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Cell Injury, Repair, Aging and Apoptosis

Fingolimod (FTY720) Enhances Remyelination Following Demyelination of Organotypic Cerebellar Slices

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Remyelination, which occurs subsequent to demyelination, contributes to functional recovery and is mediated by oligodendrocyte progenitor cells (OPCs) that have differentiated into myelinating cells. Therapeutics that impact remyelination in the CNS could be critical determinants of long-term functional outcome in multiple sclerosis (MS). Fingolimod is a S1P receptor modulator in MS clinical trials due to systemic anti-inflammatory properties, yet may impact cells within the CNS by crossing the blood-brain barrier. Previous studies using isolated dissociated cultures indicate that neural cells express S1P receptors and respond to receptor engagement. Our objective was to assess the effects of fingolimod on myelin-related processes within a multicellular environment that maintains physiological cell-cell interactions, using organotypic cerebellar slice cultures. Fingolimod treatment had no impact on myelin under basal conditions. Fingolimod treatment subsequent to lysolecithin-induced demyelination enhanced remyelination and process extension by OPCs and mature oligodendrocytes, while increasing microglia numbers and immunoreactivity for the astrocytic marker glial fibrillary acidic protein. The number of phagocytosing microglia was not increased by fingolimod. Using S1P receptor specific agonists and antagonists, we determined that fingolimod-induced effects on remyelination and astrogliosis were mediated primarily through S1P3 and S1P5, whereas enhanced microgliosis was mediated through S1P1 and S1P5. Taken together, these data demonstrate that fingolimod modulates multiple neuroglial cell responses, resulting in enhanced remyelination in organotypic slice cultures that maintain the complex cellular interactions of the mammalian brain. (Am J Pathol 2010, 176:2682–2694; DOI: 10.2353/ajpath.2010.091234)

In the CNS, oligodendrocytes produce the myelin sheath, which forms an insulating layer around axons to enhance conduction velocity of electrical impulses.1–3 The maintenance of myelin requires continuous turnover of its components throughout life.4–5 Demyelinating insults contribute to axonal degeneration, functional impairments, and clinical disability in the autoimmune disease multiple sclerosis (MS).6–7 Neuropathological and magnetic resonance-based observations indicate that re-ensheathment of demyelinated axons, termed remyelination, occurs in demyelinated lesions in MS and may contribute to clinical remittance.8–9 Remyelination is critical to restore electrical impulse conduction and protect axons from degeneration or further injury.10–14 Experimental animal models of CNS demyelination indicate that remyelination is mediated by oligodendrocytes generated from oligodendrocyte progenitor cells (OPCs).15 Demyelinating models show that both astrocytes and microglia are critical for remyelination, with microglia contributing to clearance of debris and both cell types participating in creating an environment permissive for remyelination.16–19

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The eventual failure of remyelination that occurs as MS progresses may result from the generation of a nonpermissive environment or intrinsic deficits in the remyelinating cells. Therapeutic agents that impact on myelin maintenance and remyelination are predicted to be important determinants of long-term functional outcome in MS. Fingolimod (FTY720) is currently in clinical trials due to its anti-inflammatory properties, yet readily crosses the blood brain barrier. Analysis of the distribution of fingolimod orally administered to mice for 1 week indicated that the drug is sequestered in the brain parenchyma relative to the blood. Once fingolimod has entered the brain, it is phosphorylated to its biologically active form which then binds sphingosine-1-phosphate (S1P) receptors.

Previous experiments have demonstrated that isolated neuroglial cell types in dissociated cultures can respond to S1P receptor engagement. Both human and rodent cells of the oligodendroglial lineage are able to respond to fingolimod due to the expression of S1P receptor isoforms 1, 3, and 5. S1P or fingolimod treatment of rodent pre-OLGs (O4+/Galactocerebroside−) or human fetal OPCs (A2B5+), respectively, leads to transient retraction of processes via S1P5. High doses of fingolimod synergistically enhance process branching and elongation by rat OPCs when co-applied with neurotrophic factors. Fingolimod can either promote or inhibit rodent and human OPC differentiation in a dose-dependent manner. Whereas the cytoskeletal dynamics of rat mature OLGs in dispersed cell culture are not influenced by fingolimod, this agent induces a cyclical regulation of S1P1-dependent membrane elaboration and S1P3/5-dependent membrane retraction in human mature OLGs over time. Astrocytes and microglia also respond to S1P-directed signaling. S1P receptor engagement in astrocytes leads to extracellular signal-regulated kinase-1/2 activation, resulting in proliferation and neurotrophic factor production. Astrocytes additionally produce platelet-derived growth factor (PDGF), which can alter S1P receptor levels. Fingolimod can inhibit macrophage infiltration following traumatic brain injury.

Although the effects of S1P receptor signaling on individual neuronal cell types has been described, the direct influence of fingolimod on neuronal cells and myelin-related processes in a physiological environment containing all of the resident CNS elements, and the contribution of individual S1P receptors to these responses, have not yet been investigated. Here, we delineate the consequence of fingolimod treatment on myelin maintenance and remyelination, using an in vitro model that recapitulates myelin-axon and cell-cell interactions. Rodents orally administered 0.3 mg/kg of fingolimod daily over the long-term were shown to have micromolar concentrations of the active form of the drug within the brain parenchyma, whereas subnanomolar concentrations were measured in the cerebrospinal fluid. Newborn mouse organotypic cerebellar slice cultures were treated with the active phosphorylated form of fingolimod at physiological doses representative of those measured in the brain (1 μmol/L) and cerebrospinal fluid (100 pmol/L). We have previously demonstrated that human neural cell properties are modulated in vitro by fingolimod doses within the range of 100 pmol/L to 1 μmol/L. We had observed that cells responded similarly when treated with 100 pmol/L to 1 nmol/L of fingolimod, or with 10 nmol/L to 1 μmol/L of the drug. We report that fingolimod influences cells of the oligodendroglial lineage, microglia, and astrocytes, resulting in enhanced remyelination in the absence of a systemic immune system.

**Materials and Methods**

**Animals and Tissue Preparation**

Newborn (P0) wild-type CD1 mouse pups were obtained from Charles River (Quebec, Canada). All procedures were performed in accordance with the Canadian Council on Animal Care guidelines. Slices were obtained and cultured as previously described. Cerrebellum and attached hindbrain were extracted in L15 media and cut into 300 μm sagittal sections using a McIlwain tissue chopper. Slices were separated and plated on a Millipore-Millicel-CM culture insert (Fisher Scientific, Ottawa, ON, Canada) in 6-well cell culture plates, with 6 slices per insert. Culture media was composed of 50% minimal essential media, 2.5% heat-inactivated horse serum, 2% Earle’s balanced salt solution, 6.5 mg/ml glucose (Sigma-Aldrich, St-Louis, MO), and penicillin and glutamine supplements (All from Invitrogen, Carlsbad, CA). Membranes were transferred into fresh media every 2 to 3 days. Slices were left untreated for 21 days in vitro (DIV) to allow clearance of debris and myelination to occur.

**Pharmacological Treatment**

The active phosphorylated form of fingolimod was provided by Novartis Pharma (Basel, Switzerland) and was reconstituted in dimethylsulfoxide/50 mmol/L HCl. Fingolimod was diluted in basal culture media and replaced every 2 to 3 days. For myelin maintenance studies, cultures were treated for three subsequent weeks with fingolimod. For remyelination studies, cultures were demyelinated with lysolecithin overnight (0.5 mg/ml), washed with PBS twice for 10 minutes, allowed to recover in basal culture media for 2 days, then treated with fingolimod for the next 2 weeks (14 DIV). To investigate the response to fingolimod independent from demyelination, control cultures were allowed to recover for 14 DIV postlysolecithin, then either left untreated for another 14 DIV (control 28 DIV postlysolecithin), or treated with fingolimod for 14 DIV (total 28 DIV postlysolecithin). Cultures treated with the vehicle used to reconstitute fingolimod showed no significant effects on myelin. To determine the mechanism of fingolimod-induced effects, demyelinated cultures were co-treated with fingolimod (100 pmol/L) and a S1P antagonist W123 (1 μmol/L; Cayman Chemicals, Ann Arbor, MI) or a S1P3/5 pathway antagonist suramin (100 nmol/L; EMD Bioscience, San Diego, CA). More specific S1P receptor agonists were used to determine whether activation via specific S1P receptors could mimic the fingolimod-induced effects.

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mod-induced effects, including a S1P1 agonist SEW2871 (100 nM; EMD Biosciences)\textsuperscript{27-29} and a S1P5 agonist (100 nM; provided by Novartis Pharma). The experiments were performed on six different preparations.

**Whole Mount Immunohistochemistry**

Following two quick washes in PBS, slices were fixed in 1% paraformaldehyde (PFA) for 15 minutes for node of Ranvier staining (Caspr and sodium channel), or 4% PFA for 1 hour for all other stains, then washed twice in PBS. Slices were cut from the membrane and blocked for 3 hours at room temperature in 1 mM HEPES, 2% heat-inactivated horse serum, 10% heat-inactivated goat serum, 1% bovine serum albumin, and 0.25% Triton X-100 in Hank’s Balanced Salt Solution. Primary antibodies diluted in block solution were applied for 48 hours at 4°C. These include mouse anti-myelin basic protein (MBP, 1:500; Sternberger Monoclonals Inc., Lutherville, MD), rabbit anti-platelet-derived growth factor receptor α (PDGFαR, 1:50; Santa-Cruz Biotechnology, Santa Cruz, CA), mouse anti-neurite outgrowth inhibitor protein A (NogoA, 1:1000; provided by Dr. Martin Schwab, University of Zurich), polyclonal chicken anti-neurofilament medium chain (NFM, 1:2000; EnCor Biotech, Alachua, FL), mouse anti-β tubulin III (β-TubIII, 1:400; Sigma), rabbit anti-ionized calcium binding adaptor molecule 1 (IBA-1, 1:500; Wako Chemicals USA, Richmond, VA), rabbit anti-glia fibrillary acid protein (GFAP, 1:200; NeoMarkers, Fremont, CA), mouse monoclonal pan anti-sodium channel channel (1:1000; Sigma), and rabbit anti-contactin associated protein (Caspr, 1:10 000; provided by Dr. David Colman, McGill University). Slices were washed thrice in PBS with 0.05% Triton X-100 for 1 hour on a rotary shaker then incubated in secondary antibody diluted in block solution overnight at 4°C. Secondary antibodies include goat anti-mouse Cy3 (1:500; Jackson Immunoresearch, West Grove, PA), donkey anti-chicken-Alexa-488 (1:70; Molecular Probes, Eugene, OR), goat anti-mouse Alexa-488 (1:500; Molecular Probes), and goat anti-rabbit Cy3 (1:250; Jackson Immunoresearch). Slices were counterstained with nuclear dye Topro-3 (1:1000; Molecular Probes), washed thrice, and mounted on glass slides (Fisher Scientific) using Fluoromount G (Southern Biotech, Birmingham, AL).

**Immunohistochemical Analysis**

Single optical \(\times 63\) or \(\times 100\) objective images were acquired using a Zeiss LSM 510 microscope (Peabody, MA). Two to three representative images were acquired per stain per slice. The range of the slice was determined using Z-stack imaging at 1- to 3-μm intervals. By Z-stack analysis, we found that slices thinned to approximately 30 μm in culture. Analysis demonstrated that a fully myelinated unmanipulated slice had three distinct regions in the Z plane that differed in their cellular composition and appearance: the first \(\sim 1\) to 5 μm (‘top’) is where the majority of the myelin, axons, OLGs, OPCs, and astrocytes reside; the next \(\sim 6\) to 23 μm (‘middle’) have the same composition as the ‘top’ layer however at lesser amounts; the last \(\sim 24\) to 30 μm (‘bottom’) was mostly devoid of all cellular components except microglia, which were also found throughout other depths of the slice albeit at lower levels. Quantification was performed separately in the top and middle of the slice where the myelin was located, to control for sampling error and variation in thickness of each slice.

The area of MBP or GFAP immunostaining (μm\(^2\)) per \(\times 63\) objective image was determined using Scion Image software (Scion Corporation, Frederick, MD). The numbers of PDGFαR, NogoA, and IBA-1 immunopositive cells were counted in a blinded fashion either from representative \(\times 63\) objective images or a series of images derived from Z-stack imaging. The number of nodes of Ranvier per \(\times 100\) objective image was determined by counting areas of condensed sodium channel immunostaining flanked by Caspr immunostaining. The diameter of the Node of Ranvier (μm) was evaluated by calibrating \(\times 100\) objective images and measuring the length of condensed sodium channel immunostaining using ImageJ software (U. S. National Institutes of Health, Bethesda, Maryland, http://rsb.info.nih.gov/ij).

**Phagocytosis Assay**

Phagocytosis of E. Coli-coated beads was used to measure phagocytic potential of microglia in the slices. A 2 μl drop of rhodamine-conjugated E. Coli bioparticles (1 mg/ml; Invitrogen) reconstituted in culture media was added directly on top of the slice. Slices were incubated for 3 hours at 37°C, then washed in PBS twice for 10 minutes, and fixed in 4% PFA for 1 hour. Slices were then stained against a microglial marker, IBA-1. Slices were kept in the dark as much as possible during the fixation and immunohistochemical staining processes. Absolute numbers of phagocytic microglia were taken to be IBA-1 immunopositive cells with red signal within the cell body, and were counted in a blinded manner in images obtained from Z-stacks. Phagocytosis assays were performed using one slice culture preparation.

**Bromodeoxyuridine Incorporation**

Bromodeoxyuridine (BrdU) incorporation and immunohistochemistry was performed as previously described.\textsuperscript{38} Briefly, BrdU was diluted in culture media to a concentration of 10 μmol/L and added to slices for 24 hours before fixation in 4% PFA for 1 hour. Slices were denatured with 2N HCl for 45 minutes, rinsed in PBS for 1 minute, then incubated with mouse anti-BrdU for 48 hours (1:40; Sigma) and goat anti-mouse secondary antibody overnight at 4°C.

**Statistical Analysis**

Values are represented as means ± SEM, and some are normalized to control cultures at the respective time point. Statistical analyses were performed using one-way analysis of variance (Newman-Keuls and Dunns multiple
comparison tests), unpaired two-tailed Student’s t-test, and Mann-Whitney test. P values less than 0.05 were taken to represent significant differences between treatment groups.

**Electron Microscopy**

Electron microscopy was used to confirm the integrity of myelin and to assess actual remyelination as documented by thin myelin sheaths. Cerebellar slices were fixed overnight with 2.5% glutaraldehyde in 0.1 mol/L sodium cacodylate buffer then cut out of the surrounding membrane, subsequently osmicated with potassium ferrocyanide-reduced 1% osmium tetroxide solution for 1 hour, and dehydrated with successive 10-minute washes in increasing concentrations of ethanol. Slices were infiltrated for 1 hour with 1:1, 1:2, and 1:3 ethanol: Epon blends, followed by infiltration with pure Epon. Slices were embedded in Epon in a plastic BEEM capsule. Sections of 70 to 100 μm were cut onto 200 mesh copper grids and stained with 4% uranyl acetate and Reynold’s lead citrate. Semithin sections of 1 μm were stained with toluidine-blue for neuropathological analysis.

**Results**

**Characterization of Organotypic Cerebellar Slice Cultures**

To assess the role of fingolimod on remyelination, we elected to use organotypic cerebellar slice cultures due to previous studies demonstrating robust and reproducible myelination along definable axons in slices derived from this brain region, and enhanced plasticity and resistance to mechanical trauma at this age.36,37,39 Organotypic slices retain the astrocyte and microglia elements that allow us to identify our contribution to myelination-associated processes, but avoid the complex systemic immune system interactions observed in intact animals. Confluent analysis of untreated slice cultures at the temporal onset of pharmacological manipulation (21 DIV) indicated that cultures were comprised of myelin (MBP+), mature oligodendrocytes (NogoA+), neurons (NFM+), βTubIII+, OPCs (PDGFαR+), astrocytes (GFAP+), and microglia (IBA-1+) (see Supplemental Figure 1A at http://ajp.amipathol.org). There was a significant co-localization between myelin (MBP) and axons (NFM) in the X, Y, and Z axes (see Supplemental Figure S1, A and B, at http://ajp.amipathol.org). Cross-sectional analysis confirmed that MBP+ fibers ensheathed NFM+ axons (see Supplemental Figure S1B at http://ajp.amipathol.org). Not all NFM immunopositive fibers were associated with MBP staining, consistent with not all axons being myelinated in organotypic cerebellar slices.37 Ultrastructural analysis demonstrated that the myelin was compact and organized appropriately into nodes of Ranvier (see Supplemental Figure S1C at http://ajp.amipathol.org), and was therefore mature and of good integrity.

**Effects of Fingolimod on Myelin Maintenance and Oligodendroglia**

Following recovery from dissection, cultures were treated with the active phosphorylated form of fingolimod for the subsequent 21 DIV then analyzed immunohistochemically. Fingolimod treatment (100 pmol/L, 1 μmol/L) did not significantly affect the area of MBP staining per field relative to untreated cultures (0.97 ± 0.3, 0.96 ± 0.2 fold over control, respectively), consistent with no deleterious effects of fingolimod on myelin maintenance (Figure 1A). We next investigated the effect of fingolimod on mature OLGs, labeled with the OLG cell body marker NogoA.40–41 and on OPCs identified by PDGFαR expression. Fingolimod treatment (100 pmol/L, 1 μmol/L) induced membrane elaboration and process outgrowth by OLGs (Figure 1B) and OPCs (Figure 1C), respectively, relative to control. Fingolimod (100 pmol/L, 1 μmol/L) had no significant impact on the absolute number of PDGFαR+ cells, although a trend toward an increase was observed with the lowest dose (12 ± 2 and 6 ± 1 per field, respectively, relative to 8 ± 0 for control) (Figure 1D). Fingolimod treatment for 21 DIV (100 pmol/L) caused a significant decrease in numbers of microglia (IBA-1+) from 7 ± 1 to 1 ± 0 per field, and a significant decrease in the area of GFAP immunostaining (from 3886 ± 579 μm² to 2192 ± 492).

**Effects of Fingolimod on Remyelination and Oligodendroglial Lineage Cells Following Demyelination: Characterization of Lysolecithin-Induced Demyelination**

The bioactive lipid lysolecithin (lysophosphatidylcholine) induces demyelination when focally injected into white matter tracts such as the cerebellar peduncle and spinal cord,42–43 or applied to organotypic slice cultures.37 Following 21 DIV of postisolation recovery, slices were treated with 0.5 mg/ml lysolecithin for 16 hours, which induces demyelination of organotypic slices by 2 days postapplication while minimizing concomitant axonal injury.37 Analysis of cultures 2 days after application of lysolecithin indicated a notable decrease in the area of MBP staining (0.23 ± 0.18-fold over normally myelinated control at 21 DIV) such that immunopositive staining was associated with myelin debris and ensheathed axons were rarely observed (Figure 2, A and B). In comparison with toluidine-blue stained semithin sections from fully myelinated controls (Figure 2C), sections from lysolecithin-treated cultures demonstrated many demyelinated fibers, few residually myelinated fibers, and accumulation of myelin debris (Figure 2D). NFM staining was not adversely affected by lysolecithin treatment indicating a sparing of axons (Figure 2A). Fully myelinated cultures at 21 DIV demonstrated appropriately formed nodes of Ranvier characterized by paranodal Caspr immunostaining flanking sodium channels localized to the node itself (Figure 2E). Demyelination with lysolecithin caused a lengthening of the distribution of sodium channels mark-
Effects of Fingolimod on Recovery of Myelin Following Lysolecithin-Induced Demyelination

Analysis of control cultures at 14 DIV postlysolecithin (total 35 DIV) indicated that some recovery of MBP staining and ensheathment had occurred (1.4 ± 0.42 compared with 2 days postlysolecithin; Figure 3A); however, remyelination was incomplete at this time point, as has been previously documented in this culture system. Application of fingolimod (100 pmol/L, 1 μmol/L) for 14 DIV following lysolecithin-induced demyelination caused a significant increase in the area of MBP staining per field relative to remyelinating control, indicative of enhanced remyelination (1.97 ± 0.33 and 2.00 ± 0.40-fold over control at 35 DIV, respectively) (Figure 3, A and B). Ultrastructural evaluation of fingolimod-treated cultures revealed many thin myelin sheaths, some of which were associated with paranodes suggestive of functional remyelination (Figure 3C). These cultures maintained appropriate cell–cell contacts and cyto-architecture as demonstrated by synaptic associations and adjacent localization of astrocytes (Figure 3D). Whereas control semithin toluidine-blue stained sections were characterized by many unmyelinated yet apparently healthy fibers (Figure 3E), sections from fingolimod-treated cultures revealed the presence of many thin myelin sheaths, some thicker myelin sheaths, and healthy neurons and oligodendrocytes (Figure 3F). As apparent in toluidine blue stained sections, both control and fingolimod-treated slices were characterized by the presence of phagocytosed myelin debris; we could not appreciate any qualitative difference in the amount of MBP-immunoreactive debris found in fingolimod-treated cultures compared with control cultures at 14 DIV postdemyelination.

A time course analysis of remyelination was determined with various fingolimod treatment durations by MBP immunohistochemistry. Remyelination was initiated as early as 5 DIV postdemyelination in both control and fingolimod-treated cultures. Although the area of MBP immunostaining was not significantly different between fingolimod treatment and control at both 5 and 12 DIV (data not shown), it was only significantly enhanced with fingolimod treatment at 14 DIV postlysolecithin (Figure 3, A and B). This finding indicates that the fingolimod-induced increase in remyelination occurred between 12 and 14 DIV following demyelination.

Effects of Fingolimod on Oligodendroglia During Remyelination

Fully myelinated cultures at 21 DIV were characterized by many NogoA+ mature OLGs (Figure 4A). Lysolecithin-induced demyelination was associated with a loss of NogoA immunostaining. Cells positive for another mature OLG-associated cytoplasmic protein, adenomatous polyposis coli/CC1, could still be observed under demyelinating conditions (data not shown) suggesting that demyelination eliminated myelin-associated NogoA but did not kill all OLGs (Figure 4A). Control remyelinating cultures and fingolimod-treated cultures demonstrated a recovery of No-
These numbers decreased by 5 DIV postlysolecithin and postlysolecithin-induced demyelination. Demyelination induced a significant decrease in the number of nodes of Ranvier.

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bar (pre-lysolecithin) or 2 DIV postlysolecithin-induced demyelination. Demyelination induced a significant increase in the length of the node due to lateral migration of fibers (MBP staining). Immunostaining is not associated with axons and likely reflects myelin debris. Scale bar 5 mm. Immunostaining of cultures pre-lysolecithin at 21 DIV demonstrates paranodal protein Caspr (red) flanking nodal aggregations of sodium channels (green), suggestive of proper node of Ranvier formation. Lysolecithin treatment caused a significant increase in microglia at 5 DIV postlysolecithin (5 ı 1 cells/field) compared with 2 DIV postlysolecithin (average 10 per field). Fingolimod-treated cultures (100 pmol/L) had a significant increase in microglia at 5 DIV postlysolecithin (5 ± 1 IBA-1+ cells/field compared with 2 ± 1 for control) (Figure 5B). This was further increased by 9 DIV postlysolecithin (14 ± 2 compared with 2 ± 1 for control), and maintained by 14 DIV postlysolecithin (8 ± 0 relative to 3 ± 0 for control; Figure 5A, B). To determine whether this microglial response in the presence of fingolimod could be attributed to an enhanced response to demyelination, cultures were left untreated for 14 DIV postlysolecithin then subjected to another 14 DIV of fingolimod treatment (100 pmol/L; total 28 DIV postlysolecithin). These cultures still demonstrated a significant increase in microglia cell numbers (13 ± 1 IBA-1+ cells/field) similar to that observed when fingolimod was applied for 14 DIV immediately postlysolecithin (10 ± 1) (Figure 5B). Significantly more microglia were noted in this condition than in control cultures at 28 DIV postlysolecithin (6 ± 1; Figure 5B), thereby dissociating this fingolimod-induced effect from a response to demyelination.

When assessing astrocytic reactivity, the area of GFAP immunostaining also peaked at 2 DIV postlysolecithin were maintained at this level by 9 DIV and 14 DIV postlysolecithin (Figure 5B), and may reflect the migration of microglia out of the slice. Accordingly, there was a slight increase in the number of IBA-1+ cells at the edge of the slice in these cultures (average 14 per field) compared with 2 DIV postlysolecithin (average 10 per field).

Fingolimod-treated cultures (100 pmol/L) also induced a robust outgrowth and elaboration of NogoA+ membranes compared with remyelinating control (Figure 4A).

Lysolecithin treatment caused a significant increase in absolute numbers of PDGFRα+ OPCs (17 ± 2 cells/field), as compared with before lysolecithin treatment (control at 21 DIV; 3 ± 0 cells/field) (Figure 4, B and C). The absolute number of OPCs was significantly decreased in control cultures at 14 DIV postlysolecithin (5 ± 0 cells/field; Figure 4C), suggestive of initial differentiation and loss of PDGFRα expression. The low dose of fingolimod (100 pmol/L) applied during 14 DIV postlysolecithin induced process extension in OPCs (Figure 4B) and caused a trend toward a relative increase in OPC numbers (9 ± 1 cells/field; Figure 4C).

Effects of Fingolimod on Microglia and Astrocytes During Remyelination

To further evaluate the effects of fingolimod on neuralglial cells in the context of enhanced remyelination, effects on microglia and astrocytes were investigated. A time course analysis demonstrated that average numbers of IBA-1+ microglia peaked at 2 DIV postlysolecithin, from 1 ± 0 to 4 ± 1 IBA-1+ cells/field (Figure 5, A and B). These numbers decreased by 5 DIV postlysolecithin and

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goa immunostaining over 14 DIV of postlysolecithin recovery (Figure 4A). At 14 DIV postlysolecithin, fingolimod treatment (100 pmol/L, 1 µmol/L) also induced a robust outgrowth and elaboration of NogoA+ membranes compared with remyelinating control (Figure 4A).

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Figure 2. Characteristics of lysolecithin-induced demyelination of slices. A: Representative images of slices at 21 DIV (pre-demyelination) and 2 DIV postlysolecithin induced demyelination (23 DIV total), immunostained against myelin (MBP; red) and axons (NFM; green). Following lysolecithin treatment, MBP immunostaining is not associated with axons and likely reflects myelin debris. Scale bar 20 µm. B: Quantification of amount of myelin expressed as area of MBP immunostaining (µm²) per ×63 objective image normalized to untreated control culture values at 21 DIV. Treatment with lysolecithin decreased the area of MBP staining. *P < 0.05. C: Representative toluidine-blue stained section of culture at 21DIV before demyelination. Arrows indicate myelinated fibers. Scale bar 5 mm. D: Representative toluidine-blue stained section of lysolecithin-treated culture 2 DIV postdemyelination demonstrates some residually myelinated fibers (short arrow), many demyelinated axons (arrowheads), and some neurons showing some degenerative changes (long arrow). Myelin debris in the form of small blue stained collections and granules is seen both within and outside of cells (MD). Scale bar 5 mm. E: Immunostaining of cultures pre-lysolecithin at 21 DIV demonstrates paranodal protein Caspr (red) flanking nodal aggregations of sodium channels (green), suggestive of proper node of Ranvier formation (arrows). Lysolecithin-induced demyelination induced a dispersion of Caspr immunostaining (red) and expansion of sodium channels along the axon (green; arrows). Scale bar 10 µm. F: Average diameter of nodes of Ranvier (µm) as measured by length of sodium channel aggregation ± SEM, in cultures at 21 DIV (pre-lysolecithin) or 2 DIV postlysolecithin-induced demyelination. Demyelination induced a significant increase in the length of the node due to lateral migration of sodium channels. ***P < 0.001. G: Average number of nodes of Ranvier per ×63 objective image ± SEM, in cultures at 21 DIV (pre-lysolecithin) or 2 DIV postlysolecithin-induced demyelination. Demyelination induced a significant decrease in the number of nodes of Ranvier. *P < 0.05.
(2759 ± 591 μm²) compared with before demyelination (624 ± 220 μm²) and subsided over the subsequent 14 DIV (1354 ± 550 μm²; Figure 5, C and D). Similar to the effect on microglia cell numbers, fingolimod treatment (100 pmol/L) for 14 DIV postlysolecithin was associated with an increased area of GFAP immunostaining (4870 ± 83 μm²; Figure 5, C and D). Consistent with the finding that both astrocytes and microglia were increased in cultures treated with fingolimod for 14 DIV postlysolecithin, we observed a significant increase in cells that had incorporated BrdU relative to remyelinating control (from 0.3 ± 0.2 to 5 ± 1 BrdU+ cells/field). The increased numbers of microglia and increased immunoreactivity for an astrocytic marker observed with fingolimod at 14 DIV postlysolecithin treated with 100 pmol/L fingolimod for 14 DIV postlysolecithin indicates thin myelin sheaths suggestive of remyelination. Arrows indicate demyelinated axons. Arrows indicate thin myelin sheaths suggestive of remyelination. Scale bar = 0.5 μm.

Electron micrograph of demyelinated slice allowed to recover for 14DIV postdemyelination. Arrowheads indicate demyelinated axons. Arrowheads indicate demyelinated axons. Arrows indicate thin myelin sheaths suggestive of remyelination. Scale bar = 0.5 μm. MD, myelin debris. E: Toluidine-blue stained image of demyelinated slice treated with 100 pmol/L fingolimod for 14 DIV postdemyelination demonstrates a healthy neuron (N) and oligodendrocyte (O), as well as many thin myelin sheaths (arrowheads) indicating remyelination. Scale bar = 5 μm.

**Figure 3.** Effects of fingolimod on remyelination following lysolecithin-induced demyelination. A: Representative images of demyelinated slices, control or treated with fingolimod (100 pmol/L) for 14 DIV postlysolecithin, immunostained against myelin (MBP; red) and axons (NFM; green). Fingolimod increases the amount of myelin associated with axons compared with control. Scale bar = 20 μm. B: Quantification of amount of myelin expressed as area of MBP immunostaining (μm²) per ×65 objective image normalized to control culture values at 14 DIV postlysolecithin. Fingolimod treatment (100 pmol/L, 1 μmol/L) for 14 DIV postlysolecithin significantly increases the amount of myelin compared with control cultures. **P** < 0.05. C: Electron micrograph of demyelinated slice treated with 100 pmol/L fingolimod for 14 DIV postlysolecithin. Fingolimod treatment (100 pmol/L) significantly enhanced remyelination in demyelinated slice cultures.

**Functional Basis for the Effects of Fingolimod During Remyelination**

To investigate the mechanism through which fingolimod enhanced remyelination in demyelinated slice cultures, we compared the effects of co-treating cultures with fingolimod and antagonists of specific S1P receptor subtypes on remyelination efficiency. Co-treatment of cultures with fingolimod (100 pmol/L) and a S1P1 receptor antagonist W123 only slightly decreased remyelination as assessed by MBP immunohistochemistry (from 12.9 ±
supplementation significantly reversed values compared with fingolimod alone (1.3 ± 0.5 fold over control; Figure 7, A and B). These findings suggest that whereas S1P3/5 signaling promotes astrogliosis, this is inhibited by S1P1 signaling, and blocking the latter further increases the astrocytic response induced by fingolimod. In support of this, SEW2871 applied to demyelinated cultures significantly decreased the area of GFAP immunostaining compared with control (0.5 ± 0.1 fold), whereas a S1P5 agonist induced a trend increase in values (1.3 ± 0.5 fold over control; Figure 7B).

The increase in microglia cell numbers observed at 14 DIV postdemyelination with fingolimod treatment alone (6.4 ± 0.8 fold over control) were slightly yet not significantly decreased with W123 co-treatment (5.1 ± 0.5), yet were increased when suramin was supplemented to the cultures (10.7 ± 1.0). We observed that treatment with SEW2871 or the S1P5 agonist was able to induce a significant increase in microglia cell numbers (6.2 ± 0.2 and 5.4 ± 0.4 fold over control, respectively). Together, the data suggests that signaling through S1P1 and S1P5 can contribute to enhanced microgliosis, yet the increase in cell numbers consequent to S1P3/5 blockade via suramin implies that S1P3 signaling may suppress this response. The increase in microglia cell numbers in the SEW2871-treated condition, where remyelination was inhibited, demonstrates that S1P receptor-driven enhancement of microgliosis is not sufficient to drive the myelin repair response.

**Discussion**

**Effect of Fingolimod on Myelin Maintenance**

Long-term treatment (21 DIV) of fully myelinated organotypic cerebellar slice cultures with both low (100 pmol/L) and high (1 μmol/L) doses of fingolimod did not induce any deleterious effect on myelin maintenance. Fingolimod promoted the elaboration of membranous lamellipodial extensions by OPCs and OLs, and increased OPC cell number. The elaboration of processes is critical for myelination to proceed and precise regulation of process extension by human-derived OPCs and mature OLs under basal culture conditions. We demonstrate that fingolimod-induced oligodendroglial membrane elaboration was not sufficient to increase the amount of MBP in the cultures. We hypothesize that all fibers that could be myelinated within the culture may have already been ensheathed thereby preventing any increase in myelination. In contrast with the findings we present here using organotypic slices, rodent OLs in dispersed cell cultures do not modulate their cytoskeleton in response to the drug and fingolimod alone has no effect on rodent and human OPC cell numbers. These findings indicate that OLs and OPCs may either modulate their S1P receptor levels in an environment containing multiple cell types, or that they are responding to factors being re-
leased by other fingolimod-stimulated neuroglial cells in the slice.

Effects of Fingolimod on Remyelination

Demyelination was induced in cerebellar slice cultures by overnight treatment with lysolecithin, inducing a loss of MBP associated with axons, a decrease in MBP immunostaining resembling debris, a decrease in the frequency of nodes of Ranvier, and a lengthening of the nodal region.

The adult CNS undergoes at least partial repair following injury. Myelin repair occurs via remyelination, and is considered to be associated with recovery of axonal function and clinical remittance. Studies of organotypic cerebellar slice cultures indicate that following a transient insult with lysolecithin, OPC differentiation and re-expression of myelin proteins occurs in the same temporal succession that would be observed in vivo. These studies demonstrated that the remyelination process is initiated within 1 week of demyelination, yet remains limited at 11 DIV postdemyelination. To document remyelination, we used MBP immunohistochemistry, which has previously been used to monitor this process in slice cultures, and co-labels with the late differentiation marker myelin oligodendrocyte glycoprotein by 11 DIV postdemyelination. Consistent with previous studies, the cerebellar slice cultures used in our current study initiated remyelination within 5 days of demyelination and underwent observable yet incomplete remyelination by 14 DIV postdemyelination. At 14 DIV postdemyelination, treatment with both high and low doses of fingolimod increased the amount of MBP immunostaining in demyelinated cultures in comparison with control. Toluidine-blue stained sections and ultrastructural analysis revealed the presence of many thin myelin sheaths in fingolimod-treated cultures suggestive of remyelination. Our findings using S1P receptor specific agonists and antagonists suggest that fingolimod enhanced remyelination primarily via S1P3/5 signaling; engagement of S1P5 was sufficient to induce a trend increase in remyelination. S1P5 is highly expressed on mature oligodendrocytes, and S1P-

Figure 5. Effects of fingolimod on microglia and astrocytes under remyelinating conditions. A: Representative images of slices at 55 DIV immunostained against the microglial marker IBA-1 (red). Microglia are increased at 2 DIV postlysolecithin and subsequently decrease by 14 DIV. Cultures treated with fingolimod (100 pmol/L) for 14 DIV postlysolecithin show relatively increased numbers of microglia relative to control cultures. Scale bar = 20 μm. B: Quantification of numbers of microglia represented as average IBA-1 immunopositive cells per ×63 objective image ± SEM. Microglia cell numbers are increased by 2 DIV postlysolecithin, which subsides by 5 DIV. Fingolimod-treated cultures (Fing; black bars) showed a significant increase in microglia cell numbers at 5, 9, and 14 DIV postdemyelination compared with controls (Ctrl; white bars). Delaying 14 DIV fingolimod treatment for 2 weeks following lysolecithin treatment still induced an increase in microglia cell numbers in comparison with control cultures at 28 DIV postlysolecithin. *P < 0.05, **P < 0.01, ***P < 0.001. C: Representative images of slices at 5 DIV immunostained against the astrocytic marker GFAP (red). Astrocytes are increased at 2 DIV postlysolecithin and subsequently decrease by 14 DIV. Cultures treated with fingolimod (100 pmol/L) for 14 DIV postlysolecithin show relatively increased GFAP immunostaining relative to control cultures. Scale bar = 20 μm. D: Quantification of astrocytes represented as area of GFAP immunostaining (μm²) ± SEM. Demyelination with lysolecithin (2 DIV postlysolecithin) induces an increase in GFAP staining compared with pre-demyelination controls (21 DIV). In 14 DIV postlysolecithin control cultures, GFAP staining shows a relative decrease compared with 2 DIV postlysolecithin. Fingolimod treatment (100 pmol/L) for 14 DIV postlysolecithin induces an increase in GFAP staining, as compared with control. **P < 0.01, ***P < 0.001.
mediated signaling has pro-survival effects on human and rodent mature oligodendrocytes.28,29 Whereas our studies using dissociated cultures of human OPCs demonstrated that S1P1 signaling could promote process extension, differentiation, and survival,27 here we observed that signaling through this receptor inhibited remyelination. One postulate for this unexpected result is that the growth factor environment within the slice may have modulated responses to fingolimod. Previous studies have demonstrated how responses to S1P receptor modulators may be influenced by growth factors, for instance PDGF and S1P receptor signaling are coordinated and this growth factor regulates expression levels of S1P receptors.34

The increase in absolute numbers of PDGFαR positive OPCs following lysolecithin treatment is consistent with the previously documented incorporation of BrdU into PDGFαR+ cells in lipopolysaccharide-treated organotypic cultures49 and increased OPC proliferation following demyelination in vivo.15 Whereas OPC numbers subsequently subsided over time in control cultures, by 14 DIV postlysolecithin, fingolimod treatment induced a trend increase in absolute numbers of OPCs. Increased OPC numbers may have reflected enhanced survival or proliferation, which have been documented to occur in dissociated cultures.27,30,34 More OPCs may have been available to participate in remyelination compared with control conditions. We hypothesize that the number of PDGFαR+ cells may not have reached significance with fingolimod treatment (100 nmol/L) due to a putative acceleration of OPC differentiation and consequent loss of PDGFαR expression. Fingolimod also induced process extension by both OPCs and OLGs under remyelinating conditions. Low doses of fingolimod induce differentiation of rat OPCs into mature OLs in dissociated cultures whereas high doses have the opposite effect.30,34 In our slices, OPCs may not have been exposed to high enough doses of the drug required to inhibit differentiation, perhaps as a result of sequestration by OLs/myelin.22

**Effect of Fingolimod on Remyelination**

DIV postlysolecithin, fingolimod treatment induced a trend increase in absolute numbers of OPCs. Increased OPC numbers may have reflected enhanced survival or proliferation, which have been documented to occur in dissociated cultures.27,29,30,34 More OPCs may have been available to participate in remyelination compared with control conditions. We hypothesize that the number of PDGFαR+ cells may not have reached significance with fingolimod treatment due to a putative acceleration of OPC differentiation and consequent loss of PDGFαR expression. Fingolimod also induced process extension by both OPCs and OLs under remyelinating conditions. Low doses of fingolimod induce differentiation of rat OPCs into mature OLs in dissociated cultures whereas high doses have the opposite effect.30,34 In our slices, OPCs may not have been exposed to high enough doses of the drug required to inhibit differentiation, perhaps as a result of sequestration by OLs/myelin.22

**Effects of Fingolimod on Microglia and Astrocytes Under Remyelinating Conditions**

We observed that microglial cell numbers peaked at 2 DIV postlysolecithin and declined by 14 DIV postlysolecithin, fingolimod treatment observed a significant increase in GFAP immunostaining compared with control cultures, suggesting a disinhibition of S1P3/5-associated signaling. Accordingly, suramin supplementation significantly decreased the area of GFAP immunostaining compared with fingolimod alone. Scale bar = 20 μm. **B:** Quantification of the astrocytic response represented as area of GFAP immunostaining (μm²) normalized to control at 14 DIV postdemyelination. Fingolimod-induced increase in GFAP immunostaining was significantly increased by W123 and reversed with suramin. The S1P1 agonist SEW2871 (100 nmol/L) significantly decreased GFAP immunostaining, as compared with control, whereas a S1P5 agonist (100 nmol/L) had no significant effect. *P < 0.05, **P < 0.01.
cithin in control cultures. Previous studies have demonstrated the abundance of microglia in these cultures and their activation consequent to demyelination. Microgloia are increased in numbers by 3 days following ethidium bromide-induced demyelination in vitro with concomitant increases in PDGFαR+ OPCs at the lesion border. Fingolimod-treated cultures demonstrated an increase in microglia cell numbers by 5 DIV following demyelination that was maintained to at least 14 DIV postdemyelination. When the drug application (for 14 DIV) was delayed by 2 weeks, the increase in microglia numbers was still detected, suggesting that its occurrence was not a consequence of a response to demyelination. One key mechanism whereby macrophages/microglia contribute to remyelination involves their capacity to phagocyte myelin debris, as confirmed by depletion of these cells with clodronate-liposomes. Using fluorescently-conjugated bacteria as an index of phagocytic activity, we observed a peak in numbers of phagocytic microglia shortly after demyelination (2 and 5 DIV) in control cultures. In cultures exposed to fingolimod, a decrease in the absolute numbers of actively phagocytic cells was detected at 5 and 14 DIV postlysolecithin. Although fingolimod treatment may have reduced the number of phagocytic microglia, the increase in total numbers of microglia could have been sufficient to result in myelin debris ingestion such that remyelination could occur. Ultrastructural analysis of fingolimod-treated slices indicated the presence of phagocytosed myelin debris; we could not appreciate any qualitative difference in the amount of MBP-immunoreactive debris found in fingolimod-treated cultures compared with control cultures at 14 DIV postdemyelination. Our studies using S1P receptor specific agonists and antagonists suggest that signaling via S1P1 and S1P5 contribute to microgliosis, whereas S1P3-mediated signaling may inhibit this response. The increase in microglia cell numbers under treatment conditions that either promoted remyelination (fingolimod, S1P5) or inhibited remyelination (SEW2871) suggest that S1P receptor engagement on astrocytes can induce growth factor production and growth factors can modulate S1P receptor gene expression in rat astrocytes in an autocrine manner such that S1P5 levels are increased. This suggests that within a physiological environment, astrocytes may primarily respond to fingolimod via S1P5, rather than through S1P1 as predicted from isolated cultures. Interestingly, treatment conditions under which astrogliosis was enhanced (fingolimod, S1P5) coincided with those sufficient to ameliorate remyelination, further supporting the beneficial role of astrocytes in the remyelination process.

**Conclusion**

Our studies suggest that fingolimod enhanced remyelination, which was well-correlated with an astrocytic response, both mediated via S1P3/S1P5, whereas a S1P1- and S1P5-dependent increase in microglia cell numbers and effects on phagocytosis did not appear sufficient to enhance the myelin repair process. These data establish that S1P receptor-specific signaling on multiple neuroglial cell types in a physiological environment can influence remyelination, in a manner that may not have been predicted from dissociated culture studies.

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