

Dual Effect of CTCF Loss on Neuroprogenitor Differentiation and Survival

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An increasing number of proteins involved in genome organization have been implicated in neurodevelopmental disorders, highlighting the importance of chromatin architecture in the developing CNS. The CCCTC-binding factor (CTCF) is a zinc finger DNA binding protein involved in higher-order chromatin organization, and mutations in the human *CTCF* gene cause an intellectual disability syndrome associated with microcephaly. However, information on CTCF function *in vivo* in the developing brain is lacking. To address this gap, we conditionally inactivated the *Ctcf* gene at early stages of mouse brain development. Cre-mediated *Ctcf* deletion in the telencephalon and anterior retina at embryonic day 8.5 triggered upregulation of the p53 effector PUMA (p53 upregulated modulator of apoptosis), resulting in massive apoptosis and profound ablation of telencephalic structures. Inactivation of *Ctcf* several days later at E11 also resulted in PUMA upregulation and increased apoptotic cell death, and the *Ctcf*-null forebrain was hypocellular and disorganized at birth. Although deletion of both *Ctcf* and *Puma* in the embryonic brain efficiently rescued *Ctcf*-null progenitor cell apoptosis, it failed to improve neonatal hypocellularity due to decreased proliferative capacity of rescued apical and outer radial glia progenitor cells. This was exacerbated by an independent effect of CTCF loss that resulted in depletion of the progenitor pool due to premature neurogenesis earlier in development. Our findings demonstrate that CTCF activities are required for two distinct events in early cortex formation: first, to correctly regulate the balance between neuroprogenitor cell proliferation and differentiation, and second, for the survival of neuroprogenitor cells, providing new clues regarding the contributions of CTCF in microcephaly/intellectual disability syndrome pathologies.

Key words: apoptosis; CTCF; differentiation; mouse model; neurogenesis; *Puma*

Introduction

CTCF is a multifunctional DNA binding protein that regulates higher-order chromatin structure to influence transcriptional regulation, genomic imprinting, X chromosome inactivation, and chromatin insulation (Holwerda and de Laat, 2013). It binds to a variety of highly divergent target sequences throughout the genome using a combination of its 11 zinc finger motifs (Nakashiki et al., 2013). CTCF partners with a number of chromatin-related proteins such as the cohesin complex (Parelho et al., 2008; Wendt et al., 2008), nucleophosmin, and CTCF itself (Yusufzai and Felsenfeld, 2004; Yusufzai et al., 2004). These interactions may allow CTCF sites to contact one another and/or to be tethered to subnuclear domains. CTCF-mediated chromatin interactions detected by Chromatin interaction analysis by paired-end tag sequencing (ChIA-PET) correlate with ~10% of all CTCF

binding sites (CBSs), however, indicating that CTCF likely plays additional roles within the cell (Handoko et al., 2011).

De novo mutations in *CTCF* have been identified previously in patients with varying degrees of intellectual disability and microcephaly (Gregor et al., 2013), highlighting the importance of chromatin organization for the normal development and function of the CNS. Whereas CTCF function has been studied extensively in cell culture systems, its function in an *in vivo* context remains to be completely resolved (Ohlsson et al., 2010). Ubiquitous deletion of CTCF in the mouse leads to lethality before embryonic day 3.5 (E3.5), suggesting that it is essential for early developmental processes (Fedoriw et al., 2004; Heath et al., 2008; Moore et al., 2012). Conditional deletion of *Ctcf* in specific tissues causes either reduced proliferation or apoptotic cell death, depending on the tissue targeted for *Cre* recombination. For example, deletion of *Ctcf* in thymocytes resulted in increased *p21* and *p27* expression and cell cycle arrest, whereas reduced CTCF in mouse oocytes induced meiotic and mitotic defects and apoptotic cell death before the blastocyst stage (Fedoriw et al., 2004; Heath et al., 2008; Wan et al., 2008). Deletion of *Ctcf* in the developing limb bud resulted in massive apoptosis and near-complete loss of limb structures, accompanied by increased p53 upregulated modulator of apoptosis (PUMA), a known activator of caspase-mediated apoptosis (Nakano and Vousden, 2001; Yu et al., 2001; Soshnikova et al., 2010). In human cancer cells, CTCF binds to the *Puma* gene, and its depletion results in increased

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Table 1. Primer sequences used for genotyping, gene expression, and qChIP analyses

Primer Name	Forward sequence (5'-3')	Reverse sequence (5'-3')
<i>Ctcf</i>	CTAGGAGTGTAGTTCAGTGAGGCC	GCTCTAAAGAAGTTGTGAGTTC
<i>NestinCre</i>	TGACCAGAGTCATCCTTAGCC	AATGCTTCTGTCGGTTTGCC
<i>Sry</i>	GCAGGTGGAAAAGCCTTACA	AAGCTTTGCTGTTTTTGGGA
<i>Puma</i> genotyping WT	AGGCTGTCCCTGCGGTATCC	GGACTGTGCGGGGTAGACCTCTA
<i>Puma</i> genotyping KO	AGGCTGTCCCTGCGGTATCC	ACCGCGGGTCCGAGTAGC
<i>Ctcf</i> expression	CGATATGCTCTATCCAGCA	TCCCACACTTGGAAACAGACA
<i>Puma</i> expression	CGTGTGGAGGAGGAGGAGT	GGAGGAGTCCCATGAAGAGA
β -actin expression	CTGTCCAGTCCGCTCCACCC	ACATGCGGAGCCGTGTGCG
5' 5 kb <i>Puma</i> CBS	ACCTCTGTTGGAGTGAC	CTCTGCTCTGTGCTCTAAAG
<i>Puma</i> CBS1	GCTCTCCAGGCTCTACTA	CAGCTTTCATCACTGGGACT
<i>Puma</i> CBS2	AGGAATGGATCTGCTGGATG	GTTGCTGACTACCCGGCTAT
3' 5 kb <i>Puma</i> CBS	GAGCCCTGCCTAGTAGGAT	TAGTCCCTGTGTGCTTGT
5' 1 kb p53BS	AAAAATGGCTGGAGAGC	CCACCCTGTCACAGCTTGT
<i>Puma</i> p53BS	CTGTCCACAGCTGC	GCTTGTCTGCTGGTGTG
3' 1 kb p53BS	AGCCAGGGCTACACAGAGAA	CTGAGCCATCTCCAGTCC

KO, Knock-out; WT, wild type.

Puma transcript and rapid apoptosis, indicating that *Puma* transcription can be directly influenced by the presence or absence of CTCF (Gomes and Espinosa, 2010a,b).

Given the deleterious effects of *CTCF* mutations in the human CNS, we specifically inactivated *Ctcf* in the developing mouse brain. CTCF loss of function using two different *Cre* driver lines in mice triggered apoptosis in dividing neuroprogenitor cells (NPCs) of the forebrain. Despite prevention of apoptosis by *Puma* deletion, rescued *Ctcf/Puma* double-null apical and outer radial glia (oRG) progenitors exhibited decreased proliferative capacity. Furthermore, loss of CTCF caused premature neurogenesis, resulting in depletion of the progenitor pool and a microcephaly phenotype at birth. These findings highlight the complexity of CTCF activities during neurogenesis.

Materials and Methods

Mouse husbandry and genotyping. Mice were exposed to 12 h light/dark cycles and fed tap water and regular chow *ad libitum*. The *Ctcf^{loxP}* mice, in which *loxP* sites flank exons 3–12, have been described previously (Heath et al., 2008). Mice conditionally deficient in CTCF were generated by crossing *Ctcf^{loxP/+}* females (C57BL/6 background) with heterozygous *Foxg1Cre* knock-in male mice (129/sv background) or with *NestinCre* heterozygous male mice (C57BL/6 background; Hébert and McConnell, 2000; Bérubé et al., 2005). To account for decreased *Foxg1* expression due to knock-in of the *Cre* recombinase gene, *Cre⁺* males were used as controls (*Ctcf^{+/+} Foxg1-cre^{+/-}*) unless stated otherwise. Using the *NestinCre* driver line, *Ctcf^{loxP/loxP}* mice were crossed with *Ctcf^{loxP/+}; Nestin⁺* mice to generate *Ctcf^{loxP/loxP}* or *Ctcf^{loxP/+}* (controls) and *Ctcf^{loxP/loxP}; Nestin⁺* (*Ctcf^{Nes-cre}*). *Puma^{-/-}* (*Bbc3^{tm1Ast}*) mice were obtained from The Jackson Laboratory (stock #011067; Villunger et al., 2003). DNA from tail biopsies of newborn pups or yolk sac from embryos was genotyped by PCR. Primer sequences are provided in Table 1.

Immunostaining and histology. For immunofluorescence staining, cryosections were incubated with the primary antibody overnight at 4°C, washed in PBS/0.3% Triton-X 100, and incubated with the secondary antibody for 1 h. Sections were counterstained with DAPI (Sigma-Aldrich) and mounted in SlowFade Gold (Invitrogen). Primary antibodies used were as follows: rabbit anti-CTCF (1:400; Cell Signaling Technology), rabbit anti-cleaved caspase 3 (AC3; Asp175; 1:400; Cell Signaling Technology), mouse anti-BrdU (1:50; BD Biosciences), rabbit anti-PUMA (1:200; Cell Signaling Technology), rabbit anti-TBR2 (1:200; Abcam), goat anti-SOX2 (1:400; Santa Cruz Biotechnology), goat anti-PAX6 (1:200; Santa Cruz Biotechnology), rabbit anti-SOX2 (1:100; Millipore Bioscience Research Reagents), rabbit anti-Ki67 (1:200; Abcam), rabbit anti-TBR1 (1:200; Abcam), mouse anti-SATB2 (1:200; Abcam), and rabbit anti-CTIP2 (1:200; Abcam). Secondary antibodies used were

as follows: goat-anti-rabbit Alexa 594 (1:800; Invitrogen), goat-anti-mouse Alexa 488 (1:800; Invitrogen), donkey-anti-sheep Alexa 594 (1:800; Invitrogen), and donkey-anti-mouse Alexa 488 (1:800; Invitrogen). Sections were subjected to antigen retrieval (incubated in 0.1 M sodium citrate, pH 6.0, heated to ~95°C and microwaved on low for 10 min) before overnight incubation (for BrdU, SOX2, PAX6, TBR2, STAB2, TBR1, and CTIP2). Terminal deoxynucleotidyl nick end labeling (TUNEL) was performed according to the manufacturer's instructions (Roche). For histological studies, slides were stained with hematoxylin and eosin (H&E).

BrdU labeling. Pregnant mice were injected intraperitoneally with cell proliferation labeling reagent [10 mM bromodeoxyuridine (BrdU) and 1 mM fluorodeoxyuridine (FdU) in H₂O] at 1 ml/100 g body weight, or 0.3 mg/g body weight (GE Healthcare Life Sciences). Animals were killed after 1 h by CO₂ asphyxiation, and the embryos were recovered in ice-cold PBS, pH 7.4, and were fixed in 4% paraformaldehyde. Tissue was equilibrated in 30% sucrose/PBS and frozen in OCT (Tissue Tek). For cell cycle exit analysis, pregnant female mice (E13) were injected with cell proliferation labeling reagent at 1 ml/100 g body weight (GE Healthcare Life Sciences), and embryos were collected and processed for immunofluorescence analysis 24 h later. Before immunofluorescence analysis, cryosections (8 μ m) were treated with 2N HCl to denature the DNA, and neutralized with 0.1 M Na₂B₄O₇, pH 8.5.

Primary NPC cultures and manipulation. Cortical progenitor cultures were prepared as described previously (Slack et al., 1998; Gloster et al., 1999; Watson et al., 2013) using cortices dissected from E12.5 embryos. Cells were seeded on polyornithine-coated (Sigma-Aldrich) plastic plates or glass coverslips. Cells were fixed in 4% paraformaldehyde for 10 min, washed in PBS, and processed for immunofluorescence. Cell viability was measured using the trypan blue dye exclusion method. Cell counts were determined with a hemacytometer.

Western blot analysis. Nuclear protein was extracted from the E16.5 telencephalon using a standard extraction kit (Thermo Scientific) and quantified using the Bradford assay. Protein (20 μ g) was resolved on a 6% SDS-polyacrylamide gel and transferred onto a nitrocellulose membrane (Bio-Rad). The membrane was probed with rabbit anti-p53 (1:500; Santa Cruz Biotechnology) and rabbit anti-inner centromere protein (INCENP) (1:10000; Sigma Aldrich) antibodies followed by the appropriate horseradish peroxidase-conjugated secondary antibodies (1:4000; GE Healthcare Life Sciences). The membrane was incubated in ECL before exposure to x-ray film. Densitometry analysis was performed using ImageJ software (version 1.47).

Real-time PCR. Total RNA was isolated using the RNeasy Mini kit (Qiagen). First-strand cDNA was synthesized from 1 μ g of total RNA using the SuperScriptTM II Reverse Transcriptase kit (Invitrogen) with deoxyribonucleotide triphosphates (dNTPs) (1 mM final concentration; GE Healthcare), porcine RNAGuard (GE Healthcare), and 0.3 μ g random primers (GE Healthcare). Amplification was performed using a Chromo-4 Continuous Fluorescence Detector (Bio-Rad) in the presence of iQ SYBR Green supermix (Bio-Rad) and recorded using Opticon Monitor 3 software (Bio-Rad). Results were normalized to β -actin expression, and relative gene expression levels were calculated using GeneX software (Bio-Rad). Samples were amplified as follows: 95°C for 10 s, annealed for 20 s, 72°C for 30 s. After amplification, a melting curve was generated, and samples were resolved on a 1.5% agarose gel (75 V for 1 h) to verify amplicon purity. Primer oligomers were designed using Primer3 software (<http://bioinfo.ut.ee/primer3-0.4.0/primer3/>) and were obtained from Integrated DNA Technologies. Sequences are provided in Table 1.

Chromatin immunoprecipitation. Chromatin immunoprecipitation (ChIP) experiments were performed as described previously (Kernohan et al., 2010). Briefly, mouse forebrain tissue was dissected and processed to single-cell suspension. Cells were cross-linked in 1% formaldehyde, lysed in SDS buffer and sonicated. Immunoprecipitation was performed with a rabbit anti-CTCF antibody (Cell Signaling Technology) and a rabbit anti-p53 antibody (Santa Cruz Biotechnology). Input samples represent 1/25 of total chromatin input. ChIP products were amplified in duplicate with iQ SYBR Green Master Mix (Bio-Rad) on a Chromo-4 thermocycler under the following conditions: 95°C for 5 min, followed by 35 cycles of 95°C for 10 s, 55°C for 20 s, 72°C for 30 s, and a final

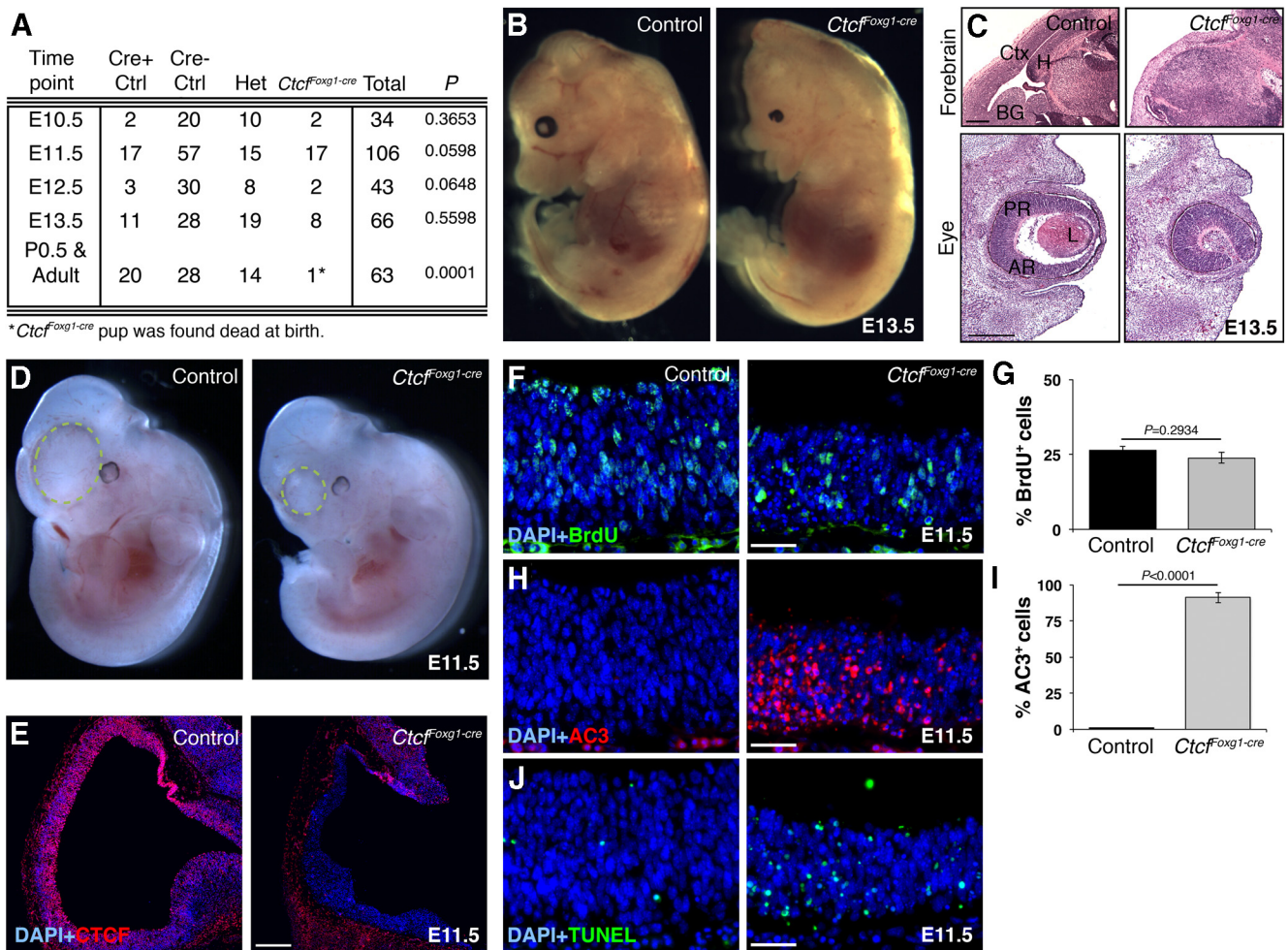


Figure 1. *Foxg1Cre*-mediated deletion of *Ctcf* results in a massive increase in apoptosis. **A**, Table of genotypes obtained during the embryonic and postnatal periods. Ratios at each time point were analyzed using a χ^2 test. Het, *Ctcf^{lox/WT};Foxg1-cre^{+/-}*. **B**, Dark-field images of control and *Ctcf^{Foxg1-cre}* embryos at E13.5. (Please note limbs were taken for genotyping). **C**, H&E staining of E13.5 sagittal cryosections demonstrates complete loss of cortex (Ctx), hippocampal hem (H), basal ganglia (BG), lens (L), and anterior retina (AR), but not the posterior retina (PR), in *Ctcf^{Foxg1-cre}* embryos. **D**, Dark-field images of control and *Ctcf^{Foxg1-cre}* embryos at E11.5. The dashed circle outlines the telencephalon, which is visibly reduced in size in the *Ctcf^{Foxg1-cre}* embryos compared to littermate controls. **E**, Immunodetection of CTCF (red) in E11.5 sagittal cryosections confirms specific loss of CTCF expression in the forebrain neuroepithelium of *Ctcf^{Foxg1-cre}* embryos. **F**, Pregnant females were subjected to a 1 h BrdU pulse before being killed. Immunodetection of BrdU in E11.5 control and *Ctcf^{Foxg1-cre}* cortical neuroepithelium is shown. **G**, BrdU⁺ cells were counted and expressed as a percentage of the total number of DAPI⁺ cells ($n = 3$). **H**, Immunodetection of activated caspase-3 (red) in control and *Ctcf^{Foxg1-cre}* cortical neuroepithelium at E11.5. **I**, AC3⁺ cells were counted and expressed as a percentage of the total number of DAPI⁺ cells ($n = 3$). **J**, TUNEL (green) detection in E11.5 control and *Ctcf^{Foxg1-cre}* cortical neuroepithelium. Error bars represent the SEM. Original magnification: **C**, 25 \times ; **E**, 50 \times ; **F**, **H**, **J**, 200 \times . Scale bars: **C**, top, 1 mm; bottom, 400 μ m; **E**, 200 μ m; **F**, **H**, **J**, 100 μ m.

melting curve generated from 55°C to 95°C in increments of 1°C per plate read. Fold change and percent input formulas were adapted from (Mukhopadhyay et al., 2008) as follows: percent input = $100 \times \frac{2^{-(\Delta C_t \text{Input} - \Delta C_t \text{Input})} - (\Delta C_t \text{Input} - \Delta C_t \text{Ab})}{25}$. Error bars represent the SEM. Primer sequences are provided in Table 1.

Microscopy. Images were captured with a digital camera (ORCA-ER; Hamamatsu) using an inverted microscope (DMI 6000b; Leica). Openlab imaging software (PerkinElmer) was used for manual image capture, and processing was performed using Velocity software (PerkinElmer). For quantification of AC3⁺ cells per area, AC3⁺ cells were counted in a defined area in at least six serial cortical cryosections, and the ratio of AC3⁺ cells to area (square millimeters) was calculated. For BrdU, SOX2, TBR2, BrdU/Ki67, TBR1, SATB2, and CTIP2 quantification, at least two serial cortical cryosections were assessed for positive cells within the indicated regions per embryo. DAPI morphology was used to bin cortex into the ventricular/subventricular zone (VZ/SVZ), intermediate zone (IZ), and cortical plate (CP).

Statistical analyses. Statistical analysis was performed using GraphPad Prism software (GraphPad Software; version 4.02), and all results are expressed as the mean \pm SEM. Unless indicated otherwise, *p* values were generated using Student's *t* test (unpaired, two-tailed) to compare be-

tween two independent data sets. Genotype ratios (Figs. 1A, 2A) were compared using a χ^2 test. To compare cell viability data (see Fig. 4E), a one-way ANOVA was used with Dunnett's multiple comparison post-test to compare the cell viability of each genotype (*Ctcf^{Nes-cre}*, *Ctcf^{Foxg1-cre}*, *Puma^{-/-}*, and *Puma^{-/-}*) to control.

Study approval. All procedures involving animals were conducted in accordance with the regulations of the Animals for Research Act of the Province of Ontario and approved by the University of Western Ontario Animal Care and Use Committee.

Results

Ctcf^{Foxg1-cre} mice exhibit widespread apoptosis and profound loss of telencephalic and anterior retinal tissue

To study the function of CTCF in the embryonic brain, *Ctcf* floxed mice were intercrossed with the previously characterized *Foxg1Cre* mice (Hébert and McConnell, 2000; Bérubé et al., 2005; Heath et al., 2008). *Foxg1* expression is first detected between E8 and E9 in the telencephalic neuroepithelium, the basal ganglia, the olfactory bulbs, and the anterior retina (Dou et al., 1999). *Foxg1-cre⁺ Ctcf^{loxP/loxP}* progeny resulting from this cross are *Ctcf*

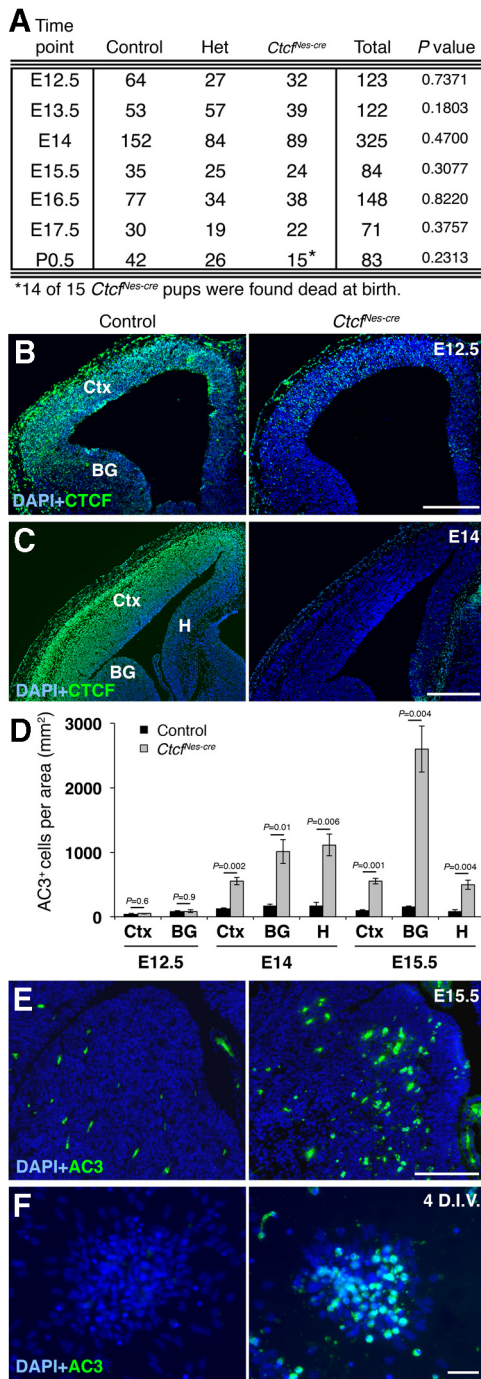


Figure 2. *NestinCre*-mediated deletion of CTCF results in activation of caspase-mediated apoptosis. **A**, Table of genotype ratios obtained during the embryonic period. Ratios at each time point were analyzed by a χ^2 test. **B**, Immunodetection of CTCF in E12.5 control and *Ctf^{Nes-cre}* coronal forebrain sections. **C**, Immunodetection of CTCF in E14 control and *Ctf^{Nes-cre}* coronal forebrain sections. **D**, Quantification of AC3 immunostaining in E12.5, E14, and E15.5 forebrain tissue ($n = 3$). AC3⁺ cells were counted and expressed per unit area (square millimeter). **E**, Immunodetection of AC3 in E15.5 control and *Ctf^{Nes-cre}* basal ganglia. **F**, Immunodetection of AC3 in control and *Ctf^{Nes-cre}* at 4 DIV. Het, *Ctf^{flox/WT};Nestin-cre*; Ctx, cortex; BG, basal ganglia; H, hippocampal hem. Error bars represent the SEM. Original magnification: **B**, **C**, 50 \times ; **D**, 100 \times ; **E**, 200 \times . Scale bars: **B**, 220 μ m; **C**, 300 μ m; **E**, 100 μ m; **F**, 25 μ m.

null in the anterior retina and forebrain, and will be referred to as *Ctf^{Foxg1-cre}* throughout this text.

We failed to recover live *Ctf^{Foxg1-cre}* pups at birth (Fig. 1A). At E13.5, the size of telencephalic and retinal structures was greatly

diminished in mutant embryos (Fig. 1B, C). At E11.5, the telencephalon of mutant embryos was already noticeably smaller compared to littermate-matched controls (Fig. 1D, hatched circle). CTCF immunostaining demonstrated nuclear expression of the protein throughout the forebrain in control tissue and its absence in mutant embryos (Fig. 1E). To determine the underlying cause of cell loss, we measured levels of proliferation and cell death in E11.5 *Ctf^{Foxg1-cre}* and control embryos. Acute (1 h) BrdU labeling of control and *Ctf^{Foxg1-cre}* embryos revealed that the percentage of cells in S phase does not vary significantly between mutant and control forebrain tissue (Fig. 1F, G). To measure apoptosis, we stained sections with an antibody against activated caspase-3 as a marker of cell death (Fig. 1H). Quantification of the results revealed a large increase in the proportion of cells undergoing apoptotic cell death in *Ctf^{Foxg1-cre}* embryos compared to littermate-matched controls (Fig. 1I). TUNEL also showed increased apoptosis in *Ctf^{Foxg1-cre}* embryos compared to control (Fig. 1J). Since we already observed profound cell loss by E11.5, we wanted to analyze whether a similar mechanism was occurring earlier in this system. We observed increased AC3 immunostaining in *Ctf^{Foxg1-cre}* forebrain cryosections at E10.5, confirming that cell death occurs at this earlier time point (data not shown). We conclude that deletion of *Ctf* in the mouse forebrain at approximately E8.5 causes extensive apoptosis resulting in profound loss of telencephalic and anterior retinal tissue by E13.5.

***NestinCre*-driven inactivation of *Ctf* decreases cell survival**

To determine the outcome of *Ctf* deletion at a later embryonic time point in the telencephalon, we mated *Ctf^{flox/flox}* mice to the previously characterized *NestinCre* transgenic mice (Bérubé et al., 2005; Heath et al. 2008). The resulting mutant is hereafter referred to as *Ctf^{Nes-cre}* for simplicity. *Cre* is expressed after pre-plate formation in this system, resulting in specific deletion of *Ctf* in neural progenitor cells at approximately E11 (Bérubé et al., 2005). We obtained the expected ratio of *Ctf^{Nes-cre}* mice at birth; however, almost all mutant pups were already dead and blue in color (Fig. 2A). One live *Ctf^{Nes-cre}* pup was found struggling to breathe, appeared weak, and was killed. We conclude that neonatal lethality of *Ctf^{Nes-cre}* mice is likely due to asphyxiation at birth.

We first established that the CTCF protein was lost in the basal ganglia and many cells of the cortex at E12.5 (Fig. 2B), and expression was undetectable in the E14 *Ctf^{Nes-cre}* cortex, basal ganglia, and hippocampal hem (Fig. 2C). To determine whether *NestinCre*-mediated deletion of *Ctf* also induces apoptosis, we analyzed *Ctf^{Nes-cre}* forebrain cryosections at E12.5–E15.5 for evidence of AC3. At E12.5, we observed no increase in apoptotic cells in the mutant telencephalon (Fig. 2D). However, at E14, we detected an increase in AC3⁺ cells in the *Ctf^{Nes-cre}* cortex, basal ganglia, and hippocampal hem compared to controls (Fig. 2D). The increase in caspase activation was also observed at E15.5 in all three forebrain regions, with the highest level of AC3⁺ cells in the basal ganglia (Fig. 2D, E). Increased apoptotic cell death was also observed *in vitro* in primary cortical progenitor cultures established from E12.5 control and *Ctf*-null telencephalon. Levels of AC3 staining were low in both control and *Ctf*-null NPCs after 2 d *in vitro* (DIV), and we only observed a detectable increase in staining at 4 DIV (Fig. 2F). Together, the data suggest that when using the *NestinCre* driver line of mice, *Ctf* deficiency does trigger the activation of caspase-mediated apoptosis in NPCs, but the effect is delayed and less severe than that observed in the *Ctf^{Foxg1-cre}* embryos, suggesting that early neuroepithelial

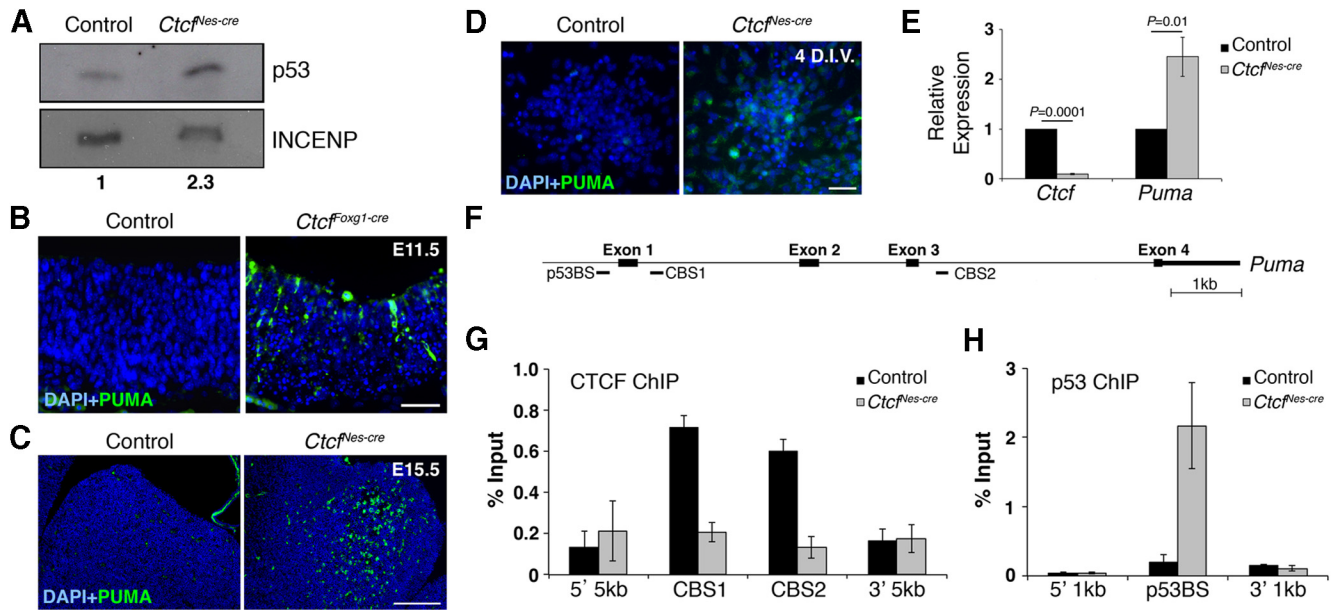


Figure 3. Loss of CTCF causes p53-dependent transcriptional activation of *Puma*. **A**, Western blot analysis of nuclear protein extracts obtained from E16.5 control and *Ctcf^{Nes-cre}* telencephalon. Densitometry analysis of blots revealed a 2.3-fold increase in *Ctcf^{Nes-cre}* p53 levels compared to control. **B**, **C**, INCENP was used as a loading control. Immunodetection of PUMA in E11.5 control and *Ctcf^{Foxg1-cre}* cortical neuroepithelium (**B**), E15.5 control and *Ctcf^{Nes-cre}* forebrain (**C**), and control and *Ctcf*-null NPCs (**D**) at 4 DIV. **E**, Quantitative real-time PCR analysis of *Ctcf* and *Puma* expression in control and *Ctcf^{Nes-cre}* E16.5 forebrain using primers spanning *Puma* exons 3–4 ($n = 3$). β -actin was used as an internal control. **F**, Schematic representation of the mouse *Puma* gene. PCR products used to detect sites of CTCF binding (CBS1 and CBS2) and p53 binding (p53BS) are shown. **G**, Quantitative chromatin immunoprecipitation of CTCF at the *Puma* gene in E16.5 control and *Ctcf^{Nes-cre}* telencephalon tissue ($n = 3$). Primer pairs 5 kb upstream and downstream of binding sites were used as negative controls. **H**, Quantitative chromatin immunoprecipitation of p53 at the *Puma* gene in E16.5 control and *Ctcf^{Nes-cre}* telencephalon shows increased p53 occupancy at the *Puma* promoter in the *Ctcf^{Nes-cre}* telencephalon ($n = 3$). Primer pairs 1 kb upstream and downstream of the binding site were used as negative controls. Error bars represent SEM. Original magnification: **A**, **C**, 200 \times ; **B**, 100 \times . Scale bars: **B**, **C**, 50 μ m; **D**, 25 μ m.

cells are more sensitive to CTCF loss than the neuroprogenitors present slightly later in development.

Increased p53 and PUMA levels in the *Ctcf*-null telencephalon

Both the *Foxg1Cre* and *NestinCre* models of CTCF loss exhibit increased levels of apoptosis, although the timing of onset differs between the two mutants. To investigate the molecular mechanism responsible for neuronal cell death due to CTCF loss, we assessed activation of the p53/PUMA pathway. Using Western blot analysis, we found that neuronal cell death correlated with an increase in p53 protein levels in the *Ctcf^{Nes-cre}* forebrain compared to control, suggesting activation and stabilization of the protein (Fig. 3A). We next investigated the levels of PUMA, which has been demonstrated previously to mediate NPC death downstream of p53 (Jeffers et al., 2003). Moreover, *Puma* is one of the most upregulated genes in response to CTCF loss in the limb bud (Soshnikova et al., 2010). CTCF binding sites in the *Puma* gene demarcate an intragenic chromatin boundary that is abolished upon CTCF knockdown in human cells, resulting in increased PUMA expression (Gomes and Espinosa, 2010b). To determine whether PUMA is involved caspase-mediated cell death in the *Ctcf* mutant embryos, we examined PUMA protein levels in the *Ctcf^{Foxg1-cre}* and *Ctcf^{Nes-cre}* mice. In the E11.5 *Ctcf^{Foxg1-cre}* forebrain, we observed an increase in PUMA immunostaining compared to littermate controls, corresponding to the high levels of AC3 staining in these embryos (Fig. 3B). In the *Ctcf^{Nes-cre}* embryos, PUMA was increased in the E15.5 basal ganglia, again correlating with the highest levels of cell death (Fig. 3C). In primary cultures, *Ctcf*-null NPCs exhibited increased PUMA immunostaining at 4 DIV when compared to control NPCs obtained from littermate embryos (Fig. 3D). We also quantified *Puma* transcript levels by quantitative reverse-transcriptase PCR and observed a significant increase in the E16.5 *Ctcf^{Nes-cre}* forebrain

compared to controls (Fig. 3E). These observations demonstrate that increased PUMA transcript and protein levels occur as a consequence of CTCF loss in NPCs and correlate with p53 activation and the onset of caspase-mediated cell death.

PUMA upregulation in the *Ctcf*-deficient embryonic brain could result from decreased CTCF occupancy within the *Puma* gene body, or because of p53-dependent activation of the gene. To investigate these possibilities further, we performed quantitative CTCF and p53 ChIP using E16.5 telencephalon isolated from control and *Ctcf^{Nes-cre}* embryos (Fig. 3F–H). ChIP analysis confirmed CTCF binding sites found downstream of *Puma* exon 1 (CBS1) and exon 3 (CBS2) at the *Puma* gene in control tissue, similar to the binding profile of CTCF in human cancer cells (Fig. 3G; Gomes and Espinosa, 2010a). As expected, CTCF binding was diminished in the *Ctcf^{Nes-cre}* tissue (Fig. 3G). ChIP for p53 demonstrated specific enrichment of the protein at the *Puma* promoter in *Ctcf^{Nes-cre}* forebrain compared to control in E16.5 tissue (Fig. 3H). Together, loss of CTCF in NPCs causes increased *Puma* transcription and protein levels due to increased p53-dependent transcriptional activation, likely combined with the loss of CTCF-dependent repression.

Deletion of *Puma* in a *Ctcf*-null context rescues cell death, but does not improve viability or brain defects at birth

To investigate whether increased PUMA levels cause cell death in the *Ctcf*-deficient embryonic brain, we introduced a mutant *Puma* allele (*Bbc3^{tm1Ast}*) in the *Ctcf^{loxP}* and *NestinCre* mice to generate mice that lack both *Ctcf* and *Puma* expression in the brain (hereafter referred to as *Ctcf^{Nes-cre};Puma^{-/-}*). Histological analysis of E16.5 *Ctcf^{Nes-cre}* cryosections showed thinning of the VZ/SVZ, hypocellularity of the intermediate zone, and a dramatic reduction in the size of the hippocampal hem (Fig. 4A). *Ctcf^{Nes-cre};Puma^{-/-}* embryos at E16.5 showed rescue in the VZ/

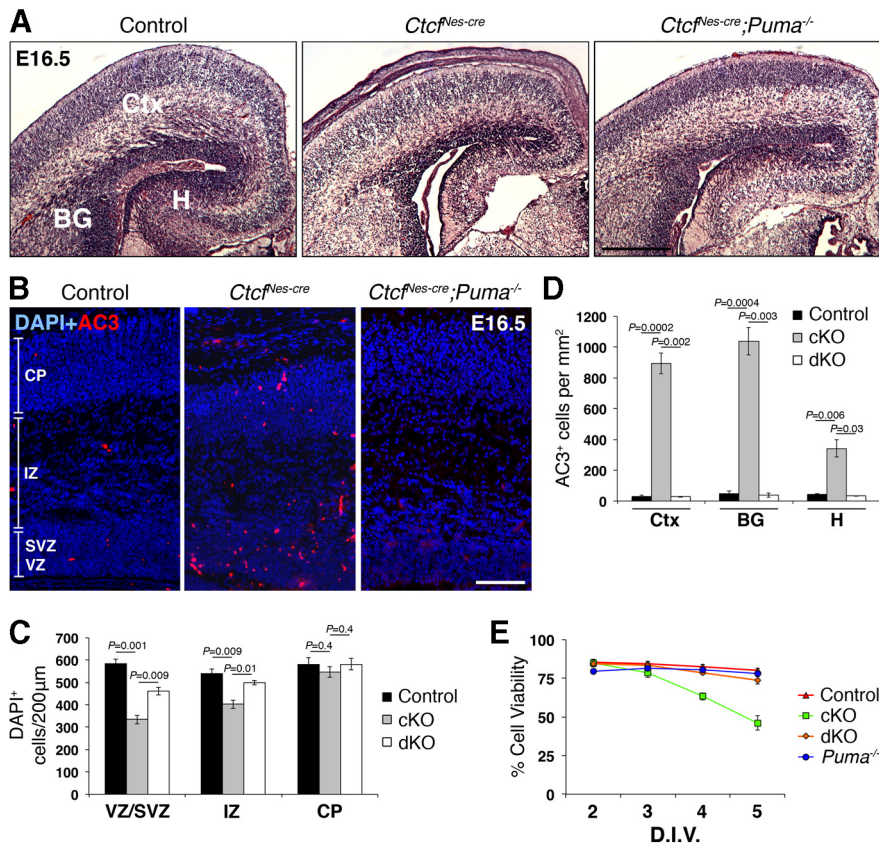


Figure 4. Apoptosis is abolished upon deletion of PUMA in the *Ctf*-deficient embryonic brain. **A**, H&E staining of E16.5 control, *Ctf^{Nes-cre}*, and *Ctf^{Nes-cre};Puma^{-/-}* coronal cortical cryosections. Skin and skull cap were not dissected from *Ctf^{Nes-cre}* brain. **B**, AC3 immunostaining in E16.5 control, *Ctf^{Nes-cre}*, and *Ctf^{Nes-cre};Puma^{-/-}* neocortex. **C**, DAPI⁺ cells were quantified in 200- μ m-wide regions of the control, *Ctf^{Nes-cre}* (conditional knock-out), and *Ctf^{Nes-cre};Puma^{-/-}* (double knock-out) CP, IZ, and VZ/SVZ ($n = 3$). **D**, AC3⁺ cells were quantified per unit area in the cortex (Ctx), basal ganglia (BG), and hippocampal hem (H; $n = 3$). **E**, Cell viability of NPC cultures was measured by trypan blue dye exclusion after 2–5 DIV ($n = 3$). Data were analyzed by a one-way ANOVA followed by Dunnett's multiple comparisons test to determine which means were significantly different from the control. No significant difference in cell viability was found at 2 and 3 DIV ($p = 0.2122$ and 0.2062 , respectively); however, viability was significantly different between *Ctf^{Nes-cre}* and control at 3 and 4 DIV ($p < 0.01$), but not between *Ctf^{Nes-cre};Puma^{-/-}* and control or *Puma^{-/-}* and control ($p > 0.05$). Error bars represent SEM. Original magnification: **A**, 100 \times ; **C**, 50 \times . Scale bars: **A**, 200 μ m; **B**, 100 μ m.

SVZ thickness, as well as hippocampal size (Fig. 4A). We confirmed these observations by partitioning the E16.5 cortex into the CP, IZ, and VZ/SVZ based on cytoarchitecture and quantifying the number of DAPI⁺ cells per region (Fig. 4B,C). We observed a restoration in the number of IZ cells and partial restoration in the number of VZ/SVZ cells in the *Ctf^{Nes-cre};Puma^{-/-}* compared to *Ctf^{Nes-cre}* cortex (Fig. 4C). There was no significant difference in the number of CP cells between genotypes at E16.5 (Fig. 4C).

We performed AC3 immunostaining in E16.5 brain sections to investigate whether apoptotic cell death was rescued in the double mutant brain. AC3⁺ cells were elevated in the *Ctf^{Nes-cre}* cortex, basal ganglia, and hippocampal hem compared to control (Fig. 4B,D). AC3⁺ cells were largely localized to the ventricular zone of the neocortex, indicating that *Ctf*-null proliferating cells are more susceptible to caspase-dependent cell death than differentiated cells of the cortical plate. Deletion of *Puma* in the *Ctf^{Nes-cre}* mouse was sufficient to abolish apoptotic cell death, indicating that PUMA mediates increased caspase-dependent cell death in *Ctf*-null neuroprogenitor cells (Fig. 4B,D). Together, these data indicate a specific decrease in intermediate zone cellularity in the E16.5 *Ctf^{Nes-cre}* cortex that is fully restored upon inhibition of

cell death. The number of cells in the ventricular/subventricular zone is only partially restored, suggesting that loss of cells cannot completely be explained by increased cell death.

To further confirm that cell viability is rescued by deletion of *Puma* in a *Ctf*-null context, we established primary cortical progenitor cultures from E12.5 control, *Ctf^{Nes-cre}*, *Ctf^{Nes-cre};Puma^{-/-}*, and *Puma^{-/-}* telencephalon and measured viability via trypan blue dye exclusion at 2–5 DIV. Each time point was analyzed by a one-way ANOVA followed by Dunnett's multiple comparisons post-test to determine which genotypes had a mean percentage of viability that was significantly different from control. No difference in viability was observed at 2 and 3 DIV; however, at 4 and 5 DIV, *Ctf^{Nes-cre}* NPCs exhibited a marked reduction in viability, correlating with increased levels of AC3 (Fig. 4E). NPCs obtained from *Ctf^{Nes-cre};Puma^{-/-}* embryos did not show a significant decrease in viability at any of the time points analyzed (Fig. 4E).

Since we observed a rescue in caspase-mediated cell death in the embryonic *Ctf^{Nes-cre};Puma^{-/-}* brain, we predicted that the brain size at birth would also be recovered and that the mice would survive into the postnatal period, allowing for more extended analyses. Surprisingly, we failed to recover live *Ctf^{Nes-cre};Puma^{-/-}* mice at birth. Histological staining of postnatal day 0.5 (P0.5) *Ctf^{Nes-cre};Puma^{-/-}* sections demonstrated severe hypocellularity and disorganization of the cortical plate and hippocampus, and an overall similarity to the *Ctf^{Nes-cre}* neonatal brain (Fig. 5A). Histological cell counts confirmed that there are fewer cells in *Ctf^{Nes-cre}* and *Ctf^{Nes-cre};Puma^{-/-}* cortical plate compared to controls; however, no significant difference in cell number was detected between *Ctf^{Nes-cre}* and *Ctf^{Nes-cre};Puma^{-/-}* (Fig. 5B). These findings indicate that although deletion of *Puma* prevents *Ctf*-null cells from undergoing apoptosis, other factors come into play to prevent cortical size expansion in the double mutant brain.

Ctf-null apical and oRG progenitors rescued from death by *Puma* deletion fail to proliferate

Given that cell death at E16.5 was rescued in the double mutant embryos but the brain hypocellularity at birth was not, we speculated that the cells rescued from apoptosis might become arrested in the cell cycle, resulting in decreased proliferation. To test this hypothesis, we performed acute BrdU labeling at E16.5 and quantified the proportion of cells in S phase by BrdU immunostaining. To compare the cortical distribution and number of NPCs in S phase, the E16.5 cortex was partitioned into the CP, IZ, and VZ/SVZ based on cytoarchitecture (Fig. 6A). The number of BrdU⁺ cells was significantly reduced in the *Ctf^{Nes-cre}* and *Ctf^{Nes-cre};Puma^{-/-}* intermediate zones and ventricular zone/subventricular zones compared to control (Fig. 6A,B). Despite

substantial rescue in cellularity of the $Ctcf^{Nes-cre};Puma^{-/-}$ cortex compared to $Ctcf^{Nes-cre}$ at E16.5, the proliferative capacity of progenitors was not significantly different (Fig. 6A,B). Our results demonstrate that *Puma* deletion rescues cell death in the *Ctcf*-null brain, but that the rescued cells display reduced proliferative capacity.

Three NPC subtypes exist in the embryonic cortex: apical radial glia (apical progenitors), basal progenitors, and outer radial glia. Apical progenitors can be identified by their expression of SOX2 and PAX6 transcription factors (Götz et al., 1998; Tarabykin et al., 2001; Bani-Yaghoob et al., 2006; Shitamukai and Matsuzaki, 2012). They undergo interkinetic nuclear migration and divide at the ventricular surface to either self-renew or differentiate into basal progenitors, outer radial glia, or cortical neurons (Shitamukai et al., 2011; Shitamukai and Matsuzaki, 2012). Basal progenitors reside in the subventricular zone and uniquely express the T-box transcription factor TBR2 (Englund et al., 2005). They are reported to have the potential to self-renew; however, the majority of their divisions are neurogenic (Haubensak et al., 2004; Miyata et al., 2004; Noctor et al., 2004). Outer radial glia are similar to radial glia in that they have a basal process and express SOX2 and PAX6; however, oRG cells are located outside of the VZ in the outer SVZ. They are able to divide asymmetrically, producing one oRG and one neuron-committed cell with each division (Reillo et al., 2011; Shitamukai et al., 2011; Wang et al., 2011a,b; Shitamukai and Matsuzaki, 2012). We performed acute BrdU-labeling for 1 h and coimmunostained E16.5 cortical sections with antibodies against BrdU and TBR2 or SOX2 to examine the behavior of different progenitor subtypes after loss of CTCF (Figs. 7, 8).

The total number of TBR2⁺ basal progenitor cells was reduced in the $Ctcf^{Nes-cre}$ cortex compared to control (Fig. 7A,B). The number of basal progenitors was restored to control levels in the $Ctcf^{Nes-cre};Puma^{-/-}$ cortex, and most rescued cells were still able to enter S phase (Fig. 7A,C). Despite the correct overall number of TBR2⁺ cells in the $Ctcf^{Nes-cre};Puma^{-/-}$ cortex, the localization of basal progenitors appeared to be shifted apically compared to control (Fig. 7A,B). This is perhaps due to an inability of the rescued basal progenitors to correctly delaminate from the ventricular surface.

Quantification of SOX2⁺ cells indicated a dramatic reduction in the number of apical progenitors and oRGs in the $Ctcf^{Nes-cre}$ cortex (Fig. 8A,B). The number of SOX2⁺ apical progenitors was only partially rescued in the $Ctcf^{Nes-cre};Puma^{-/-}$ cortex, indicating that CTCF may control the size of the apical progenitor pool independently of apoptosis (Fig. 8A,B). Conversely, oRG (SOX2⁺ cells in the intermediate zone) numbers were restored to control levels in the $Ctcf^{Nes-cre};Puma^{-/-}$ cortex, indicating that these cells are lost via *Puma*-mediated apoptotic cell death upon deletion of *Ctcf* (Fig. 8A,B). The proliferative capacity of apical progenitors (SOX2⁺/BrdU⁺ cells in the VZ/SVZ) was severely

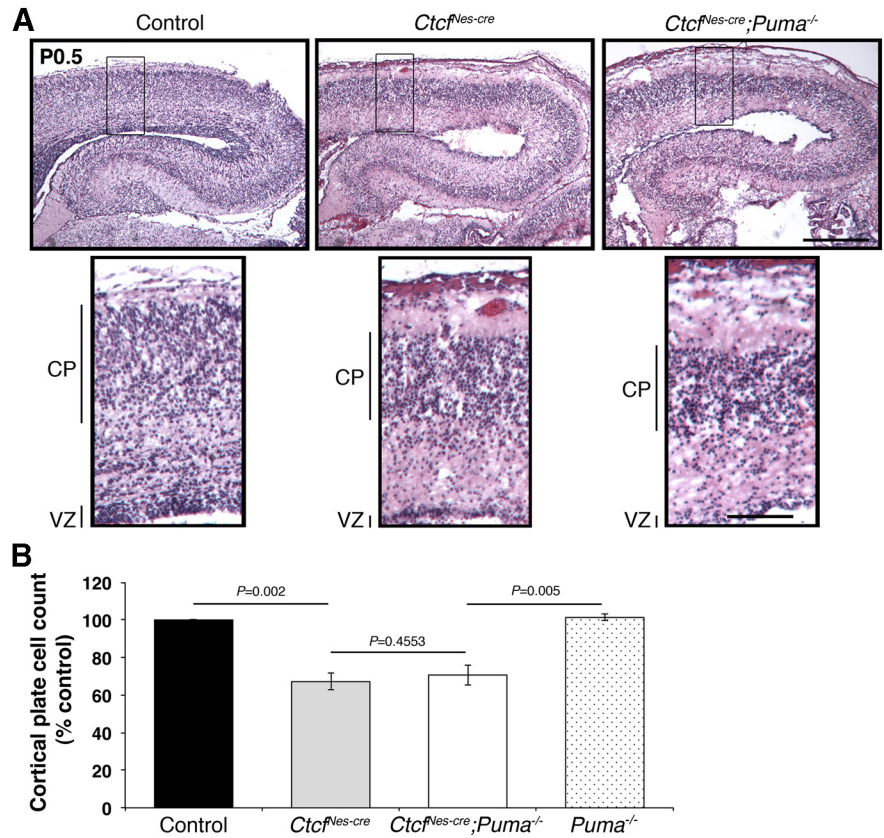


Figure 5. *Puma* deletion in the *Ctcf*-deficient brain fails to restore hippocampal size or hypocellularity of the cortical plate. **A**, H&E staining of neonatal (P0.5) control, $Ctcf^{Nes-cre}$, and $Ctcf^{Nes-cre};Puma^{-/-}$ cortex and hippocampus. Scale bars: top, 800 μ m; bottom, 200 μ m. Original magnification, 25 \times . **B**, Graph depicting cortical plate cell counts as a percentage of control ($n = 3$). Error bars represent SEM.

diminished in the $Ctcf^{Nes-cre}$ cortex compared to control, and was not restored in the $Ctcf^{Nes-cre};Puma^{-/-}$ cortex (Fig. 8A,C). Similarly, BrdU incorporation in the rescued $Ctcf^{Nes-cre};Puma^{-/-}$ oRG cells was extremely low, indicating that they are likely arrested in the cell cycle and fail to correctly enter S phase (Fig. 8A,C). To confirm that these cells are oRG progenitors, we performed SOX2 and PAX6 coimmunostaining and indeed observed a decreased number of SOX2⁺PAX6⁺ cells in the intermediate zone of $Ctcf^{Nes-cre}$ embryos compared to control and $Ctcf^{Nes-cre};Puma^{-/-}$ at E16.5 (Fig. 8D).

Ctcf loss causes premature differentiation of apical progenitors

Incomplete restoration of the apical progenitor population in the $Ctcf^{Nes-cre};Puma^{-/-}$ cortex suggests that loss of CTCF leads to an apoptosis-independent reduction of these cells (Fig. 8A,B). To identify the cause of reduced apical cell numbers, we analyzed the progenitor pool composition, cell cycle exit indices, and postmitotic projection neuron subtypes in control and $Ctcf^{Nes-cre}$ cortex at E14.

Apical progenitors begin to produce projection neurons at approximately E11.5. At the same time, they generate neuron-committed basal progenitors and oRG progenitors. Since the majority of basal progenitor divisions are terminal and result in the production of two neurons, this would result in increased generation of neurons at the expense of the progenitor pool (Haubensak et al., 2004; Miyata et al., 2004; Noctor et al., 2004). The relative abundance of TBR2⁺ basal progenitors was significantly increased in the E14 $Ctcf^{Nes-cre}$ cortex compared to control, whereas the number of SOX2⁺ apical progenitors was not af-

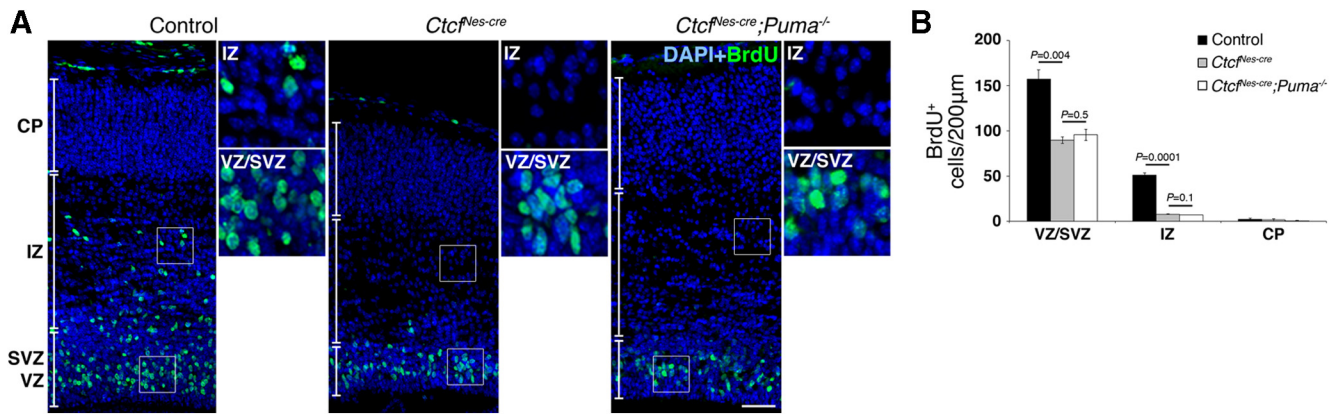


Figure 6. *Ctcf*-deficient cells that are rescued from apoptotic death display reduced proliferative capacity. Pregnant females were subjected to a 1 h BrdU pulse before being killed. **A**, BrdU (green) immunostaining of E16.5 control, *Ctcf^{Nes-cre}*, and *Ctcf^{Nes-cre};Puma^{-/-}* cortical cryosections. The inset demonstrates fewer BrdU⁺ cells in the *Ctcf^{Nes-cre}* and *Ctcf^{Nes-cre};Puma^{-/-}* IZ compared with control. Scale bar, 50 µm. Original magnification, 100×. **B**, The number of cells in S phase was quantified by counting BrdU⁺ cells in 200-µm-wide cortical images (*n* = 3). Error bars represent SEM.

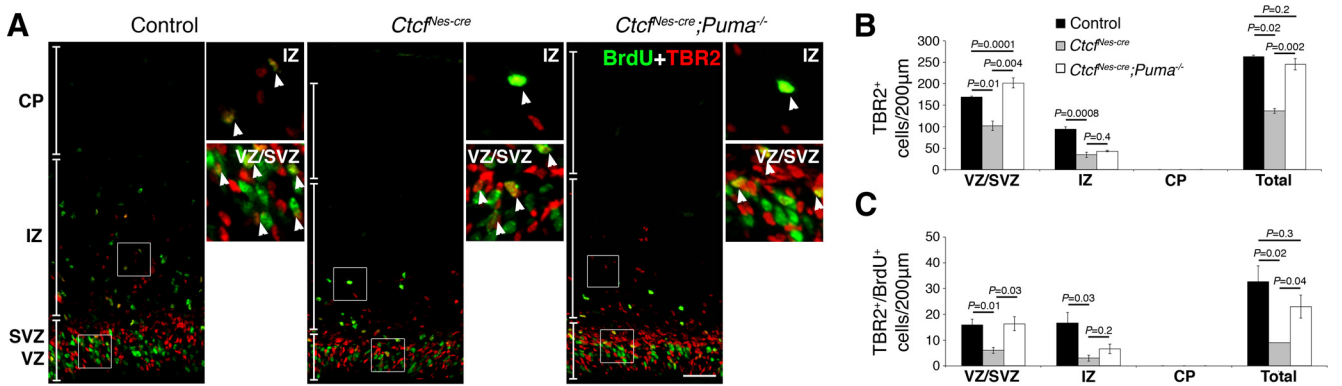


Figure 7. *Ctcf*-deficiency results in PUMA-dependent apoptosis of basal progenitor cells. Pregnant females were subjected to a 1 h BrdU pulse before being killed. **A**, TBR2 (red) and BrdU (green) coimmunostaining of E16.5 cortical cryosections. Arrowheads indicate TBR2⁺/BrdU⁺ cells in the IZ and VZ/SVZ. Scale bar, 50 µm. Original magnification, 100×. **B**, TBR2⁺ cells were quantified in 200-µm-wide cortical images (*n* = 3). **C**, TBR2⁺/BrdU⁺ cells were quantified in 200-µm-wide cortical images (*n* = 3). Error bars represent SEM.

ected (Fig. 9A,B), suggesting that differentiation of apical progenitors into basal progenitors is increased. Next, we measured the proportion of cells exiting the cell cycle by labeling embryos with BrdU for 24 h and analyzing Ki67⁺ and BrdU⁺ cells in E14 control and *Ctcf^{Nes-cre}* cortex. Cell cycle exit (BrdU⁺Ki67⁻ cells/total BrdU⁺ cells) was significantly higher in *Ctcf^{Nes-cre}* cortex compared to controls (Fig. 9C,D).

The cortical plate is established by projection neurons that organize themselves in an “inside-out” manner, such that early-born neurons populate deeper cortical layers (layer VI, then layer V), and late-born neurons migrate past the early-born neurons to populate the more superficial layers of the cortex (layer IV, then layer II/III; Greig et al., 2013). Increased cell cycle exit and differentiation at E13–E14 is predicted to result in an increased number of layer VI corticothalamic projection neurons (TBR1⁺), layer V subcerebral projection neurons (CTIP2⁺), and callosal projection neurons (SATB2⁺). We found that the relative generation of these neuronal subtypes was increased in the *Ctcf^{Nes-cre}* cortex compared to control (Fig. 8E–G), confirming premature differentiation of *Ctcf*-null apical progenitors, which is predicted to result in a reduced number of this progenitor pool by late neurogenesis.

Discussion

This study provides evidence that CTCF is required at very early stages of telencephalon development for the maintenance and

survival of neuroprogenitor cells. We found that ablation of CTCF leads to *Puma*-dependent apoptosis of NPCs using two different conditional deletion strategies. Increased apoptosis correlated with p53-dependent *Puma* transcription, suggesting that CTCF loss results in p53 stabilization and transcriptional activation of its downstream targets. CTCF loss might also result in a more open chromatin environment at the *Puma* gene, facilitating p53-dependent activation of transcription and elongation of RNA polymerase II (Gomes and Espinosa, 2010a,b). *Ctcf* inactivation in postmitotic cortical and hippocampal neurons does not induce apoptosis (Hirayama et al., 2012), pointing to a specific survival role for CTCF in proliferating cells. However, CTCF loss of function may not induce apoptosis in all types of proliferating cells *in vivo*, as *Ctcf* deletion in thymocytes was shown previously to induce cell cycle arrest without induction of apoptosis (Heath et al., 2008). It is possible that the outcome of CTCF deficiency leads to either p53-dependent cell cycle arrest or apoptosis, depending on varying cell-specific or temporal cues. This could be similar to the outcomes described upon *Nbs1* (*Nijmegen breakage syndrome 1/Nibrin*) deletion in the CNS, which leads to p53-dependent apoptosis in the cerebellum, but causes p53-dependent cell cycle arrest in the neocortex (Li et al., 2012).

In the present report, we demonstrate that deletion of *Puma* effectively rescues apoptotic cell death observed at E16.5 in the *Ctcf^{Nes-cre}* brain (Fig. 4). Despite this apparent recovery at E16.5,

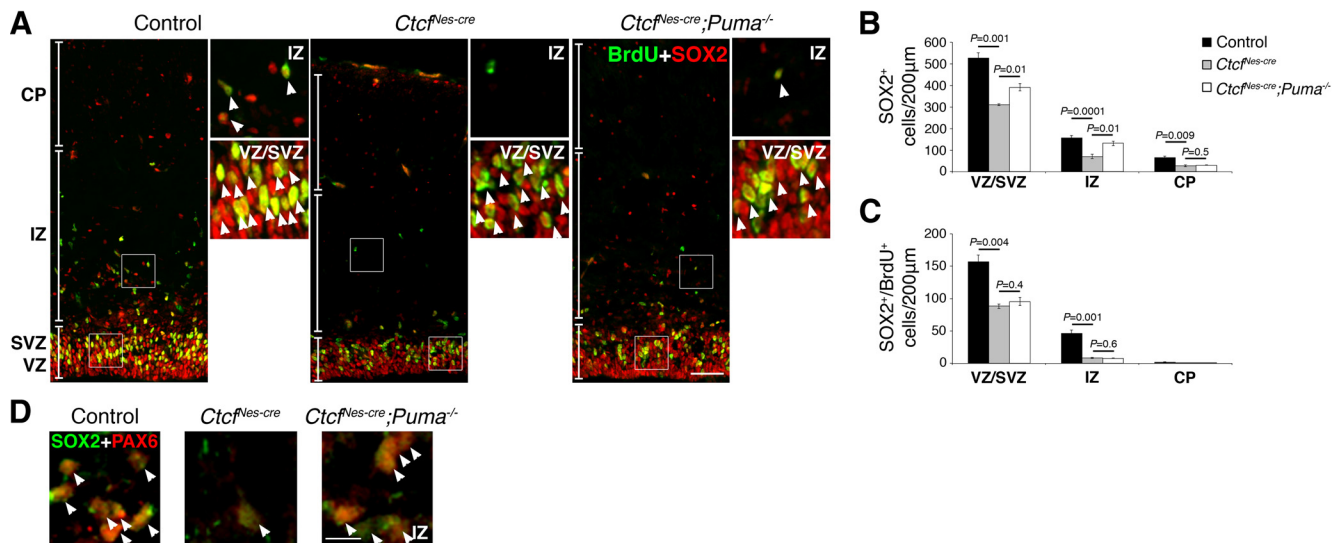


Figure 8. *Ctcf*-deficiency results in PUMA-dependent apoptosis of apical and outer radial glia progenitors. Apical and outer radial glia progenitors that are rescued from apoptotic death fail to proliferate. Pregnant females were subjected to a 1 h BrdU pulse before being killed. **A**, SOX2 (red) and BrdU (green) coimmunostaining of E16.5 cortical cryosections. The inset demonstrates fewer SOX2⁺ cells in the *Ctcf*^{Nes-cre} IZ than control or *Ctcf*^{Nes-cre};*Puma*^{-/-}. **B**, SOX2⁺ cells were quantified in 200- μ m-wide cortical images ($n = 3$). **C**, SOX2⁺/BrdU⁺ cells were quantified in 200- μ m-wide cortical images ($n = 3$). **D**, Immunodetection of SOX2 (green) and PAX6 (red) in the E16.5 cortical IZ demonstrates restoration of oRG progenitors in *Ctcf*^{Nes-cre};*Puma*^{-/-} cortex compared to *Ctcf*^{Nes-cre}. Arrowheads indicate SOX2⁺/PAX6⁺ oRG cells. Error bars represent SEM. Original magnification: **A**, **D**, 100 \times . Scale bars: **A**, 50 μ m; **D**, 10 μ m.

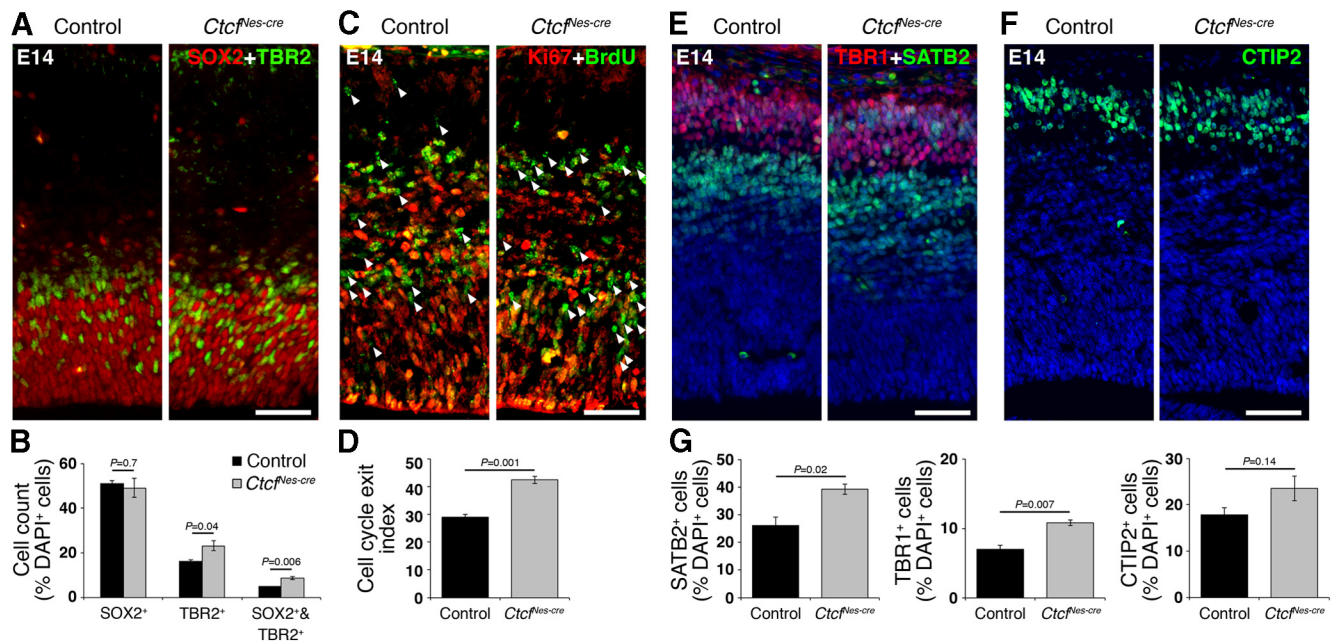


Figure 9. *Ctcf*-deficiency causes premature neurogenesis. **A**, Immunodetection of SOX2 (red) and TBR2 (green) in E14 control and *Ctcf*^{Nes-cre} cortex. **B**, SOX2⁺, TBR2⁺, and SOX2⁺/TBR2⁺ cells were quantified in 150- μ m-wide cortical images and expressed as a percentage of total DAPI⁺ cells ($n = 3$). **C**, Pregnant female mice were subjected to a 24 h BrdU pulse before being killed. Ki67 (red) and BrdU (green) immunostaining was used to determine the percentage of cells exiting the cell cycle in control and *Ctcf*^{Nes-cre} E14 cortex. Arrowheads indicate BrdU⁺/Ki67⁻ cells that have exited the cell cycle. **D**, Cell cycle exit indices were calculated by measuring the ratio of BrdU⁺/Ki67⁻ cells to total BrdU⁺ cells in control and *Ctcf*^{Nes-cre} cortex at E14 ($n = 3$). **E**, Immunodetection of TBR1 (red) and SATB2 (green) in E14 control and *Ctcf*^{Nes-cre} cortex. **F**, Immunodetection of CTIP2 (green) in E14 control and *Ctcf*^{Nes-cre} cortex. **G**, SATB2⁺, TBR1⁺, and CTIP2⁺ cells were quantified and expressed as a percentage of DAPI⁺ cells ($n = 3$). Error bars represent SEM. Original magnification: **A**, **C**, **E**, **F**, 100 \times . Scale bars: **A**, **C**, 50 μ m; **E**, **F**, 50 μ m.

the double mutant brain at birth appeared hypocellular and was nearly undistinguishable histologically from the *Ctcf*^{Nes-cre} brain (Fig. 5). We showed that *Ctcf*^{Nes-cre};*Puma*^{-/-} apical and oRG progenitors that cannot activate the apoptotic pathway fail to incorporate BrdU, which might be explained by a p53-mediated cell cycle arrest. Further investigations into the mechanism underlying increased p53 levels in the *Ctcf*-null brain will be important to fully elucidate the role of CTCF in progenitor cell survival.

Given that oRG cells are implicated in neocortical expansion in humans, their reduced ability to proliferate in the double mutant cortex might partly explain the failure to recover cortical size at birth, and may be relevant to microcephaly caused by CTCF mutations in humans (Hansen et al., 2010; Lui et al., 2011; Reillo et al., 2011).

A key observation is that deletion of *Puma* did not completely restore the number of apical cells in the *Ctcf*^{Nes-cre};*Puma*^{-/-} cor-

tex, suggesting that while a proportion of this progenitor population undergoes *Puma*-dependent death in the absence of CTCF, apical cells exhibit an independent defect. We found that *Ctcf*-null apical progenitors differentiate prematurely, causing an initial increase in the production of basal progenitors and early-born postmitotic neurons (Fig. 9). However, the increased number of basal progenitors is later counteracted by a reduction of the progenitor pool from which they are derived, and exacerbated by increased levels of apoptosis. Given that CTCF colocalizes with the cohesin complex and is required for cohesin localization to specific genomic sites to influence gene expression (Parelho et al., 2008; Wendt et al., 2008), it is conceivable that premature neurogenesis in the *Ctcf*^{Nes-cre} cortex results from dysregulation of CTCF target genes.

The number of basal progenitor cells marked by TBR2 expression was completely restored in the double mutant brain, indicating that PUMA activation causes basal progenitor cell death in the absence of CTCF. Simultaneous labeling of brain sections with TBR2 and BrdU showed that rescued basal progenitors are still able to proliferate, unlike rescued apical and oRG cells. Despite the overall equivalent number of TBR2⁺ cells in the control and double mutant cortex, several rescued basal progenitors did not move basally out of the ventricular zone, perhaps due to a defect in apical radial glia or an inability to correctly delaminate from the apical surface of the *Ctcf*-deficient cortex.

In summary, we demonstrated that CTCF is required in the early developing mouse brain for neuroprogenitor cell survival and that its deletion induces p53- and PUMA-dependent apoptosis. Independent from its role in promoting cell survival, CTCF is required for the correct balance of proliferative versus differentiative divisions and maintenance of the apical progenitor pool. Together, these functions of CTCF contribute to the normal development of the mammalian neocortex.

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