Peripherally derived FGF21 promotes remyelination in the central nervous system

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Demyelination in the central nervous system (CNS) leads to severe neurological deficits that can be partially reversed by spontaneous remyelination. Because the CNS is isolated from the peripheral milieu by the blood-brain barrier, remyelination is thought to be controlled by the CNS microenvironment. However, in this work we found that factors derived from peripheral tissue leak into the CNS after injury and promote remyelination in a murine model of toxin-induced demyelination. Mechanistically, leakage of circulating fibroblast growth factor 21 (FGF21), which is predominantly expressed by the pancreas, drives proliferation of oligodendrocyte precursor cells (OPCs) through interactions with β-klotho, an essential coreceptor of FGF21. We further confirmed that human OPCs expressed β-klotho and proliferated in response to FGF21 in vitro. Vascular barrier disruption is a common feature of many CNS disorders; thus, our findings reveal a potentially important role for the peripheral milieu in promoting CNS regeneration.

Results
OPC proliferation exhibits kinetics similar to those of vascular barrier disruption. We investigated the time changes in vascular permeability and OPC proliferation in the spinal cord following injection of lysophosphatidylcholine (LPC), which perturbs vascular barrier-
er (18) and myelin structures (14, 25) with no change of neuronal number (Supplemental Figure 1A; supplemental material available online with this article; https://doi.org/10.1172/JCI94337DS1). Histological analysis revealed the presence of extravasated cadaverine around the site of LPC injection 3 and 7 days after the operation (Figure 1A). In mice that underwent LPC injection, similar kinetics were observed for extravasation of endogenous IgG around the LPC injection site (Figure 1A and Supplemental Figure 1B) and Evans blue leakage into the spinal cord (Figure 1B). Consistent with this, IHC analysis revealed that the number of PDGFRα+ cells (proliferating OPCs) around the site of LPC injection increased significantly 3 and 7 days after the operation (Figure 1, C and D).

Because OPC proliferation is correlated with remyelination (26, 27), we investigated the change over time in the size of the myelin basic protein–positive (MBP+) area in the dorsal column of the spinal cord following LPC injection. IHC evaluation revealed loss of myelin in the spinal cord 3 and 7 days after LPC injection; however, this loss was partially restored 7 and 14 days after LPC injection, respectively (Figure 1, E and F). Consistent with this, remyelination after removal of the cuprizone diet, which does not cause BBB disruption (28, 29), was less efficient than remyelination after LPC injection (Figure 1, G and H, and Supplemental Figure 1C). These data suggest that vascular barrier disruption is synchronized with OPC proliferation, which precedes remyelination.

**Circulating FGF21 promotes OPC proliferation.** Because vascular barrier disruption causes leakage of circulating factors into the CNS (16), we hypothesized that the leaked circulating factors stimulate OPC proliferation, which in turn contributes to remyelination. To test this idea, we collected adult mouse serum and asked whether serum treatment would promote OPC proliferation in vitro. BrdU incorporation assays revealed that serum treatment promoted DNA synthesis in OPCs (Figure 2A and Supplemental Figure 2A, A and B), suggesting that adult serum contains an OPC proliferation factor(s). The BrdU incorporation activity was almost completely abolished by heat or Pronase treatment (Figure 2B), indicating that the OPC proliferation factor in adult mouse serum must be a heat-sensitive protein. A pharmacological screen revealed that 2 FGF receptor (FGFR) inhibitors, PD173074 and NF449 (30, 31), blocked serum–promoted OPC proliferation (Figure 2, C and D, Supplemental Figure 2C, and Supplemental Table 1), suggesting that the OPC proliferation factor in adult mouse serum belongs to the FGF family. We then sought to determine which FGFRs are involved in serum–promoted OPC proliferation. Knockdown of FGFR1 and FGFR3 in OPCs prevented serum–mediated enhancement of BrdU incorporation (Figure 2E and Supplemental Figure 2D).

As the candidate molecule appears to act on both FGFR1 and FGFR3, we focused on FGF21, which is involved in FGFR1 and FGFR3 signaling (23, 32). Because FGFR signaling mediated by FGF21 requires β-klotho, an essential receptor of FGF21, we asked whether β-klotho was required for serum–mediated OPC proliferation. β-Klotho knockdown in OPCs shows lower proliferation activity after serum treatment compared with control OPCs (Figure 2E and Supplemental Figure 2D), and this inhibitory effect was comparable to that in FGFR1 knockdown (P value for interaction = 0.8828) and FGFR3 knockdown (P value for interaction = 0.855). These data indicate that the receptor for FGF21 is involved in serum–mediated OPC proliferation. To test whether FGF21 promotes OPC proliferation directly, we used recombinant FGFs (FGF15 [the mouse ortholog of human FGF19], FGF21, and FGF23), which act as endocrines (33). FGF21 treatment promoted BrdU incorporation into OPCs (Figure 2F), whereas FGF15 and FGF23 did not (Figure 2F). Moreover, neutralizing antibody against FGF21 abolished the serum–mediated increase in BrdU incorporation by OPCs (Figure 2G). Furthermore, FGF21 did not promote BrdU incorporation in astrocytes (Supplemental Figure 3A). We detected no significant difference in MTT absorbance in OPCs following β-klotho knockdown (Supplemental Figure 3B). Differentiation into MBP+ cells was not altered by β-klotho knockdown (Supplemental Figure 3C) and FGF21 treatment (Supplemental Figure 3D). We detected no significant difference in OPC proliferation cultured in astrocyte supernatant with or without serum pretreatment (Supplemental Figure 3E). These data suggest that FGF21 in adult mouse serum acts directly as an OPC proliferation factor.

**Pancreas-derived FGF21 mediates OPC proliferation.** Next, we investigated the expression pattern of FGF21 in adult mice. Quantitative mRNA and protein analysis revealed prominent FGF21 expression in pancreas (Figure 3A), as reported previously (34, 35). IHC analysis revealed abundant FGF21 expression in pancreatic islets, especially in glucagon-producing α cells (Figure 3, B and C), which are endocrine cells. To determine whether pancreatic FGF21 is involved in serum–mediated OPC proliferation, we specifically knocked down FGF21 in vivo by transfecting FGF21 siRNA into pancreas (36) (Supplemental Figure 4A), and then collected serum from the transfected mice. BrdU incorporation analysis revealed that serum from pancreas–specific FGF21-knockdown mice exerted weaker proliferative activity than control serum (Figure 3D and Supplemental Figure 4B). These data indicate that pancreas–derived FGF21 is involved in promotion of OPC proliferation mediated by circulating factors in vitro.

**Circulating FGF21 drives remyelination.** We next asked whether circulating FGF21 promotes OPC proliferation and remyelination in vivo. To test this idea, we monitored the FGF21 level around demyelinating lesions following LPC injection, because leakage of circulating factor into the CNS could be detected around the lesion site with BBB disruption (37). Following LPC injection, FGF21 protein in the spinal cord was present at a higher level than in the control (Figure 4A), although the level of FGF21 mRNA in the spinal cord was unchanged (Supplemental Figure 5A). Fluorescent dye–labeled FGF21 by i.v. administration was detected in the spinal cord of mice after LPC injection (Supplemental Figure 5B). In addition, in FGF21-KO mice, LPC injection caused FGF21 detection in the spinal cord of mice that received recombinant FGF21 intravenously (Supplemental Figure 5C). These results indicate that the elevated FGF21 level around demyelinating lesions is a result of influx from circulating FGF21 into the CNS.

To determine whether in vivo OPC proliferation is promoted by circulating FGF21, we injected LPC into the spinal cords of FGF21-KO mice. The absence of FGF21 prevented the increase in PDGFRα+ Ki67+ cell number around the site of LPC injection (Figure 4B). Furthermore, the MBP+ demyelination area in the dorsal column of the spinal cord was larger in FGF21-KO mice than in control littermates (Figure 4C). Under intact conditions, there was no significant difference between FGF21-KO mice and control lit-
OPCs in the spinal cord tissues from individuals with multiple sclerosis (Figure 6A). We also found that treatment with recombinant human FGF21 increased BrdU incorporation in human OPCs in vitro (Figure 6B), implying that FGF21-mediated OPC proliferation is conserved in humans.

**Discussion**

Peripheral tissue–derived factors promote OPC proliferation in the CNS (Supplemental Figure 9). Our findings suggest a novel paradigm for the mechanism of CNS regeneration, which has previously been considered to primarily involve the CNS microenvironment. Because OPC proliferation is just one process of CNS regeneration, the influence of peripheral tissue–derived factors on other regen-

**OPC proliferation is promoted under vascular disruption. (A)** Representative images of spinal cord sections visualized with Alexa Fluor 488–conjugated cadaverine (top) and endogenous IgG (bottom). Spinal cord sections were prepared at the indicated times after LPC injection. **(B)** Quantitation of Evans blue leakage in the spinal cord (n = 3); **P < 0.01.** **(C)** Representative images of spinal cord sections labeled for PDGFRα (top) and Ki67 (middle). Spinal cord sections were prepared at the indicated days after LPC injection. Arrows indicate cells that are colabeled with PDGFRα and Ki67. **(D)** Graph shows quantitation as indicated in each image (n = 3 for control, 3 for d3, 4 for d7, 4 for d14); **P < 0.05.** **(E)** Representative images of spinal cord sections labeled for MBP. Spinal cord sections were prepared at the indicated days after LPC injection. **(F)** Graph shows quantitation as indicated in each image (n = 3 for control, 3 for d3, 4 for d7, 4 for d14); P = 0.002, 0.0424, 0.0018 (left to right); **P < 0.05.** **(G)** Representative images of brain sections visualized with MBP (top) and endogenous IgG (bottom) the indicated number of days after LPC injection. **(H)** Graph shows the time course of demyelination after LPC injection (n = 3 for control, 6 for d3, 7 for d7, 4 for d14); **P < 0.05.** **(I)** Error bars represent SEM. Scale bars: 200 μm (A and B); 50 μm (C); 100 μm (G).

terminates in PDGFRα+ OPC number, PDGFRα+ BrdU+ cell number, or myelin formation (Supplemental Figure 5, D–F). We confirmed that FGF21-KO mice did not exhibit a significant increase in demyelination after LPC injection (Supplemental Figure 6A). There was no significant difference between groups in GFAP+ cell number in the spinal cord 7 days after LPC injection (Supplemental Figure 6B). These data imply that peripheral tissue–derived factors on other regen-

To confirm that these observations rely on the degree of structural remyelination, we calculated the g-ratio, which is positively correlated with the thickness of myelin sheaths. A typical g-ratio for a normally myelinated axon is between 0.6 and 0.8, whereas a g-ratio of 1.0 indicates complete demyelination. FGF21-knockdown mice exhibited few signs of remyelination relative to con-

We also investigated the possibility that circulating FGF21 acts directly on remyelination in vivo. To this end, we injected LPC into the spinal cord of FGF21-KO mice, with or without intrathecal FGF21 treatment. IHC analysis revealed that the mice subjected to intrathecal administration of FGF21 had reduced demyelination around the site of LPC injection, relative to control mice (i.e., mice not subjected to FGF21 treatment) (Figure 4F). We also found that FGF21 treatment promoted reduction of demyelination around the site of LPC injection in WT mice (Figure 4G). These data raise the possibility that circulating FGF21 acts directly on OPCs in the CNS, resulting in remyelination.

**β-Klotho in OPCs is involved in remyelination.** We next inves-

Because FGF21-mediated signaling is required for the expression of β-klotho (21–24), which forms a complex with FGFRs, we examined β-klotho expression in OPCs after LPC injection. IHC analysis revealed that β-klotho expression in PDGFRα+ OPCs was elevated after LPC injection (Figure 5A and Supplemental Figure 7A). β-Klotho expression in proliferating OPCs was higher than that in quiescent OPCs (Supplemental Figure 7B). β-Klotho expression was also detected in O4– immature and mature oligodendrocytes and in Adenomatous polyposis coli (APC)–positive mature oligodendrocytes in the spinal cord after LPC injection (Supplemental Figure 7C).

To determine whether OPC proliferation depends on β-klotho in OPCs, we generated a conditional knockout mouse in which β-klotho is knocked down specifically in PDGFRα+ OPCs (Klb−/− Pdgfra-CreERT mice; Supplemental Figure 7D). We conducted LPC injection into the spinal cord of the conditional knockout mice and performed histological analysis to evaluate OPC number and demyelination area in the spinal cord. There were fewer PDGFRα+ cells in the spinal cords of conditional knockout mice than in spinal cords of control mice 7 days after LPC injection (Figure 5B). These conditional knockout mice also had a larger MBP area in the dorsal spinal cord than control littermates (Figure 5C), indicating that β-klotho expression in OPCs is involved in OPC proliferation and subsequent remyelination. We also found that FGF21 admin-

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Because demyelination is a common feature of many CNS disorders, including traumatic brain injury (39), we conducted controlled cortical injury to evaluate the suitability of FGF21 as a therapeutic target with broad pathological significance. Intracerebroventricular administration of FGF21 increased the number of PDGFRα+Ki67+ cells around the lesion 7 days after brain injury (Figure 5D and Supplemental Figure 8B). The MBP area in the corpus callosum was smaller in FGF21-treated mice than in control mice (Figure 5E). Furthermore, electron microscopy also revealed an increase in structural remyelination by FGF21 treatment after brain injury (Figure 5F). These results suggest that FGF21 promotes remyelination in many CNS pathological states that are characterized by demyelination.

Finally, we confirmed that our findings may apply to humans. IHC analysis revealed enhanced β-klotho expression in PDGFRα+ OPCs in the spinal cord tissues from individuals with multiple sclerosis (Figure 6A). We also found that treatment with recombinant human FGF21 increased BrdU incorporation in human OPCs in vitro (Figure 6B), implying that FGF21-mediated OPC proliferation is conserved in humans.
Thus, CNS entry of peripheral FGF21 is limited in normal adult subjects. Indeed, we observed no significant difference in OPC number between WT and Fgf21−KO mice, indicating that FGF21 does not control OPC proliferation under normal conditions. In the CNS, FGF21 regulates metabolism, circadian behavior, and neuroendocrine control of female reproduction via direct action on hypothalamus (32, 45), which structurally lacks a BBB (46). This anatomical feature also supports our hypothesis: peripheral FGF21 acts on the CNS directly under certain situations, such as pathological lesions with vascular barrier disruption or normal tissues with high vascular permeability.

Our culture experiments suggested that β-klotho is involved in OPC proliferation. β-Klotho is widely believed to be a simple coreceptor required for FGF21 binding to FGFRs (21–24), which are involved in signal transduction. All FGFRs mediate cell proliferation (47), and conditional knockout mice lacking FGFR1/FGFR2 in OPCs show decreased oligodendrocyte repopulation in a chronic demyelination model (48). This finding is consistent with our results that FGFR1 is involved in serum-mediated OPC proliferation. Regarding FGFR3, FGFR3-null mice exhibit reduced oligodendrocyte number during development, whereas this change is not due to reduction of oligodendrocyte progenitor proliferation mechanisms, including oligodendrocyte development and neuronal plasticity, remains an open question. Regarding the duration of vascular barrier disruption after CNS injury, we observed that vascular permeability was significantly recovered 14 days after LPC injection, relative to 7 days after injection. These kinetic observations are supported by other reports that restoration of vascular barrier in CNS lesions begins within a week after injury (40, 41). By contrast, our in vivo experiment revealed extension of the MBP-expressing area 2 weeks after LPC injection. Considering the relative kinetics of vascular barrier disruption and remyelination, it is conceivable that peripheral tissue–derived factors play a strongly supportive role during early regeneration mechanisms such as OPC proliferation, at a time when there is abundant extravasation of blood components into the CNS.

In this context, we found that FGF21, which is predominantly expressed from pancreas, is crucial for peripheral tissue–mediated OPC proliferation. The results of this study reveal an unexpected role of FGF21, which has been previously characterized as a metabolic regulator (42). In reviewing previous findings regarding FGF21 function in the CNS, we noted that FGF21 can cross the BBB (43), but the FGF21 level in the cerebrospinal fluid of healthy patients is approximately 40% of that in the plasma (44). Thus, CNS entry of peripheral FGF21 is limited in normal adult subjects. Indeed, we observed no significant difference in OPC number between WT and Fgf21−KO mice, indicating that FGF21 does not control OPC proliferation under normal conditions. In the CNS, FGF21 regulates metabolism, circadian behavior, and neuroendocrine control of female reproduction via direct action on hypothalamus (32, 45), which structurally lacks a BBB (46). This anatomical feature also supports our hypothesis: peripheral FGF21 acts on the CNS directly under certain situations, such as pathological lesions with vascular barrier disruption or normal tissues with high vascular permeability.

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As developmental myelination occurs at postnatal weeks 2–3 when the BBB is already formed, FGF21 in blood does not penetrate into the CNS, and therefore FGF21 level in the CNS should be much lower than that in the blood. In contrast, our in vitro experiments mimic the pathological condition in the CNS that suffered exposure of blood components (including FGF21) by disruption of BBB, which is formed during the embryonic stage. The seemingly contradictory findings in the effects of FGFR3 inhibition on OPC proliferation during the developmental stage in vivo and in our in vitro condition may be due to the absence or presence of ligands for FGFR3, including FGF21. We note that FGFR1 and FGFR3 are expressed in OPCs in vivo (50–52), and our in vitro experiments reveal that serum-mediated OPC proliferation is dependent on FGFR1 and FGFR3. In addition, FGFR3 expression in the white matter of the spinal cord is elevated in the experimental demyelinating disease (53), and FGFR3 is considered to promote proliferation of oligodendrocyte progenitors (54). Therefore, we do not exclude the possibility that FGFR3 might be another component necessary for the signal transduction of FGF21 specifically in the pathological condition in the CNS.
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**Figure 4. Circulating FGF21 drives OPC proliferation.** (A) Quantitation of FGF21 protein in the spinal cord 1 day and 3 days after LPC injection (n = 5 for control, 5 for d1, 4 for d3). (B) Representative images of spinal cord sections double-labeled for PDGFRα and Ki67. Sections were obtained from FGF21-KO mice and control littermates 7 days after LPC injection. Graph shows quantitations as indicated in the images (n = 5). Arrows indicate Ki67+ cells colabeled with PDGFRα; arrowheads indicate Ki67− cells not labeled with PDGFRα. (C) Representative images of spinal cord sections labeled for MBP. Sections were obtained from mouse spinal cord 14 days after LPC injection. Graph shows quantitations as indicated in the images (n = 5). (D) Representative immunoelectron microscopy images of myelin in the spinal cord. Sections were obtained from FGF21-KO mice and control littermates 14 days after LPC injection. Graphs show quantitations of g-ratio indicated in the images (n = 3). (E) Motor function was assessed by ladder-walk test (n = 11 for control littermates, 9 for FGF21-KO mice). (F) Representative images of spinal cord sections labeled for MBP. Graph shows quantitations as indicated in the images (n = 5 for control littermates + vehicle, 5 for FGF21-KO mice + vehicle, 4 for FGF21-KO mice + FGF21). (G) Representative images of spinal cord sections labeled for MBP. Graph shows quantitations as indicated in the images (n = 4); **P < 0.01 as determined by Student’s t test or by ANOVA with Tukey’s post hoc test or Dunnett’s test. *P < 0.05, **P < 0.01. Error bars represent SEM. Scale bars: 50 μm (B); 200 μm (C, F, and G); 2 μm (D).

We should note that FGF21-mediated OPC proliferation is one of the mechanisms of remyelination. In terms of molecular mechanism, we just focused on the direct action of FGF21 on OPC proliferation; however, FGF also regulates expression of VEGF receptor 2 (55). Because VEGF signaling is related to brain homeostasis, including OPC migration, a process that involves remyelination (56, 57), an indirect effect of FGF21 on OPCs may also contribute to oligodendrocyte development and remyelination. There is also some possibility that other molecules, which are not targeted by the inhibitors that use pharmacological screening, are involved in the serum-mediated OPC proliferation. Future studies should seek to reveal novel molecular mechanisms of remyelination based on the concepts proposed in this study.

Meanwhile, FGF21-associated drugs for treating diabetes have recently been developed by pharmaceutical companies, and some of these compounds have reached the stage of clinical trials (58). In this study, we observed that FGF21 treatment also promotes human OPC proliferation. Therefore, we believe that these FGF21-associated drugs may exert FGF21-mediated remyelination effect and provide clinical benefits in patients with CNS demyelination.

The findings of this study emphasize the novel concept that the peripheral milieu controls CNS regeneration. It is widely believed that CNS regeneration occurs rarely in adults because of the age-dependent decline in regenerative capacity (2), which can be partially explained by a reduction in cell-intrinsic plasticity (59). Several recent studies suggested that age-related decline of neuronal plasticity depends on an aged systemic milieu (60–62). The results of these studies predict that a young peripheral milieu should enhance regenerative capacity in the adult CNS. Although our study is currently limited to focusing on a circulating molecule in the adult systemic milieu, infiltrating peripheral macrophages in the youthful systemic environment are known to enhance spinal cord remyelination (17). Future studies focusing on the young systemic milieu may contribute to the development of methods for CNS regenerative medicine.

**Methods**

**Mice.** All experimental procedures were approved by the Institutional Animal Care Committee of Osaka University. All experiments were run in a blinded fashion. Female C57BL/6 mice were obtained from Charles River Japan, Japan SLC, or CLEA Japan. Fgf21-KO mice (C57BL/6 background) were previously described (63). *Pdgfra-CreERT* mice (C57BL/6 background) were purchased from the Jackson Laboratory. β-Klotho-floxed mice (Klbfl/fl; C57BL/6 background) were provided by Steven Kliewer (University of Texas Southwestern Medical Center). Mice were housed in an air-conditioned room at 23°C ± 2°C with a 12-hour light–dark cycle, and had free access to water and food.

The OPC-specific β-klotho deletion mice were obtained by crossing of the β-klotho-floxed mice with the *Pdgfra-CreERT* mice. Cre recombination in the generated mice (7–8 weeks) was induced by administration of tamoxifen (75 mg/kg, i.p.; Sigma-Aldrich) daily for 5 consecutive days. Ten days after the first tamoxifen administration, mRNA in PDGFRα cells in brain was obtained from the Cre+/−:floxed/floxed mice and −/−:floxed/floxed mice to assess the efficiency of β-klotho deletion. Relative β-klotho expression in PDGFRα-labeled cells was assessed by immunohistological analysis.

**Surgical procedure.** Focal demyelination in the brain was induced by stereotactic infusion of 1% (wt/vol) lysophosphatidylcholine (LPC, Sigma) dissolved in PBS. LPC was slowly injected into the corpus callosum of adult female mice (2 mm posterior to bregma, midline, depth of 2 mm, 2 μl per site) using a glass capillary attached to a microsyringe.

To induce demyelination in the spinal cord, adult female mice underwent laminectomy at Th11–12 and received injections of 2 μl of 1% (wt/vol) LPC in the dorsal column midline at a depth of 0.5 mm. For administration of recombinant FGF21, a cannula from an Alzet osmotic pump (model 1007D, Alzet Corp.) was placed under the dura at the thoracic spinal cord immediately after LPC injection. The pump was filled with vehicle solution (PBS containing 0.5% BSA) or recombinant mouse FGF21 (50 ng/kg of body weight per day for 1 week) and implanted s.c. into the back. Administration was continued until the end of the study.

To create traumatic brain injury, adult female mice were stabilized in a stereotactic frame (Narisilige) after deep anesthesia. A midline incision was made in the scalp, and the fascia was retracted to expose the cranium. A circular craniotomy opening 4 mm in diameter was created on the left side, with the center at 0 mm anteroposterior and 2 mm lateral to the bregma. Controlled cortical impact was produced with a Pneumatic Impact Device (AmScien Instruments) using a 3-mm-diameter flat-tip, as described previously (64). The impact parameters were 4.0–4.5 mm/ms velocity, 1 mm depth, and 120 ms duration time. The scalp was then sutured and closed, and the mice were left to wake from the anesthesia. For infusion of recombinant FGF21, a 28-gauge stainless steel cannula connected to an Alzet osmotic pump (model 1002, Alzet Corp.) was guided to the lateral ventricle (coordinates: 0 mm bregma, 1 mm lateral, and 2.5 mm ventral). The pump was filled with vehicle solution (PBS containing 0.5% BSA) or recombinant mouse FGF21 (50 ng/kg of body weight per day for 2 weeks) and implanted s.c. into the back.

**Demyelination by cuprizone diet.** Mice were placed on a diet of 0.2% (wt/wt) cuprizone mixed into chow pellets. Mice were exposed to continuous cuprizone feeding for 12 weeks, and then returned to normal chow pellets. Mice were maintained in sterile, pathogen-free conditions.

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Vascular barrier permeability assays. Mice were transcardially perfused with 4% paraformaldehyde (PFA) in PBS. Spinal cord tissues were immersed in 30% sucrose in PBS overnight at 4°C. Cryosections (30-μm thickness) were mounted on Matsunami adhesive silane-coated slides (Matsunami Glass) and permeabilized in PBS containing 0.05% Tween-20 and 2% BSA for 1 hour at room temperature. To detect IgG extravasation, sections were incubated with Alexa Fluor 568-conjugated goat anti-mouse IgG (1:500; Invitrogen, A-11004) for 1 hour at room temperature. In another set of experiments, mice were injected i.v. with 0.1 ml of Alexa Fluor 488-conjugated cadaverine at a concentration of 1 mg/ml (Life Technologies, A30676) 2 hours before fixation. Spinal cord tissues were immersed in 30% sucrose in PBS for 24 hours at 4°C, after which they were sectioned at 30-μm thickness on a cryostat and mounted on Matsunami adhesive silane-coated slides. All samples were analyzed by laser scanning confocal microscopy (Olympus FV1200).

Evans blue leakage. Evans blue dissolved at 5 mg/ml in saline was injected into the tail vein of mice at a dose of 10 mg/kg 24 hours before the end of breeding. Mice were intracardially perfused with PBS, and thoracic spinal cord at the level of Th11–12 was dissected out following immersion for 48 hours at 4°C in 200 μl of formamide. The supernatants from the samples were collected, and the concentration of Evans blue in the supernatants was determined spectrophotometrically at 620 nm.

Immunohistochemistry. At 0, 3, 7, or 14 days after LPC injection, mice were transcardially perfused with PBS followed by 4% PFA in 0.1 M phosphate buffer (PB). Brain, spinal cord, and pancreas were removed and were postfixed with 4% PFA in PB for 1 hour (for PDGFRα staining) or overnight at 4°C following immersion in 30% sucrose in PBS. Tissues were embedded in OCT compound (Tissue-Tek), and then 30-μm sections were cut and mounted on Matsunami adhesive silane-coated slides (Matsunami Glass). To detect BrdU-labeled cells, the sections were pretreated with 2N HCl for 30 minutes at 37°C and washed in 0.1 M borate buffer (pH 8.5) for 10 minutes. The sections were permeabilized with PBS containing 0.3% Triton X-100 and 10% Goat Serum (Sigma-Aldrich) (for PDGFRα staining) or PBS containing 0.1% Triton X-100 and 5% BSA for 1 hour at room temperature. The sections were then incubated with primary antibodies overnight at 4°C, and then incubated with fluorescently labeled secondary antibody for 1 hour at room temperature. The primary antibodies used were as follows: rat anti-mouse PDGFRα (1:400; 558774, BD Biosciences), rabbit anti-human Ki67 (1:200; ab16667, Abcam), goat anti-human MBP (1:500; sc-13914, Santa Cruz Biotechnology), mouse anti-BrdU (1:100; M0744, Dako), mouse anti-bovine O4 (1:250; MAB345, Millipore), rabbit anti-human Adenomatous polyposis coli (APC) (1:250; sc-896, Santa Cruz Biotechnology), rabbit anti-human β-klotho (1:300; HPA021136, Sigma-Aldrich), rabbit anti-human FGF21 (1:200; ab64857, Abcam), mouse anti-porcine glucagon (1:500; G2654, Sigma-Aldrich), mouse anti-human insulin (1:500; ab6995, Abcam), and rat anti-human somatostatin (1:200; MAB354, Millipore). Alexa Fluor 488- or 568-conjugated goat antibody against rabbit IgG, goat antibody against rat IgG, goat antibody against mouse IgG, goat antibody against mouse IgM, and donkey antibody against goat IgG were used as secondary antibodies. In another set of experiments, we also used Alexa Fluor 647-conjugated mouse anti-human Ki67 antibody (1:100; 558615, BD Biosciences) and Alexa Fluor 488-conjugated anti-NeuN antibody (1:100; MAB377X, Millipore). We validated all antibodies for use in this study. To estimate the numbers of proliferating and total OPCs, the sections were immunostained with PDGFRα-specific and Ki67-specific antibodies. The numbers of PDGFRα+ and Ki67+ cells were counted in sections around the LPC lesion (segments Th11–12, just caudal to the lesion center). The means were calculated from 3 sections spaced 100 μm apart. All samples were analyzed by laser scanning confocal microscopy (Olympus FV1200).

Primary culture of OPCs and cell proliferation assay. Primary cultures of OPCs were obtained from C57BL/6J mice at postnatal day 1. Whole brains were dissected in HBSS (Life Technologies) and dissociated into single-cell suspensions using the Neural Tissue Dissociation Kit (Miltenyi Biotec). OPCs were isolated by A2B5- and PDGFRα-specific antibody-coated magnetic beads (Miltenyi Biotec). The cells were plated on poly-L-lysine– precoated (PLL- precoated) 96-well plates (Greiner Bio-One) at a density of 1 × 10^4 cells per well. The cells were maintained at 37°C with 5% CO_2_ and cultured in DMEM containing 1 mM sodium pyruvate (Sigma-Aldrich), 0.1% BSA (Sigma-Aldrich), 50 μg/ml apo-transferrin (Sigma-Aldrich), 5 μg/ml insulin (Sigma-Aldrich), 30 nM sodium selenite (Sigma-Aldrich), 10 nM biotin (Sigma-Aldrich), 10 nM hydrocortisone (Sigma-Aldrich), 10 ng/ml PDGF-AA (PeproTech), 10 ng/ml basic FGF (PeproTech), and 1% penicillin/streptomycin (Life Technologies). The culture conditions we used were suitable for OPC culture, as previously described (65). Immunocytochemistry revealed that 93.4% ± 0.01% of cells in A2B5+ cell culture were labeled with PDGFRα, an OPC marker (data not shown). To conduct pharmacological screening, we used the Inhibitor Select 384-well Protein Kinase Inhibitor Library 1 (Calbiochem). Cells were pretreated with various compounds in the library for 30 minutes, and then cultured in the presence of mouse serum. To inhibit FGFR signaling, cells were pretreated with PD173074 (Sigma-Aldrich) and
Figure 6. FGF21 promotes human OPC proliferation. (A) Representative image of β-klotho expression in an autopsied sample from healthy patient and a patient with multiple sclerosis. Graphs show quantitations as indicated in the images (n = 4 for healthy patients, 3 for multiple sclerosis patients); *P < 0.01. (B) BrdU incorporation in human OPCs after stimulation with recombinant FGF21 (n = 6 for control, 4 for FGF21); *P < 0.05 as determined by Student's t test. Error bars represent SEM. Scale bar: 20 μm.

NF449 (Calbiochem) for 30 minutes, and then incubated in the presence of mouse serum. To assess the role of FGFRs, we used recombinant mouse FGF15 (Abcam), recombinant mouse FGF21 (R&D Systems), and recombinant mouse FGF23 (R&D Systems). For neutralization experiments, neutralizing antibodies against mouse FGF21 (final concentration, 6 μg/ml; sc-16842, Santa Cruz Biotechnology) were added to the culture 30 minutes before serum treatment. Control samples were treated with normal goat IgG (Sigma-Aldrich).

To assess cell proliferation, cells were incubated in DMEM supplemented with or without 1% (vol/vol) serum obtained from adult mice. BrdU was added into the culture 24 hours after the beginning of culture, and cell proliferation was estimated 48 hours after stimulation by measurement of BrdU incorporation into newly synthesized cellular DNA using the Cell Proliferation ELISA and BrdU (colorimetric) kit (Roche). Absorbances were measured by microplate reader (Molecular Devices SpectraMax 5).

BrdU detection by immunocytochemistry. OPCs were cultured in medium containing BrdU (10 μM; Sigma-Aldrich) for 2 hours. After culture, cells were fixed with 4% PFA in PBS for 30 minutes at room temperature, and then cells were pretreated with 2N HCl for 30 minutes at 37°C and washed in 0.1 M borate buffer (pH 8.5) for 10 minutes. After permeabilization with PBS containing 0.3% Triton X-100 and 10% goat serum (Sigma-Aldrich), cells were incubated with primary antibodies overnight at 4°C. Cells were incubated with fluorescent dye-conjugated secondary antibody for 1 hour at room temperature, and then cells were pretreated with 2N HCl for 30 minutes, and the tissue was lysed after 48 hours, and the lysates were subjected to real-time PCR.

To assess cell proliferation, cells were incubated in DMEM supplemented with or without 1% (vol/vol) serum obtained from adult mice. BrdU was added into the culture 24 hours after the beginning of culture, and cell proliferation was estimated 48 hours after stimulation by measurement of BrdU incorporation into newly synthesized cellular DNA using the Cell Proliferation ELISA and BrdU (colorimetric) kit (Roche). Absorbances were measured by microplate reader (Molecular Devices SpectraMax 5).

Quantitative reverse transcriptase PCR. Total RNA was isolated using the RNeasy Mini kit (Qiagen), and cDNA was synthesized using the High Capacity cDNA Reverse Transcriptase Kit (Applied Biosystems). cDNA fragments were amplified using the following primer pairs: FGF21 forward, GCTGCTGGAGGACGTCTAC; FGF21 reverse, CACAGAAGAGATCCTACAG; FGF21 forward, CACAGAAGAGATCCTACAG; FGF21 reverse, CGTGGGTGAGATCCAAGTA; FGF21 forward, GCCAGAUGGAGCCUCUCUATT-3′ (sense) and 5′-AUAGAGACCGAUCUCUGGTG-3′ (antisense); β-klotho siRNA, 5′-GGAUUAACUGAAACGAT-3′ (sense) and 5′-UCGUAUUCGGAGUAUACG-3′ (antisense); FGFR1 siRNA, 5′-GCAUCUGUGAGAGAUGUAUATT-3′ (sense) and 5′-UACACCUUCUCAGACG-3′ (antisense); FGFR2 siRNA, 5′-CUCUCACGCUAUUGAGA-3′ (sense) and 5′-CUACUACGCUAAUUGAGA-3′ (antisense); FGFR3 siRNA, 5′-GGAUUAAUGUUUGCUUUU-3′ (sense) and 5′-AAAAGACCAAACACUCAU-3′ (antisense); FGFR4 siRNA, 5′-CAUUGAAGCACUAUAGAATT-3′ (sense) and 5′-UUUUGAUAUAUGACAGT-3′ (antisense). Transfection of cultured OPCs with siRNAs targeting FGFRs and β-klotho was performed using Lipofectamine RNAiMAX (Invitrogen). The cells were lysed after 48 hours, and the lysates were subjected to real-time PCR. For knockdown of FGF21 in pancreas, FGF21 siRNAs were mixed with in vivo–jetPEI reagent (Polyplus Transfection). Mice received intrapancreatic injection of either PEI-conjugated FGF21 siRNA or PEI-conjugated nontargeting siRNA (0.02 μg/mouse, 3 points of pancreas) as previously described with slight modifications (36) just after LPC injection. The tissue was lysed after 48 hours, and the lysates were subjected to real-time PCR. Injection of siRNA into the pancreas did not result in any signs of toxicity at the behavioral level (data not shown).

Preparation and transfection of siRNA. Mouse FGF21 siRNA and β-klotho siRNA were synthesized by Ambion (Life Technologies). Mouse FGFR siRNAs were synthesized by Thermo Fisher Scientific. The sense and antisense strands of siRNA were as follows: FGF21 siRNA, 5′-GCCAGAUGGAGGCUCUCUATT-3′ (sense) and 5′-CUCAUGAGACCGAUCUCUGGTG-3′ (antisense); β-klotho siRNA, 5′-GGAUUAACUGAAACGAT-3′ (sense) and 5′-UCGUAUUCGGAGUAUACG-3′ (antisense); FGFR1 siRNA, 5′-GCAUCUGUGAGAGAUGUAUATT-3′ (sense) and 5′-UACACCUUCUCAGACG-3′ (antisense); FGFR2 siRNA, 5′-CUCUCACGCUAUUGAGA-3′ (sense) and 5′-CUACUACGCUUAUUGAGA-3′ (antisense); FGFR3 siRNA, 5′-GGAUUAAUGUUUGCUUUU-3′ (sense) and 5′-AAAAGACCAAACACUCAU-3′ (antisense); FGFR4 siRNA, 5′-CAUUGAAGCACUAUAGAATT-3′ (sense) and 5′-UUUUGAUAUAUGACAGT-3′ (antisense).
GAPDH forward, TCACCACCATGGAGAGGC; GAPDH reverse, GCTAAGCATTGGTTGCTCA. Samples for SYBR Green assays consisted of a 1× final concentration of PowerSYBR Green PCR Master Mix (Applied Biosystems), 200 nM gene-specific primers, and 10 ng cDNA (ABI ViaA7 real-time PCR system; Applied Biosystems). PCR conditions included 1 cycle at 95°C for 10 minutes, followed by 40 cycles of 95°C for 15 seconds and 60°C for 60 seconds. A melting analysis was carried out following PCR to monitor amplification specificity. Relative mRNA expression was normalized against GAPDH mRNA levels in the same samples, and calculated by the Δ/Δ Ct method.

**MTT assay**. PDGFRe OPCs were plated on PLL-coated 96-well plates at a density of 1×10⁴ cells per well. Cells were transfected with β-klotho siRNA using Lipofectamine RNAiMAX. After 72 hours of culture, MTT solution was added to each well, and the cultures were incubated for an additional 4 hours. Crystalline formazan was solubilized with 100 μl of a 10% (wt/vol) SDS solution for 24 hours. Absorbance at 595 nm was read spectrophotometrically using a microplate reader.

**Differentiation assay**. Primary cultures of OPCs were obtained from C57BL/6J mice at postnatal day 1. Whole brains were dissected in PBS and dissociated into single-cell suspensions by incubation at 37°C for 15 minutes in 0.25% trypsin-PBS. After neutralization by DMEM containing 10% FBS, cells were centrifuged at 300 g for 5 minutes, suspended in 10% FBS-DMEM, and filtered through a 100-μm cell strainer. Single cells were plated on PLL-precoated 10-cm dishes (Greiner Bio-One) and maintained at 37°C with 7% CO₂ in DMEM containing 10% FBS, cells were centrifuged at 300 g for 15 minutes in 0.25% trypsin-PBS at 35°C for 4 minutes. Collected cells were centrifuged at 300 g for 5 minutes, and the pellets were suspended in OPC medium (DMEM containing 1 mM sodium pyruvate [Sigma-Aldrich], 0.1% BSA [Sigma-Aldrich], 50 μg/ml apo-transferrin [Sigma-Aldrich], 5 μg/ml insulin [Sigma-Aldrich], 30 nM sodium selenite [Sigma-Aldrich], 10 nM biotin [Sigma-Aldrich], 10 nM hydrocortisone [Sigma-Aldrich], 10 ng/ml PDGF-AA [PeproTech], 10 ng/ml basic FGF [PeproTech], and 1% penicillin/streptomycin [Life Technologies]). Cells were plated on noncoated dishes for 1 hour to separate cell types based on differences in adhesion. Cells that did not adhere after incubation (OPCs) were collected and plated on PLL-precoated 96-well plates at a density of 1×10⁴ cells per well, and then maintained at 37°C with 5% CO₂ in OPC medium.

To induce oligodendrocyte differentiation, cells were cultured for an additional 3 days in DMEM containing 1 mM sodium pyruvate (Sigma-Aldrich), 0.1% BSA (Sigma-Aldrich), 50 μg/ml apo-transferrin (Sigma-Aldrich), 5 μg/ml insulin (Sigma-Aldrich), 30 nM sodium selenite (Sigma-Aldrich), 10 nM biotin (Sigma-Aldrich), 10 nM hydrocortisone (Sigma-Aldrich), 30 ng/ml Triiodo-L-Thyronine (T₃; Sigma-Aldrich), and 1% penicillin/streptomycin (Life Technologies) with or without recombinant mouse FGF21 (R&D Systems).

**Primary culture of astrocytes**. Primary cultures of OPCs were obtained from C57BL/6J mice at postnatal day 1. Whole brains were dissected in PBS and dissociated into single-cell suspensions by incubation at 37°C for 15 minutes in 0.25% trypsin-PBS. After neutralization by 10% FBS-DMEM, cells were centrifuged at 300 g for 5 minutes, suspended in 10% FBS-DMEM, and filtered through a 100-μm nylon cell strainer. Single cells were plated on PLL-coated 10-cm dishes and maintained at 37°C with 5% CO₂ in 10% FBS-DMEM. Ten days after culture, cells were detached by treatment with 0.25% trypsin-PBS and neutralized with an equal volume of 10% FBS-DMEM. The cells were pelleted by centrifugation at 1,000 g for 3 minutes, and then resuspended in 10% FBS-DMEM. The cells were plated on culture plates precoated with PLL. After 2 passages, the cells were plated on PLL-coated 96-well plates at a density of 1×10⁴ cells per well. To collect astrocyte supernatant, the cells were cultured in DMEM with or without 1% (vol/vol) serum obtained from adult mice. One day after incubation, supernatant was collected and added to the OPC culture.

**ELISA**. FGF21 levels in mouse serum or tissue lysates were examined using a Mouse/Rat FGF21 Quantikine ELISA (R&D Systems). The tissues were homogenized in the lysis buffer (25 mM Tris-HCl [pH 7.5], 150 mM NaCl, 1% Triton X-100, and 1 mM EDTA containing protease inhibitor [Roche]). The lysates were clarified by centrifugation at 13,000 g at 4°C for 20 minutes, and the supernatants were used for assay.

**FGF21 labeling**. Recombinant mouse FGF21 was conjugated with HiLyte Fluor 555 labeling kit (Dojindo Molecular Technologies). To detect leakage of FGF21 into the spinal cord, mice received fluorescence-labeled FGF21 (200 pg/mouse) i.v. 2 hours before sampling. Spinal cord tissues were isolated after PBS perfusion and were homogenized in lysis buffer. After centrifugation (13,000 g at 4°C for 20 minutes), fluorescence in the supernatant was measured by microplate reader.

**BrdU labeling in vivo**. BrdU (1 mg/ml) was added to the drinking water (supplemented with 1% [wt/vol] sucrose) for 5 weeks before sacrifice of the mice. The water bottles containing BrdU were protected from light and changed every 3 days during labeling periods.

**Electron microscopy**. Mice were transcardially perfused with 0.1 M PB containing 2.5% glutaraldehyde and 2% PFA. Spinal cord tissues were removed and were postfixed in the same fixative at 4°C overnight. The spinal cords were sliced into 50-μm pieces using a Vibrating Blade Microtome (VT1000s, Leica) and washed in 0.1 M PB to prepare the sample for electron microscopy as described previously (66). The samples were incubated in 0.1 M PB containing 1% OsO₄ (TAAB) in 0.1 M PB for 1 hour and dehydrated in an ascending ethanol series, and then embedded in epoxy resins through propylene oxide (Wako). The samples were cut into 80-nm sections using a Reichert-type ultramicrotome (Ultracut N, Leica), and ultrathin sections on formvar-coated 1-hole Cu grids (Nisshin EM Corp.) were observed by transmission electron microscopy (H-7650, Hitachi). Nonoverlapping obtained images of nerve fiber cross sections were analyzed using ImageJ version 1.44p (NIH). The diameter of axons, myelin thickness, and g-ratio were determined from about 600 fibers in the corticospinal tract.

**Ladder-walk test**. We observed the walking of mice on a horizontal ladder (whole length of 1 m) with stainless steel rungs spaced 1–4 cm apart. Mice were habituated to the apparatus before the surgery. The number of faulty placements of the injured hind paw was counted when the mice walked through the 1-m stretch. Deep slips/misses, minor slips, and placement errors (correction, replacement) were considered as faults (67). The tests were performed during the light cycle. The control baseline scores were obtained just before the injury. The spacing between rungs was changed accordingly to prevent the animal from learning the rung locations.

**IHC staining of human tissues and cell proliferation assay of human OPCs**. This study was approved by the Local Ethical Committee, Toneyama National Hospital (Toyonaka, Japan), with research number 1135-1 (Research Resource Network, Japan), and by the Local Ethical Committee, Aichi Medical University (Nagakute, Japan), with research number 15-017 (Aichi Medical University, Kareiken Brain Resource Center). Informed consent was obtained at the occasion...
of autopsy (brain banking). We obtained autopsy brain tissues from 4 control individuals (3 men; median age: 68 years; range: 63–70 years) and 3 individuals with multiple sclerosis (1 man; median age: 66 years; range: 63–85 years). Informed consents were obtained from family member of patients. Formalin-fixed spinal cord samples were embedded in paraffin and cut into 10-μm-thick sections for IHC. Sections were deparaffinized and were permeabilized with PBS containing 0.3% Triton X-100 and 5% BSA for 1 hour at room temperature. Sections were incubated with primary antibodies against mouse anti-human PDGF-Rα (1:50; 0100-0220, Serotec) and rabbit anti-human β-klotho (1:50; AV53325, Sigma-Aldrich) and Alexa Fluor 488–conjugated goat anti-mouse IgG and/or Alexa Fluor 647–conjugated goat anti-rabbit IgG (Invitrogen) secondary antibodies. The sections were treated with a solution of Sudan Black B. All images were captured by laser scanning confocal microscopy. The research protocol was approved by the Human Use Review Committees of the Graduate School of Medicine, Osaka University. Informed consent was obtained from all subjects.

Primary cultures of human OPCs were obtained from ScienCell. The cells were plated on PLL precoated 96-well plates at a density of 1 × 10⁴ cells per well, and then maintained at 37°C with 5% CO₂ in DMEM containing 1 mM sodium pyruvate (Sigma-Aldrich), 0.1% BSA (Sigma-Aldrich), 50 μg/ml apo-transferrin (Sigma-Aldrich), 5 μg/ml insulin (Sigma-Aldrich), 30 mM sodium selenite (Sigma-Aldrich), 10 mM biotin (Sigma-Aldrich), 10 mM hydrocortisone (Sigma-Aldrich), 10 ng/ml PDGF-AA (PeproTech), 10 ng/ml basic FGF (PeproTech), and 1% penicillin/streptomycin (Life Technologies). To assess cell proliferation, cells were cultured in DMEM supplemented with or without recombinant human FGF21 (R&D Systems) at a final concentration of 6 μg/ml. BrdU was added to the culture 24 hours after the beginning of culture, and cell proliferation was estimated 48 hours after stimulation by measurement of BrdU incorporation into newly synthesized cellular DNA using the Cell Proliferation ELISA and BrdU (colorimetric) kit.

**Statistics.** Data are presented as means ± SEM. Statistical significance between groups was determined by unpaired Student’s t-test, or repeated-measures ANOVA followed by Bonferroni post hoc test. No randomization was used to assign experimental groups or collect data, but samples were assigned to specific experimental groups without bias. No data points were excluded. Histopathological analyses and behavioral tests were performed in a blinded manner.

**Author contributions**

M Kuroda performed almost all experiments and analyzed the data. RM developed the concept, designed the experiments, and wrote the manuscript. NM performed immunohistochemical experiments. YK carried out electron microscopy analyses. MH helped with tamoxifen injection and cuprizone feeding. HF, MY, and HM provided autopsied samples. M Konishi and NI contributed to FGF21-KO mouse experiments. TY supervised all aspects of this project.

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