Mllt11 Regulates Migration and Neurite Outgrowth of Cortical Projection Neurons during Development

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The formation of connections within the mammalian neocortex is highly regulated by both extracellular guidance mechanisms and intrinsic gene expression programs. There are two types of cortical projection neurons (CPNs): those that project locally and interhemispherically and those that project to subcerebral structures such as the thalamus, hindbrain, and spinal cord. The regulation of cortical projection morphologies is not yet fully understood at the molecular level. Here, we report a role for Mllt11 (Myeloid/lymphoid or mixed-lineage leukemia; translocated to chromosome 11/All1 Fused Gene From Chromosome 1q) in the migration and neurite outgrowth of callosal projection neurons during mouse brain formation. We show that Mllt11 expression is exclusive to developing neurons and is enriched in the developing cortical plate (CP) during the formation of the superficial cortical layers. In cultured primary cortical neurons, Mllt11 is detected in varicosities and growth cones as well as the soma. Using conditional loss-of-function and gain-of-function analysis we show that Mllt11 is required for neuritogenesis and proper migration of upper layer CPNs. Loss of Mllt11 in the superficial cortex of male and female neonates leads to a severe reduction in fibers crossing the corpus callosum (CC), a progressive loss in the maintenance of upper layer projection neuron gene expression, and reduced complexity of dendritic arborization. Proteomic analysis revealed that Mllt11 associates with stabilized microtubules, and Mllt11 loss affected microtubule staining in callosal axons. Taken together, our findings support a role for Mllt11 in promoting the formation of mature upper-layer neuron morphologies and connectivity in the cerebral cortex.

Key words: axonogenesis; callosal projection neurons; microtubules; Mllt11/All1q; neuritogenesis; neurodevelopmental disorders

Significance Statement

The regulation of cortical projection neuron (CPN) morphologies is an area of active investigation since the time of Cajal. Yet the molecular mechanisms of how the complex dendritic and axonal morphologies of projection neurons are formed remains incompletely understood. Although conditional mutagenesis analysis in the mouse, coupled with overexpression assays in the developing fetal brain, we show that a novel protein called Mllt11 is sufficient and necessary to regulate the dendritic and axonal characteristics of callosal projection neurons in the developing mammalian neocortex. Furthermore, we show that Mllt11 interacts with microtubules, likely accounting for its role in neuritogenesis.

Introduction

The mammalian neocortex is a complex structure, underlying the capacity for executive function, sensory processing, emotion, motor output, and cognition. Its laminated organization of excitatory pyramidal cortical projection neurons (CPNs) and inhibitory interneurons (INs) emerges from the coupling of neurogenesis with neuronal migration. Functional circuits emerge progressively during development as newborn neurons acquire molecular identities, dendritic morphologies, and projections characteristic of their laminar position. A fundamental question concerns how migration is controlled during sequential generation of more superficial cortical layers. While it is clear that the cytoskeleton plays a key role in neuronal translocation, its role in CPN subtype-specific neurite morphogenesis and the
maintenance of laminar transcriptional programs is not well understood. The sorting of differentiating CPN subtypes into discrete layers occurs as cells migrate radially from progenitor domains populating the ventricular zone (VZ) and adjacent subventricular zone (SVZ) along the apicobasal axis to invade the cortical plate (CP). The earliest-born CPNs migrate along radial glial fibers projecting from the basal surface of VZ progenitors to the pia via somal translocation as forces exerted on the cytoskeleton by motor proteins push and pull the nucleus toward the pia until a neuron reaches its terminal location (Bellion et al., 2005; Tsai et al., 2007). As the somas of these cells approach the pia, the trailing apical pole becomes the nascent axon projecting to form white matter (WM) tracts (Nadarajah et al., 2001; Marin and Rubenstein, 2003; Ayala et al., 2007; Sakakibara et al., 2014). Consequently, the intermediate zone (IZ) and CP become densely populated, necessitating multiple modes of motility in later-born upper layer (UL). These events rely on coordinated cytoskeletal reorganization as cells transiently acquire multipolarity to weave through the dense IZ and then reacquire bipolarity on invasion of the CP (Sakakibara et al., 2014). The growth of axons during brain formation also involves dynamic cytoskeletal reorganization at the growth cone in response to external guidance cues or cell-cell contacts (Hirokawa and Takemura, 2004; Ayala et al., 2007).

While axonogenic mechanisms are conserved across CPN subtypes, the targeting and wiring of each cortical layer is dependent on its unique molecular characteristics. Each cortical layer expresses a unique transcription factor program that provides the molecular instructions for synapse formation with appropriate target regions of the brain. Beginning with the earliest-born Cajal–Retzius (CR) cells of layer 1 (L1), which secrete Reelin to guide the radial migration of nascent neurons (Frotscher, 1998; Chai et al., 2009, 2016; Gil-Sanz et al., 2013), neurogenesis and laminarization proceeds in an inside-out, cross-repressive manner. Transcription factors Tbr1 (Hevner et al., 2001; Thomson, 2010) and Bcl11a/Ctip2 (Arlotta et al., 2005) of deep layer (DL)6 and DL5, respectively, suppress UL-specific factors until DL formation is complete and subcerebral projections are initiated (Toma et al., 2014). UL neurogenesis follows with expression of Satb2 in thalamocortically-targeted L4 CPNs (Hisaoka et al., 2010; Leone et al., 2015; Lopez-Bendito and Molnar, 2003), and co-expression of Satb2 (Alcama et al., 2008; Britanova et al., 2008; Leone et al., 2015), Cux1 (CDP), and Cux2 (Nieto et al., 2004; Zimmer et al., 2004) in UL2/3 corticocortical CPNs, 80% of which project across the corpus callosum (CC; Leone et al., 2015).

In an effort to identify regulators of neural migration and differentiation, we previously reported the expression of Mllt11/Af1q (Myeloid/lymphoid or mixed-lineage leukemia; translocated to chromosome 11/All1 Fused Gene From Chromosome 1q) in the developing CP (Yamada et al., 2014). Mllt11 is a vertebrate-specific, 90 amino acid protein first identified as a fused protein with Mll in infant acute myeloid leukemia (Tse et al., 1995). Here, we report that Mllt11 is a neural-specific tubulin-associating protein and its inactivation in the superficial layers of the mouse brain results in improper migration and formation of callosal projections. Loss of Mllt11 led to neurite outgrowth defects coupled with loss of UL-specific transcriptional programs characteristic of neurodevelopmental disorders. In contrast, Mllt11 overexpression promoted the invasion of the CP by differentiating neurons. We provide evidence that Mllt11 interacts with stabilized microtubules, consistent with Mllt11 mutants phenocopying brain tubulinopathies (Bahi-Buisson et al., 2014). Altogether, these findings contribute significantly to our understanding of the genetic regulation of CPN development, dendritic complexity, and axonal connectivity.

Materials and Methods

Animals

All animal experiments were done according to approved protocols from the IACUC at Dalhouse University. Mice (Mus musculus) carrying a null mutation in the Mllt11 gene were generated using embryonic stem (ES) cell clones obtained from the mouse knock-out consortium project (UCDavis KOMP repository, Mllt11tm1a(KOMP)Mgpbp). The targeting construct is a “knock-out, conditional second” approach, which inserts the gene encoding β-galactosidase (β-gal) into exon 2, the protein-coding sequence of the Mllt11 gene. Two independently targeted clones were injected into blastocysts, and the resulting chimeras were mated to BL/6 females to achieve germ-line transmission. Offspring were genotyped by PCR using the following primers: wild-type forward (F) 5'-CGGTCGTGCCCTTGTAGGTTCTCAGC-3' and reverse (R) 5' -GCTACTGCACAAGGCTTCTTTGG-3' (expected product size: 379 bp), Mutant F5'-GAGATGGCGGCAACAGATTATG-3' and R5'-AAGCAGATTGTTGCTACTGGCCTTG-3' (expected product size: 274 bp). Heterozygotes (Mllt11tm1a(KOMP)Mgpbp) were maintained on a C57BL/6 background and crossed with FlpO+ /F6 (B6.Cg-Tg(Pkg1-flp)105ykr/), 011065, The Jackson Laboratory) mice and converted to a conditional allele via germ line Flp recombinase expression. Resulting Mllt11Flox/Flox offspring were crossed with Ai9 Rosa26.TdTomato (B6.Cg-Gt(Rosa26)Sor1MmU1-cre(CAG-Tomato)Ai9/J, 007909, The Jackson Laboratory) mice and Cre F5'/-Galactosidase (β-galactosidase) into exon 2, the protein-coding sequence of the Mllt11 gene. Two independently targeted clones were injected into blastocysts, and the resulting chimeras were mated to BL/6 females to achieve germ-line transmission. Offspring were genotyped by PCR using the following primers: wild-type forward (F) 5'-CGGTCGTGCCCTTGTAGGTTCTCAGC-3' and reverse (R) 5' -GCTACTGCACAAGGCTTCTTTGG-3' (expected product size: 379 bp), Mutant F5'-GAGATGGCGGCAACAGATTATG-3' and R5'-AAGCAGATTGTTGCTACTGGCCTTG-3' (expected product size: 274 bp). Heterozygotes (Mllt11tm1a(KOMP)Mgpbp) were maintained on a C57BL/6 background and crossed with FlpO+ /F6 (B6.Cg-Tg(Pkg1-flp)105ykr/), 011065, The Jackson Laboratory) mice and converted to a conditional allele via germ line Flp recombinase expression. Resulting Mllt11Flox/Flox offspring were crossed with Ai9 Rosa26.TdTomato (B6.Cg-Gt(Rosa26)Sor1MmU1-cre(CAG-Tomato)Ai9/J, 007909, The Jackson Laboratory) mice. The resulting Cux2iresCre+ /- or Cux2iresCre+ /+ offspring were crossed with the conditional knockout line to conditionally knock out Mllt11. Offspring of these crosses were genotyped using the following primers: wild-type forward (F) 5'-CGGTCGTGCCCTTGTAGGTTCTCAGC-3' and R5'-GCTACTGCACAAGGCTTCTTTGG-3' (expected product size: 379 bp), Post Flp/Cre F5'-CGGTCGTGCCCTTGTAGGTTCTCAGC-3' and R5'-AAGCAGATTGTTGCTACTGGCCTTG-3', and Cre F5'-GCTACTGCACAAGGCTTCTTTGG-3', and R5'-AAGCAGATTGTTGCTACTGGCCTTG-3', and Cre F5'-GGCATGACACACCGGAAATG-3', and R5'-GGCATGGAAAAACCTATCCGACAGA-3'. We genotyped for the presence of TdTomato primers F5'-TACGGCATGAGGACGGTCGAGATAAA-3' and R5'-CAGGGCAGACGACGCAAGAAA-3' (expected product size: 500bp) and allelism was determined with primers F5'-TACTAATGGCCGGGGTGTGTT-3', R5'-TCTACTGGTGCTTCTCAGG-3', and R5'-TACTAATGGCCGGGATCGGAGAATA-3' (expected product size: 250 pb for wild-type and 300 bp for mutant). Similar numbers of control and Mllt11cko mutant phenotype were observed in pilot studies. Statistical analysis (see below) was conducted on combined numbers of male and female offspring.
GST protein expression, mass spectrometry, and immunoprecipitation

GST-tagged protein expression

The entire Mllt11 sequence was cloned into pGEX-4T-2 vector for GST pulldown assays. BL21 competent cells (Sigma) were transformed with pGEX-4T-2 (GST) and pGEX-4T-2-MLLT11 (GST-MLLT11). Colonies were picked and incubated overnight in 2 ml 2xYT media with ampicillin at 37°C with shaking. The following day, overnight culture was added to 150 ml 2xYT media with ampicillin and incubated at 37°C with shaking until OD600 reached 0.6. Protein expression was induced by adding IPTG (Invitrogen) to the culture to a final concentration of 0.1 mM and shaken at 28°C for ~4 h. Cells were pelleted by centrifugation and resuspended in 8 ml NP-40 lysis buffer (50 mM Tris, pH 8.0, 150 mM NaCl, and 1% NP-40) with protease inhibitor cocktail (Sigma). Cells were incubated at 4°C for 10 min, sonicated on ice (30 s on/30 s off, for 3 min total, level 6 intensity), and again pelleted at 4°C for 10 min. Lysed cells were pelleted and supernatant collected (protein lysate).

Immobilization of GST or GST-MLLT11 bait protein

GST or GST-MLLT11 proteins were immobilized to Glutathione Sepharose 4B beads (Pierce). GST or GST-MLLT11 protein lysates were added to equilibrated 50% bead slurry and incubated at 4°C for ~3 h. Beads with immobilized protein were collected, washed, and resuspended in PBS to make a 50% slurry.

GST pull-downs and mass spectrometry

Whole embryonic mouse brains were harvested at embryonic day (E) 15.5 and lysed in NP-40 lysis buffer with protease inhibitor cocktail (Sigma). Whole-brain lysates from one embryo were added to either 6-µg GST or 6-µg GST-MLLT11 bead slurries and incubated overnight at 4°C. Beads with bound lysate were pelleted, washed, resuspended in sample buffer. Samples were heated for 15 min at 37°C and run on an SDS-PAGE gel, then stained with Coomassie Blue (Pierce). Bands were cut from the SDS-PAGE gel and processed by Dalhousie’s Biological Mass Spectrometry Core Facility.

Co-immunoprecipitation (co-IP) and Western blotting

HEK293 cells were transfected with myc or myc-MLLT11 vectors using Lipofectamine 2000 (Invitrogen). Twenty-four hours later, cells were lysed on ice with Tris-HCl lysis buffer containing 50 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1 mM EDTA, 1% Triton X-100, and protease inhibitor cocktail (Sigma). Lysed cells were collected, centrifuged, and supernatant used for co-IP. Anti-c-Myc Agarose resin (Pierce) was used, as per manufacturer’s protocol, to co-immunoprecipitate mhc or mhc-MLLT11 and their binding partners. Proteins were eluted from the resin with 50 mM NaOH, neutralized with 1 M Tris, pH 9.5, and added to nonreducing sample buffer for Western blot analysis. Whole-brain lysates, E18.5 cortical protein samples were separated on 8% SDS-PAGE gels for 1 h at 120 V and transferred overnight at 20 V on to PVDF membranes (Bio-Rad). Blots were probed with mouse anti-actin acetylated tubulin (1:20,000, Sigma), and rabbit anti-Mllt11 (1:1,000, Abcam). Secondary antibodies were goat anti-rabbit HRP (1:5,000, Invitrogen) and goat anti-mouse HRP (1:5,000, Invitrogen). Blots were developed with Clarity Western ECL Substrate (Bio-Rad) and imaged on a ChemiDoc Touch Gel Imaging System (Bio-Rad). Band densitometry was done using Image Lab Software (Bio-Rad).

qPCR

To confirm the loss of Mllt11 transcripts, E18.5 RNA was extracted from cortices of three genotypic conditional knockout (kOs) and four controls using the RNAeasy Micro kit (QIAGEN). RNA was reverse transcribed to cDNA using the SuperScript II Reverse Transcriptase kit (Invitrogen). qPCRs were conducted using the SensiFAST SYBR No-ROX Kit (Bioline) with the following primers for Mllt11 and internal control GAPDH: Mllt11 F 5'-GAACTGGATCTGTCGGAGCT-3' and R 5'-GCCGCTTCTCCAGAAGTGTGAAG-3', GAPDH F 5'-ACCAACG TCCATGCCCATAC-3' and R 5'-TCCACACCCCTGTGGTGTGA-3' (Weng et al., 2014). Reactions were performed in triplicates.

Histology

Fetal brains were dissected out and fixed in 4% paraformaldehyde (PFA) for 4–8 h, depending on embryonic stage, before being equilibrated in surose, embedded in Optimum Cutting Temperature (OCT) compound (Tissue-Tek), and cryosectioned at 12 µm. Cortex morphology was assessed by DAPI staining. β-Gal Tissue Stain kit (Millipore). Immunohistochemistry was conducted on E14.5–E18.5 brains as described previously (Yang et al., 2008).

Immunohistochemistry was conducted using the following antibodies: rabbit anti-CFP/CAx1 (1:100, Santa Cruz), mouse anti-Satb2 (1:250, Abcam), rat anti-Bcl11b/Ctip2 (1:500, Abcam), rabbit anti-Tbr1 (1:200, Abcam), rabbit anti-Cleaved Caspase 3 (CC3; 1:500, Cell Signaling Technology), rabbit anti-Pax6 (1:500, Abcam), goat anti-Sox2 (1:200, Santa Cruz), rat anti-Tbr2 (1:200; ebioscience), mouse anti-neurofilament 2H3 (1:200; DSHB, University of Iowa), rabbit anti-Tuj1 (Tubb3; 1:1000, Biologend), and mouse anti-acetylated α-tubulin (1:1000, Sigma). Species-specific Alexa Fluor 488, 568, 594, and/or 647-conjugated IgG (1:1500; Invitrogen) secondary antibodies were used to detect primary antibodies.

In situ hybridization was performed on 30-mm frozen sections obtained from E18.5, postnatal day (P7), P14, P21, and P28 Control and cKO brains fixed overnight as previously described (Yamada et al., 2014) using an Mllt11 riboprobe.

For EDU birth dating studies, dams were injected intraperitoneally with 30 mg/kg body weight of Edu (Invitrogen) at E14.5, E16.5, and E18.5 and killed at E14.5 or E18.5. Sections were immunostained using the Click-It kit according to the manufacturer’s protocol (Invitrogen).

For Golgi stains, brains were harvested from mice at P28 and subjected to the FD Rapid GolgiStain kit (FD Neurotechnologies) as per manufacturer instructions. Crude sections were cut and mounted on slides with Permount Mounting Medium (Fisher Scientific). Four brains were analyzed per genotype and 30 neurons per individual were analyzed.

Microscopy

Images were captured using a Zeiss AxioObserver fluorescence microscope equipped with an Apotome 2 structured illumination device, 10 ×, 20 ×, and a Hamamatsu Orca Flash v4.0 digital camera. β-Gal and Mllt11 in situ staining was captured using an upright Zeiss Primostar compound microscope with an ERCCs color camera. Images were processed using Zen software (Zeiss) and Photoshop CS6 (Adobe).

Primary cortical cell culture and immunocytochemistry

Cortices were microdissected from E18.5 embryos, digested in trypsin (Pierce), manually triturated and plated on 35-mm well onto poly-D-lysine coated coverslips at a density of 150,000 cells for the neurite outgrowth assay or 500,000 for cellular localization experiments. Cells were plated in medium containing DMEM with 10% FBS and 1% penicillin/streptomycin and 4 h after plating, media was completely removed and replaced with Neurobasal media containing B-27 and 1% penicillin/streptomycin. Cells were cultured for 24 h or one week in a 37°C incubator containing 5% CO2, then fixed for 4 h in a 37°C incubator containing 5% CO2, then fixed for 10 min in 4% paraformaldehyde. Immunocytochemistry was conducted using the following antibodies: mouse anti-Tau (1:200, Abcam), rabbit anti-α-Tubulin (1:1000, Sigma), and mouse anti-acetylated α-tubulin (1:1000, Sigma).

In utero electroporation and cDNA constructs

In utero electroporation was conducted using standard methodology under sterile surgical conditions (Salto, 2006). Endotoxin-free DNA was prepared according to the instructions of the manufacturer (QIAGEN) and injected at 1.5 μg/μl into the telencephalic vesicles of embryos in time-staged pregnant females anesthetized under inhalable isoflurane (5 l/min). A small incision was made on the ventral midline of anesthetized pregnant FVB females under a sterile field treatment. Single uteri containing the E13.5 fetuses were extruded and electric current was delivered across the fetal brains as five pulses for 50 ms at 900-ms
intervals using tweezer-style electrodes linked to the pulse generator CUY21 Vivo SQ (Sonidel). The embryos were returned in the body cavity, the peritoneum was sutured and the skin was stapled. Experimental plasmids used were Mllt11ires-eGFP, and control plasmids included pIRE2-EGFP (Clontech) or pCIG (Addgene). To ensure comparable development staging, for each dam one uterus was electroporated with the experimental construct and the other with the control vector. Fetuses were allowed to survive for 2 d until E15.5 after which they were processed for cryosectioning to evaluate GFP expression and cortical layer development by immunostaining. N = 3 Mllt11-eGFP and N = 4 eGFP control fetal brains were analyzed.

**Dil tracing of callosal projections**  
Brains of E18.5 embryos were removed and embedded in 7% low gelling temperature agarose in DMEM medium with 1% penicillin/streptomycin with 0.5 µl of Dil (Invitrogen) was injected into the cortical WM tracts at rostral and caudal axial levels and incubated for 8 h in Tyrode’s solution. Tissues were then fixed overnight at 4°C in 4% paraformaldehyde before incubating for two weeks at room temperature in PBS with 1% penicillin/streptomycin. Images were captured using a Zeiss V16 Axiozoom fluorescent stereomicroscope equipped with an AxioCam 506 mono digital camera (Zeiss).

**Image sampling, quantification, and analysis**  
For analysis of immunostaining markers and EdU, counting frames (100 x 100 µm) were placed in a vertical strip along the somatosensory (S1) cortex with the first counting frame along the edge of the ventricle. At least three histological sections within the somatosensory cortex from three to eight different animals were analyzed for each immunostain, Dil or electroporated vector. Cells that were positively labeled for both DAPI and the marker were counted within each frame using ImageJ (FIJI; Schindelin et al., 2012). For analysis of primary cortical cell culture and Golgi stain, neurites were traced and measured and Sholl analyses were conducted using the “simple neurite tracer” plugin in FIJI. Analysis of radial glial parameters, E14.5, E16.5, and E18.5 cortical slices were immunostained with goat anti-Nestin antibodies (1:250, Santa Cruz). Control and Mllt11 KO mutant images were captured with identical acquisition parameters and radial glial morphology was assessed using the “directionality” plugin in FIJI to calculate the percentage of fibers in the image aligned in the same direction. Automatic thresholds were then applied and the area covered by Nestin staining was measured and expressed as a percentage of the total image area (Extended Data Fig. 3-1).

To ensure consistency among samples, cell counts were restricted to the presumptive S1 (somatosensory cortex) of the embryonic brain. In all experiments, three to six Mllt11 KO and control embryos were used for quantification analysis using the unbiased and systematic sampling method we described previously (Yamada et al., 2015). Counts for Chip2, Br1, Cux1, Cux2, Sox2, Pax6, CC1, and Sath2 are represented as line graphs, quantifying the proportion of DAPI stained cells expressing those markers. For EdU, TdTomato, and DAPI distribution analyses, proportion of total stained cells expressing markers within each cortical bin was quantified to ensure counting methods were consistent with previous studies (Chen et al., 2008; Lawson et al., 2017; Reyes et al., 2019). Image montages were assembled using Photoshop CS6.

**Statistical analysis**  
Statistical differences were determined with Student’s t tests (two-tailed) with Welch’s correction. Sholl analyses were performed with paired t tests (two-tailed) to compare branch points within comparable radii. Bar charts, line graphs, and statistical testing were conducted using GraphPad Prism V5.04d software, with results shown as mean ± SD. In all quantification studies, significance level was set at p ≤ 0.05 (*p ≤ 0.05, **p ≤ 0.01, ***p ≤ 0.001, ****p ≤ 0.0001).

**Results**  
**cKO of Mllt11 from UL2/3 CPNs**  
Mllt11 is expressed in developing neurons of the central nervous system, including the neocortex, but its role in cortical neurogenesis is unknown (Yamada et al., 2014). To investigate the role of Mllt11 in neurogenesis, we generated mice carrying a null mutation in the Mllt11 gene using ES cell clones obtained from the mouse knock-out consortium project (UCDavis KOMP repository, Mllt11tm1a(KOMP)Mbp). We used the targeted (unflipped) allele to visualize Mllt11 by β-galactosidase (β-gal) staining (Fig. 1A) because of the LacZ gene inserted into the Mllt11 locus (Extended Data Fig. 1-1A). Initial Mllt11 locus activity begins in the pallial MZ at E12.5, showing stronger expression in the IZ of the fetal cortex at E14.5 coinciding with the birth of basal progenitors fated to become UL2/3 CPNs, intensifying in the superficial cortical layers by E16.5–E18.5 (Fig. 1A). The localization of β-Gal staining reflected the activity of the Mllt11 locus and confirmed our previous report of Mllt11 mRNA and protein localization in differentiating neurons of the neocortex (Yamada et al., 2014). A detailed examination of Mllt11 transcript expression by in situ hybridization revealed that cortical expression was maintained until P21, when it began to taper to levels indistinguishable from background at P28 (Extended Data Fig. 1-1B,C). In the hippocampus, Mllt11 transcript levels decreased progressively from P7 to P21 and remained low at P28 (Extended Data Fig. 1-1C).

Given the temporal and spatial overlap between Mllt11 expression and neurogenic regions of the cerebrum, we set out to investigate the role of Mllt11 in the development cerebral cortex, and specifically in the formation of the UL2/3 callosally projecting CPNs that bridge communication interhemispherically (Fame et al., 2011) where Mllt11 expression appeared highest during corticogenesis (Fig. 1A). We used a targeted Mllt11 null allele to generate cKO Mllt11 mice, in which the entire protein coding sequence in Exon 2 is flanked by loxp sites (Extended Data Fig. 1-1A). To avoid nonspecific effects of the selection cassette, mice encoding germline expressing fpl recombinase were crossed to mice harboring the Mllt11 targeted allele, leaving only loxp sites used to excise the entire protein-coding exon (exon 2) to create a conditional allele. Subsequently, to inactivate the Mllt11 in UL2/3 of the cerebral cortex, we crossed the Mllt11lox/lox+ mice to the Cux2iresCre reporter mice to visualize recombined neurons, creating Mllt11 cKO mutant (cKO) fetuses and neonates. Cux2 is highly expressed in developing UL2/3 projections neurons, and mosaically expressed in the SVZ in the fetal pallial cortex (Nieto et al., 2004; Zimmer et al., 2004; Cubelos et al., 2008a,b). We confirmed Mllt11 loss in UL CPNs by qPCR of E18.5 cortices (Extended Data Fig. 1-2A,B) as well as by ISH at P7 (Extended Data Fig. 1-2C,D) from Cux2iresCre-driven Mllt11 cKO brains.

**Loss of Mllt11 leads to a progressive decrease in cortical thickness**  
Brain and body weight did not differ significantly between Mllt11 cKO mutants and controls at E18.5 (Fig. 1B–D). Mutants displayed a thinning of the cortex visible in the CP and WM beneath it (Figs. 1E, 5A,E). To explore the origin of this cortical thinning, we examined the morphology of the cortex beginning at E14.5 (Fig. 1F) when UL CPNs are born (Zimmer et al., 2004; Cubelos et al., 2008b). A thinning of the mutant cortex was observed beginning at E16.5 (Fig. 1G), increasing in severity by E18.5 (Fig. 1H). This was reflected by significant changes in the distribution of DAPI+ nuclei in deeper bins of the Mllt11 cKO, with a larger proportion of DAPI+ nuclei localized apically (toward the VZ) in E18.5 mutants relative to controls (Fig. 1H). We also evaluated whether programmed cell death could account for
the cortical thinning, but did not find any cells positive for the apoptotic marker CC3 in the Mllt11 cKO mutant neonatal cortex (Extended Data Fig. 1-3A,B), while retrosplenial regions, which typically show enhanced apoptosis, showed similar CC3+ staining in both controls and mutants (Extended Data Fig. 1-3C).

Mllt11 expression was restricted to the developing CP and not in the apical progenitor regions (VZ) of the cortex (Fig. 1A).
To rule out potential cell nonautonomous contributions from neural progenitors to the Mllt11 mutant cortical phenotype we quantified cells positive for progenitor markers from E14.5 to E18.5, corresponding to the period of UL neurogenesis. Pax6, a radial glial cell (RGC) marker (Göttz et al., 1998), was unaltered on Mllt11 loss at all observed time points (Extended Data Figs. 1-4A-C, 1-5A-C). Sox2, a less restrictive marker for neural progenitors and quiescent RGCs (D’Amour and Gage, 2003; Ellis et al., 2004), also exhibited no significant differences in expression in the Mllt11 cKO mutants compared with controls (Extended Data Figs. 1-4D-F, 1-5D-F). Tbr2+ intermediate (basal) progenitors, which give rise to UL 2/3 CPNs, did not differ markedly between control and Mllt11 cKO groups, but displayed fewer cells within deep (nonprogenitor) regions of the mutant cortex at E18.5, suggesting alterations in the formation and/or migration of nascent neurons from the SVZ (Extended Data Figs. 1-4G-I, 1-5G-I).

Mllt11 is required for the maintenance of UL CPN molecular identity

The transcriptional de-repression loop that specifies cell types in discrete cortical laminae coincides with the birth and migration of projection neurons (Kumamoto et al., 2013; Toma et al., 2014). Thus, a loss of Mllt11 in UL progenitors can potentially affect neuronal birth, migration, and/or specification. We therefore investigated whether Mllt11 loss had any role in regulating the molecular identity of UL CPNs. Expression of Sath2 in CPNs of L2-4 exhibited a decrease at E18.5 (Fig. 2C; Extended Data Fig. 2-1C) as did CDP/Cux1, another marker of UL CPNs (Fig. 2F; Extended Data Fig. 2-1F). Investigation at earlier developmental time points revealed that this phenotype was progressive, beginning at E16.5 and increasing in severity by E18.5 (Fig. 2A,B,D,E; Extended Data Fig. 2-1A,B,D,E). While the extent of CDP/Cux1 and Sath2 staining displayed variability across individual mutants, there was a consistent decrease in expression levels an apical shift in expression domain in all Mllt11 cKOs (Fig. 2C,F; Extended Data Fig. 2-1C,F). TdTomato labeling was used as a short-term lineage tracer of Cux2+ intermediate progenitors that give rise to UL2/3 CPNs, and while TdTomato levels were largely unaltered at all time points (Fig. 2G-I), there was a consistent apical shift in expression domain at E18.5 (Fig. 2I). Collectively, these findings suggested that Mllt11 loss in nascent UL neurons did not affect their neurogenesis, or the activation of UL gene expression programs, but may have affected their migration to contribute to L2/3 formation.

The cortical de-repression loop functions on a cellular level by alternately repressing DL and UL fate (Toma et al., 2014) so we wanted to determine whether the loss of UL CPN fate was coupled with a complementary upregulation of DL CPN fates in these cells. We examined the localization of DL6-specific and DL5-specific markers Tbr1 and Ctip2, respectively, in Mllt11 cKOs and controls at E14.5–E18.5 (Fig. 2J–O; Extended Data Fig. 2-1G–L). Mllt11 cKO mutants displayed an apical shift in the expression domains of both Tbr1 and Ctip2 from E16.5 to E18.5, consistent with decreased cortical thickness and impaired UL formation (Fig. 2K–O; Extended Data Fig. 2-1H–L).

Mllt11 is required for the migration of UL CPNs

Given the thinning of the cortex and shift of cortical layer markers throughout development, we hypothesized that Mllt11 may regulate CPN migration into the CP. To explore the role of Mllt11 in CPN migration, we pulsed pregnant dams with EdU to label UL CPNs at E14 or E16 and observed the localization of EdU at E18.5. CPNs labeled with EdU at E14 exhibited an apical shift of roughly 100 µm in the Mllt11 cKO mutants relative to controls (Fig. 3A), consistent with a decrease in cortical thickness in the cKOs (Figs. 1, 2). A greater proportion of E14-born CPNs remained restricted to the deeper bins of Mllt11 cKO cortices relative to controls (Fig. 3A). As CPNs labeled with EdU at E16 were still in transit by E18, with only a few control CPNs reaching the outermost layers of the cortex, the discrepancy in UL cell migration was less pronounced in cKOs (Fig. 3B). To examine whether Mllt11 loss affected the neurogenesis of UL neurons, we pulsed dams with EdU for 2 h at E14, which efficiently labeled nuclei within the basal progenitor region (Fig. 3C). We observed no difference in the numbers or distribution of EdU+ nuclei in Mllt11 mutants versus control cortices after a short pulse at E14 (Fig. 3C). To determine whether Mllt11 loss caused a delay in the timing of UL neurogenesis, we repeated this short pulse at E18 when UL neurogenesis is nearly complete and noted no differences in EdU+ cells in the superficial cortex of Mllt11 mutant versus control brains (Fig. 3D). We did observe a slight increase of EdU+ cells in the deepest 100-µm bin of the Mllt11 mutant cortex relative to controls, possibly reflecting aberrant migration of later born neurons (Fig. 3D). Overall, the EdU birth dating analysis suggested that Mllt11 primarily regulates the migration, but not neurogenesis, of UL CPNs into the CP.

Guidance of migratory CPNs is dependent on the earliest born CR cells of L1, which secrete Reelin to attract and halt neuronal migration at the pial interface (Tueting et al., 1999; Hirota and Nakajima, 2017). However, there were no differences in the expression of CR markers p73 or Reelin at E14.5 (Fig. 3E) or E18.5 (Fig. 3F) following Mllt11 loss in UL progenitors and nascent neurons, suggesting that the apparent migration defect was not because of abrogated Reelin signaling. Additionally, Mllt11 loss did not result in the perturbation of the radial glial scaffold on which migrating nascent neurons rely to reach their respective laminae, as evidenced by the comparable distribution of Nestin+ radial glia fibers in the Mllt11 cKO mutant and control fetal cortices (Extended Data Fig. 3-1A,E,I). We quantified the radial glial angle (Extended Data Fig. 3-1B,F,J), average area covered (Extended Data Fig. 3-1C,G,K), and dispersion of fibers (Extended Data Fig. 3-1D,H,I,J) at E14.5, E16.5, and E18.5, respectively, and found no significant differences between controls and cKOs. We conclude that the migratory defects in Mllt11 mutant cortices were not likely caused by alterations in the radial glial scaffold.

Given the migration deficit in Mllt11 mutants, we examined whether its overexpression in fetal brains is sufficient to promote migration to the superficial regions of the developing cortex. We used in utero electroporation to express a bicistronic vector containing Mllt11 and GFP cDNAs, or control GFP vector alone, in wild-type mouse embryonic forebrains. We performed electroporations in anesthetized pregnant dams at E13.5, by injecting a short-term lineage tracer of Cux2+ intermediate (basal) progenitors, which give rise to UL 2/3 CPNs, did not differ markedly between control and Mllt11 cKO groups, but displayed fewer cells within deep (nonprogenitor) regions of the mutant cortex at E18.5, suggesting alterations in the formation and/or migration of nascent neurons from the SVZ (Extended Data Figs. 1-4G-I, 1-5G-I).

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**Figure 2.** Mllt11 loss progressively perturbed the formation of UL CPNs. **A–O,** Coronal cortical slices stained for cortical layer markers. **A, B,** Coronal cortical slices stained for Satb2 expression were similar between Mllt11 cKOs and controls at E14.5 (**A**) but displayed decreased numbers of Satb2+ cells in upper bins, and an apical shift at E16.5 (**B**). **C,** Mllt11 cKO cortices displayed decreased Satb2+ cell numbers and apical shift at E18.5 compared with controls. **D, E,** Cdp/Cux1+ cell numbers were largely normal at E14.5 (**D**), but began to decrease at E16.5 (**E**). **F, G,** Ctip2 levels were severely decreased in Mllt11 cKOs compared with controls at E18.5. **G–I,** Distribution of TdTomato expression in the cortex was comparable between controls and cKOs at E14.5 (**G**), E16.5 (**H**), and E18.5 (**I**). **J–L,** Tbr1 levels and localization were comparable between control and cKO at E14.5 (**J**) and E16.5 (**K**), but its expression domain exhibited an apical shift at E18.5 (**L**). **M–O,** Ctip2+ cell numbers and localization were largely unaltered in cKO at E14.5 (**M**) and E16.5 (**N**), but exhibited an apical shift in expression domain at E18.5 (**O**). Line charts represent percentage of positive cells normalized to DAPI+ nuclei per 100×100 μm bin (**A–F, J–O**) and percentage of total TdTomato+ cells per 100×100 μm bin (**G–I**). Student’s t test with Welch’s correction, (**A, E, J, L–O**) N = 4, (**F–I**) N = 5, (**G**) N = 5 controls, 6 mutants, (**K**) N = 4 controls, 5 mutants. Data presented as mean ± 50 n.s., not significant; *p ≤ 0.05, **p ≤ 0.01, ***p ≤ 0.001, ****p ≤ 0.0001. Scale bar: 50 μm (**A–O**). VZ, ventricular zone. UL CPNs, upper layer cortical projection neurons. See Extended Data Figure 2-1.
Mllt11 loss leads to reduced callosal crossing fibers and a thinning of WM tracts

The migratory defects alone may be insufficient to explain the decreased cortical thickness of Mllt11 cKO brains at E18.5 (Fig. 1E,H). Given that reduction in UL CPNs (Fig. 2A,B) would likely affect interhemispheric projections (Aboitiz and Montiel, 2003; Alcamo et al., 2008; Fame et al., 2011), we examined whether Mllt11 loss impacted formation of projection fibers contributing to the cortical WM tracts by staining for NF in coronal brain slices. We observed notable decreases in the thickness of NF+ fibers in the region of the fetal cortex containing WM tracts at E16.5 (Fig. 5A–D) and E18.5 (Fig. 5E–H) following Mllt11 loss. Importantly, while NF+ fiber staining in the cc was greatly reduced in Mllt11 cKOs at E18.5, NF+ fibers in the internal capsule, reflecting corticothalamic projections, remained unchanged in the mutant brains (Fig. 5I). Taken together, our findings

Figure 3. EdU birthdating demonstrated a CPN migratory defect in Mllt11 cKO mutants. A, B, EdU birthdating of UL CPNs by injection at E14 showed altered distribution in Mllt11 cKOs at E18.5, reflecting an apical shift in expression correlating with the decreased cortical thickness. B, Apical shift in migrating CPNs in cKOs relative to controls following an EdU pulse at E16 (B). C, Proliferating cells labeled with a short pulse of EdU at E14 showed no significant differences in numbers or distribution between controls and cKOs. D, No significant difference in numbers of nuclei labeled by a short pulse of EdU at E18 populating most bins in cKOs versus controls, but a greater proportion of cells were retained in the cko VZ region. E, F, Pial sections at E14.5 (E) and E18.5 (F) showed comparable levels of expression of CR markers P73 and Reelin in L1 of cKOs versus controls. Line charts represent percentage of positive cells per 100 × 100 μm bin as a proportion of total EdU+ cells. Total co-labeled cells are shown as bar graphs. Student’s t test with Welch’s correction, (A, C, E) N = 4, (B, D, F) N = 3. Data presented as mean ± SD n.s., not significant; *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001. Scale bar: 50 μm (A–F). VZ, ventricular zone. CPN, cortical projection neuron. See Extended Data Figure 3-1.
revealed a critical role for Mllt11 in UL CPN neurite morphology and axonogenesis of callosal fibers connecting the developing cerebral hemispheres.

The reduction of NF+ fibers crossing the cc in mutant brains suggested that Mllt11 loss impacted the ability of UL2/3 CPNs to extend fibers across the telencephalic midline. To address the potential inability of Mllt11 cKOs to extend axonal projections, we injected the lipophilic fluorescent dye DiI into the WM tracts of cortical slices at E18.5 and allowed it to diffuse through the tissue for two weeks before visualization. DiI labeling of projections was not detectable in the cc of Mllt11 cKOs, while controls exhibited clearly labeled crossing callosal fibers (Fig. 5 J). In contrast, the corticothalamic projections of the internal capsule were intact and traced by the DiI in both controls and cKOs, demonstrating that corticothalamic projections from DL6 were unaffected by the Cux2iresCre driven Mllt11 cKO strategy (Fig. 5 J). Altogether, our findings demonstrated that Mllt11 loss in UL neurons specifically impacted the development of callosal projections.

**Mllt11 interacts with cytoskeletal proteins**

Our birthdating analysis suggested that the migratory defects observed in Mllt11 cKOs are likely independent of the maintenance of UL transcriptional programs. To identify potential pathways through which Mllt11 might be exerting its effects on CPN migration, we performed a GST pulldown assay to identify interacting proteins using lysate from E18.5 brains. Potential Mllt11-interacting proteins are listed in Table 1 and include multiple α and β tubulin isoforms (>70% coverage) as well several atypical myosins, such as Myosin 5a (36%), Myosin 9 (24%), and Myosin 10 (8%). A potential association between Mllt11 and tubulin is consistent with a recent whole cell proteomic study, which identified Mllt11 as a likely interactor of both α-tubulins and β-tubulins (Go et al., 2021). Given that acetylation of α-tubulin is associated with stabilized microtubules and is crucial for outgrowth of stable neurites, we decided to focus on validating interactions between Mllt11 and acetylated-α-tubulin by immunoprecipitation analysis. We overexpressed a myc-tagged Mllt11 in Hek293 cells and pulled down acetylated-α-tubulin.
Probing the blots with anti-Mllt11 antibodies confirmed this interaction was dependent on heterologous Mllt11 expression in the cells (Fig. 6B), validating an association between Mllt11 and stabilized tubulin isoforms.

To evaluate whether Mllt11 localized to growing neurites, we used immunocytochemistry to probe the subcellular distribution of Mllt11, Tubb3, and acetylated α-tubulin in cultures of primary fetal cortical neurons. Cortical neurons were extracted from wild-type embryos at E18.5 and cultured for 24 h or one week, then immunostained to reveal the localization of Tubb3, acetylated α-tubulin, and Mllt11. After 24 h in vitro, corresponding to the extension of a primary neurite, Mllt11/Tubb3 co-localized in discrete punctate patterns in varicosities along the distal portion of the developing neurite, as well as in the growth cone with limited overlap in the soma (Fig. 6C,c'). After one week in culture, neurons displayed a more elaborate neurite arborization pattern and acetylated α-tubulin and Mllt11 were both detected along proximal portion of neurite (Fig. 6D).
### Table 1. Potential Mllt11-interaction targets from GST pull-downs in E18.5 whole-brain lysates

| Band | Accession | Description | Sum PEP score | Coverage (%) | # Peptides | # PSMs | # AAs | MW (kDa) | Reference |
|------|-----------|-------------|---------------|--------------|------------|--------|-------|---------|-----------|
| 1    | D32435    | Unconventional myosin-Va (Myo5a) | 201.776       | 36           | 53         | 240    | 1855  | 215.4   | Li et al. (2014) |
| 2    | Q8VD05    | Myosin-9 (Myh9) | 181.542       | 24           | 36         | 142    | 1960  | 226.2   | Li et al. (2014) |
| 3    | Q09JH59   | Myosin-10 (Myh10) | 31.272        | 8            | 10         | 22     | 2013  | 233.3   | Li et al. (2014) |
| 4    | P63017    | Heat shock cognate 71-kDa protein OS = Mus musculus | 199.247       | 59           | 32         | 303    | 646   | 70.8    | Li et al. (2014) |
| 5    | P36475    | Stress-70 protein, mitochondrial (Hspa9) | 20.395        | 12           | 6          | 12     | 679   | 73.4    | Go et al. (2021) |
| 6    | P20029    | Endoplasmic reticulum chaperone BiP (Hsp90) | 23.74         | 10           | 5          | 50     | 655   | 72.4    | Go et al. (2021) |
| 7    | Q3THK7    | GMP synthase [glutamine-hydrolyzing] (Gmps) | 8.67          | 5            | 3          | 6      | 693   | 76.7    | Go et al. (2021) |
| 8    | Q70553    | Dihydropyrimidinase-related protein 2 OS = Mus musculus | 12.297        | 15           | 4          | 7      | 572   | 62.2    | Go et al. (2021) |

**Mllt11** expression displayed a high degree of overlap with acetylated α-tubulin and β-tubulin isoforms differentially over neurite development, consistent with a role in neurite development. We next evaluated microtubule levels in the CC of Mllt11 mutants and found that both Tubb3 and acetylated α-tubulin isoforms differentially over neurite extension because of the cycling of microtubule stability (Sudo and Baas, 2010; Lin and Smith, 2015; Wei et al., 2018). If Mllt11 loss affected the relative amounts of stabilized microtubules, we would expect to see significant changes in neurite morphogenesis of UL neurons. We therefore evaluated the morphology of cultured primary cortical neurons from Mllt11 cKO mutants. We took advantage of the Ai9 tdTomato+ reporter introduced into the Cux2iresCre/+; Mllt11flox/flx conditional mouse to efficiently label and isolate UL primary neurons in culture. After 24 h in vitro, the length of MAP2+ primary neurites was reduced in cKOs compared with controls (Fig. 7A,B), as was the total length of all neurites (Fig. 7C). Moreover, the proportion of total length of extended neurites represented by the primary neurite at 24 h was slightly reduced in cKOs compared with controls (Fig. 7D). Quantification of the average number of neurites per neuron revealed a significant decrease in cKOs compared with controls (Fig. 7E,F), further suggesting that initiation of neurite outgrowth was abrogated on Mllt11 loss.

After one week of culture in vitro, the total neurite length was drastically reduced in Mllt11 cKO CPNs (Fig. 7G). Since Mllt11 loss progressively attenuated primary neurite outgrowth in...
cultured primary UL CPNs, we next evaluated a requirement for Mllt11 in the elaboration of dendritic arborization patterns in cultured CPNs. An advantage of using cultured neurons is that they are free of apical and basal contacts, allowing their neurites to radiate uniformly from the soma. This feature of two-dimensional neuronal culture allows for the evaluation of dendritic complexity by Sholl analysis. Mllt11 mutant primary neurons displayed much less complexity in the branching of their neurites (Fig. 7H,I), thus Mllt11 loss greatly impacted the arborization patterns of cultured UL CPNs.

Given the prevalence of neurite growth and pruning during postnatal development, we evaluated the requirement of Mllt11 in the formation of dendritic arborization morphology characteristic of UL CNs in vivo at a time when mature
arborization patterns have fully developed. Golgi staining was performed on control and Mllt11 cKO brains harvested at P28 to capture mature CPN morphologies using the Golgi staining method. While control UL CPNs displayed complex dendritic morphologies typical of cortical pyramidal neurons (Fig. 8Aa–Ad), Mllt11 cKOs UL CPNs displayed greatly attenuated dendritic arborization morphologies (Fig. 8Aa’–Ad’). We quantified the length of neurites in the Golgi-stained UL CPNs of cKO and control brains and observed a severe decrease in total neurite length in the Mllt11 mutants (Fig. 8B). The extent of decrease of neurite arborization in vivo was similar to that observed in cultured primary neurons (Fig. 7G–I). Sholl analysis revealed that the overall neuronal morphology was also much less complex with less elaborately-branched neurites in Mllt11 cKO UL CPNs (Fig. 8C). Taken together, these findings confirmed a critical requirement of Mllt11 in the growth and elaboration of
neurites during the development of mature superficial CPN morphologies and projection phenotypes.

Discussion
We described the role of a novel vertebrate-specific and neuronally restricted protein, Mllt11, in the regulation of neurite extension and migration of superficial CPNs. The Mllt11 locus has been identified in human genomic regions potentially affected in schizophrenia patients (Brzustowicz et al., 2002, 2004; Su et al., 2017), suggesting a role in CNS development. Our study is the first to demonstrate that Mllt11 regulates key aspects of cortical neuron morphogenesis and connectivity. Using the

Figure 8. Decreased neurite outgrowth and branching complexity in UL CPNs in the Mllt11 cKO cortex. A, Corresponding 100 × (AA, AA') and 200 × (AB–AD') magnification images of coronal sections of P28 Golgi-stained control (AA–AD) and Mllt11 cKO cortices (AA'–AD'). B, C, Quantification of neurite length (B) and branching complexity (C) within UL2/3 showed reduced neurite length and branching complexity in cKOs compared with controls. Length measurements analyzed with Student’s t test with Welch’s correction, layer-specific Sholl analysis compared with paired t test, N = 4 (controls and cKOs), 30 neurons/individual. Data presented as mean ± SD; *p ≤ 0.05, **p ≤ 0.01. Scale bar: 100 μm (AA, AA') and 50 μm (AB–AD').
Cux2iresCre driver to excise Mllt11 in developing UL CPNs, we revealed that conditional Mllt11 loss decreased cortical WM tracts and callosal fibers, and abrogated extension and maintenance of mature arborecent dendrites. Moreover, Mllt11 conditional mutants did not affect the formation of Reelin+ CR cells nor radial glial scaffolding, both of which are crucial for guiding the migration of CPNs. This suggests that Mllt11 loss acted cell autonomously to hinder radial migration of CPN progenitors into the CP. The inefficient migration of UL CPNs in Mllt11 mutant brains likely reflected a role for Mllt11 in regulating the cytoskeleton, as it associated with stabilized microtubules, and was required for their maintenance in growing neurites. Neurons lacking Mllt11 displayed deficits in neurite complexity both in vivo and in vitro, demonstrating that Mllt11 is a critical regulator of neurite outgrowth.

We show that Mllt11 interacts with tubulin isoforms, confirming a recent proteomic screen (Go et al., 2021), which provides a molecular rationale for the basis of the Mllt11 mutant phenotype. Cortical thinning and increased ventricular lumen surface area are common phenotypes associated with neurodevelopmental disorders, and were observed in Mllt11 mutant brains, suggesting cytoskeletal dysregulation may underlie a subset of these disorders (Bahli-Buisson et al., 2014; Mensen et al., 2017; Li et al., 2019). Moreover, changes in microtubule dynamics can affect neuronal migration and the generation of functional connections in the brain. These are processes whose dysfunction may underlie a wide spectrum of neurodevelopmental disorders. Given that Mllt11 mutant mice displayed both migratory and neurite outgrowth defects, they are a model to study brain disorders arising from aberrant regulation of the neuronal cytoskeletal during development.

The microtubule cytoskeleton generates polarized force and provides a functional railway along which motor proteins migrate, whether to drive actin polymerization in generating protrusive forces at the leading edge of the growth cone, or integrate extracellular cues into regulation of microtubule activity (Dent and Kalil, 2001; Buck and Zheng, 2002; Lee et al., 2004; Zhou et al., 2004). Tubulin mutations have been shown to impair neuronal migration into the cortex (Aiken et al., 2017), and specifically affect UL CPNs (Keays et al., 2007) reliant on dynamic, multipolar cytoskeletal reorganization to extend short processes, “crawling” through established DL CPNs (Miyata et al., 2001; Sakakibara et al., 2014). The dynamic expression of Mllt11 during UL CPN development also supports its role as a critical regulator of UL migration. Interestingly, mutations in α-tubulin have been shown to cause axonal trafficking defects that impair synaptic stability and function without impacting axonal degeneration or neuronal survival (Buscaglia et al., 2020). The interaction of Mllt11 with microtubules could therefore provide a mechanistic explanation for the impaired migration and reduced CPN dendritic complexity seen in Mllt11 mutants. Specifically, we showed that Mllt11 associates with acetylated α-tubulin in fetal brain lysates, suggesting that it may be required for the stabilization of the cytoskeleton in growing neurites, and account for the similarities between the Mllt11 mutant phenotype with those targeting tubulin isoforms.

The inefficient extension of neurites may also be attributed to impaired trafficking of organelles and ribosomes for local protein synthesis, or vesicles carrying secreted cues and growth cone machinery to the axonal periphery (Kennedy and Ehlers, 2006; González et al., 2016). Specifically, a potential Mllt11 interactor Myosin 10 (Table 1) is known to regulate radial migration of CPNs through its effect on the localization of N-cadherin (Lai et al., 2015), and has been implicated in tumor invasion by regulating process extension (Ropars et al., 2016). It is tempting to speculate that altered neuronal trafficking may underlie morphologic and transcriptional deficits of Mllt11 mutant CPNs. Loss of Mllt11 from the superficial cortex altered levels of UL2/3-specific transcription factors Satb2 and CDP/Cux1, which function in establishment and maintenance of the CC as well as mature arborecent neuronal morphology and somal packing (Alcami et al., 2008; Cubelos et al., 2015; Rodríguez-Tornos et al., 2016; He et al., 2021). Importantly, the loss of layer-specific morphology and identity has been demonstrated in postmortem brains from autistic humans (Casanova et al., 2013; Stoner et al., 2014; Fujimoto et al., 2021). These clinical data are reminiscent of the severe dysplasia in the cortex of our conditional Mllt11 mutants, revealed by the Golgi staining method. It is presently unclear whether the loss of UL-specific gene expression we observed in our Mllt11 mutants, and tubulopathies more generally, is primarily because of cytoskeletal dysregulation, or other uncharacterized gene expression pathways functioning downstream of Mllt11. Some neuronal subtypes exhibit altered expression of subpopulation-specific markers when cytoskeletal regulation is altered such that connections to target regions cannot be established (Hippenmeyer et al., 2005), implying that there may be uncharacterized feedback mechanisms between establishment of connectivity and maintenance of expression of CPN subtype-specific transcription factors.

Other potential roles for Mllt11 may depend on binding to chaperon protein HSPA8 and regulating protein export from the nucleus required for protein degradation (Li et al., 2014). Yet another possible function for Mllt11 may include regulating the Wnt signaling cascade via T cell factor 7, which in turn regulates CD44 to promote cell migration and metastasis (Park et al., 2015; Li et al., 2018). As ours is the first study to explore Mllt11 function in the developing brain, additional studies are needed to determine whether Mllt11 could be exerting its effect on transcriptional regulation directly or indirectly via cytoskeletal interactions and intracellular trafficking, or both.

In light of this, we showed that the overexpression of Mllt11 promoted migration into the CP, consistent with a cytoskeletal regulatory function. This finding is also consistent with in vitro studies, which showed that the overexpression of Mllt11 in cancer stem cell lines promotes proliferation and invasiveness (Tse et al., 2017). It is possible that Mllt11-dependent cytoskeleton reorganization may favor symmetric divisions of organelles, nuclear components, and polarity protein complexes, therefore enhancing invasiveness of cells (Nance and Zallen, 2011; Piroli et al., 2019). On the other hand, the transcriptional regulation of Mllt11 by REST has been associated with promoting neuronal differentiation, with high REST and low Mllt11 expression in undifferentiated neuronal tissue, and increasing Mllt11 levels correlated with decreased REST activity during terminal neuronal differentiation (Hu et al., 2015). These previously published findings, in conjunction with the complexity of the Mllt11 mutant phenotype we reported here, imply that there are potentially two discrete pathways through which Mllt11 regulates neuronal development: a cytoskeletal regulatory mechanism, and a transcriptional regulatory mechanism.

Much of what is known about Mllt11 has been discerned from overexpression studies in immortalized cell lines and oncogenic clinical case studies, revealing pathogenic roles in cell process extension, invasiveness, and secretion of factors that promote efficient cellular motility (Lin et al., 2004; Tse et al., 2004; Li et al., 2006; Chang et al., 2008; Tiberio et al., 2017; Park et al.,
Oncogenesis and neurogenesis share commonalities at the level of cytoskeletal regulation of somatic and nuclear morphology, as well as extension of and trafficking along cytoplasmic processes, allowing for invasion of and migration through tissues. However, it is important to make the distinction between pathologic studies and our current findings on the role of Mllt11. Until the current study, the physiological role for Mllt11 was unclear. In all the published in vitro and oncogenic case studies, Mllt11 is aberrantly overexpressed in non-neuronal tissues, either by itself or as a fusion with the chromatin remodeling protein MII. We previously reported that Mllt11 protein and mRNA was exclusively localized to the developing CNS, and not in any other tissues (Yamada et al., 2014). The expression pattern of β-gal in the targeted Mllt11 allele confirmed that it is normally expressed only within the developing CNS, with the highest levels reflecting UL cortical neurogenesis. Consistent with this expression profile, we now show that Mllt11 is required for proper migration but not neurogenesis of UL CPNs. Furthermore, we also revealed that Mllt11 regulates neurite outgrowth and the maintenance of UL gene expression programs. The loss of Mllt11 led to a reduction of interhemispheric connectivity via reduced crossing callosal fibers. We also characterized a severe dysplasia of CPNs in Mllt11 EKO mutant neonatal brains; a phenotype found in severe neurodegeneration diseases. Whether the regulatory role of Mllt11 exerts its effect by altering cytoskeletal organization during development of UL CPNs using a genetic KO and labeling strategy to target the superficial cortex. Numbers, laminar distribution, and morphology of UL CPNs were assessed over development. By combining in vitro and in vivo neurite outgrowth and morphology assays, we demonstrate that Mllt11 is required for neuronal invasion in the CP, formation of mature dendritic morphologies characteristic of UL CPNs, and the extension of axons across the CC. Mllt11 interacts with the microtubule cytoskeleton and likely exerts its effect by altering cytoskeletal organization during development. Whether the regulatory role of Mllt11 is exerted through the stabilization of cytoskeletal architecture or by trafficking of cellular machinery along neurites will be the subject of future investigations.

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