



Defective myelination in an RNA polymerase III mutant leukodystrophic mouse

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RNA polymerase (Pol) III synthesizes abundant short noncoding RNAs that have essential functions in protein synthesis, secretion, and other processes. Despite the ubiquitous functions of these RNAs, mutations in Pol III subunits cause Pol III-related leukodystrophy, an early-onset neurodegenerative disease. The basis of this neural sensitivity and the mechanisms of disease pathogenesis are unknown. Here we show that mice expressing pathogenic mutations in the largest Pol III subunit, *Polr3a*, specifically in *Olig2*-expressing cells, have impaired growth and developmental delay, deficits in cognitive, sensory, and fine sensorimotor function, and hypomyelination in multiple regions of the cerebrum and spinal cord. These phenotypes reflect a subset of clinical features seen in patients. In contrast, the gross motor defects and cerebellar hypomyelination that are common features of severely affected patients are absent in the mice, suggesting a relatively mild form of the disease in this conditional model. Our results show that disease pathogenesis in the mice involves defects that reduce both the number of mature myelinating oligodendrocytes and the ability of these cells to produce a myelin sheath of normal thickness. The findings suggest unique sensitivities of oligodendrogenesis and myelination to perturbations of Pol III transcription.

RNA polymerase III | leukodystrophy | oligodendrocyte | hypomyelination

Leukodystrophies constitute a group of genetically heterogeneous inherited diseases that affect myelin sheath production and/or function in the central nervous system (CNS) (1). They include disorders, such as Pelizaeus-Merzbacher disease, where the causal mutation involves a myelin-specific gene, and many other disorders where the causal mutations affect gene functions that are required more broadly (1). With the increasing functional diversity of leukodystrophy-associated genes, it has become apparent that these diseases are not restricted to cells of the oligodendrocyte (OL) lineage, which ultimately produce myelin but may involve astrocytes, microglia, and neurons (2). RNA polymerase (Pol) III-related leukodystrophy, an autosomal recessive disorder, is one of the most frequent leukodystrophies based on exome sequencing (3). The condition was identified when five initially distinct syndromes with overlapping clinical and radiological features were found to have a common molecular basis (4–6). Pol III-related leukodystrophy is typically a hypomyelinating disorder and is known for its high phenotypic variability (4–10). Patients are diagnosed between birth and adolescence with neurological and nonneurological deficits, and are classified according to distinct brain magnetic resonance imaging (MRI) features and confirmed by sequencing (11). Neurological deficits include developmental delay with a progressive decline in cognitive and motor function, ataxia, cerebellar atrophy, and thinning of the corpus callosum. Nonneurological deficits may include hypodontia, myopia, and endocrine abnormalities, such as hypogonadotropic hypogonadism (8). An understanding of the disease mechanisms underlying these clinical characteristics is currently lacking.

Pol III is the largest eukaryotic RNA polymerase and is composed of 17 subunits. Ten of these subunits are unique to Pol III, five are common to all three nuclear RNA polymerases, and the remaining two subunits are shared with Pol I. The enzyme transcribes abundant noncoding RNAs, such as tRNA, 5S rRNA, U6 snRNA, and 7SL RNA that are involved in essential cellular processes, including protein synthesis, RNA processing/splicing, and protein secretion, among others (12–14). Pol III-related leukodystrophy mutations were initially identified in the two largest subunits, *Polr3a* and *Polr3b*, which form the catalytic core of the enzyme. Subsequently, disease-causing mutations have been found in three other subunits: *Polr1c*, *Polr3k*, and *Polr3gl* (15–17). In most cases, biallelic, compound heterozygous mutations have been reported, although several homozygous mutations have also been identified. Mapping of >100 Pol III-related leukodystrophy disease mutations on the cryoelectron microscopy (cryo-EM) structure of human Pol III suggests a variety of molecular defects from disrupted subunit folding and subunit–subunit interactions that likely impact enzyme assembly (15) to impaired functions involving structural elements important for catalysis and surface-exposed sites (18, 19). How these hypomorphic alleles alter the Pol III transcriptome to cause disease is a major unanswered question. Studies with patient-derived

Significance

Pathogenic mutations in subunits of RNA polymerase (Pol) III cause a prevalent autosomal recessive form of leukodystrophy. However, understanding of the mechanisms of pathogenesis, including how ubiquitously expressed Pol III mutations affect primarily the central nervous system, has been limited by the absence of an animal model of the disease. We show that conditional knockin of pathogenic *Polr3a* mutations in the *Olig2* lineage in mice results in growth, neurobehavioral, and hypomyelination phenotypes, reflecting a subset of clinical features of Pol III-related leukodystrophy patients. Myelination defects in the mice identify neural-specific roles for Pol III transcription. The phenotypes of Pol III-related leukodystrophic mice enable genetic and pharmacological approaches aimed at mitigating the consequences of this disease in humans.

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fibroblasts and other cell-based systems suggest a range of possible effects from a global reduction in translation due to decreased synthesis of tRNA and 5S rRNA to more limited or even gene-specific effects (e.g., reduced BC200 and 7SL RNAs) (16, 20, 21). However, the molecular mechanisms of pathogenesis remain poorly characterized and further progress is significantly limited by the absence of an animal model of the disease.

In this study, we report the characterization of a mouse model of Pol III-related leukodystrophy that is unique in exhibiting features of the disease found in patients. Conditional knockin of a *Polr3a* double-mutant allele in Olig2-expressing cells results in developmental delay in neonates and cognitive, sensory, and fine sensorimotor deficits in adult mice. These neurobehavioral phenotypes are consistent with hypomyelination of multiple regions in the CNS as revealed by MRI and diffusion tensor imaging (DTI), immunohistochemistry, and electron microscopy. Along with an analysis of OL cell populations and myelin-specific proteins, the results reveal defects in the number and function of myelinating OLs in the mutant mice. The work highlights the importance of Pol III transcription for the normal functioning of cells in the OL lineage and provides a model for further exploration of the role of Pol III in diseases of the CNS and other tissues.

Results

A Neural-Specific Defect in *Polr3a* Leads to Reduced Growth and Delayed Postnatal Development. We engineered mice with whole body and conditional knockin mutations (W671R/G672E) at adjacent positions in *Polr3a*. Homozygosity for either mutation causes Pol III-related leukodystrophy in humans (8). However, mice that are homozygous or hemizygous for the G672E mutation alone have no apparent phenotype (22, 23). The corresponding mutation in the homologous yeast subunit also had no phenotype (24). In contrast, mutation of both sites in the yeast *Polr3a* homolog results in temperature-sensitive growth and defects in Pol III transcription (24). The properties of this double mutant in yeast provided the foundation for the present study. Mice that are heterozygous for the whole-body double-mutant knockin (*Polr3a*^{KI/+}) have a normal body weight compared to WT mice (*SI Appendix, Fig. S1A and B*) and express WT and mutant *Polr3a* mRNAs at approximately equal levels (*SI Appendix, Fig. S1C*). POLR3A protein and total tRNA levels are unchanged in the brain and Northern blotting for a precursor tRNA that serves as an indicator of Pol III transcription also showed no change (*SI Appendix, Fig. S1E–J*). Nonetheless, homozygosity for the whole-body knockin (*Polr3a*^{KI/KI}) was lethal prior to embryonic day 12.5 (E12.5) (*SI Appendix, Fig. S1D*). Since the major clinical feature of Pol III-related leukodystrophy is hypomyelination (4–8), we used an *Olig2*-Cre driver and conditional *Polr3a* mutant knockin (*Polr3a*^{cKI/cKI}) mice in an effort to bypass embryonic lethality and investigate the effects of the mutation on myelination. Olig2, a CNS-specific basic helix–loop–helix transcription factor, becomes active at E11.5 and is expressed throughout the OL lineage, as well as in a subset of motor neurons and GFAP⁺ astrocytes (25–27). *Olig2*^{Cre+/-} *Polr3a*^{cKI/cKI} mice (hereafter referred to as *Polr3a*-cKI mice) are viable and exhibit recombination frequencies of ~83% in females and ~88% in males based on FACS analysis of O4⁺ OLs from brain homogenates of mice expressing a dual tdTomato-EGFP reporter (*SI Appendix, Fig. S2*).

Polr3a-cKI mice are often diagnosed at an early stage in development (early childhood to adolescence); therefore, we investigated the mice for phenotypes as neonates (postnatal day 1 [P1] to P21), adolescents (3 to 5 wk of age), and adults (≥8 wk of age). For neonates, we tracked body weight and monitored the acquisition of developmental milestones that serve as indicators of neurodevelopmental disorders. *Polr3a*-cKI neonates of both sexes have significantly lower body weights than the corresponding WT mice (Fig. 1A and B). This difference is more pronounced in males and increases over time from P5 to P21, in contrast to females that maintain a relatively constant and smaller

body weight difference. Male and female *Polr3a*-cKI mice also show a delay in achieving developmental milestones that reflect deficits in body-righting mechanisms, strength, coordination, locomotion, and the extinguishing of pivoting behavior (Fig. 1C and D) (28). Notably, impaired growth and developmental delay is also seen in patients with Pol III-related leukodystrophy (8).

Impaired Growth and Myelination in Adolescent and Adult *Polr3a*-cKI Mice. In humans and in rodents, postnatal growth impairment (e.g., due to malnutrition) can be followed by a period of catch-up growth. However, the body-weight difference of *Polr3a*-cKI mice was maintained in males and enhanced in females between 3 and 5 wk of age (Fig. 1G and J). These changes correlated with a smaller body size and lower brain weight throughout adolescence (Fig. 1E, F, H, I, K, and L). Differences in body weight and size were also evident in adult animals at 12 and 16 wk of age (*SI Appendix, Fig. S3A–D*).

To investigate myelin deposition in the CNS, we harvested brains and spinal cords from adolescent and adult WT and *Polr3a*-cKI mice (Fig. 1F) and performed immunohistochemistry targeting the lipid and protein components of myelin. In coronal sections, Luxol fast blue (LFB) staining of the phospholipid portion of myelin revealed reduced intensity in the corpus callosum splenium, hippocampus, and cerebral cortex of adolescent *Polr3a*-cKI mice (Fig. 2A–D). Similar observations were made for LFB staining of these regions in adult mice (*SI Appendix, Fig. S3E and J*). Additionally, FluoroMyelin staining of the total lipid portion of myelin was reduced in the corpus callosum genu (*SI Appendix, Fig. S3G and K*). Comparable findings were obtained for myelin basic protein (MBP), which showed reduced staining in the corpus callosum splenium, hippocampus, and the cerebral cortex of adolescent and adult *Polr3a*-cKI mice (Fig. 2E–H and *SI Appendix, Fig. S3F and J*). Changes in myelination were also apparent in the spinal cord where LFB and MBP staining of the ventral and dorsal horns was reduced and there was a reduction in the size of the white matter tract due to the projection of the ventral horns into the ventrolateral funiculus (Fig. 2I–L). Surprisingly, given the important role of the cerebellum in motor control and the frequent ataxia reported in Pol III-related leukodystrophy patients (8), LFB and MBP staining in sagittal sections of the cerebellum did not show any difference compared to WT (Fig. 2M–P). Additionally, we counted Purkinje cells, which express Olig2 (25), after Nissl staining, and found no difference (*SI Appendix, Fig. S3H, L, and N*). A quantitative assessment of immunofluorescence intensity confirmed the reduction in MBP signal in the cortex, corpus callosum, hippocampus, and the spinal cord (Fig. 2Q and *SI Appendix, Fig. S3M*). A small difference in MBP intensity in the thalamus of adolescent mice was absent in adult animals, and no difference was measured in the cerebellum. Altogether the histochemical data reveal impairments to myelination in distinct anatomical regions of the cerebrum and the spinal cord but no apparent changes in the cerebellum.

Adult *Polr3a*-cKI Mice Exhibit Multiple Neurobehavioral Deficits. Pol III-related leukodystrophy patients present with neurological deficits, including impaired cognitive and motor functions (8). To determine whether neurological deficits are present in *Polr3a*-cKI mice, we performed a battery of behavioral tests. Initially, we utilized a behavioral spectrometer with video tracking and pattern-recognition analysis to quantify numerous home cage-like locomotion and repetitive behaviors. No significant differences were observed, although mutant male mice showed a trend toward a shorter center track length, suggesting an anxiety-like behavior (*SI Appendix, Fig. S4A–F*). However, further investigation of this behavior using an elevated plus maze did not reveal any difference compared to WT mice (*SI Appendix, Fig. S4I*). Cognitive functions involving visual and spatial memory were assessed by object recognition (OR) and object placement (OP) tests, respectively, using a preference score

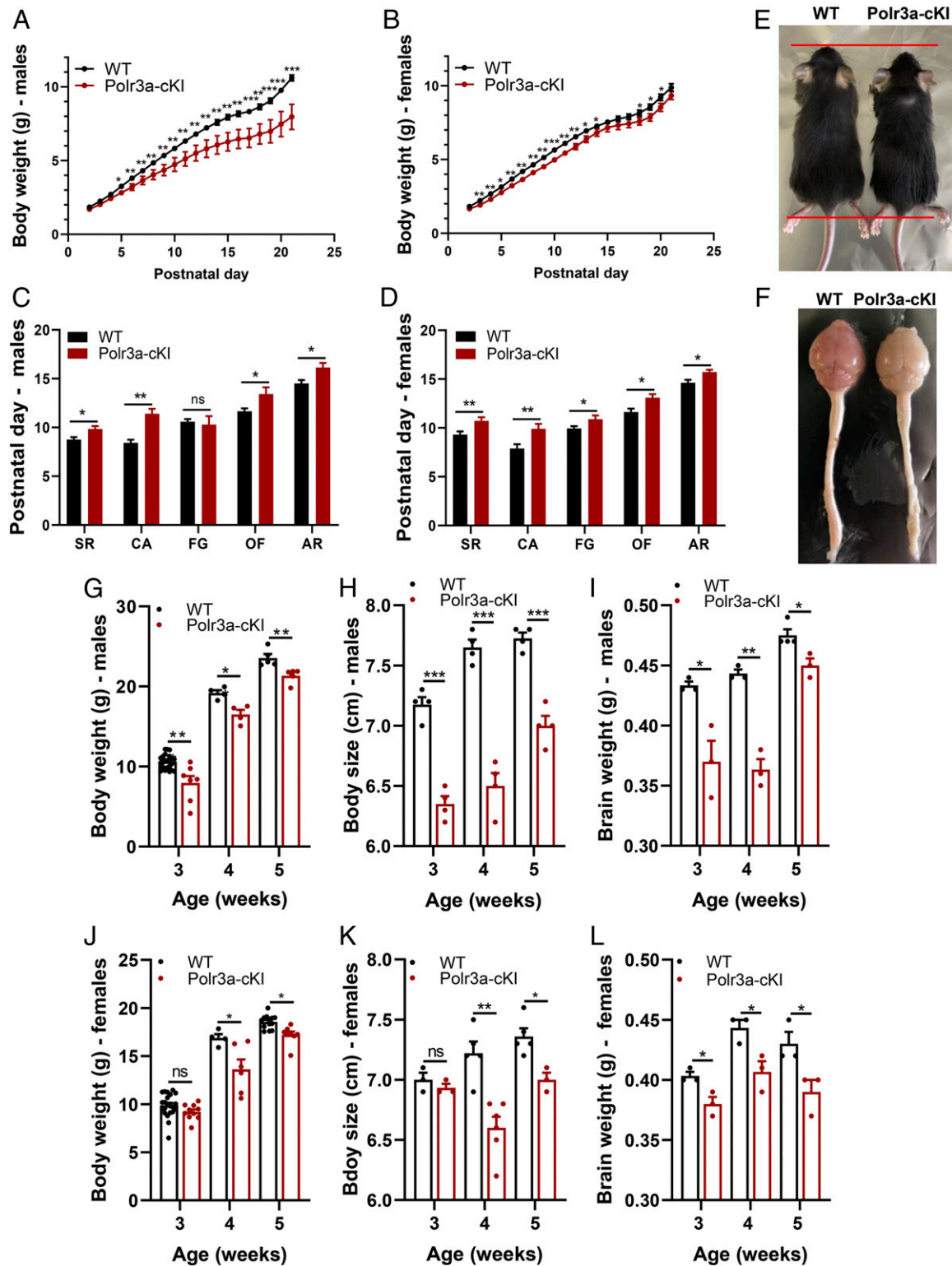


Fig. 1. Body metrics and developmental milestone assessment of neonates and adolescent WT and Polr3a-cKI mice. (A) Neonate male body weight curves (WT $n = 22$, Polr3a-cKI $n = 7$, nested t test $P = 0.0011$). (B) Neonate female body weight curves (WT $n = 22$, Polr3a-cKI $n = 11$, nested t test $P = 0.01$). (C) Male developmental milestones (WT $n = 20$ (SR, OF, and AR), 19 (CA), and 17 (FG); Polr3a-cKI $n = 6$ (SR, FG, OF, and AR), and 5 (FG); nested t test $P = 0.0061$). (D) Female developmental milestones (WT $n = 19$ (SR, OF, and AR), 18 (CA), and 17 (FG); Polr3a-cKI $n = 11$ (SR, CA, OF, and AR), 9 (FG); nested t test $P = 0.0028$). AR, air righting; CA, cliff aversion; FG, forelimb grasp; OF, open field; SR, surface righting. For A–D, the same mice were analyzed at multiple data points or tests (repeated measures). (E) Image of WT and Polr3a-cKI mice highlighting the body size difference at 3 wk of age. (F) Image of WT and Polr3a-cKI brain and spinal cord. (G–I) Body weight, body size, and brain weight of WT and Polr3a-cKI adolescent males, respectively. (J–L) Body weight, body size and brain weight of WT and Polr3a-cKI adolescent females, respectively. Values are presented as the mean \pm SEM. Groups were compared using two-sided Student's t test at each time point (multiple t test), * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$.

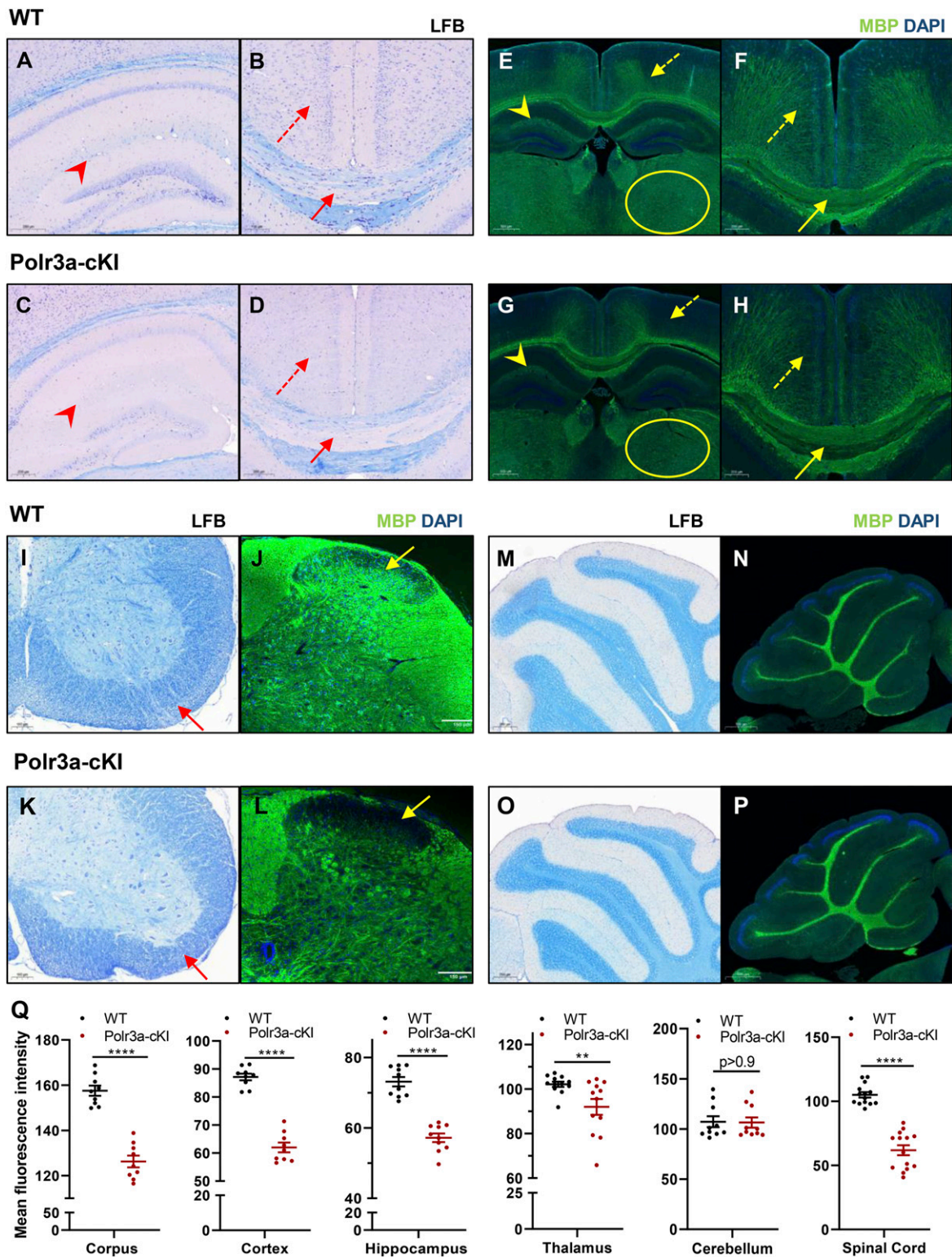


Fig. 2. Immunohistochemical staining of cerebral, cerebellar, and spinal cord sections from adolescent WT and Polr3a-cKI mice. (A–D) LFB staining of the hippocampus (arrowhead), corpus callosum splenium (solid arrow), and cortex (dashed arrow). (E–H) MBP staining of the hippocampus (arrowhead), corpus callosum splenium (solid arrow), cortex (dashed arrow), and thalamus (dashed oval). (I and K) LFB staining of spinal cord sections (solid arrow highlights differences in the ventral horn). (J and L) MBP staining of the spinal cord (solid arrow highlights differences in the dorsal horn). (M and O) LFB staining of cerebellar sagittal sections. (N and P) MBP staining of cerebellar sagittal sections. Images are representative of three mice per condition/sex. (Q) Quantification of mean MBP fluorescence intensity in different regions of the CNS in adolescent mice. Individual values are shown with their mean \pm SEM. Multiple sections from three mice per group/sex were compared using multiple *t* tests, ***P* < 0.01, *****P* < 0.0001.

cutoff of 55 (time exploring a new object or location/total time of exploration \times 100) to demonstrate the animals' inherent preference for novelty. With this as the criterion, 90 to 100% of male and female WT mice scored above the cutoff on both the OP and OR tests. In contrast, >60% of male and ~80% of female Polr3a-cKI mice scored below the cutoff in OP (i.e., failed to differentiate between familiar and novel locations) (Fig. 3 *A* and *B*) and ~60% of female Polr3a-cKI mice failed to differentiate between familiar and novel objects in OR (SI Appendix, Fig. S4H). These data point to impaired cognitive functions involving the hippocampus and other brain regions (29).

The Hargreaves test was used to investigate the reflex response to thermal sensation. Both WT and Polr3a-cKI mice showed increased latency for hind paw withdrawal as a function of stimulus intensity (Fig. 3 *C* and *D*). However, the relative response was significantly delayed for Polr3a-cKI mice at lower intensities for males and at all intensities for females, suggesting an impaired sensation and reflex response. Auditory and sensorimotor reflexes were investigated using an acoustic startle test. Male Polr3a-cKI mice exhibited a significant reduction in the magnitude of the startle reflex at all stimulus intensities, whereas the corresponding female mice exhibited a trend at lower stimuli and a significant reduction at 115 dB (Fig. 3 *G* and *H*). Given the differences observed for male and female mice at 115 dB, we examined the latency to respond at this stimulus intensity and found a significant delay, relative to WT, for both sexes to elicit a startle reflex (Fig. 3*I*). Together, these data indicate that Polr3a-cKI mice have impaired auditory and thermal sensorimotor reflexes.

To determine if the ataxic phenotype observed in patients is evident in the mice, balance beam and rotarod experiments were used to assess gross motor coordination and motor memory. The number of slips made by Polr3a-cKI mice on four balance beams of varying diameter was not statistically different from WT mice (Fig. 3 *J* and *K*). Similarly, Polr3a-cKI mice exhibited no difference in latency to fall from an accelerating rod compared to WT mice using protocols to assess short-term or long-term memory (SI Appendix, Fig. S4J and *K*). The lack of any difference in these tests was consistent in both sexes, suggesting an absence of gross motor coordination and motor memory dysfunction. In contrast, significant differences in fine sensorimotor function were revealed using a tape removal test. Polr3a-cKI mice showed a significant delay in active removal latency over 2 consecutive days (Fig. 3 *E* and *F*). The delay was present for both days and in both sexes, suggesting impaired fine sensorimotor function. Thus, along with impaired growth and hypomyelination, these data indicate that adult Polr3a-cKI mice exhibit deficits in cognitive, sensory, fine sensorimotor, auditory, and reflex functions compared to WT mice.

Impaired Myelin Integrity Revealed by MRI and DTI. Quantitative MRI is an important diagnostic tool for detecting and monitoring lesions, inflammation, and white matter changes in the CNS caused by traumatic injury and a wide range of neurodegenerative diseases (30). In patients with Pol III-related leukodystrophy, diffuse hypomyelination is typically identified by MRI as hyperintensity in T2-weighted images of white matter regions (4, 5, 7, 8). We performed brain MRIs on adult WT and Polr3a-cKI mice and determined T1 and T2 relaxation times for different white and gray matter regions (Table 1 and SI Appendix, Fig. S5A). In agreement with the immunohistochemical findings for adolescent and adult Polr3a-cKI mice, T2 relaxation times increased in the corpus callosum, consistent with hypomyelination. To further explore the myelination defect in the mice, we utilized DTI, a high-resolution methodology that provides microstructural information about white matter fibers based on the diffusion properties of water (31). Parallel bundles of axons promote water diffusion along their length (axial diffusivity, AD) and restrict it perpendicular to the tract (radial diffusivity, RD), resulting in high fractional anisotropy (FA). Interestingly, in Polr3a-cKI mice, FA was reduced significantly in the

anterior commissure, corpus callosum genu and splenium, and the optic tract due to increased RD in the absence of effects on AD (Table 2 and SI Appendix, Fig. S5 *B* and *C*). These data suggest a myelin deficiency in these regions, analogous to those reported in Shiverer mice, which are mutant for MBP (32). In contrast, the optic nerve showed no change in FA but reduced RD, AD, and mean diffusivity (MD) (Table 2). These changes recapitulate observations made in a mouse model of retinal ischemia that were interpreted to represent axonal degeneration and myelin fragmentation in the optic nerve (33). The DTI data also revealed high but unchanged FA in the fimbria, the pons, and structures of the cerebellum, with no changes in diffusivities (SI Appendix, Fig. S5 *C* and *D*), suggesting that myelin integrity in these regions is not altered. The absence of DTI changes in the cerebellum is consistent with the lack of immunohistochemical changes in this region and the absence of gross motor defects in the mice.

Ultrastructural Analysis Reveals Defects in Myelination. Transmission electron microscopy (TEM) was performed on the corpus callosum to directly examine axonal myelination in the mice. Images were collected at multiple magnifications (Fig. 4 *A–D*) for the quantitation of axon size, myelin thickness (i.e., g-ratio, axon radius/axon plus myelin radius), and the number of unmyelinated axons per unit volume. Myelin sheath thickness of Polr3a-cKI mice was substantially reduced compared to WT mice as indicated by the higher average g-ratio (Fig. 4*E*). Reduced myelin thickness was observed across a range of axon calibers, as seen from a linear fit of the g-ratios versus axon diameter (Fig. 4*G*). These results reveal a defect in the capacity of mature OLs to produce a myelin sheath of normal thickness in the mutant mice. Additionally, myelinated small and large diameter axons appeared to be underrepresented in Polr3a-cKI mice (Fig. 4*G*). To examine this further, we calculated the frequency distributions of myelinated axon diameters. The peak of the distribution was unchanged. However, myelinated axons in Polr3a-cKI mice showed a narrower size range (Fig. 4*H*). This was especially notable for large diameter axons (1.4 to 1.8 μ m), which were present in the WT but absent in the mutant. Interestingly, the frequency distributions of unmyelinated axons were markedly different (Fig. 4*I*): The peak of the distribution for Polr3a-cKI mice was broader and shifted to larger diameters compared to WT mice. The largest myelinated and unmyelinated axons in the mutant mice were comparable in size and considerably smaller than the largest WT myelinated axons. Consistent with these observations, the mutant mice exhibited a marked increase in the number of unmyelinated axons with cross-sectional areas greater than the smallest myelinated axons observed in WT mice (Fig. 4*F*). The increase in unmyelinated axons suggests that Polr3a-cKI mice may have fewer mature OLs and fewer myelinating processes per mature OL (34).

The origin of these changes in myelination seemed likely to involve cells of the OL lineage, given their ubiquitous expression of Olig2, which drives Cre expression in the mice. In addition, neurons derived from embryonic Olig2-expressing progenitors might also play a role (35). To examine this possibility, we analyzed confocal z-stacked images of the corpus callosum for colocalization of NeuN-stained neurons and EGFP fluorescence, which serves as a marker for Olig2-Cre recombination (SI Appendix, Fig. S6A). Of the 204 neurons examined, no colocalization was observed. Additionally, we note that Olig2-expressing GFAP⁺ astrocytes are not found in the corpus callosum (27). These observations support the view that the myelination defects in the corpus callosum reside in cells of the OL lineage.

We also examined TEM images of the dentate gyrus region of the hippocampus (SI Appendix, Fig. S6 *B* and *C*). Although variability in axon orientation relative to the sectioning plane limits the morphometric analysis, an examination of multiple images supports the qualitative conclusion that the number of myelinated

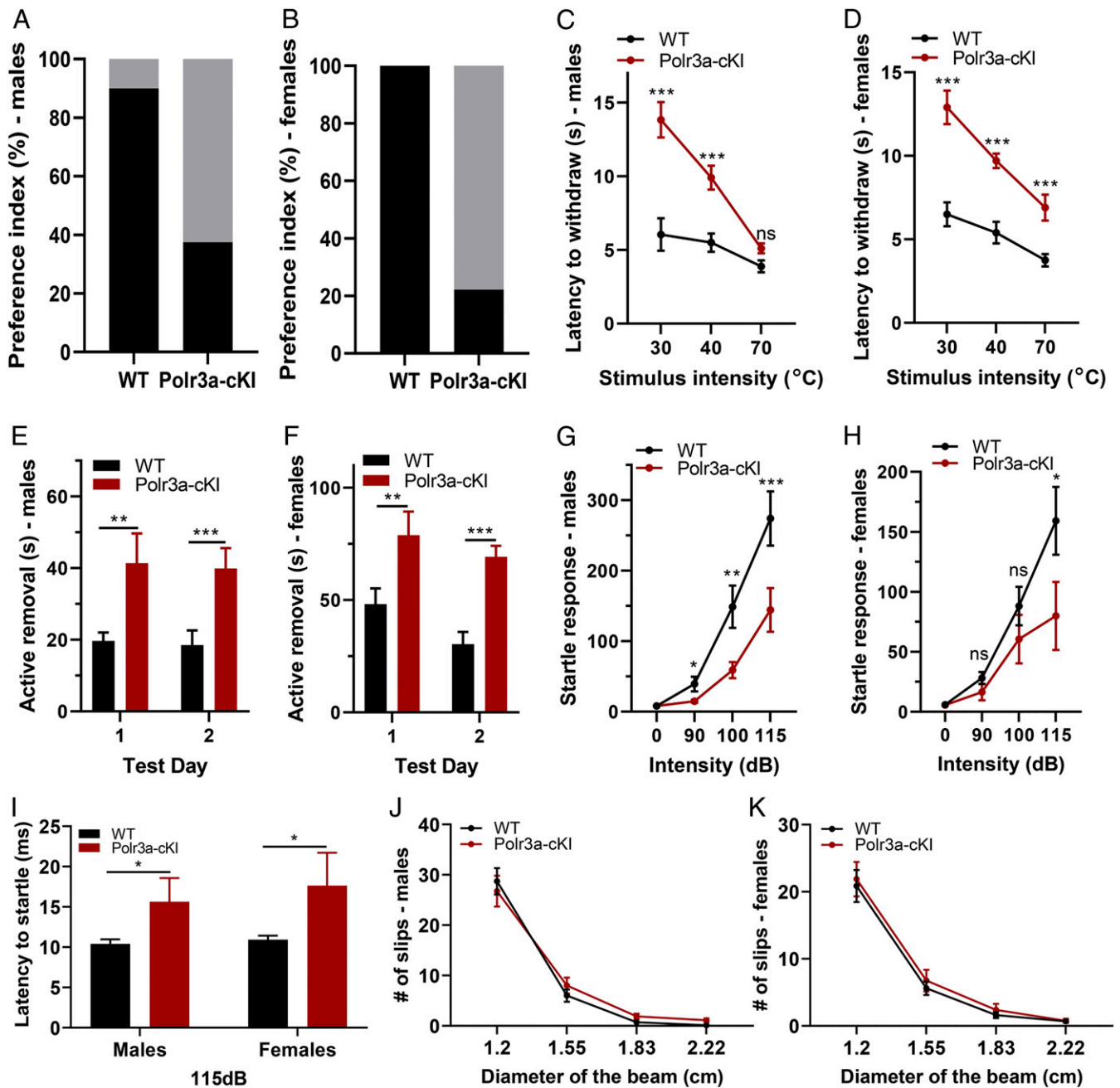


Fig. 3. Behavioral testing of adult WT and Polr3a-cKI mice reveals cognitive, sensory, and fine sensorimotor deficits. (A and B) Cognitive assessment of object placement for males and females. Data are plotted as the percentage of animals that achieve a preference score >55 (black bar, animals scoring above the cutoff) and those that fail to reach this cutoff (i.e., no preference) (gray bar, χ^2 test for both sexes $P < 0.001$). (C and D) Hargreaves test for males and females. The latency for hind paw withdrawal is plotted as a function of stimulus intensity (repeated-measures ANOVA, $*P < 0.05$, $**P < 0.01$, $***P < 0.001$). (E and F) Tape removal test for males and females. The time of active tape removal is reported for tests performed on consecutive days (one-way ANOVA, $**P < 0.01$, $***P < 0.001$). (G and H) Acoustic startle test for males and females. The startle reflex (millivolts per gram body weight) is reported as a function of stimulus intensity (repeated-measures ANOVA as for panels C and D). (I) Acoustic startle latency at a stimulus intensity of 115 dB (one-way ANOVA for each sex; both sexes $*P < 0.05$). (J and K) Balance beam reporting the number of slips from four different beams ranging from hard (1.2 cm) to easy (2.22 cm) for males and females, respectively (repeated-measures ANOVA $P > 0.85$). ns, not significant.

axons and myelin thickness is reduced in this region, consistent with the findings in the corpus callosum (Fig. 4).

Reduced Numbers of Myelinating OLs in Adolescent and Adult Polr3a-cKI Mice. The production of mature myelinating OLs from OL precursor cells (OPCs) begins at birth, reaches a peak during postnatal development (~P14 to P28) and then declines, but continues into adulthood where newly formed OLs play multiple important

roles, including neuronal activity-based myelination during task-learning (35). Ultrastructure analysis of Polr3a-cKI mice suggested that, in addition to the diminished function of myelinating OLs, myelination might be further limited by reductions in the number of mature OLs. To examine this possibility, we used flow cytometry to quantify OPCs and mature OLs in single-cell suspensions of dissociated whole brain (36). As a gating strategy, we utilized Alexa-Fluor 647-conjugated antibodies against O4, a marker of

Table 1. MRI T1 and T2 relaxation times of different brain regions

Structure	T1 (s)	T2 (ms)
Corpus callosum genu		
WT	1.84 ± 0.22	34.5 ± 2.1
Polr3a-cKI	1.88 ± 0.14	37.5 ± 2.7
<i>P</i>	0.72	0.038
Corpus callosum splenium		
WT	1.69 ± 0.06	34.7 ± 1.7
Polr3a-cKI	1.72 ± 0.06	38.7 ± 1.9
<i>P</i>	0.46	0.0014
Cortex		
WT	1.83 ± 0.03	43.9 ± 3.2
Polr3a-cKI	1.86 ± 0.03	45.7 ± 3.0
<i>P</i>	0.06	0.32
Hippocampus		
WT	1.86 ± 0.03	45.7 ± 2.8
Polr3a-cKI	1.87 ± 0.05	46.8 ± 2.7
<i>P</i>	0.71	0.44

All values are presented as the mean ± SD, combining data from males and females *n* = 7.

OL progenitors, and PE-Cy7-conjugated antibodies against MOG, a marker of terminally differentiated myelinating OLs (36). Additionally, we used mice containing the dual tdTomato-EGFP reporter and gated EGFP-expressing cells in Polr3a-cKI samples, so that only OL lineage cells that had undergone Cre-mediated recombination would be counted (*SI Appendix, Fig. S7*). Representative scatter plots of samples from adolescent female mice show the cell populations, including O4 and MOG double-positive premyelinating OLs (36) (Fig. 5A). From these data we calculated the fractional representation of the three OL populations (Fig. 5B–E). The results for adolescent and adult mice of both sexes show reduced numbers of MOG-only expressing OLs. In addition, there was a significant reduction in double-positive pre-OLs among adolescent male Polr3a-cKI mice and a similar trend among adolescent female mice that, in both cases, was absent in the adults. In keeping with the lower number of OLs

in Polr3a-cKI mice, this observation is consistent with a defect in oligodendrogenesis.

To confirm these findings, we performed Western blot analyses to assess changes in the levels of proteins (CNP, MOG, QKI, and MBP) that serve as markers for premyelinating and mature myelinating OLs (37). Protein extracts were prepared from microdissected regions of the cerebrum (cortex, corpus callosum, and thalamus) along with the cerebellum and spinal cord and the signals from the resulting blots were quantified and normalized to vinculin. Similar to the immunohistological findings for MBP (Fig. 2), Polr3a-cKI mice exhibited region-specific changes in myelin protein abundance. The CNP, MOG, and MBP signals were all reduced in the cortex and corpus callosum, while in the spinal cord, CNP, MBP, and QKI were lower, although the latter did not reach statistical significance (Fig. 6). In the cerebellum, the MOG signal was reduced. Otherwise, no significant changes were seen in this region or in the thalamus (*SI Appendix, Fig. S8*). Finally, since previous work has shown that apoptosis of premyelinating OLs can control OL maturation and myelination (37, 38), we looked for evidence of caspase 3 activation by Western blotting but found none (*SI Appendix, Fig. S8*). While this does not exclude a role for apoptosis in the hypomyelinating phenotype of Polr3a-cKI mice, the findings from Western analysis and flow cytometry remain consistent with the idea (discussed below) that a defect in oligodendrogenesis may reduce the number of mature myelinating OLs.

Discussion

Previous efforts to develop a mouse model of Pol III-related leukodystrophy have been unsuccessful, either because a known homozygous disease-causing *Polr3a* G672E mutation had no phenotype in mice or because homozygosity for one mutation (*Polr3b* R103H) from a compound heterozygous variant was embryonic-lethal (22, 23). We reasoned that enhancing the defect of the *Polr3a* G672E mutation might produce a viable disease model. However, homozygosity for the *Polr3a* W671R/G672E mutation, a construct informed by patient data and by structure–function studies in yeast (8, 24), resulted in embryonic lethality prior to E12.5. In other studies, early embryonic lethality between E3.5 and E6.5 has been documented for deletions of the Pol III transcription factor *Brfl* (39) and for *Polr3g*, a Pol III subunit isoform that is expressed in

Table 2. DTI of different brain regions

Structure	FA	MD	AD	RD
Anterior commissure				
WT	0.64 ± 0.04	0.64 ± 0.03	1.18 ± 0.06	0.36 ± 0.03
Polr3a-cKI	0.59 ± 0.04	0.66 ± 0.03	1.17 ± 0.05	0.41 ± 0.03
<i>P</i>	0.044	0.134	0.729	0.017
Optic nerve				
WT	0.88 ± 0.06	0.78 ± 0.10	1.79 ± 0.09	0.27 ± 0.11
Polr3a-cKI	0.89 ± 0.03	0.62 ± 0.03	1.54 ± 0.08	0.16 ± 0.04
<i>P</i>	0.848	0.043	0.0002	0.035
Corpus callosum genu				
WT	0.65 ± 0.03	0.75 ± 0.09	1.33 ± 0.08	0.42 ± 0.07
Polr3a-cKI	0.58 ± 0.05	0.76 ± 0.06	1.31 ± 0.05	0.48 ± 0.08
<i>P</i>	0.009	0.880	0.631	0.161
Corpus callosum splenium				
WT	0.77 ± 0.03	0.66 ± 0.03	1.39 ± 0.09	0.28 ± 0.02
Polr3a-cKI	0.70 ± 0.06	0.67 ± 0.01	1.32 ± 0.09	0.34 ± 0.06
<i>P</i>	0.024	0.352	0.137	0.017
Optic tract				
WT	0.78 ± 0.04	0.65 ± 0.07	1.34 ± 0.09	0.26 ± 0.02
Polr3a-cKI	0.73 ± 0.03	0.64 ± 0.02	1.30 ± 0.04	0.31 ± 0.03
<i>P</i>	0.0244	0.598	0.337	0.0071

Values for MD, AD, and RD are 10^{-5} cm²/s. All values are presented as the mean ± SD, combining data from males and females, *n* = 7.

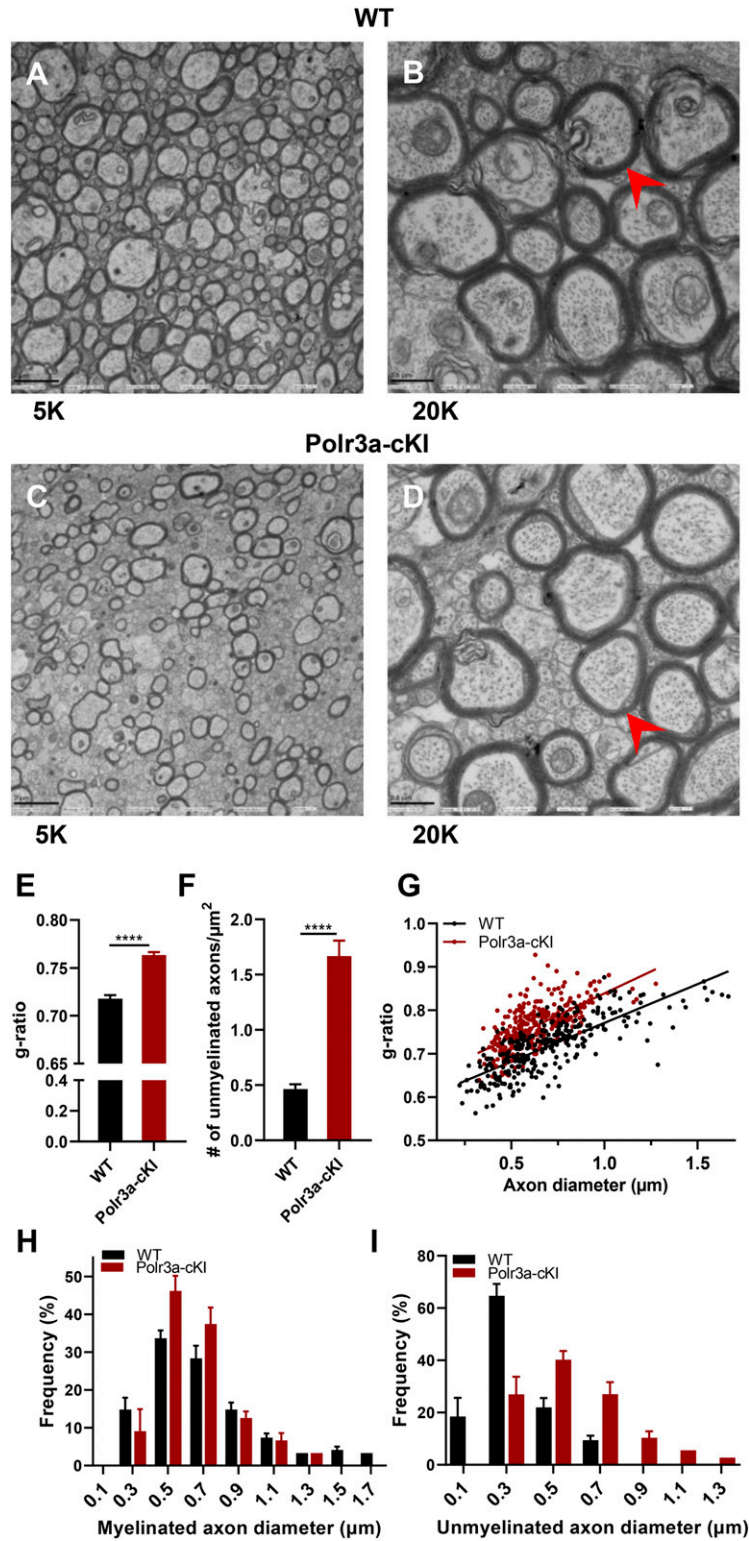


Fig. 4. TEM images of the corpus callosum in adult male mice. (A and B) Representative WT images at 5K and 20K magnification, respectively. (C and D) Representative Polr3a-cKI images at 5K and 20K, respectively. Arrowhead highlights differences in myelin thickness for axons of similar size. (E) Average g-ratio (radius of axon/radius of axon+myelin), $n = 300$ per group. (F) Number of unmyelinated axons/ μm^2 , $n = 10$ fields per group. An axon is counted as unmyelinated if the area of the axon is equal to or larger than the smallest myelinated axon in the corresponding WT field. (**** $P < 0.0001$) (G) g-ratio as a function of axon diameter, $n = 300$ per group. The data show a uniformly higher g-ratio for Polr3a-cKI mice based on the similar slopes of the linear regressions and the different y intercepts. (H) Frequency distribution of myelinated axon diameters, $n = 300$ per group. (I) Frequency distribution of unmyelinated axon diameters, $n = 10$ fields per group. Quantitation was performed on 10 independent images/group at 10K magnification. Values are presented as the mean \pm SEM. Groups were compared using two-sided Student's t test unless otherwise indicated, **** $P < 0.0001$.

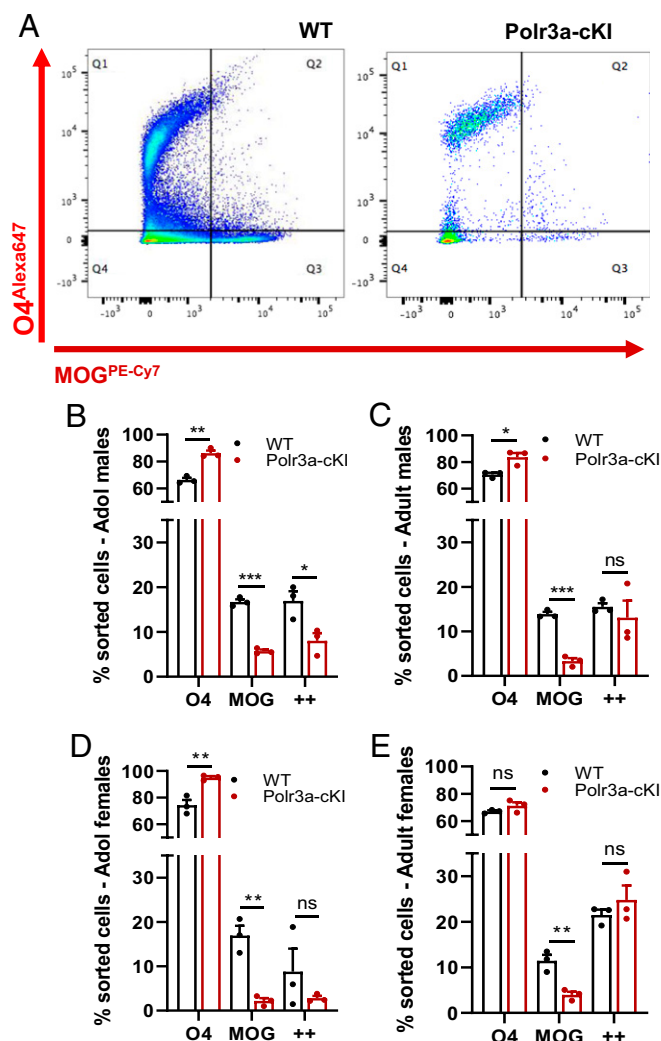


Fig. 5. Flow cytometry of OL cell populations from WT and *Polr3a*-cKI mice. (A) Representative scatter plots from flow cytometry show gating of O4, MOG, and double-positive OLs in whole brain from adolescent female mice. (B and D) Quantitation of O4, MOG and double-positive (++) OLs in adolescent male and female samples. (C and E) Quantitation of O4, MOG, and double-positive (++) OLs in adult male and female samples. EGFP-expressing cells in *Polr3a*-cKI samples were gated so that only OL lineage cells that had undergone Cre-mediated recombination would be counted. Values are presented as the mean \pm SEM. Groups of $n = 3$ mice per condition/sex were compared using multiple *t* tests, * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$.

mouse embryonic stem cells at a level sixfold higher than its *Polr3gl* paralog (40). This implies that viability during embryonic development requires $>15\%$ of WT Pol III activity in the critical period of rapid growth prior to gastrulation. Consistent with this, biochemical studies on a yeast Pol III mutant corresponding to *Polr3a* W671R/G672E show that the mutations reduce Pol III activity to approximately one-third of the WT level (24). Conversely, mice that are homozygous or hemizygous for the *Polr3a* G672E mutation and thus are likely to have no more than 50% of WT Pol III function, appear to be normal up to at least 12 mo of age (22). Accordingly, there appears to be a narrow functional window in mice versus humans for *Polr3a* mutations to both support embryogenesis and generate leukodystrophy phenotypes.

In mice, the production of OPCs begins around E12.5, with the first myelin-expressing OLs appearing just before birth (35). Given the timing of oligodendrogenesis relative to the rapid early cell divisions that precede gastrulation, we hypothesized that a conditional

knockin of the *Polr3a* W671R/G672E mutation in OLs might support viability and exhibit disease phenotypes. Indeed, *Polr3a*-cKI mice display multiple phenotypes similar to those seen in Pol III-related leukodystrophy patients. The developmental delay in neonates and the growth rate reduction that continues through adulthood are notable examples (8). The existence of a growth rate phenotype in a Pol III hypomorph that is restricted to the Olig2 lineage is intriguing and suggests a cell-nonautonomous effect. Interestingly, the short stature of many Pol III-leukodystrophy patients correlates with a growth hormone deficiency in a small number of patients evaluated by a growth hormone stimulation test (8, 41). How this reduction in growth hormone might be achieved is not known. However, the patient data together with our findings may indicate a role for OLs or other Olig2-expressing cells in the regulation of growth hormone production by the pituitary gland and control of the hypothalamic-pituitary axis. Such effects could also account for the observed hypogonadotropic hypogonadism of Pol III-related leukodystrophy patients (42).

Polr3a-cKI mice exhibit impairments in cognition, auditory and thermal sensorimotor reflexes, and fine sensorimotor function, consistent with hypomyelination in multiple regions of the cerebrum (hippocampus, cortex, corpus callosum) and the spinal cord (43). Diffuse hypomyelination, assessed by MRI, is also a common phenotype of Pol III-related leukodystrophy patients and typically includes regions in the cerebrum, notably the corpus callosum, as well as the cerebellum, where it is often associated with cerebellar atrophy and gross motor defects (8). Interestingly, even though Olig2 is expressed throughout the OL lineage, *Polr3a*-cKI mice showed no signs of hypomyelination in the cerebellum by histology, DTI, or Western blotting and did not exhibit a gross motor defect or ataxia. The basis for this difference in hypomyelination between the cerebellum and other CNS regions is unknown. However, it is well-known that the survival and differentiation of OPCs, along with the function of OLs and the maintenance of myelin, is dependent on the actions of neighboring astrocytes and microglia and on the ensheathed axons (2). These observations raise the possibility that cell-intrinsic changes in OLs may be suppressed in our model by cell-extrinsic factors. A possible mechanism is the ability of astrocyte-derived lipids to supplement lipid synthesis in OLs (44), although it is unclear how the cerebellar selectivity would be achieved in this case. Alternatively, functional heterogeneity within the OL population may play a role (45). Whatever the mechanism, targeting the *Polr3a* mutation to additional cell types in the CNS may provoke hypomyelination in the cerebellum, may exacerbate it in other regions and is likely to produce more severe disease phenotypes (2).

Our experiments suggest that pathogenic *Polr3a* mutations expressed under *Olig2*-Cre control cause distinct defects that contribute in different ways to changes in myelination. One defect reduces the number of mature myelinating OLs, while the other diminishes the capacity of mature OLs for myelin deposition (i.e., reduces myelin thickness). These changes likely reside in cells of the OL lineage, although neurons derived from Olig2-expressing progenitors and Olig2⁺, GFAP⁻ astrocytes may also make phenotypic contributions in some regions of the CNS. However, these cell populations appear to be absent in the corpus callosum where the hypomyelination phenotype is relatively robust. Additionally, we note that Olig2-expressing Purkinje neurons in the cerebellum of *Polr3a*-cKI mice do not appear to negatively affect myelination in this region.

Genetic labeling of myelinating OLs in mice has shown these cells to have an exceptionally long lifespan, with 90 to 100% survival in all CNS regions, except for the optic nerve, over the first 8 mo of life (46). Thus, the low and relatively constant number of mature (MOG⁺) OLs in adolescent and adult *Polr3a*-cKI mice, determined by flow cytometry, is unlikely due to the reduced viability of this cell population and more readily explained by a defect in oligodendrogenesis. Lineage tracing of OPCs in the cerebral cortex of

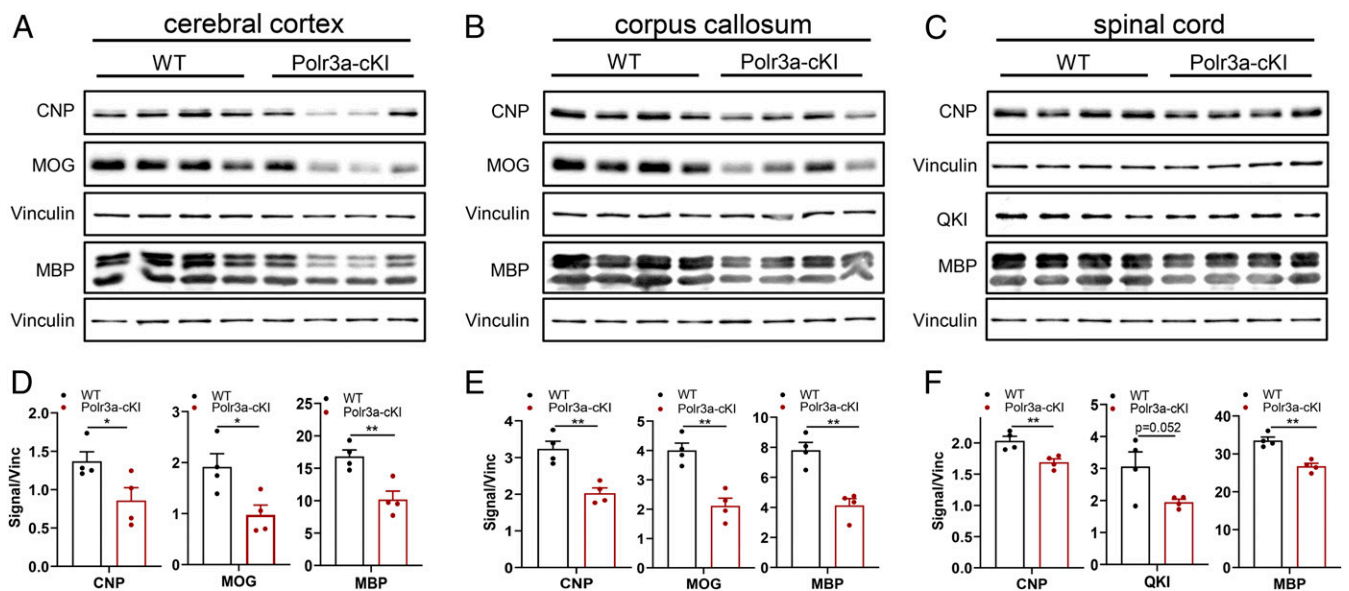


Fig. 6. Western blot analysis of the cortex, corpus callosum, and spinal cord extracts from adolescent male mice. (A) Extracts of dissected cerebral cortex were blotted for MOG, CNP, MBP, and vinculin. (B) Extracts of dissected corpus callosum were blotted as in A. (C) Spinal cord extracts were blotted for CNP, QKI, MBP, and vinculin. For A–C, vinculin blots were used for normalization and are shown below the corresponding myelin-specific proteins. (D–F) Myelin-specific protein signals from A–C, respectively, were quantified and normalized to the corresponding vinculin signals. Values are presented as the mean \pm SEM. Groups of $n = 4$ mice were compared using multiple t tests, * $P < 0.05$, ** $P < 0.01$.

young mice has shown that differentiation is followed by substantial (~80%) cell death prior to commitment to myelination (47). Limited survival of uncommitted OL precursors has also been reported during early postnatal development (48). These observations suggest that decreased levels of Pol III transcription may affect differentiation and survival functions during oligodendrogenesis to reduce the number of mature OLs in *Polr3a-cKI* mice. Currently, there is little knowledge about causal roles for Pol III transcription as an effector of cell differentiation or survival under conditions that are independent of other perturbations (e.g., cell transformation, DNA damage, viral infection) (13). One notable exception is the finding that genetic or chemical inhibition of Pol III transcription can promote adipogenesis in primary cells cultured from the stromal vascular fraction of inguinal fat pads (49). Conversely, increased Pol III transcription in mouse embryonic stem cells, achieved by knockdown of its master negative regulator MAF1, was shown to inhibit formation of the mesodermal germ layer and the terminal differentiation of mesoderm into adipocytes (49). Based on this work, further studies of *Polr3a-cKI* mice are needed to assess the impact of decreasing Pol III transcription on oligodendrogenesis.

In summary, we describe an animal model of Pol III-related leukodystrophy that demonstrates clinical features of the disease. The complexity of the disease and the ability of Pol III mutations to broadly impact cellular function is reflected by the identification of defects that reduce both the number and the function of myelinating OLs. Further studies of the mice will enhance our molecular understanding of disease pathogenesis and enable testing of genetic and pharmacological approaches aimed at mitigating the consequences of Pol III-related leukodystrophy.

Materials and Methods

Animals. All experiments involving mice were performed using protocols approved by the Institutional Animal Care and Use Committee of the Albert Einstein College of Medicine. Mice were housed under standard laboratory conditions on a 12-h light/dark cycle with constant access to food and water. The C57BL/6J-*Polr3a^{tm1wvl}* mouse line was generated by homologous recombination in C57BL/6J blastocysts (Ozgene Pty. Ltd.). This conditional *Polr3a* knockin mouse (*Polr3a^{cKI/+}*) carries an insertion of a *Polr3a* cDNA encoding

exons 15 to 31 and a downstream heterologous polyadenylation signal flanked by *loxP* sites followed by intronic sequences and a mutant exon 15 encoding the leukodystrophy mutations W671R and G672E. Cre-mediated recombination removes the WT cDNA to allow expression of the *Polr3a* mutant. Crosses of this mouse to one expressing a germline Cre recombinase were performed to generate mice heterozygous for the whole-body *Polr3a* knockin (*Polr3a^{KI/+}*). B6.129-*Olig2^{tm1.1(cre)Wdr/J}* (*Olig2Cre*, stock no. 025567) (50) mice and B6.129(Cg)-*Gt(ROSA)26Sor^{tm4(ACTB-tTomato,EGFP)Lox/J}* (tdTomato-EGFP, stock no. 007676) (51) mice were purchased from the Jackson Laboratory. Conditional *Polr3a* mutant mice and WT controls were obtained by breeding *Olig2Cre* male mice heterozygous for the floxed *Polr3a* allele to homozygous floxed *Polr3a* females. All experiments were performed with male and female mice except TEM and Western blots, which were performed using males.

Developmental Milestones. The protocol from Hill et al. (28), was followed with two investigators participating in the execution and documentation of the tests (*SI Appendix*). Briefly, prior to daily testing, pups were removed from the home cage and placed with a small amount of nesting material on a heating pad set at 37 °C. On P2, each mouse pup was weighed and given a unique tail tattoo mark. Tests were repeated daily until completed successfully on 2 consecutive days. The first day of passing each test is reported.

Western Analysis. All tissues were snap-frozen in liquid nitrogen or frozen on dry ice immediately after harvesting. For POLR3A detection, whole-brain samples were homogenized in RIPA lysis buffer supplemented with protease inhibitors (cOmplete mini + EDTA, Roche, 1 mM PMSF, 10 μ g/mL leupeptin) and phosphatase inhibitors (10 mM sodium fluoride, 20 mM glycerophosphate- β , 2 mM sodium orthovanadate and 25 mM sodium pyrophosphate). Proteins were quantified by a bicinchoninic acid assay (Pierce). Protein (50 μ g) was resolved by SDS/PAGE (6%) and transferred to nitrocellulose in 1 \times Tris-glycine buffer with 20% methanol. POLR3A and vinculin were detected by ECL (LAS-4000, GE Healthcare) and quantified using ImageQuant software. Extracts from cerebrum, cerebellum, and spinal cord were prepared by homogenization in RIPA buffer with protease inhibitors, as above and phosphatase inhibitors (mixtures 2 and 3; Sigma). Proteins were quantified by Bradford assay (Bio-Rad). Protein (20 μ g) was resolved by SDS/PAGE (10% gels for CNP, MOG, and caspase-3 and 12% gels for QKI and MBP) and transferred to nitrocellulose using 1 \times Tris-glycine 0.1% SDS buffer with 20% methanol using a wet transfer apparatus. Proteins were detected by ECL and quantified as described above. Antibodies and dilutions were: POLR3A (Abcam 96328; 1:750), vinculin (Santa Cruz 73614; 1:2,000), CNP (EMD MAB326R, 1:1,000), MOG (EMB MAB5680; 1:1,000), CC1 which recognizes the QKI protein (Abcam 16794; 1:1,000), caspase-3

(Cell Signaling 9662; 1:1,000), cleaved caspase-3 (Cell Signaling 9664; 1:1,000), and MBP (Abcam 7349; 1:1,000).

RNA Analysis. Tissue samples (50 to 100 mg, flash-frozen in liquid N₂) were homogenized into TRIzol lysis reagent (Thermo Fisher), and RNA was purified according to the manufacturer's directions. RNA was reprecipitated, quantified, and resolved by denaturing polyacrylamide electrophoresis before electrophoretic transfer to Nytran Plus membranes (GE Healthcare) and hybridization with [³²P]-end-labeled oligonucleotide probes at 42 °C or direct staining. Precursor tRNA^{Leu}(TAT) transcripts detected by phosphorimaging were quantified and normalized to U3 snRNA (52). Mature tRNAs visualized with ethidium bromide were quantified and normalized to 5.8S rRNA.

Behavioral Studies. Behavioral studies were carried out by blinded operators on two separate cohorts of mice at 12 wk of age. The tests included video monitoring and analysis in a Behavioral Spectrometer (BIOBSERVE), object placement, and object recognition tests, Hargreaves, acoustic startle and tape removal tests, elevated plus maze, balance beam, and rotarod (*SI Appendix*).

Immunohistochemistry. Brain and spinal cord were harvested after isoflurane anesthesia and cardiac perfusion (4% paraformaldehyde, PFA) and were stored at 4 °C in 4% PFA overnight before long-term storage in 25% sucrose. Frozen tissue was sectioned into 10- μ m slices using a cryostat and stained with the BrainStain imaging kit (Invitrogen B3645) according to the manufacturer's protocol. Paraffin-embedded tissue was sectioned into 15- μ m slices for staining with LFB and Nissl, which followed standard procedures. Purkinje cell counts were performed as previously reported (53). For MBP staining, 5- μ m sections were deparaffinized before heat-mediated antigen retrieval in citrate buffer. Blocking, primary, and secondary antibody steps used 0.1% Triton X-100, 5% goat serum, 5% BSA in 1 \times TBS. MBP primary antibody (SMI99 EMD NE1019; 1:1,000) was detected with anti-mouse Alexa-Fluor 488 (1:1,000 dilution). Slides were mounted with ProLong diamond antifade with DAPI (Invitrogen P36966) and images were acquired using a 3D Hitec p250 high capacity slide scanner. For EGFP and NeuN staining, 20- μ m sections were fixed in 4% PFA for 10 min at room temperature and rinsed in 1 \times PBS 3 \times 5 min before permeabilization and blocking in 0.25% Triton X-100, 5% goat serum, 5% BSA in 1 \times PBS for 2 h at room temperature with mouse blocking reagent (Vector MKB-2213, 1 drop/mL). Slides were incubated overnight at 4 °C with primary antibodies, GFP (Novus NB1001770, 1:100) and NeuN (EMD Millipore MAB377, 1:150) in 0.1% Triton X-100, 5% goat serum, 5% BSA in 1 \times PBS, washed with PBS 3 \times 5 min each, and further incubated with anti-goat Alexa-Fluor 488 (Invitrogen A32814; 1:500) and anti-mouse Alexa-Fluor 647 (Invitrogen A31571; 1:500). Images were acquired using a Leica SP8 inverted DMI8 confocal microscope (40 \times N.A.1.3).

Immunofluorescence Image Analysis and Quantification. For the measurement of MBP fluorescence intensity, micrographs were first converted to TIF images in CaseViewer and uploaded to ImageJ. The area of interest was selected on the TIF image and the mean fluorescence intensity value was directly measured by the ImageJ "Measure" tool, using the readout of "mean gray value." Mean intensity values were recorded for adjacent brain sections from multiple animals (at least three mice per genotype and sex) to calculate the average MBP fluorescence intensity for each brain region. For the analysis the NeuN and EGFP signals, the LIF micrograph was first extracted and converted in FIJI. A composite image was generated and then split into the individual channels. The threshold for the green and red channels was set uniformly across all sections according to signal intensity. Individual neurons were counted in the midline of the corpus callosum and colocalization with EGFP was interrogated in the merged image throughout the z-stack and confirmed in the individual split channels. Neurons were examined and counted in two to four adjacent brain sections from two Polr3a-*ckl* animals that express the tdTomato-EGFP dual reporter.

Transmission Electron Microscopy. Brains, harvested after anesthesia and sequential cardiac perfusion with 15 mL per mouse of 0.09% NaCl containing heparin (25 mg/mL) and 4% PFA in 0.1 M cacodylate buffer, were postfixed in 2% PFA 2.5% glutaraldehyde in 0.1 M cacodylate buffer for 30 min at room temperature. A brain matrix was used to dissect the region encompassing the midline of the corpus callosum. Next, 1-mm sections were oriented to generate cross-sections of axons within the corpus callosum. Ultrathin sections were cut and stained with Toluidine blue. A minimum of 10 images that showed axons cut in cross-section were obtained at 5K, 10K and 20K magnification for quantitation. Images were analyzed by Photoshop software using the free hand selection tool and manually thresholded to determine the number of myelinated axons, axon cross-sectional area, axon diameter (caliber), and g-ratios. Three-hundred axons per mouse were analyzed from 10 nonserial sections from 10K images. An axon was counted as unmyelinated if the area of the axon was equal to or larger than the area of the smallest myelinated axon in the corresponding field. Images were acquired on a JEOL 1200 EX transmission electron microscope.

Flow Cytometry. Following cardiac perfusion with ice-cold 1 \times PBS, the brain was harvested and cut into small pieces in a Petri dish filled with PBS on ice. An adult brain dissociation kit (Miltenyi 130-107-67) was used to obtain a single-cell suspension per the manufacturer's protocol. Briefly, tissues were dissociated mechanically and enzymatically followed by debris and red blood cell removal steps. The cell suspension was passed through a 0.35- μ m filter before staining with Trypan blue to assess cell viability and determine cell counts. The suspension was centrifuged at 300 \times g for 10 min at 4 °C and resuspended (up to 10⁷ cells) in 900 μ L of FACS buffer (ice-cold 1 \times PBS). The suspension was blocked with 100 μ L of FcR blocking reagent (Miltenyi 130-092-575) for 10 min on ice. Primary conjugated antibodies were diluted according to the manufacturer's protocol; O4 Alexa-Fluor 647 (RND FAB1326R; 6 μ L/10⁶ cells), mGalC-FITC (EMD FCMA312F; 10 μ L/10⁶ cells), MOG (EMD MAB5680; 10 μ L/10⁶ cells) conjugated with a lightning-link PE-Cy7 tandem conjugation kit per the manufacturer's protocol (Novus 7620005). Primary antibodies were incubated on ice for 20 min in the dark, washed with 2 mL of FACS buffer, centrifuged at 350 \times g for 7 min at 4 °C. The pellet was resuspended in 1 mL of FACS buffer and analysis was performed on a LSRII-U flow cytometer (BD Biosciences). Single-color controls were used to determine proper gating prior to sample analysis.

MRI/Proton MRS (¹H MRS). All MRI and ¹H MRS data were acquired in a 9.4 T Varian Direct Drive system (Agilent Technologies). A 14-mm diameter receive-only surface RF coil (Doty Scientific) along with a 7-cm ID ¹H transmit and receive body coil (M2M Imaging) was used. Anesthesia was achieved with 1.25% isoflurane mixed with room air. Respiration was monitored with a pressure pad (SA Instruments). Temperature was maintained at ~37 °C using warm air with feedback from a thermocouple placed in the rectum (SA Instruments). Imaging parameters and details of the analyzes are provided in *SI Appendix*.

Data Availability. All study data are included in the article and *SI Appendix*.

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