each fraction was immunoblotted to detect the presence of CK2 α and CK2 β proteins (lower panels). (D) Two-dimensional (2D) phosphopeptide mapping analysis of CK2 β -dependent Olig2(1-177) phosphorylation. The major phosphopeptide (arrowhead; upper panel) was subjected to automated Edman degradation and released derivatives from each cycle were spotted onto plates and autoradiographed (lower panel). Deduced amino acids in the Olig2 sequence are indicated above or below the spots. (E) Recombinant GST fusion proteins containing different Olig2 fragments (1 to 177, 1 to 108, and 109 to 177) were incubated with [³⁵S]methionine-labeled CK2 β . The ability of the GST fusion proteins to pull down the CK2 β subunit was detected by autoradiography after electrophoresis. On the left, one-tenth of the input is shown. Loading controls were checked after Coomassie blue staining of the gel (*). (F) Coimmunoprecipitation of Flag-CK2 β and HA-Olig2(1-323) in Cos7 cells. Flag-CK2 β was immunoprecipitated (IP) with anti-Flag antibody, and the immunoprecipitated complexes were analyzed by Western blotting (WB) with anti-HA and anti-CK2 β antibodies. Hc, heavy chain; Lc, light chain. (G) Schematic representation of the suggested CK2-Olig2 interaction.

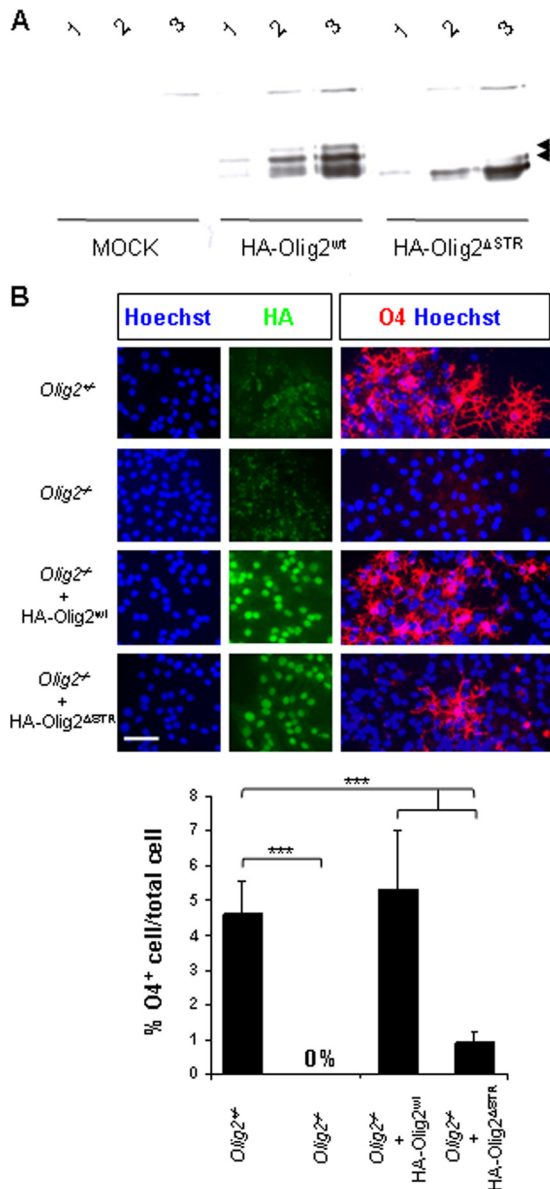


FIG. 9. The Olig2 STR domain is required for NSC oligodendroglial potency in neurosphere cultures. (A) Transient expression analysis of the HA-Olig2^{ΔSTR} mutant protein in Cos7 cells. Comparative pattern expressions of full-length (mouse; amino acids 1 to 323) HA-Olig2^{wt} and HA-Olig2^{ΔSTR} in transiently transfected Cos7 cells. Anti-HA Western blots revealed shifted-up bands within the HA-Olig2^{wt}-transfected protein extracts (arrowheads), and these were not detected in the HA-Olig2^{ΔSTR} version. This observation is reminiscent of the STR sequence as phosphorylated sites. "MOCK" indicates a control experiment performed with the empty vector. Amounts of protein are as follows: 1, 4 μ g; 2, 8 μ g; 3, 16 μ g. (B) Olig2^{-/-} NSCs were transfected with HA-Olig2^{wt}(1-323) and HA-Olig2^{ΔSTR}. NSC-derived neurospheres were allowed to differentiate and stained with HA (green) to detect exogenous Olig2 and with O4 (red) to assess for oligodendroglial potency. Olig2^{+/-} and Olig2^{-/-} assays are shown in parallel. The ratio (percentage) of cells expressing the O4 oligodendroglial marker to the total cell number evaluated by Hoechst-stained nuclei (blue) showed a significant deficiency (***, $P = 5.6 \times 10^{-7}$ [4 neurosphere cultures per genotype]) in the oligodendroglial activity of the HA-Olig2^{ΔSTR} mutant compared to that of the HA-Olig2^{wt} control. Scale bars, 50 μ m.

drocytes (17, 21, 36, 43). Multiple regulations such as interactions (NFIA/B, SCL, and ID4) and phosphorylation (Akt/protein kinase B [PKB]) that inhibit Olig2 function dictate a loss into the oligodendroglial lineage (6, 24, 30, 31). The increase in astrocytogenesis observed in CK2 β ⁻ neurosphere cultures after differentiation (Fig. 6A) is compatible with a glial switch following loss of the oligodendroglial function of Olig2 (6, 7, 21, 24, 30, 31, 36, 43). However, whether CK2 β modulation of astrocytic instruction is direct or a consequence of a regulation of oligodendrocyte differentiation is unclear. We show that the Olig2(1-177) fragment is a strict CK2 β -dependent CK2 substrate, probably through a CK2 β -specific Olig2 recruitment by direct interaction with the bHLH domain, and that CK2 is an Olig2 kinase in forebrain extracts (Fig. 8). CK2-dependent major phosphorylation occurs in the STR domain. In neurosphere assays, we demonstrated that the CK2-targeted STR domain is required for the full oligodendroglial function of Olig2 (Fig. 9). Although we cannot exclude phosphorylation(s) of the STR domain by one or more other serine/threonine kinases, our results do not definitively prove, but do suggest, that CK2 β -dependent CK2 activity may play a role as a determinant in the combinatorial code that trigger Olig2-dependent oligodendrocyte lineage progression. Alternatively, we cannot rule out other CK2 β -dependent transcription factor substrates important for the oligodendroglial network which could act in synergy inside the same genetic pathway.

During CNS development, Olig2 is sequentially required for the generation of subsets of motor neurons (E9 to E10.5) and OPCs (E12.5 to E14.5) (21, 36, 43). If the assumption for a defect in Olig2 function in CK2 β ⁻ progenitors is correct, then it is intriguing that the process of neurogenesis is not affected. However, in CK2 β mutant brains, complete ablation of CK2 β does not occur before E18.5, by which neurogenesis is largely completed. Additional studies will be needed to address the earlier role of CK2 β in neuronal development.

The essential role of CK2 β for cell viability initially detected in ES cells (2) precluded analysis of later roles for CK2 during development. Our conditional strategy revealed unexpected roles for CK2 β in the CNS. We demonstrated that CK2 β represents a unique signaling component in NSC proliferation and self-renewal and in the oligodendroglial lineage-restricted pathway. If CK2 β -dependent CK2 activity is required for cancer stem cell proliferation and self-renewal as it is for NSCs, then inhibitors of interaction between the CK2 β regulatory subunit and CK2 catalytic subunits (α and α') could be useful chemotherapeutic agents. Conversely, because differentiation block of OPCs has been demonstrated to cause neurodegenerative diseases, such as chronic multiple sclerosis (16), activation of CK2 β -dependent CK2 activity may be useful in promoting remyelination.

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