

GPR56/ADGRG1 regulates development and maintenance of peripheral myelin

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Myelin is a multilamellar sheath generated by specialized glia called Schwann cells (SCs) in the peripheral nervous system (PNS), which serves to protect and insulate axons for rapid neuronal signaling. In zebrafish and rodent models, we identify GPR56/ADGRG1 as a conserved regulator of PNS development and health. We demonstrate that, during SC development, GPR56-dependent RhoA signaling promotes timely radial sorting of axons. In the mature PNS, GPR56 is localized to distinct SC cytoplasmic domains, is required to establish proper myelin thickness, and facilitates organization of the myelin sheath. Furthermore, we define plectin—a scaffolding protein previously linked to SC domain organization, myelin maintenance, and a series of disorders termed “plectinopathies”—as a novel interacting partner of GPR56. Finally, we show that *Gpr56* mutants develop progressive neuropathy-like symptoms, suggesting an underlying mechanism for peripheral defects in some human patients with *GPR56* mutations. In sum, we define *Gpr56* as a new regulator in the development and maintenance of peripheral myelin.

INTRODUCTION

Myelin, the lipid-rich insulation that ensures rapid nerve conduction velocity, is generated by specialized glial cells: oligodendrocytes in the central nervous system (CNS) and Schwann cells (SCs) in the peripheral nervous system (PNS; Nave and Trapp, 2008). Disruption of the myelin sheath in patients with peripheral neuropathies (e.g., Charcot-Marie-Tooth disease [CMT]) causes severe symptoms, including pain and neuronal damage that can ultimately lead to paralysis (Harel and Lupski, 2014). Proper myelination is required for a high quality of life in humans, yet the molecular mechanisms governing myelinating glial cell development and myelination are poorly understood.

Recent works from our laboratory and others have identified the adhesion class of G protein-coupled receptors (aGPCRs) as key regulators of myelinating glial cell development (Monk et al., 2009, 2011; Glenn and Talbot,

2013; Mogha et al., 2013; Shin et al., 2013a; Liebscher et al., 2014; Ackerman et al., 2015; Giera et al., 2015; Petersen et al., 2015; Küffer et al., 2016; Salzman et al., 2016). aGPCRs represent the second-largest class of GPCRs in humans, yet remain largely understudied because of their genomic and structural complexity (Hamann et al., 2015). In brief, mature aGPCRs exist at the cell membrane as bipartite receptors consisting of an N-terminal fragment (NTF) facilitating adhesion and a C-terminal fragment (CTF) containing the seven-transmembrane pass signaling domain, and modulation of NTF–CTF interactions is associated with receptor activation (Hamann et al., 2015). Using a transcription activator–like effector nuclease-induced mutant allele of the aGPCR *Gpr56/Adgrg1* in zebrafish (*gpr56^{stl13/stl13}*), we previously determined that *Gpr56* regulates proliferation of oligodendrocyte precursor cells (Ackerman et al., 2015). Analysis of published expression datasets revealed that *Gpr56* is also robustly expressed in SCs (Miller et al., 2006; Weiss et al., 2016). Given the importance of *Gpr126* (a closely related aGPCR) in regulating SC biology (Monk et al., 2009, 2011; Glenn

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and Talbot, 2013; Mogha et al., 2013; Liebscher et al., 2014; Petersen et al., 2015; Küffer et al., 2016) and the established role of GPR56 in oligodendrocyte precursor cells (Ackerman et al., 2015; Giera et al., 2015; Salzman et al., 2016), we sought to determine if GPR56 also regulates myelinating glial development in the PNS.

Here, we use zebrafish and rodent models to define the role of *Gpr56* in SC development and myelination. Whereas loss of *Gpr56* from the CNS results in hypomyelination because of a reduction in oligodendrocyte precursor cell proliferation and mature oligodendrocyte numbers, we find no apparent defect in SC proliferation or overall SC number in *Gpr56* mutants. Instead, we demonstrate that GPR56 functions through RhoA in early SC development to regulate proper radial sorting of axons by SCs and that disrupted radial sorting in *Gpr56* mutants likely results from altered cytoskeletal remodeling. Additionally, we find that loss of *Gpr56* from mature nerve results in multiple myelin structural abnormalities, including increased myelin thickness, aberrant myelin domain organization, shortened internode length, and failed myelin maintenance. Finally, we identify plectin as a novel interacting partner of GPR56 in SCs. Plectin is a large cytoskeletal linker protein that causes multiple diseases (plectinopathies) when mutated in humans (Winter and Wiche, 2013). The most common plectinopathy is epidermolysis bullosa simplex with muscular dystrophy, a rare skin-blistering disorder that presents with late-onset muscular dystrophy, but mutations in *PLECTIN* have also been linked to defects in peripheral myelin ultrastructure (Bauer et al., 2001). We find that plectin physically interacts with GPR56, that plectin levels are up-regulated in the absence of *Gpr56* (as expected for a receptor-ligand pair), and that *Plectin* mutant SCs phenocopy *Gpr56* mutant SCs, suggesting that these proteins function together to regulate SC biology. Together, our studies define GPR56 as a novel and evolutionarily conserved regulator of peripheral nerve development and function with important implications for human health and disease.

RESULTS

GPR56 regulates SC radial sorting via

$G\alpha_{12/13}$ and RhoA signaling

Radial sorting is the process by which immature SCs encompass bundles of naked axons and sort individual axonal segments into a 1:1 SC/axon segment relationship. The timing of radial sorting is essential; when radial sorting is impaired, both myelination and peripheral nerve maturation are delayed (Jessen and Mirsky, 2005; Feltri et al., 2016). Analysis of peripheral nerves from *Gpr56* mouse mutants (Li et al., 2008) revealed radial sorting delays by transmission EM (TEM; Fig. 1). At postnatal day (P) 3 in *Gpr56* mouse mutant sciatic nerves (SNs; $n = 4$ mutant mice), the number of axons sorted 1:1 with SCs was significantly reduced relative to WT ($n = 3$) and heterozygous ($n = 3$) siblings (Fig. 1 G; $P < 0.0002$, one-way ANOVA). Additionally, we observed abnormal cytoplasmic protrusions from sorting SCs (indic-

ative of disrupted cytoskeletal rearrangements, Fig. 1, I and J), basal lamina trails reflecting aberrant SC process extension and retraction during radial sorting (Fig. 1 K), along with other myelin abnormalities in the SNs of *Gpr56* mutant mice (Fig. 1, L–N; $P < 0.002$, one-way ANOVA). Deficient SC precursor proliferation can cause radial sorting delays (Feltri et al., 2016), and GPR56 is a known regulator of cell division (Wu et al., 2013; Bae et al., 2014; Ackerman et al., 2015; Giera et al., 2015); thus, we first tested if *Gpr56* mutant SCs were less proliferative than WT. Interestingly, SC proliferation and total number were unaffected in *Gpr56* mouse mutant SNs, as was overall peripheral (saphenous) nerve size (Fig. S1). Therefore, in contrast to its role in oligodendrocyte precursor cells, GPR56 is dispensable for SC proliferation.

These findings suggest that GPR56 does not regulate radial sorting by affecting SC numbers. Given the cytoplasmic defects observed on P3 (Fig. 1, I and J), we hypothesized that GPR56 may function in controlling dynamics of immature SC protrusions and/or adhesion to axonal membranes. To test this, we used a modified Boyden chamber to mimic axon–SC interactions during radial sorting (Poitelon et al., 2015). In brief, purified rat SCs were seeded within the upper compartment of a Boyden chamber in which neuronal fragments were suspended below to induce SC “pseudopod” (mimicking cytoplasmic protrusion) growth. Pseudopods were then collected, processed, and analyzed by long-gradient nano reversed-phase liquid chromatography–mass spectrometry (LC–MS; Fig. 2, A and B; see Materials and methods for full details). Indeed, we determined that GPR56 is highly enriched in SC pseudopods (Fig. 2 C and Dataset 1), supporting the hypothesis that GPR56 autonomously regulates SC cytoplasmic protrusion dynamics during radial sorting.

In the nervous system, GPR56 primarily signals through $G\alpha_{12/13}$ and RhoA (Iguchi et al., 2008; Luo et al., 2011; Ackerman et al., 2015; Giera et al., 2015), and indeed, $G\alpha_{13}$ (protein symbol GNA13) and RhoA signaling proteins are also enriched in SC pseudopods (Fig. 2, C and D; and Dataset 1). Further, we found a significant reduction in the relative concentration of active, GTP-bound RhoA in *Gpr56* mutant nerves relative to WT sibling nerves on P6. We also detected a significant decrease in active, GTP-bound RhoA in heterozygous nerves relative to WT (Fig. 2 E, $n = 12$ nerves per genotype across three replicates), consistent with the observation that *Gpr56*^{+/-} animals exhibit some mutant phenotypes on P3 (Fig. 1 H; $P < 0.039$, one-way ANOVA). To test whether GPR56 functions through $G\alpha_{12/13}$ and RhoA to promote radial sorting, we used standard genetic manipulations to either enhance (via inhibition of $G\alpha_{12/13}$) or suppress (via transient overexpression of constitutively active RhoA) radial sorting defects in *gpr56*^{sd13/sd13} mutant zebrafish (Lin et al., 2009; Ackerman et al., 2015). Strikingly, injection of antisense morpholinos that block translation of the three zebrafish transcripts encoding $G\alpha_{12/13}$ (3MO) strongly enhanced radial sorting defects in the posterior lateral line nerve (pLLn) of *gpr56*^{sd13/sd13} mutants and induced radial sorting defects in

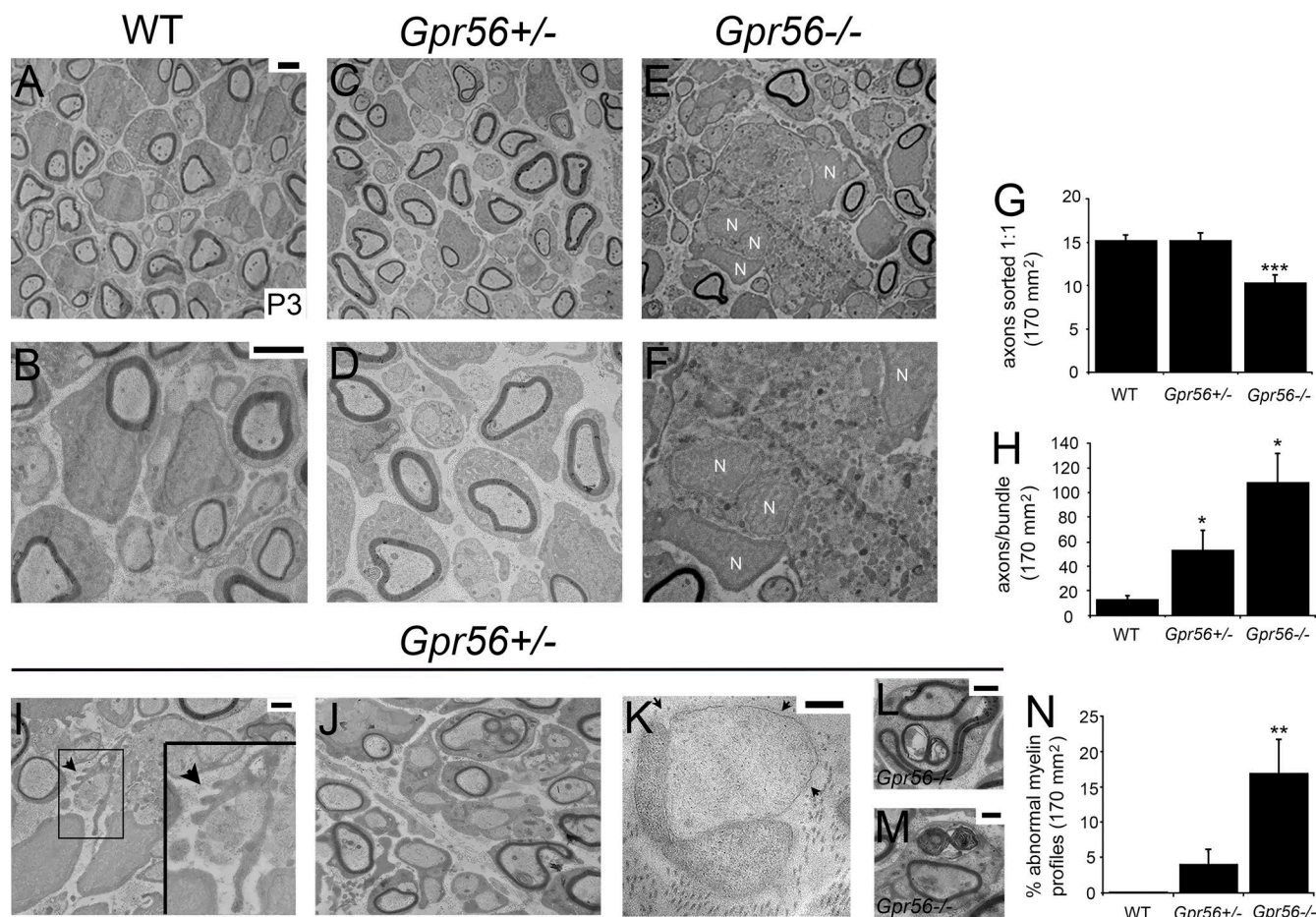


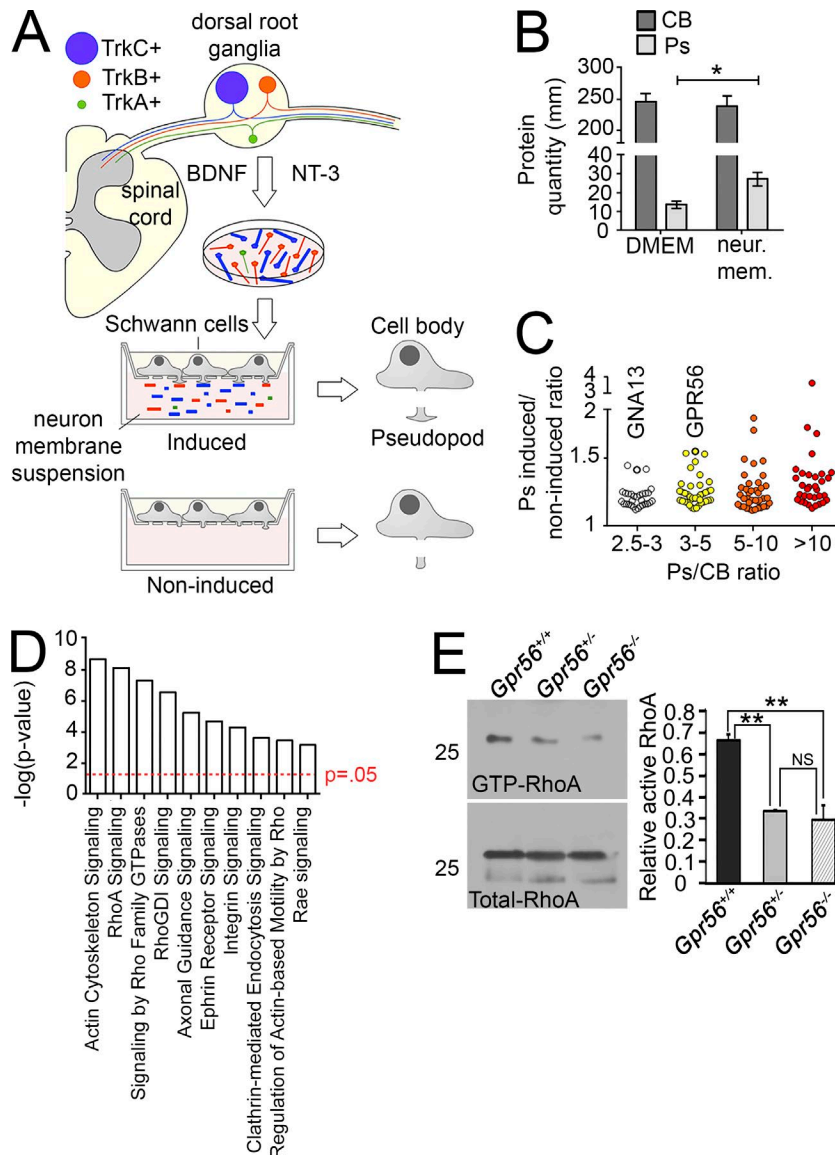
Figure 1. GPR56 is required for efficient radial sorting. (A–F) TEM images of SN from (A and B) WT ($n = 3$), (C and D) *Gpr56*^{+/-} ($n = 3$), and (E and F) *Gpr56*^{-/-} mice ($n = 4$) on P3. SC precursor nuclei (N) surrounding and within axon bundles in *Gpr56*^{-/-} nerves. Higher magnification images: B, D, and F. Bars, 2 μ m. (G) Quantification of axons sorted 1:1 with SCs (***, $P < 0.0002$, one-way ANOVA). (H) Quantification of the number of axons per unsorted bundle (*, $P < 0.04$, one-way ANOVA). (I–M) TEM images of P3 SN *Gpr56*^{-/-} ($n = 4$) animals. Bars: (I, J, L, and M) 1 μ m; (I inset and K) 500 nm. (I) Large bundles of unsorted axons surrounded by SC processes are observed in a *Gpr56*^{-/-} SN. Some SCs also exhibited abnormal cytoplasmic protrusions (arrowhead). Higher magnification of the boxed in region shown within the inset. In addition, (J, L, and M) myelin abnormalities (outfoldings) and (K) nearly naked axons associated with SCs and partially surrounded by SC basal lamina (arrows) were often observed. (N) Quantification of the percent of axons with myelin abnormalities revealed a significant increase in myelin defects in *Gpr56*^{-/-} nerves compared with WT controls (**, $P < 0.002$, one-way ANOVA). (G, H, and N) Error bars represent SEM.

heterozygote and WT siblings in a dose-dependent manner ($P < 0.001$, two-way ANOVA; Fig. 3, A–I and M). Conversely, transient overexpression of synthetic mRNA encoding constitutively active RhoA (*rhov14*) completely suppressed radial sorting defects in *gpr56*^{stl13/stl13} mutants, suggesting a positive role for RhoA in radial sorting via Gpr56 signaling (control WT vs. control *gpr56*^{stl13/stl13}; $P < 0.023$, unpaired Student's *t* test; *rhov14* injected WT vs. *rhov14* injected *gpr56*^{stl13/stl13}; $P < 0.58$, unpaired Student's *t* test; Phenol red: $n = 3$; WT: $n = 3$ *stl13*^{+/-}, $n = 4$ *stl13*^{-/-}; +*rhov14*: $n = 3$ animals per genotype; Fig. 3, J–L, O, and P). Total axon number was not affected by experimental conditions throughout (Fig. 3, N and Q–R). The importance of RhoA in maintenance of cell–cell contact points has been established (Parsons et al.,

2010), suggesting that sorting delays in *Gpr56* mutants may not result from impaired process extension but rather from an inability of SCs to maintain secure contacts with target axons during radial sorting. This hypothesis is consistent with the aforementioned signs of SC process retraction observed in P3 mouse mutant SNs (Fig. 1 K).

Myelin domain organization is impaired in *Gpr56* mutants

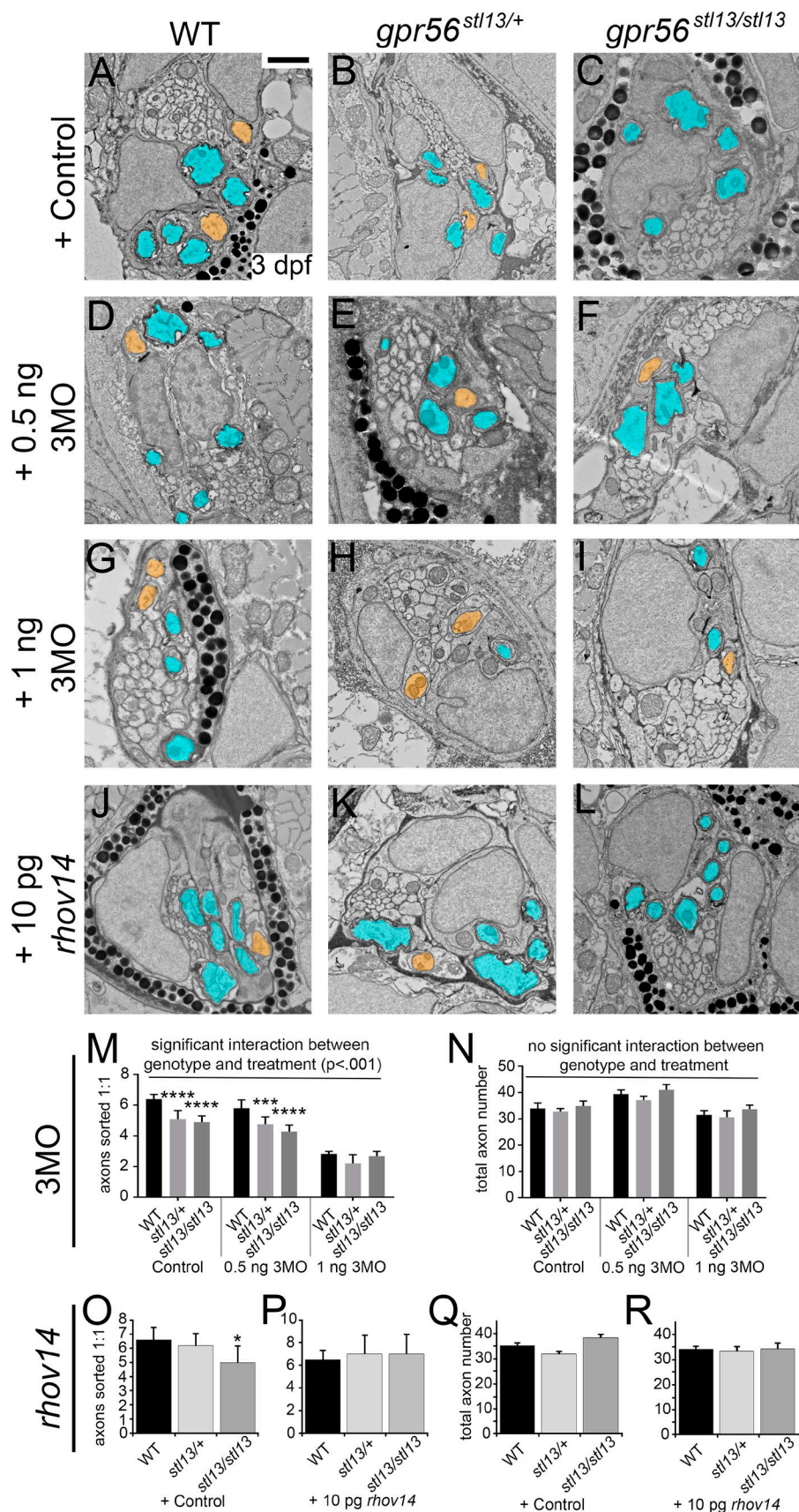
We've defined that GPR56 signaling through RhoA is required for timely radial sorting of axons during peripheral nerve development. Although this phenotype did not persist through adulthood (Fig. 4, A–D; $n = 3$ animals per genotype, $P < 0.57$, unpaired Student's *t* test; Fig. S2, A–D), we noted several other pronounced alterations in mature periph-



eral nerve. First, we observed Remak bundle defects. Remak bundles are groups of small caliber axons sorted and individually encompassed by a single nonmyelinating SC. In *Gpr56* mutants, Remak bundle morphology was impaired, such that interdigitation of SC processes between axons within a single bundle was incomplete (Fig. 4 E, $P < 0.027$, one-way ANOVA). In addition, we noted several deviations in mature myelin ultrastructure. Myelin thickness was significantly increased in *Gpr56* mouse mutant SNs (Fig. 4 F; reflected in lower mean G-ratio, ~ 180 fibers analyzed from three animals per genotype, $P < 0.002$, one-way ANOVA), and we noted myelin outfoldings commonly associated with hypermyelination (Fig. 4 B).

In the PNS, myelin internodes possess two primary structural domains: Cajal bands (longitudinal channels of SC cytoplasm) and appositions (where the outermost layer of my-

elin contacts the SC plasma membrane; Sherman and Brophy, 2005). Strikingly, we also observed defects in the suborganization of peripheral myelin into Cajal bands and appositions in *Gpr56* mouse mutant SCs (Fig. 4 G; 154 WT and 168 *Gpr56*^{-/-} fibers from three animals per genotype, $P < 0.002$, one-way ANOVA), as well as SC cytoplasmic abnormalities in *gpr56*^{st13/st13} zebrafish mutant SCs (Fig. S2 B). The function of Cajal bands is poorly understood because few proteins have been linked to Cajal band formation (Court et al., 2004, 2009; Sherman et al., 2012; Walko et al., 2013). Of these proteins, Periaxin has been best studied; Periaxin stabilizes SC appositions, and *Periaxin* mutations cause neuropathy-like phenotypes in mouse and CMT disease in humans (Guilbot et al., 2001). Although *Periaxin* is well characterized for its role in CMT, mutations in *PLECTIN* (a cytolinker protein shown to regulate SC domain organization) are similarly as-



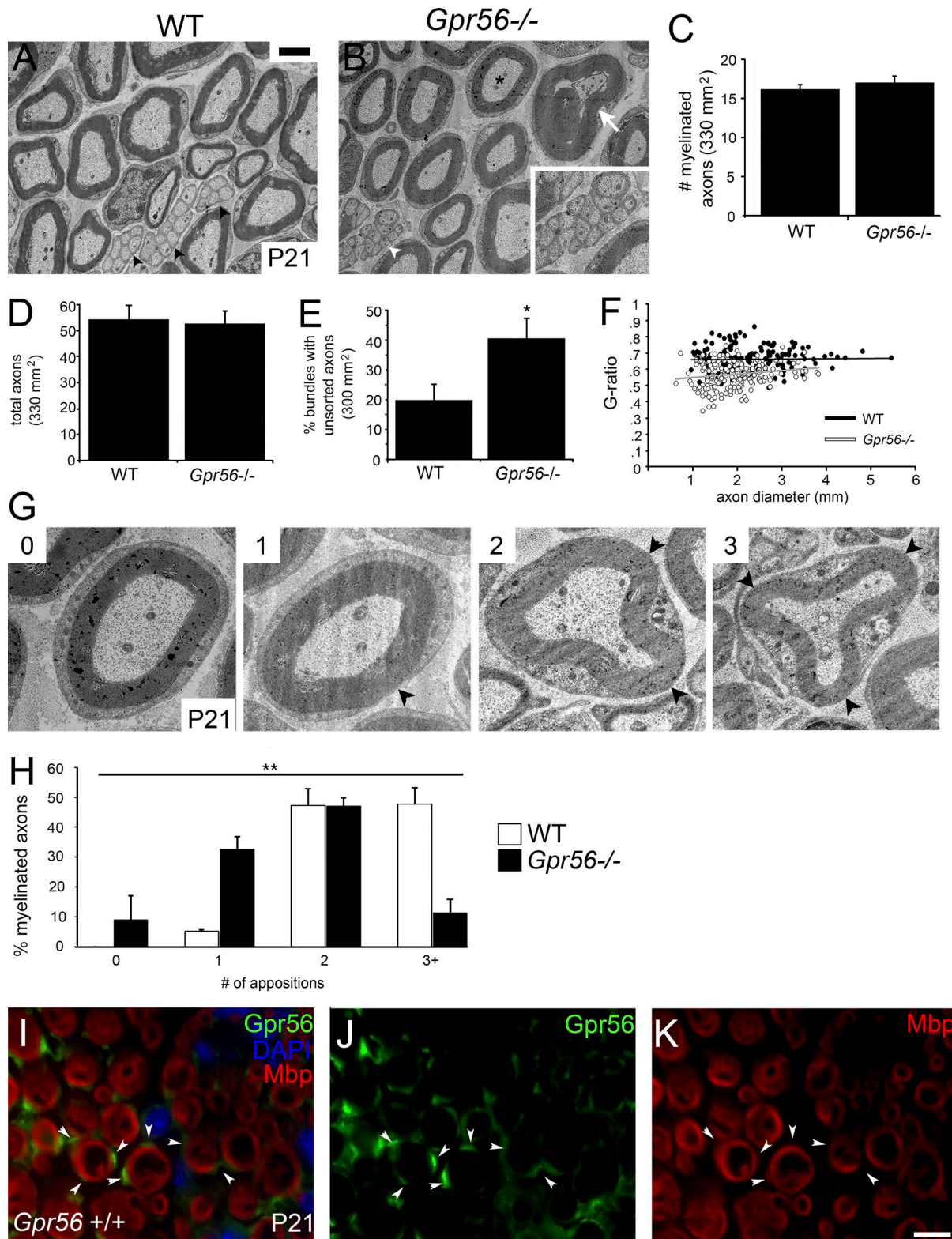


Figure 4. GPR56 regulates mature myelin ultrastructure and domain organization. (A and B) TEM images of WT ($n = 3$) and *Gpr56* mutant mouse ($n = 3$) SNs on P21. Bar, 2 μ m. White arrow denotes myelin outfolding. Black arrowheads denote morphologically WT Remak bundles, which are incompletely formed in *Gpr56* mutants (B, white arrowhead/inset). The asterisk marks an axon with no appositions. (C–E) Quantification of (C) total number of myelinated axons ($P < 0.45$, one-way ANOVA), (D) overall axon number ($P < 0.81$, one-way ANOVA), and (E) percentage of bundles containing axons that are not

sociated with myelin maintenance defects in mice and neuropathy-like symptoms in humans (Walko et al., 2013; Winter and Wiche, 2013), demonstrating the importance of myelin domain architecture for sustained PNS health in humans. Interestingly, analysis by immunohistochemistry (IHC) revealed that in mature SNs, GPR56 is specifically localized to discrete cytoplasmic domains on the abaxonal SC membrane (Fig. 4, I–K) and, thus, poised to regulate Cajal band formation.

Altered expression of cytoskeletal genes in developing and mature *Gpr56* mutant nerves

GPR56 signals via RhoA to regulate radial sorting (Figs. 1, 2, and 3), but the downstream signaling pathway(s) through which GPR56 regulates the development of mature myelin are not known. To determine what signaling pathway(s) may underlie the phenotypes observed in *Gpr56* mutants during SC maturation in contrast to early SC development (radial sorting), we performed RNA sequencing (RNA-seq) on RNA isolated from SNs of P21 and P3 mutants and WT controls, respectively ($n = 3$ per genotype on P21; $n = 4$ per genotype on P3; Fig. S3). Importantly, we observed a significant decrease in the expression of *Gpr56* at both developmental time points (Fig. S3 F). In accordance with the hypothesis that GPR56 normally functions in SC pseudopods to promote stable cell-cell contacts during radial sorting, and with the abnormal SC cytoplasmic extensions observed in *Gpr56* mutant SCs on P3 (Fig. 1, I and J), we observed a significant increase in the expression of genes associated with dynamic actin filaments in mutant SN harvested on P3 during active radial sorting (Fig. S3 D). We also found a statistically significant enrichment ($-\log[p\text{-value}] < 20$) in gene ontology (GO) processes associated with actin dynamics, including (a) actin-myosin filament sliding, (b) actin-mediated cell contraction, (c) actin filament-based movement, and (d) actin filament-based process using g:Profiler (Dataset 2).

Interestingly, we also observed significant changes in expression of genes related to the cytoskeleton on P21, when Cajal band defects are pronounced in *Gpr56* mutant nerves. Although we detected no differential expression of genes relating to the actin cytoskeleton, the expression of genes that promote microtubule assembly (*Map2* and *Ttl7*) was significantly increased in mutant nerves relative to controls, whereas expression of genes that promote microtubule disassembly (e.g., *Eml2*) was significantly down-regulated (Fig. S3 E). We also found a statistically significant enrichment ($-\log[p\text{-value}] < 4$) in the Cytoskeletal Remodeling GO pathway by g:Profiler (Dataset 2). The molecular mechanisms that guide Cajal band formation are poorly understood; however, microtubule

disorganization is a common feature of other mutants that lack Cajal bands (Court et al., 2004). These findings suggest that precise microtubule organization is necessary for Cajal band formation and represents an important area of future research into the development of these understudied domains.

In addition to changes in the expression of genes involved in microtubule dynamics, we also observed a striking dysregulation of a cluster of genes that share the SREBP-1 binding motif. Although we observed a modest increase in expression of SREBP1 target genes on P3, by P21 we noted a significant decrease in the expression of genes that share the SREBP-1 motif. Loss of *sterol regulatory element-binding factor-1c* (*Srebf1c*) results in increased peripheral myelin thickness, Remak bundle defects, and peripheral neuropathy-like symptoms in mice (Cermenati et al., 2015). Finally, although the authors did not formally examine the role of *Srebf1c* in Cajal band formation, *Srebf1c* mutants did not present with overt Cajal band defects based on published TEM images of *Srebf1c*^{−/−} peripheral nerve (Cermenati et al., 2015).

Our transcriptional profiling experiments (Fig. S3 and Dataset 2), combined with our genetic and biochemical analyses (Figs. 2 and 3) demonstrate that GPR56 signaling through $\alpha_{12/13}$ and RhoA is required to modify the actin skeleton for proper radial sorting during early SC development. These data further suggest that, during SC maturation, *Gpr56* signals through SREBP1 pathway to regulate lipid biogenesis and Remak bundle biology and through an alternative mechanism to modify microtubule dynamics for proper Cajal band assembly.

Gpr56 is required for SC myelin maintenance

Defective organization of the myelin sheath precedes neuropathy-like symptoms in all mutants (e.g., *Periaxin*, *Drp2*, and *Plectin*) previously shown to have Cajal band defects (Court et al., 2004; Sherman et al., 2012; Walko et al., 2013). *Gpr56* mutant mice present with Cajal band/apposition phenotypes, as well as *Srebf1c*-like hypermyelination and Remak SC defects (Cermenati et al., 2015); thus, we hypothesized that *Gpr56* mutants would develop neuropathy-like symptoms with age as is observed in both Cajal band and *Srebf1c* mutants. Ultrastructural analyses of mouse SNs on P90, P180, and P300 (Fig. 5, A–I) revealed small yet significant increases in the percent of axons with abnormal myelin profiles in *Gpr56* mutants and heterozygotes relative to WT siblings at each stage ($n = 3$ animals minimum per genotype/time point). Myelin abnormalities were composed of outfoldings and other defects: axon regeneration signatures (bands of Büngner), SCs myelinating other myelinating and Remak SCs, extensive

fully sheathed in *Gpr56* mutants and WT sibling nerves (*, $P < 0.03$, one-way ANOVA). Error bars represent SEM. (F) Plots of G-ratios in mutants and WT controls ($P < 0.002$, one-way ANOVA). (G) TEM images of *Gpr56*^{−/−} myelinated axons with 0, 1, 2, or 3 appositions. (H) Distribution of the percentage of myelinated axons (mean \pm SD) per apposition number in WT (white bars) and *Gpr56*^{−/−} SNs (black bars, **, $P < 0.002$, one-way ANOVA). (I–K) IHC showing GPR56 (green), DAPI (nuclear stain, blue), and MBP (marker of mature myelin) localization on P21 SNs. In WT SNs, GPR56 is detected in pockets (arrowheads) outside of MBP-positive rings but absent from mature axons. (J) GPR56 stain alone. (K) MBP stain alone. Bar, 5 μ m.

cytoplasmic extensions, minifascicle formation, myelin debris (Fig. 5, J–O), and disrupted Schmidt-Lanterman incisures (Fig. S4, A and B). In addition, we observed a significant interaction between genotype and age with respect to myelin abnormalities ($P < 0.0001$, two-way ANOVA; Fig. 5 P), indicating that these phenotypes worsen with time. Axon number remained largely unchanged with age, suggesting that loss of *Gpr56* does not induce widespread neuronal loss (Fig. 5, Q–S). Analysis of *Gpr56* mouse mutant dorsal and ventral spinal roots, composed of purely sensory and purely motor neurons, respectively, revealed similar demyelinating signatures (Fig. S4, C–L).

Given the importance of *Gpr56* in the formation of Cajal bands, structures that span the entire internode and can reach nearly 1 mm in length in adult mice (Court et al., 2004), we hypothesized that standard TEM analysis may underestimate myelin abnormalities in *Gpr56* mutants. Indeed, analysis of serial EM reconstructions of *Gpr56* mouse mutant SN on P90 revealed a cumulative average of $68 \pm 10\%$ axons showing at least one myelin defect over 15 μm of internode, compared with $15 \pm 6\%$ in WT controls (Fig. 6, Fig. S5, and Videos 1, 2, 3, and 4).

***Gpr56* mutants display neuropathy-like symptoms with age**

In addition to myelin loss, impaired Cajal band formation in *Periaxin* mutants is associated with shortened internodes, leading us to hypothesize that loss of *Gpr56* would affect internode length (Court et al., 2004). Teased nerve fiber analysis revealed a significant reduction in internode length on P6 (not depicted) and on P21 in *Gpr56* mutant mice (Fig. 7, A–C; $P < 0.05$, one-way ANOVA). Moreover, on P180, *Gpr56* mutants displayed a hindlimb clasping phenotype (Fig. 7, D and E), and both *Gpr56*^{+/-} ($n = 16$ on P180, $n = 15$ on P300) and *Gpr56*^{-/-} mutants ($n = 13$ on P180, $n = 12$ on P300) performed poorly relative to WT controls ($n = 11$ on P180, $n = 10$ on P300) on the accelerated rotarod, indicative of impaired sensory-motor coordination and reduced grip strength (Fig. 7 F; $P < 0.001$; Fig. 7 G, $P < 0.005$; two-way ANOVA analysis). To test whether behavioral deficits observed in *Gpr56* mutant and heterozygous mice reflected a reduction in nerve function, we performed electromyography (EMG) analyses in vivo and found a significant increase in EMG latency in *Gpr56* mutants ($n = 11$) relative to controls ($n = 5$ WT, $n = 12$ *Gpr56*^{+/-}) in animals aged P200–P300 (Fig. 7 H; $P < 0.001$, one-way ANOVA) with a corresponding reduction in response velocity (Fig. 7 I; $P < 0.001$, one-way ANOVA). See Fig. 7 J for example traces. Interestingly, although we observed no overt changes in Na⁺ channel structure by teased nerve analysis (Fig. 7, A and B), we detected a significant increase in the expression of genes related to K⁺ channel activity on P21 and a significant decrease in the expression of the Na⁺ channel clustering protein *Ndr4* (Fontenas et al., 2016) on both P3 and P21 (Fig. S3, C and D). GPR56 is also localized to sensory neurons (Fig. S4, M–O), and future work with

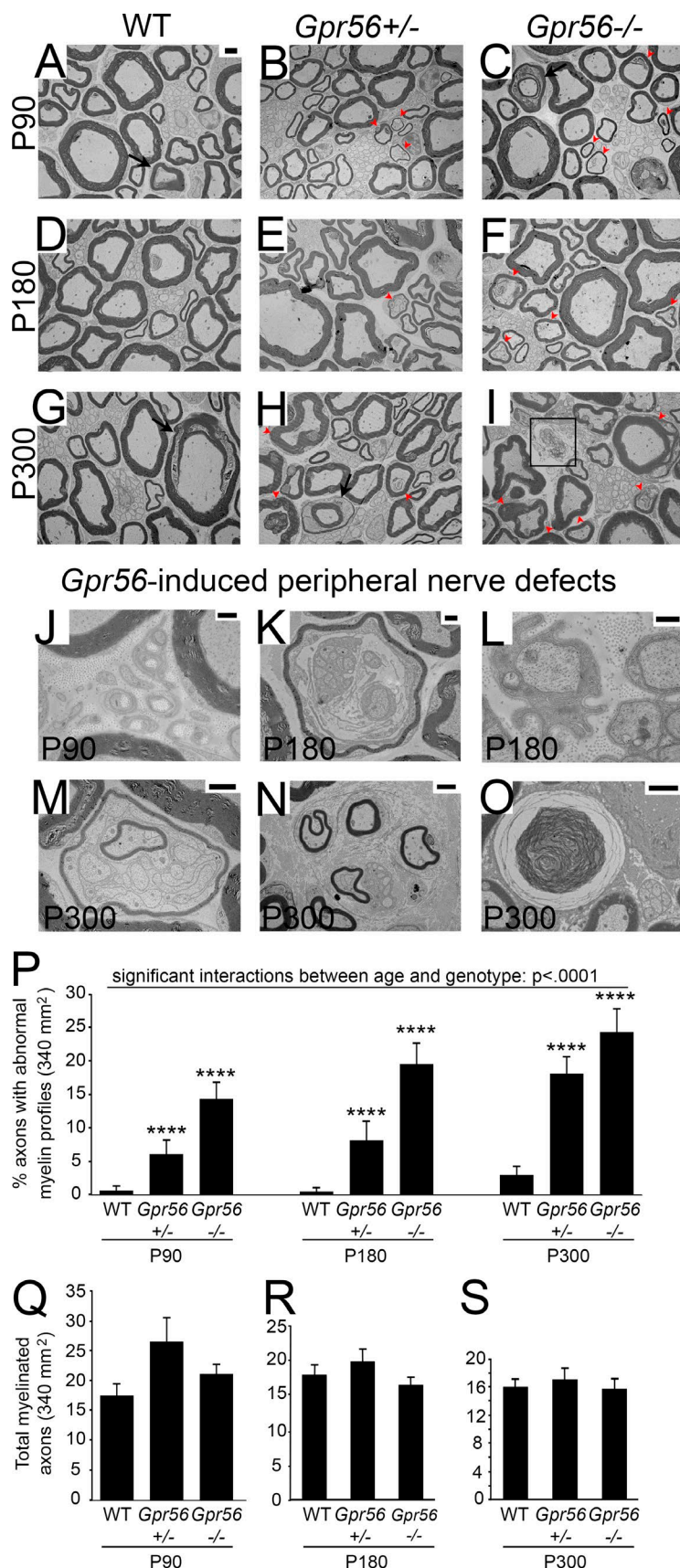
cell type-specific knockout lines will be important for understanding the relative contributions of GPR56 in neurons versus SCs to peripheral nerve function.

Plectin is a novel interacting partner of Gpr56 in peripheral nerve

To further define how GPR56 regulates SC function, we next sought to determine which protein(s) interact with GPR56 to facilitate peripheral nerve development. We recently showed that cellular prion protein (*Prnp*/Prp^C) is secreted from neurons and modulates GPR126 (an aGPCR closely related to GPR56) signaling and SC function. We used the previously published N-terminal flexible tail ([FT] residues 23–120) of Prp^C (Küffer et al., 2016), which can activate GPR126 signaling in vitro and in vivo, to test whether Prp^C can also activate *Gpr56* signaling in vitro. We did not observe any significant increase in GPR56 signaling either through RhoA or through an SRE-mediated transcriptional response to FT compared with application of collagen III, a known ligand of GPR56 (Luo et al., 2011; Fig. 8, A and B; $P < 0.03$, unpaired Student's *t* test). FT activates *Gpr126* signaling through cAMP; to test whether GPR56 may signal through this alternative pathway in response to FT, we measured cAMP levels after FT addition to GPR126 or GPR56. As previously shown, FT robustly stimulates cAMP signaling through GPR126 (Küffer et al., 2016) but had no effect on GPR56 (Fig. 8 C; $P < 0.0027$, unpaired Student's *t* test).

We next used an unbiased mass-spectrometry screen to determine what protein(s) physically interact with GPR56 (unpublished data). Interestingly, the top hit from this screen was plectin, a large cytolinker protein known to interact with and modulate cell surface signaling molecules through its coordinate reorganization of the actin and intermediate filament cytoskeletal networks (Winter and Wiche, 2013). *PLECTIN* is also mutated in a series of human diseases termed plectinopathies, including epidermolysis bullosa simplex with muscular dystrophy, a skin blistering disorder that also presents with neuropathy-like symptoms because of degeneration of intramuscular myelinated nerves (Bauer et al., 2001; Winter and Wiche, 2013). First, we verified that GPR56 and plectin physically interact in WT P5 SNs ($n = 8$ nerves from four animals) by coimmunoprecipitation (co-IP; Fig. 8 D). Interestingly, although we were able to detect binding of plectin to the CTF of GPR56 (not depicted), we observed robust binding of plectin to the extracellular NTF of GPR56 as well. Although this result most likely reflects binding of plectin to the larger GPR56 NTF-CTF complex, plectin has been shown to be secreted to the extracellular matrix in pathological contexts (Shin et al., 2013b). Future work will therefore be required to test whether plectin is secreted to serve as a direct activating ligand for GPR56.

We next tested whether loss of *Gpr56* affects plectin expression in P6 SNs and found a significant increase in plectin levels in *Gpr56* mutants ($n = 7$ replicates: 1 biological replicate = 2 pooled nerves from one animal) relative



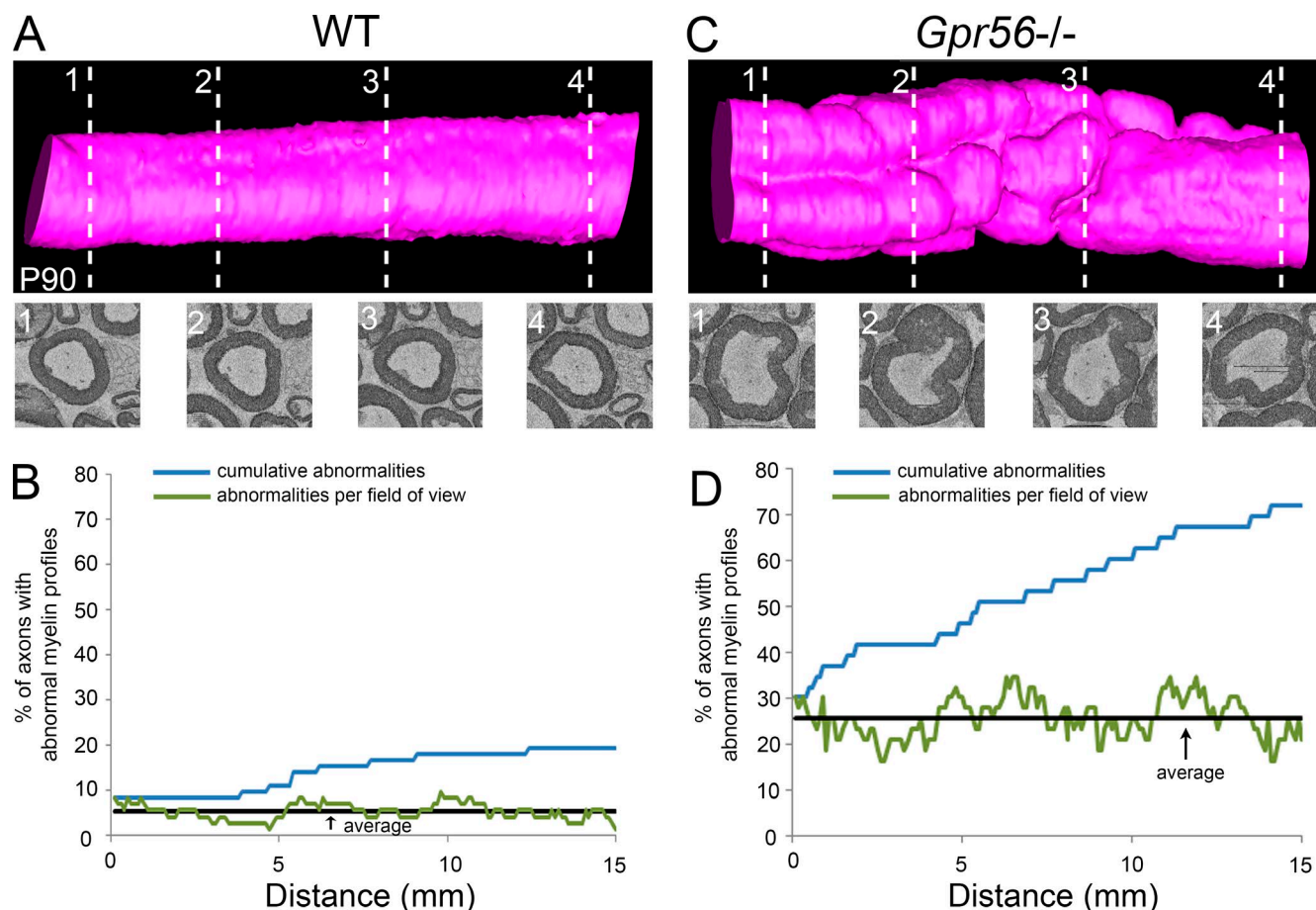


Figure 6. ***GPR56* is required for myelin stability.** (A–D) Serial scanning EM images (350, 100 nm apart) of WT and *Gpr56*^{-/-} mouse SN on P90 ($n = 3$ animals per genotype) were acquired and reconstructed to generate the WT and *Gpr56*^{-/-} models shown in A and C, respectively. Single sections taken from multiple locations along the nerve (1–4) are shown below each model (dashed lines show approximate locations). The percentage of axons with abnormal myelin profiles was calculated per section over a 15- μ m distance for the WT (B) and *Gpr56* mutant (D) reconstruction and is graphed to show the percentage of abnormalities per section (green line), the average percentage of abnormalities over all sections (black line) and the cumulative percentage of abnormalities (percentage of axons that show at least one abnormality over 15 μ m [blue line]).

to WT ($n = 8$ replicates) and heterozygous ($n = 5$ replicates) siblings, as expected for a positive interacting partner of GPR56 (Fig. 8 E; $P < 0.001$, unpaired Student's t test). Finally, plectin is strongly expressed in SCs and specifically localized to Cajal bands, and SC-specific KO of *Plectin* disrupts myelin domain organization and causes neuropathy-like symptoms with age (Walko et al., 2013). Thus, plectin is a novel binding partner for GPR56, and the presence of phenocopy coupled with both physical and genetic interactions (Fig. 8, D and E) strongly suggest that GPR56 and plectin function together to promote SC function and long-term health of peripheral nerves.

DISCUSSION

GPR56 in myelinating glial cell development

In the CNS, oligodendrocytes form the myelin sheath that both protects and insulates axons for rapid action potential propagation (Jessen and Mirsky, 2005). We previously defined

GPR56 as a novel regulator of oligodendrocyte development (Ackerman et al., 2015; Giera et al., 2015; Salzman et al., 2016). In the CNS, GPR56 regulates the timing of the transition between oligodendrocyte precursor cell proliferation and differentiation. In the absence of GPR56, oligodendrocyte precursor cells prematurely exit the cell cycle resulting in fewer numbers of myelinating oligodendrocytes; however, the myelinating oligodendrocytes that are produced generate phenotypically normal myelin (Ackerman et al., 2015; Giera et al., 2015). In the PNS, SCs ensure long-term health of peripheral axons (Jessen and Mirsky, 2005). In contrast to the role of GPR56 in regulating the numbers of myelinating glia in the CNS, GPR56 does not affect SC proliferation or overall numbers. Instead, we observe novel and distinct functions of GPR56 in SC development (radial sorting) and in SC myelination (thickness, organization, and maintenance), and our data suggest defects in cytoskeletal dynamics as underlying causes of both early and late functions of GPR56 in peripheral nerve.

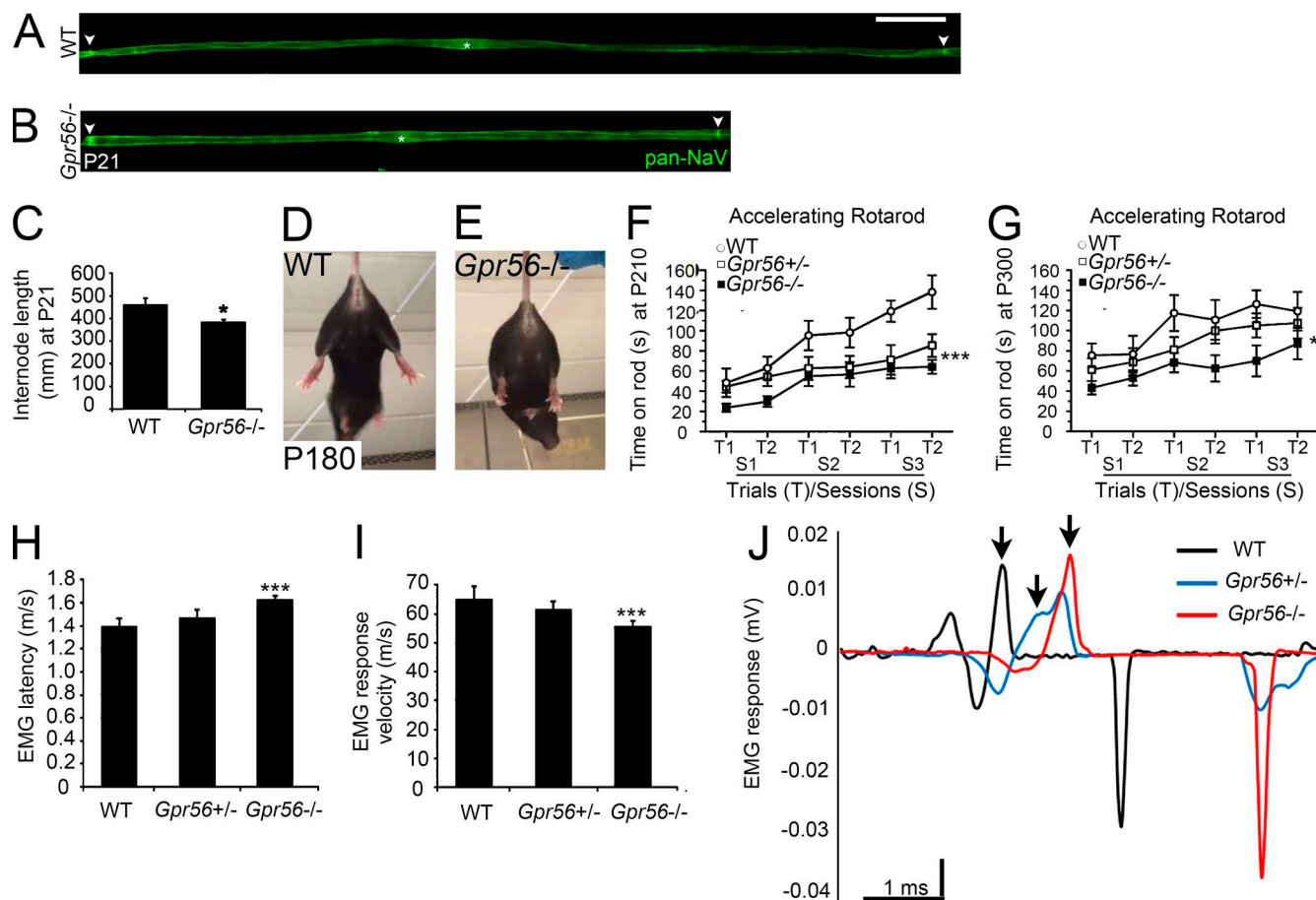


Figure 7. *Gpr56* mutants display neuropathy-like symptoms. (A and B) IHC on teased nerve fibers from WT ($n = 3$) and *Gpr56*^{-/-} ($n = 4$) mouse SNs on P21 stained for Pan-NaV to mark nodes of Ranvier (arrowheads). Bar, 50 μ m. (C) Quantification of internode length on P21 ($P < 0.05$, one-way ANOVA). (D and E) Hindlimb clasping observed in *Gpr56* mutants ($n = 8/11$) but not WT siblings on P180 ($n = 1/11$). (F and G) Quantification of time on rotarod to assess sensorimotor control on P180 (genotype effect: ***, $P < 0.001$, two-way ANOVA; $n = 11$ WT, $n = 16$ *Gpr56*^{+/-}, $n = 13$ *Gpr56*^{-/-}) and P300 (genotype effect: *, $P < 0.02$, two-way ANOVA; $n = 10$ WT, $n = 15$ *Gpr56*^{+/-}, $n = 12$ *Gpr56*^{-/-}). Error bars represent SEM. (H) Quantification of EMG latency (m/s) in animals aged P200–P300 (***, $P < 0.001$, one-way ANOVA). Error bars represent SD. (I) Quantification of EMG response velocity (m/s) in animals aged P200–P300 (***, $P < 0.001$, one-way ANOVA). Error bars represent SD. (J) Representative traces from EMG recordings from WT (black, $n = 5$), *Gpr56*^{+/-} (blue, $n = 12$), and *Gpr56*^{-/-} (red, $n = 11$) SNs. Arrows denote the point of the first positive deflection.

RhoA-dependent regulation of radial sorting by GPR56

During early SC development, we find that GPR56 signaling is required for timely sorting of axons. Many proteins have been shown to function during radial sorting, including small Rho-GTPases (Nodari et al., 2007; Pereira et al., 2009; Feltri et al., 2016). RhoA is robustly expressed during radial sorting (Taylor et al., 2003; Melendez-Vasquez et al., 2004; Nodari et al., 2007) and specifically enriched in SC pseudopods. The function of RhoA in SCs has not been thoroughly examined, however, given the established role of the related Rho GTPase Rac in radial sorting (Nodari et al., 2007; Pereira et al., 2009; Feltri et al., 2016). Rac positively regulates the formation/protrusion of lamellipodia-like structures from SCs, whereas RhoA activation typically inhibits process extension (Pereira et al., 2009). Indeed, up-regulation of Rho kinase (ROCK; a major downstream effector of RhoA signaling) in *ILK*

SC-specific KO mice results in severe radial sorting delays because of impaired process extension (Pereira et al., 2009). In contrast, we find that reduced RhoA signaling in *Gpr56* mutants likewise impairs radial sorting, which can be restored upon transient overexpression of constitutively active RhoA.

These results indicate that RhoA levels must be very tightly regulated during SC development, because both increased signaling in *ILK* mutants impairs SC process extension, and decreased signaling in *Gpr56* mutants delays radial sorting theoretically through impaired SC adhesion to axonal membranes. In accordance with this hypothesis, a previous study demonstrated that inhibition of ROCK in vitro results in aberrant actin fiber organization and reduced SC adhesion because of loss of stress fibers (Melendez-Vasquez et al., 2004). ROCK is a serine-threonine kinase that regulates actin organization through phosphorylation of proteins that modulate

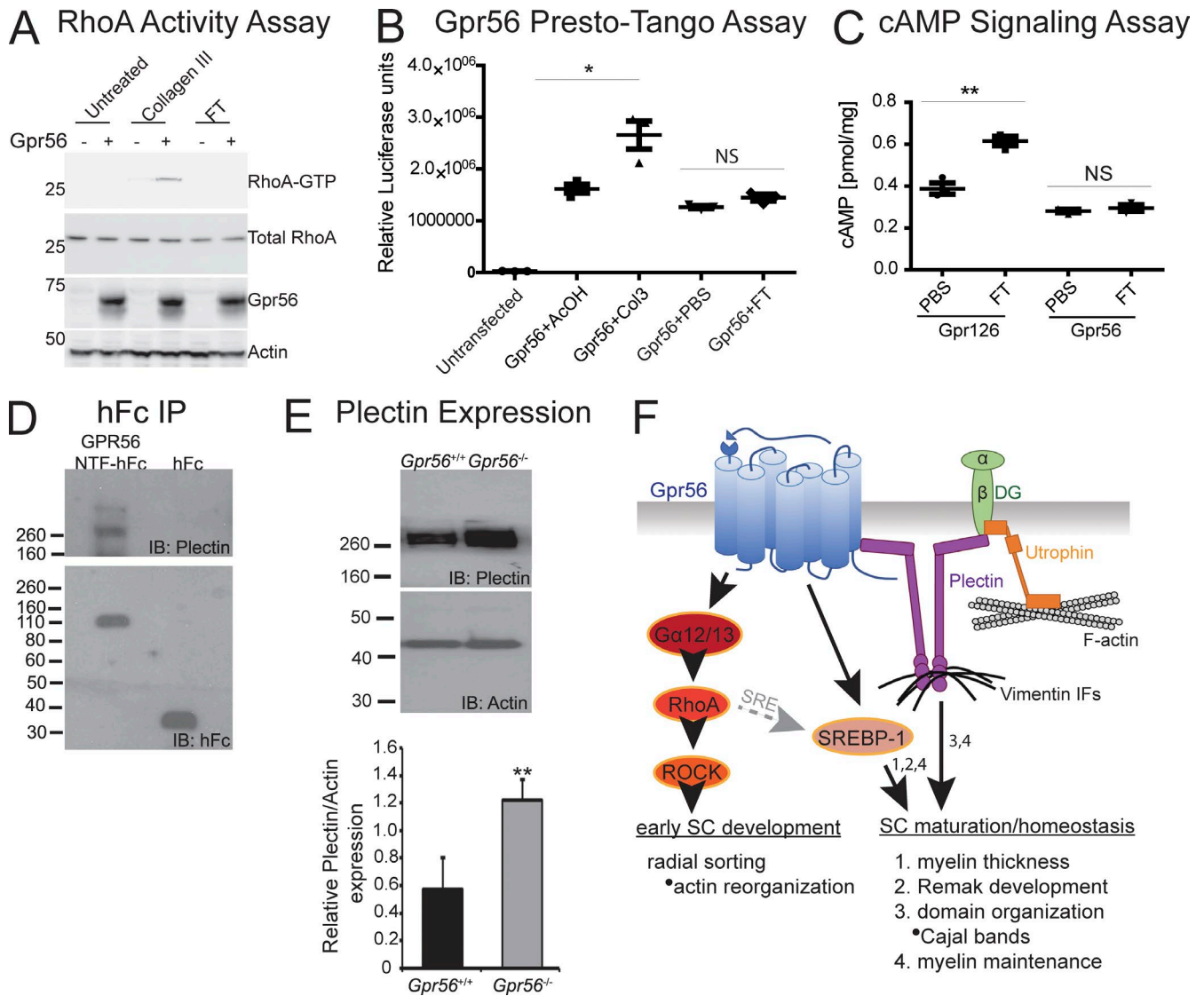


Figure 8. Plectin and GPR56 physically and genetically interact in peripheral nerve. (A) GTP-RhoA assay to assess Gpr56 signaling in response to exogenous ligands. Addition of collagen III, a known activating ligand of Gpr56, to HEK 293T cells coexpressing *Gpr56* result in elevation of GTP-RhoA levels over control (untransfected). Addition of prion protein (FT) does not induce Gpr56 signaling through RhoA. (B) Gpr56 Presto-Tango luciferase assay to assess Gpr56 signaling in HTLA cells. Addition of collagen III results in a significant elevation of luciferase compared with AcOH alone control (*, $P < 0.05$, unpaired Student's *t* test). Addition of FT did not result in a significant elevation of luciferase relative to PBS controls ($P = 0.0647$, unpaired Student's *t* test). (C) cAMP ELISA assay to assess cAMP signaling through Gpr56 and Gpr126 (positive control) after addition of FT. Addition of FT to HEK 293T cells expressing *Gpr126* results in a significant elevation of cAMP (**, $P < 0.01$, unpaired Student's *t* test) compared with PBS controls. Addition of FT to cells expressing *Gpr56* does not induce cAMP signaling ($P = 0.4575$, unpaired Student's *t* test). (A–C) Data are representative of three biologically independent experiments. Error bars represent SEM. (D) Co-IP to assess interactions between GPR56 and plectin in P5 WT SNs ($n = 8$ nerves from four animals per replicate; $n = 3$ replicates). Lane 1: plectin coimmunoprecipitates with GPR56 when the NTF of GPR56 fused to human IgG Fc (NTF-hFc) is used as a bait but not when pulling down with hFc alone (lane 2). (E) Analysis of plectin levels in P6 SNs. Plectin is significantly increased in *Gpr56*^{−/−} nerves over sibling controls (**, $P < 0.002$, unpaired Student's *t* test, $n = 12$ nerves from six animals; error bars represent SEM). (F) Graphic model: Gpr56 promotes function of developing and mature SCs. During early SC development, Gpr56 signals through Gα_{12/13} and RhoA to modulate the actin cytoskeleton for efficient radial sorting of axons. During SC maturation/homeostasis, Gpr56 functions upstream of SREBP1 to regulate myelin thickness and Remak SC biology, potentially through SRE signaling downstream of RhoA (gray dashed arrow). Concordantly, Gpr56 regulates mature myelin domain organization and myelin maintenance through direct interactions with the cytolinker protein plectin, which has been shown to physically interact with additional regulators of myelin domain organization (utrophin and β-dystroglycan). We therefore hypothesize that plectin serves as a docking center for cell signaling molecules that govern the partitioning of myelin into mature subdomains (Cajal bands and appositions) and helps generate these structures through its interactions with the cytoskeleton.

cytoskeletal dynamics (Amano et al., 1996). Myosin II light chain is a direct target of ROCK, functions to stabilize and increase the contractility of actin-myosin filaments (Tan et al., 1992), and is required for ROCK-dependent reorganization of actin fibers in SCs (Melendez-Vasquez et al., 2004). RNA-seq of *Gpr56* mutant nerves during radial sorting (P3) revealed a significant dysregulation of genes that modify the actin and actomyosin cytoskeleton, including *Myosin light chain 1* (*My11*, also known as *Mhc1/Mhc3*), *Myosin Light Chain*, *Phosphorylatable*, *Fast Skeletal Muscle* (*My1pf*, also known as *Mlc2b*), and *Mybpc1*, a myosin binding protein. These findings support the hypothesis that GPR56 signals through RhoA and ROCK to modify the actin and actomyosin cytoskeleton to establish firm SC-axon contacts during radial sorting (Fig. 8 F, working model).

Gpr56 regulates peripheral myelin structure and maintenance

Although radial sorting recovers in *Gpr56* mutants with time, we show that loss of GPR56 leads to multiple defects in mature SCs. First, we observed a significant increase in myelin thickness despite the delay in myelination onset. RNA-seq analysis of mutant nerves on P21 (postradial sorting defects) revealed a significant decrease in the expression of a cluster of SREBP1 target genes. SREBP1 is a transcription factor that regulates membrane lipid composition, and deletion of the primary isoform of SREBP1 (*Srebp1c*) results in hypermyelination, incomplete separation of axons by Remak SCs, and neuropathy-like symptoms with age (Cermenati et al., 2015). Although the signaling pathways upstream of SREBP1 activity are poorly defined, analysis of the promoter region of human and mouse *SREBP1* revealed SRE binding sites (Eberlé et al., 2004). GPR56-dependent RhoA signaling is known to function upstream of SRE to regulate neural precursor migration (Iguchi et al., 2008), suggesting that GPR56 may also signal through this cascade to regulate SREBP1 expression and in turn, myelin thickness and Remak SC dynamics (Fig. 8 F, working model).

In addition to changes in myelin thickness and Remak SC biology, we also observed a striking defect in mature myelin domain organization in *Gpr56* mutants—a significant reduction in Cajal band number. Mature myelin is longitudinally organized into two primary structural domains: Cajal bands and appositions. Cajal bands are cytoplasmic channels separated from one another by regions of completely compact SC membrane called appositions. The molecular mechanisms that drive Cajal band development are poorly understood, in part because few proteins have been linked to Cajal band development. Mutations in the large intracellular cytolinker protein *PLECTIN* in humans are linked to peripheral nerve defects (Winter and Wiche, 2013), and a recent study in mice defined plectin as a key regulator of myelin domain organization (Walko et al., 2013). In SCs, plectin is specifically localized to Cajal bands where it interacts with β -dystroglycan and utrophin to restructure vimentin intermediate filaments in the cy-

toskeleton (Walko et al., 2013). Interestingly, β -dystroglycan is also a member of the Periaxin–DRP2–dystroglycan complex that stabilizes appositions for proper formation of Cajal bands (Sherman and Brophy, 2005). Here, we define plectin as a novel interacting protein of GPR56: GPR56 and plectin physically interact, and loss of *Gpr56* results in up-regulation of plectin, as expected if plectin mediates GPR56 downstream signaling.

Plectin has now been shown to interact with proteins that are specifically enriched in Cajal bands (utrophin and GPR56) as well β -dystroglycan, which is present in both Cajal bands and in appositions as part of the Periaxin–DRP2–dystroglycan complex. Furthermore, plectin is known to function as a scaffold for receptors that mediate intracellular signaling (Castañón et al., 2013). Together, these data suggest that in myelinating SCs, plectin could serve as a docking center for both structural (Periaxin, DRP2, and utrophin) and signaling (GPR56 and β -dystroglycan) molecules required for efficient compartmentalization and long-term integrity of the myelin sheath (Fig. 8 F, working model). Although the downstream signaling pathways that promote peripheral myelin organization remain elusive, our RNA-seq analysis on P21 (when changes in Cajal bands are present in *Gpr56* mutant nerves) revealed significant dysregulation of genes that regulate microtubule dynamics. Disorganization of the microtubule cytoskeleton is present in *Periaxin* mutants, which also present with Cajal band defects (Court et al., 2004), supporting the hypothesis that reorganization of microtubules is required for mature myelin domain assembly.

Neuropathy-like symptoms in GPR56 mutants: links to human health

Several of the developmental defects observed in *Gpr56* mutants (e.g., hypermyelination and reduced Cajal bands) are known to precede disruptions in myelin integrity (Court et al., 2004, 2009; Sherman et al., 2012; Walko et al., 2013; Cermenati et al., 2015), and indeed, myelin maintenance is also defective in *Gpr56* mutants relative to controls. We demonstrate that in *Gpr56*^{−/−} mouse adults, impaired myelin stability coupled with shortened internode length results in behavioral deficits consistent with the recorded EMG latency delay in *Gpr56* mutants. Similar phenotypes are often noted in mouse models of peripheral neuropathy and in human neuropathy patients (Martini, 1999), suggesting that mutations in *GPR56* could cause neuropathy-like symptoms in humans. Mutations in *GPR56* cause several neurological disorders affecting the CNS, including bilateral perisylvian polymicrogyria (BPP or BPPR; OMIM #615752). BPP is a cortical brain malformation resulting in severe neurological impairments, including epilepsy and oromotor apraxia, and a minority of BPP patients also present with arthrogryposis multiplex congenital and exhibit EMG signatures consistent with PNS pathology (Poduri et al., 2010). Our work defines GPR56 as a novel, evolutionarily conserved regulator of SC development and myelin maintenance and suggests a mechanism underlying PNS defects observed in BPP patients.

MATERIALS AND METHODS

Zebrafish

Stocks and rearing conditions. Zebrafish (*Danio rerio*) were maintained in the Washington University Zebrafish Consortium facility. All experiments were performed in compliance with Washington University's institutional animal care and use committee. Embryos were collected from pairwise matings and reared at 28.5°C in egg water (5 mM NaCl, 0.17 mM KCl, 0.33 mM CaCl₂, and 0.33 mM MgSO₄). Embryos and larvae were staged in days postfertilization (dpf) by using morphological markers (Kimmel et al., 1995). The *gpr56*^{stl13/stl13} line was genotyped as described previously (Ackerman et al., 2015).

3MO and synthetic mRNA injection. 3MOs targeting *gna12*, *gna13a*, and *gna13b* (3MO) are previously published (Lin et al., 2009; Ackerman et al., 2015) and were provided by F. Lin (University of Iowa, Iowa City, IA). 3MOs were combined and diluted in sterile water supplemented 1:10 with phenol red dye and then injected at a final concentration of 0.5 ng each in a total volume of 1 nl or 1 ng each in a total volume of 2 nl. To control for adverse side effects from the injection process, we also injected control siblings with an equal volume of phenol red diluted 1:10 in water (phenol red-injected controls). Constitutively active Rho (*rhov14*) was provided by L. Solnica-Krezel (Washington University, St. Louis, MO) and was transcribed and injected as described previously (Lin et al., 2009; Ackerman et al., 2015).

TEM. For zebrafish larvae, TEM was performed as described (Czopka and Lyons, 2011) by using a PELCO BioWave Pro with SteadyTempTM Digital Plus water-recirculating system. Microwave-assisted fixation was used for all steps before Epon treatment. In brief, embryos were fixed in modified Karnovsky's fix (4% paraformaldehyde, 2% glutaraldehyde, and 0.1 M sodium cacodylate, pH 7.4) and left overnight at 4°C. After fixation, embryos were postfixed in 2% osmium tetroxide in 0.1M sodium cacodylate and 0.1M imidazole, pH 7.4, washed, and treated with saturated uranyl acetate. Embryos were then dehydrated by using increasing concentrations of ethanol followed by 100% acetone and then infiltrated overnight in an acetone-Epon mixture. Embryos were then transferred to 100% Epon, embedded in molds, and baked for at least 48 h at 65°C. Fish were examined at approximately the same body segment level to control for variability along the anterior/posterior axis. For preparation of pLLNs from 6-mo-old adult fish, we anesthetized animals in an ice-water bath and severed the hindbrain before dissecting a portion of the anterior pLLN, and fixation of adult tissue occurred as described previously (Monk et al., 2009). In brief, adult tissues were fixed via microwave-assisted fixation in modified Karnovsky's fix as earlier in this paragraph and then postfixed in 2% osmium tetroxide in 0.1M sodium cacodylate, washed, and dehydrated in increasing concentrations of ethanol followed by 100% propylene oxide. The tissues were then sub-

merged in a 2:1 mixture of propylene oxide/EPON for 1 h, followed by a 1:1 mixture overnight. The next day, the tissues were transferred to 100% Epon before embedding and baking at 65°C. Thin sections (70 nm) for all samples were mounted on mesh grids and stained with 8% uranyl acetate followed by Sato's lead stain. We viewed sections on a Jeol (JEM-1400) electron microscope, and images were acquired with an AMT V601 digital camera. All images were processed and analyzed by using ImageJ and Adobe Photoshop software. At 3 dpf, quantifications were performed on the entire nerve. For *rhov14* transient overexpression experiments, we examined five nerves from three WT, five nerves from three *gpr56*^{stl13/+}, and seven nerves from four *gpr56*^{stl13/stl13} larvae for phenol red controls; and six nerves from three WT, six nerves from three *gpr56*^{stl13/+}, and five nerves from three *gpr56*^{stl13/stl13} larvae injected with 10 pg *rhov14* synthetic mRNA. For 3MO experiments, we examined 10 nerves from 6 WT, 10 nerves from 6 *gpr56*^{stl13/+}, and 9 nerves from 6 *gpr56*^{stl13/stl13} larvae for phenol red control treatment. For larvae treated with 0.5 ng 3MO, we examined 10 nerves from 6 WT, 8 nerves from 5 *gpr56*^{stl13/+}, and 7 nerves from 5 *gpr56*^{stl13/stl13} animals. For larvae treated with 1 ng 3MO, we examined six nerves from three WT, five nerves from three *gpr56*^{stl13/+}, and six nerves from *n* = 3 *gpr56*^{stl13/stl13} animals. For larval studies at 5 dpf, we examined seven nerves from four WT and six nerves from three *gpr56*^{stl13/stl13} animals. For adult studies, three nerves from three individuals at 6 mo of age for WT and *gpr56*^{stl13/+}, and four nerves from three *gpr56*^{stl13/stl13} animals were analyzed.

Mice

Lines and rearing conditions. All mouse (*Mus musculus*) experiments were performed in compliance with Washington University's Institutional Animal Care and Use Committee as well as the Animal Care and Use Committee at Boston Children's Hospital. The *Gpr56* knockout mice were obtained pure on C57BL/6 (*n* = 11 generations) from the Mutant Mouse Regional Resource Centers (032342-UCD), maintained on a C57BL/6 background, and genotyped as described previously (Li et al., 2008).

SC cultures and preparation of neuronal membranes. Primary rat SCs were prepared from SN on P3 as described (Feltri et al., 1992). SCs were maintained in SC medium: DMEM (Gibco), 1% FCS, 2 mM L-glutamine, 2 μM Forskolin (Calbiochem), and 2 ng/ml recombinant human NRG1-β 1 (R&D) and passaged not more than three times.

Neuronal membranes were prepared from dorsal root ganglia (DRG) of rat embryos at 15 gestational days. In brief, 20 DRGs were plated on 9.5-cm² dishes in C media: MEM (Gibco), 10% FBS, 4 g l⁻¹ D-glucose, 2 mM L-glutamine, 25 ng/ml BDNF (PeproTech), and 10 ng/ml NT-3 (PeproTech). After 12 h, cells were switched for 2 wk in NB medium (Gibco), 1X B27 supplement (Gibco), 4 g/l D-glucose, 2 mM L-glutamine, 25 ng/ml BDNF (PeproTech), and 10 ng/ml NT-3 (PeproTech). To obtain neuron-only cultures, the cells

were treated with FUDR (10 μ M floxuridine and 10 μ M uridine) in NB medium for three cycles (2 d with FUDR, 2 d without FUDR). Neurons were collected by scraping the network with forceps and homogenized in PBS with a Dounce homogenizer. The homogenized solution was centrifuged at 300 *g* for 20 min to remove debris and collagen. The supernatant was centrifuged at 30,000 *g* for 1 h. The pellets were resuspended in DMEM (67 μ l for each 9.5-cm² dish) and stored at -80°C for 1 mo. To test their activity, membranes were centrifuged onto serum-starved, rat primary SCs, incubated at 37°C for an additional 20 min, lysed, and analyzed by Western blotting for Akt activation.

Pseudopod assay. Pseudopod assay was performed as described previously (Poitelon et al., 2015). In brief, starved SCs were plated onto the upper compartment of the Boyden chamber insert (3414; Corning). Cells were maintained in DMEM, 2 mM L-glutamine and 0.5% BSA for 4 h at 37°C . Then pseudopods were allowed to grow for 2 h at 37°C by replacing bottom-chamber DMEM with neuronal membrane suspension. After induction, inserts were washed twice in PBS, fixed with ice-cold methanol for 30 min on ice, and washed again in PBS. For each filter, either the cell bodies on the upper surface or the pseudopods on the undersurface were removed with a cotton swab.

Sample preparation for shotgun proteomics for SC pseudopod assay. Microporous filters with either pseudopods or cell bodies were manually cut into tiny pieces and suspended in 200 μ l of cold lysis buffer (50 mM Tris-FA, 150 mM NaCl, 0.5% sodium deoxycholate, 2% SDS, 2% IGEPAL CA-630 [Sigma-Aldrich], pH 8.0) supplemented with protease/phosphatase inhibitor cocktail tablets (Roche Applied Science). The samples were sonicated until the liquid turned clear and were then placed on ice for 30 min, followed by a 30-min centrifugation at 20,000 *g* under 4°C . Supernatant from all samples was carefully transferred to Eppendorf tubes, and protein concentrations were measured by bicinchoninic acid assay kit (Pierce Biotechnology, Inc.).

For each extraction, 100 μ g of protein was subjected to a surfactant-aided precipitation/on-pellet digestion procedure (An et al., 2015) to generate tryptic peptides for LC-MS analysis. Proteins first undergo reduction and alkylation by 3 mM tris(2-carboxyl)phosphine and 20 mM iodoacetamide. Both steps were performed under 37°C for 30 min in darkness, with constant oscillation in an Eppendorf Thermomixer (Eppendorf). Proteins were then precipitated by addition of eight volumes of chilled acetone, and the samples were incubated under -20°C overnight. Precipitated proteins were collected by centrifugation at 20,000 *g* under 4°C for 30 min, and the supernatant portion was removed. This step allowed the elimination of detergents and unwanted contaminants (e.g., phospholipids), which will compromise the analytical quality of the LC-MS system. Protein pellets were rinsed with 500 μ l of chilled acetone/water mixture (85:15, vol/vol percentage),

air-dried for 5 min, and resuspended in 80 μ l Tris-FA. Trypsin from porcine pancreas (Sigma-Aldrich) at a concentration of 0.25 μ g/ μ l was added to the samples in two steps, at a final enzyme/substrate ratio of 1:20 (wt/wt percentage). The samples were incubated under 37°C overnight (18–20 h) with constant oscillation to achieve complete proteolytic cleavage. The digestion step was terminated by addition of 1 μ l formic acid, after which the samples were centrifuged at 20,000 *g* under 4°C for 30 min and transferred to LC vials.

Long-gradient nano reversed-phase LC-MS for SC pseudopod assay. The nano RPLC system used in the present study consists of a Spark Endurance autosampler (Emmen) and an ultrahigh-pressure Eksigent Nano-2D Ultra capillary/nano LC system. A nano LC/nanospray setup featuring low-void-volume and high-chromatographic reproducibility was used so that comprehensive separation of complex peptide samples could be achieved. Mobile phase A was 0.1% formic acid in 2% acetonitrile, whereas mobile phase B was 0.1% formic acid in 88% acetonitrile. Peptides were first loaded onto a large inner-diameter trap (300 μ m inner diameter \times 1 cm, packed with Zorbax 3M C18 material) with 1% B at 10 μ l min⁻¹, and the trap was washed for 3 min before bringing in line with the nano LC flow path. A series of optimized gradients (250 nl/min) was used to back-flush the trapped peptides onto the nano LC column (75 μ m inner diameter \times 100 cm, packed with Pepmap 3M C18 material), which was heated homogeneously to 52°C to improve chromatographic resolution and reproducibility. The 7-h gradient profile used for extensive peptide separation was listed as follows: 3–8% B over 15 min, 8–24% B over 215 min, 24–38% B over 115 min, 38–63% B over 55 min, 63–97% B in 5 min, and finally isocratic at 97% B for 15 min.

An LTQ Orbitrap XL mass spectrometer (Thermo Fisher Scientific) was used for peptide detection. Reproducibility of ionization efficiency was maintained via rinsing and conditioning of the nanospray needle (by dripping 50% methanol after every three runs). The instrument was carefully calibrated for mass accuracy and Fourier transform transmission. One complete scan cycle consisted of one MS1 survey scan (*m/z* 310–2,000) at a resolution of 60,000 and seven subsequent MS2 scans by collision-induced dissociation activation mode to fragment the top seven most-abundant precursors in the survey scan. The automatic gain control target was set to 8×10^6 , which allowed highly sensitive detection with little compromise to mass accuracy and resolution. Dynamic exclusion was enabled with the following settings: repeat count = 1, repeat duration = 30 s, exclusion list size = 500, and exclusion duration = 40 s. The activation time was 30 ms with an isolation width of 3 D for ion trap mass spectrometry; the normalized activation energy was 35%, and the activation *q* was 0.25. All 20 samples were randomly analyzed.

Protein identification and quantification for SC pseudopod assay. LC-MS.raw files were matched to a combined

Swiss-Prot *Mus musculus* and *Rattus norvegicus* protein database because of the incomprehensiveness of the current rat protein database. Files were matched by SEQUEST by using a target-decoy search strategy, in which a concatenated protein database with both forward and reverse sequences was used (Elias et al., 2005). The searching parameters include 15 ppm for precursor ion mass, 1.0 D tolerance for fragment ion mass, 2 for maximal missed cleavages, carbamidomethylation of cysteine as fixed modification, and oxidation of methionine as dynamic modification. Peptide inference, protein grouping, and false-discovery rate control was accomplished by Scaffold (v4.3.2; Proteome Software Inc.; Searle, 2010). Corresponding parameters include for minimal peptide length, 25 ppm for parent mass tolerance; 0.1 for DeltaCn; 2.1, 2.2, 2.6, 3.5 (pseudopods); and 1.1, 1.4, 1.7, 2.5 (cell bodies) for Xcorr thresholds for singly, doubly, triply, and quadruply charged ions.

Chromatographic alignment and global intensity-based MS1 feature detection/extraction were accomplished by SIEVE (v2.1.377; Lopez et al., 2011). The major steps we took are as follows. (a) Alignment of LC-MS runs via the use of ChromAlign algorithm. The quality of LC-MS runs was monitored and benchmarked by the alignment scores and base-peak ion current intensity among all runs. (b) Determination of quantitative “frames” based on *m/z* and retention time in the aligned collective dataset. The reliability of quantification was assured by retaining only frames with high-quality area-under-the-curve data (e.g., signal/noise ratio >10). (c) Calculation of peptide ion intensities for each frame. The resulting .sdb files were incorporated with spectrum reports exported from Scaffold to associated MS2 fragmentation scans with each frame by using customized R scripts. Another in-house-developed R package was used for the normalization of ion current intensities for each protein (by quantile ion intensities among individual runs) and the aggregation of ion intensities from frame levels to protein levels. The calculation of protein-abundance ratio, as well as the evaluation of statistical significance with Student's *t* test (unpaired) between induced and noninduced (i.e., control) groups, was conducted to determine proteins with significant changes in abundance. Detailed information about the quantitative proteomics workflow is described in several pieces of our previous works (Tu et al., 2014a,b; An et al., 2015; Shen et al., 2015).

Bioinformatics for SC pseudopod assay. The pathway and signaling networks were analyzed by using Ingenuity Pathways Analysis software (Ingenuity Systems). Lists of proteins with relative enrichment in pseudopods by neuronal membrane were associated with canonical pathways in Ingenuity Pathways Analysis. The probability of association between the dataset and the canonical pathway proteins was measured by using Fisher's exact test (Dataset 1). Networks of proteins were algorithmically generated based on their connectivity.

RT-PCR. Standard RT-PCR was performed on cDNAs derived from purified mouse DRG and purified rat SCs (gifts from Y. Sasaki, J. Milbrandt laboratory, Washington University, St. Louis, MO). For mouse *Gpr56*, the following primer pair was used to amplify a 577-bp fragment that spans exons 5–10: 5'-CAG TTACAGAGCCTGGAGTC-3' and 5'-TGTGAGTGGCTT CTACCTCT-3'. For rat *Gpr56*, the following primer pair was used to amplify a 573-bp fragment that spans exons 9–12: 5'-GAGGGCTGTGAGACTGTTAG-3' and 5'-TAGAGG GGTAGATGACATGG-3'. The following primers were used to amplify a 702-bp fragment from exons 3–6 of mouse and exons 2–6 of rat *Gapdh*: 5'-GCCATCAACGACCCCTTC ATT-3' and 5'-CGCCTGCTTCACCACCTTCTT-3'.

TEM. For standard EM analyses, tissues were drop-fixed in modified Karnovsky's fix and processed as described previously (Monk et al., 2011). Semithin sections (200 nm) were stained with toluidine blue and imaged with an AxioCam MRm on a light microscope (Zeiss AxioImager M2). Thin sections (70 nm) were treated as described for zebrafish analyses. For examination of dorsal and ventral roots, two roots each were analyzed from two WT and three *Gpr56*^{−/−} animals on P180. The number of animals examined at time point for SN analyses is as follows: P3, six independent regions (4,000×) of nerve from three WT, three *Gpr56*^{+/−}, and four *Gpr56*^{−/−}; P14, four independent regions (2,000×) of nerve from three WT, three *Gpr56*^{+/−}, and three *Gpr56*^{−/−}; P21, four independent regions (2,000×) of nerve from three WT and three *Gpr56*^{−/−}; P90, three independent regions (2,000×) of nerve from three WT, three *Gpr56*^{+/−}, and four *Gpr56*^{−/−}; P180, three independent regions (2,000×) of nerve from three WT, three *Gpr56*^{+/−}, and three *Gpr56*^{−/−}; and P300, three independent regions (2,000×) of nerve from three WT, three *Gpr56*^{+/−}, and three *Gpr56*^{−/−}.

For all experiments, myelinated fibers showing myelin outfoldings, infoldings, or demyelinating profiles were considered abnormal.

Antibodies for IHC. We used the following primary antibodies: mouse anti-Gpr56-H11 (1:1,000; Jeong et al., 2012), rabbit anti-Ki67 (1:400; Abcam; Mogha et al., 2013), rabbit anti-TUJ1 (1:1,000; Covance; Mogha et al., 2013), rat anti-MBP (1:10; AbD Serotec; Mogha et al., 2013), and mouse anti-pan sodium channel (IgG1, 1:400; Sigma-Aldrich; Saito et al., 2003). Fluorescently conjugated secondary antibodies were applied at a concentration of 1:1,000 (Invitrogen). Rhodamine-phalloidin (1:100; Invitrogen; Terada et al., 2012) was used for staining of Schmidt-Lanterman incisures.

IHC. For tissue sections, nerves were harvested and drop-fixed in 4% paraformaldehyde overnight, transferred into 30% sucrose, and then embedded in optimal cutting temperature compound. IHC was done as described previously on 7–10-μm-thick sections (Jeong et al., 2012). Stained sections were imaged with an AxioCam MRm on a light microscope

(Zeiss AxioImager M2), and analyses were performed in ImageJ software. SNs and DRG from three WT were used for expression analyses. SNs from three WT and two *Gpr56*^{+/-} mice (controls) and three *Gpr56*^{-/-} mice were used for quantification of cell proliferation on P3. For teased nerve preparations, nerves were harvested and fixed for 30 min in 4% paraformaldehyde before teasing on glass slides. Samples were then permeabilized for 15 min in acetone at -20°C before antibody staining.

RNA sequencing. Mouse SNs were harvested on P3 and P21 ($n = 4$ WT and $n = 4$ *Gpr56*^{-/-} on P3; $n = 3$ WT and $n = 3$ *Gpr56*^{-/-} on P21), flash frozen in liquid nitrogen, and stored at -80°C. Total RNA was extracted by using an RNeasy Mini kit (74104; Qiagen). First, nerves were allowed to thaw on ice for at least 5 min in Buffer RLT containing β -mercaptoethanol. During this time, small spring-loaded surgical scissors were used to cut the nerves into smaller pieces while the nerves were still in the liquid. Scissors were cleaned with 100% ethanol between each sample. Once thawed, nerves were homogenized by using a TissueRuptor for ~30–60 s followed by vortexing briefly, then passing the sample through a syringe and successively smaller needles (22 gauge and then 28 gauge) at least 10 times each or until large clumps of tissue were no longer visible. We then followed manufacturer instructions for the RNeasy Mini kit to obtain total RNA for each animal. RNA was eluted in 30 μ l elution buffer and then submitted on dry ice to the Genome Technology Access Center at Washington University in St. Louis, MO, for processing and RNA-seq. Libraries were generated by using Clontech Library Prep kits, and 12 samples were run on a single lane of Illumina 3000, 1 \times 50 bp. Approximately 20–40 million mapped reads were generated per sample. All reads were mapped with STAR (Dobin et al., 2013).

R/Bioconductor was used for subsequent analysis. Differential expression analysis was performed by using the packages, DESeq (Anders and Huber, 2010), and edgeR (Robinson et al., 2010) in R (<http://www.R-project.org>). The original p-values were corrected via false-discovery rate for multiple testing errors. In addition to a false-discovery rate of <0.05, we considered genes differentially expressed if the adjusted p-value was <0.05 and fold change >1.4 \times ($\log_2[\text{fold change}] > 0.5$). Individual P3 and P21 volcano plots and comparison between the two developmental stages were generated on normalized gene expression by using R. GO analysis was performed with MetaCore and g:Profiler.

Morphometry. G-ratios were calculated from a total of ~180 myelinated fibers per genotype (~60 fibers each from three mice per genotype) by taking the ratio of axon diameter to fiber diameter. Measurements were obtained by using ImageJ software. Internodal lengths were similarly measured from IHC images in ImageJ software. For internode analyses, a total of a total of 21 WT and 64 *Gpr56*^{-/-} fibers were measured per genotype on P21 ($n = 3$ mice per genotype). For apposi-

tion quantifications from electron micrographs, a total of 154 WT and 168 *Gpr56*^{-/-} myelinated fibers were analyzed on P21 ($n = 3$ mice per genotype). For analysis of Schmidt-Lanterman incisures on P300, a total of 104 WT fibers and 93 *Gpr56*^{-/-} fibers were analyzed ($n = 3$ mice per genotype).

Serial scanning EM and reconstruction. Nerves from P90 mice (three nerves from three WT and three *Gpr56*^{-/-} animals) were harvested and drop-fixed in modified Karnovsky's fixative overnight. Nerves were then washed into 0.1 M sodium cacodylate before staining, embedding, and imaging at Renovo Neural Inc. Approximately 400 serial sections were imaged (100 nm each in thickness) per nerve, and data were analyzed and reconstructed by using IMOD software (Kremer et al., 1996). Videos depicting data and models from serial EM analyses (Videos 1, 2, 3, and 4) were generated in ImageJ software.

Motor function. Motor function tests were performed in the Animal Behavior Core Lab at Washington University School of Medicine. The accelerated rotarod was used to assess sensorimotor function and grip strength and was performed on a cohort of 11 WT, 16 *Gpr56*^{+/-}, and 13 *Gpr56*^{-/-} mice on P180, and the same cohort was used for testing on P300 ($n = 10$ WT, $n = 15$ *Gpr56*^{+/-}, and $n = 12$ *Gpr56*^{-/-}) as described previously (Idol et al., 2014).

Electrophysiology. Animals (WT, $n = 5$; *Gpr56*^{+/-}, $n = 12$; *Gpr56*^{-/-}, $n = 11$) were anesthetized by using 4% isoflurane/96% oxygen (induction) and 2% isoflurane/98% oxygen (maintenance) administered by inhalation. After preparation of the sterile field, the right SN was isolated from the sciatic notch to below the point of trifurcation, distal to the popliteal artery. Cathodic, monophasic electrical impulses (duration = 50 μ s, frequency = single, amplitude = 1 mA) were generated by a single-channel isolated pulse stimulator (model 2100; A-M Systems Inc.) and delivered to the SN proximal to the implanted device via bipolar platinum/iridium microwire electrodes (Medwire). Resulting EMGs were then differentially recorded in the tibialis anterior muscle by using intramuscular microwire electrodes (4 mil; California Fine Wire). Measured signals were band-pass filtered (low pass = 1 Hz, high pass = 5 kHz, notch = 60 Hz) and amplified (gain = 1,000–10,000 \times) by using a two-channel microelectrode AC amplifier (model 1800; A-M Systems, Inc.) before being recorded on a desktop PC (Dell Computer Corp.) equipped with a data acquisition board (DT3003/PGL; Data Translations). EMG traces were digitally recorded while peak latency and maximal amplitude were calculated by using custom Matlab software (The MathWorks, Inc.).

Western blot and GTP-Rho pull-down assay. For analysis of RhoA activity in mouse nerves, P6 litters consisting of *Gpr56*^{+/+}, *Gpr56*^{+/-}, and *Gpr56*^{-/-} mice were used in each of three separate experiments (we combined four

nerves from the same genotype for one pull-down and did three separate experiments, with a total of 12 for each genotype group). P6 SNs were dissected under a Leica stereo microscope (MZ 6; Leica Pte Ltd.); four nerves from same the genotype were combined in one group, powdered on liquid nitrogen, then lysed in ice-cold RIPA buffer (1% Nonidet P-40, 50 mM Tris, pH 7.6, 120 mM NaCl, and 1 mM EDTA) containing protease inhibitor cocktail set 1 (Calbiochem). The lysates were cleared of insoluble materials by centrifugation at 16,000 *g* for 10 min at 4°C. The protein concentration was determined by a Bio-Rad protein assay method (Bio-Rad) according to the manufacturer's protocol, and equal amounts of protein were used for each pull-down assay. The GTP-Rho pull-down assay was performed as previously described (Luo et al., 2011). In short, tissues were pulverized on liquid nitrogen, lysed in 300 μ l of ice-cold RIPA buffer containing protease inhibitors with a cell disruptor for 10 min and homogenized with a 26-gauge syringe needle. Equal amounts of total protein were incubated with 60 μ g GST-RBD beads (Cytoskeleton) at 4°C for 90 min. The beads were washed twice with lysis buffer and once with Tris-buffered saline buffer. Bound Rho proteins were eluted by Laemmli sample buffer and detected by Western blot by using mouse monoclonal anti-RhoA antibody (ARH04, 1:1000; Cytoskeleton). For analysis of GPR56-dependent RhoA signaling in vitro, HEK 293T cells were plated in 6-well plates (500,000 cells/well) and transfected with mouse *Gpr56* (Luo et al., 2014) or not. 24 h posttransfection, cells were subjected to serum starvation for 36 h followed by treatment with human collagen III (84 nM, CC054, positive control; Sigma-Aldrich) or full-length recombinant FT (res. 23–110, 10 μ M) and as a control with either 0.5 M acetic acid or PBS, respectively, for 5 min. Cells were lysed in lysis buffer (PBS + 1% Triton X-100 + 1 \times protease inhibitors) followed by pull-down of RhoA-GTP (Luo et al., 2011). We also blotted for GPR56 (ab172361, concentration of 1 μ g/ml; Abcam) to assess the success of transfection and actin (MAB1501, 200 ng/ml antibody and c concentration; Chemicon) for a loading control. Data are representative of three biologically independent experiments.

Gpr56 PRESTO-Tango assay. HTLA cells (a HEK293 cell line stably expressing a tTA-dependent luciferase reporter and a β -arrestin2-TEV fusion gene) were provided by the laboratory of B. Roth (University of North Carolina, Chapel Hill, NC; Kroeze et al., 2015). Cells were plated in six-well plates (500,000 cells/well) and transfected with either *Gpr56* Tango plasmid (Addgene) or as a control with pCDNA3 plasmid. 24 h posttransfection, cells were subjected to serum starvation for further 24 h followed by treatment with human collagen III (150 nM; CC054; Sigma-Aldrich) or full-length recombinant FT (res. 23–110, 10 μ M), and as a control, with either 0.5 M acetic acid or PBS, respectively, overnight. Cells were washed with PBS and processed with Bright-Glo luciferase

system (E2610; Promega) according to manufacturer's protocol. 200 μ l of Bright-Glo solution, diluted in the assay buffer, was added to 25 μ g of cell lysate, and followed by 20 min of incubation, luciferase activity was measured (as a function of chemiluminescence). Data are representative of three biologically independent experiments.

cAMP assay. cAMP assay was done with a colorimetric competitive immunoassay (Enzo Life Sciences) as described previously (Küffer et al., 2016). In brief, HEK 293T cells were transfected with either human *GPR126* (cloned into pCDNA3.1/V5-His-TOPO vector) or mouse *Gpr56* (cloned into pCDNA3.1 vector) and 48 h posttransfection were treated with either PBS or full-length recombinant FT (10 μ M) and then processed for quantification of cAMP. Data are representative of three biologically independent experiments.

Co-IP and immunoblotting. Co-IP and immunoblotting were done as previously described (Luo et al., 2011). In brief, P5 mouse SNs ($n = 8$ nerves from four animals for each co-IP) were lysed in RIPA buffer as described for the GTP-RhoA pull-down assay. After extraction at 4°C with rocking, insoluble material was removed by centrifugation, and lysates were precleared for 1 h at 4°C with protein G-Sepharose (101241; Invitrogen). The lysate was equally divided into two; half of the lysate was used for GPR56 IP, and the other half was used for the control IP. Protein G beads were then used to pull-down the GPR56NTF-hFc protein complex. Immuno-complexes were subjected to SDS-PAGE and Western blot by using pan-plectin antibody (MAB5674; Sigma-Aldrich), and rabbit anti-human IgG Fc antibody (31142; Thermo Fisher Scientific) following standard protocols.

Plectin expression assay. P6 mouse SNs ($n = 12$ nerves from six animals per genotype) were collected and lysed as described for the GTP-RhoA pull-down assay. Plectin levels were assessed by Western blot by using a pan-plectin antibody (MAB5674; Sigma-Aldrich). Two nerves were pooled from each animal to make up a single replicate.

Statistical analyses

Analyses were performed blinded to genotype and treatment. All data are presented as the mean \pm SEM unless otherwise noted. To represent significance, *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$; ****, $P < 0.0001$. A minimum of two technical replicates was performed for each described experiment. Although we did not perform statistical tests to predetermine sample size, our samples sizes are similar to previously published studies in the field. Statistical analyses were performed by using one-way ANOVA, two-way ANOVA, or unpaired Student's *t* test with two tails with unequal variance when required conditions were met. A departmental statistician was consulted to confirm the appropriateness of each statistical analysis performed.

Online supplemental material

Fig. S1 shows IHC analysis of cellular proliferation and the total cell number in P3 mutant and control mouse SNs as well as an assessment of overall saphenous nerve size by toluidine blue staining, which can be affected by cell number. Fig. S2 shows TEM images assessing pLLn integrity and myelin ultrastructure in *gpr56^{std13/std13}* zebrafish mutants during development and in adulthood to show conservation with mouse phenotypes. Fig. S3 shows RNA-seq analysis of the transcriptome of *Gpr56* mutant and WT control nerves on P3 and P21. Fig. S4 Shows further structural analysis of *Gpr56* mutant nerves: Schmidt Lanterman incisures quantification by IHC, toluidine blue, and TEM images of *Gpr56^{-/-}* dorsal and ventral roots compared to control roots on P180. Analysis of *Gpr56* expression in neurons versus SCs is shown. Fig. S5 shows additional TEM reconstructions to supplement observations in Fig. 6. Video 1 shows serial reconstruction of a myelinated fiber from a WT mouse SN on P90 generated from 350 successive scanning EM images (100 nm thick). Video 2 shows serial scanning EM images (210 sections shown, 100 nm thick) from a WT mouse SN on P90. Video 3 shows serial reconstruction of a myelinated fiber from a *Gpr56^{-/-}* mutant mouse SN on P90 generated from 350 successive scanning EM images (100 nm thick). Video 4 shows serial scanning EM images (210 sections shown, 100 nm thick) from a *Gpr56^{-/-}* mutant mouse SN on P90. Dataset 1 is a summary of all proteins significantly upregulated in SC pseudopods after induction and analysis of the major cellular pathways found to be enriched. Dataset 2 is a summary of gProfiler results for significantly enriched or depleted pathways in *Gpr56* mutant nerves based on RNA-seq analyses on P3 and P21. Raw data files can be accessed from Zenodo (<https://zenodo.org/record/1154250>).

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Author contributions: Y. Poitelon and M.L. Feltri designed the SC pseudopods assay, which was executed by Y. Poitelon, J. Li, and J. Qu. A.K.K. Lakkaraju and A. Aguzzi designed the PrP experiments, which were performed by A.K.K. Lakkaraju. W.Z. Ray and M.R. MacEwan designed the electrophysiology experiments, which were performed by M.R. MacEwan and P. Gamble. The remaining experiments were designed

by S.D. Ackerman, X. Piao, and K.R. Monk and executed by S.D. Ackerman, R. Luo, A. Mogha, M. D'Rozario, N.E. Sanchez, and B.L. Hartly. S.D. Ackerman and K.R. Monk wrote the manuscript, which was read and edited by all authors.

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