NEUROBIOLOGY

Oligodendrocyte Progenitor Cell Susceptibility to Injury in Multiple Sclerosis

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Accepted for publication April 8, 2013.

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The extent of remyelination in multiple sclerosis (MS) lesions is variable and often incomplete. In experimental models, oligodendrocyte progenitor cells (OPCs) rather than previously myelinating oligodendrocytes (OLs) are responsible for remyelination. This study compares the relative susceptibility of adult human OPCs and mature OLs to injury in actively demyelinating MS lesions and under in vitro stress conditions. In all lesions (n = 20), the number of OLs (Olig2 weak/NogoA positive) was reduced compared to control white matter (mean 38 ± 4% of control value). In 11 cases, OPC numbers (Olig2 strong; NogoA negative) were also decreased; in eight of these, the reduction was greater for OPCs than for OLs. In the other nine samples, OPC numbers were greater than control white matter, indicating ongoing OPC migration and/or proliferation. Analysis of co-cultures with rat dorsal root ganglia neurons confirmed that OPCs were more capable of contacting and ensheathing axons than OLs. In isolated culture under stress conditions (withdrawal of serum/glucose and/or antioxidants), OPCs showed increased cell death and reduced process extension compared to OLs. Under all culture conditions, OPCs up-regulated expression of genes in the extrinsic proapoptotic pathway, and had increased susceptibility to tumor necrosis factor–induced cell death as compared to OLs. Our data suggest that susceptibility of OPCs to injury within the MS lesion environment contributes to the limited remyelination in MS. (Am J Pathol 2013, 183: 516–525; http://dx.doi.org/10.1016/j.ajpath.2013.04.016)
the excitotoxin glutamate compared to more mature OLs.19–21 Ziabreva et al.,22 however, found that inhibitors of mitochondrial respiratory complex IV induced greater injury of OLs matured in vitro from the neonatal rat brain compared to OPCs.

In the current study, we show that relative numbers of OPCs can be more reduced than mature OLs in actively demyelinating MS lesions. Using cells isolated from noninflamed adult human brain samples, we provide evidence that OPCs (A2B5 antibody selected) rather than previously myelinating mature OLs have greater capacity to ensheathe axons. Furthermore, we demonstrate that cultured OPCs have enhanced vulnerability to injury compared to mature OLs when placed under metabolic stress conditions (withdrawal of serum/glucose and/or antioxidants); under all culture conditions, OPCs had higher expression of genes in the extrinsic proapoptotic signaling cascade and were more susceptible to subsequent TNF-α–induced injury.

Materials and Methods

Analysis of OPCs and OLs in Early MS Lesion Stages

We analyzed surgically obtained brain samples containing actively demyelinating lesions from MS patients for the expression of OL lineage transcription factor 2 (Olig2) as a marker of OPCs and a member of a family of integral membrane proteins termed reticulons (NogoA) as a marker of mature OLs.23 All actively demyelinating lesions (active) showed macrophage infiltration and/or microglial activation and presence of myelin degradation products within macrophages [2',3'-cyclic nucleotide 3'-phosphodiesterase (CNP), myelin OL glycoprotein (MOG), myelin associated glycoprotein (MAG)].23,24 Two sets of samples were evaluated, namely, samples included in an earlier study (n = 16)6 and four new samples. Each was compared to brain tissue specimens that were reactive with O1 antibody (recognizes GalC28), whereas almost all (85% to 95%) cells in the negatively selected fraction were O1+ (Supplemental Figure S1).

An Affymetrix-based microarray comparison (Affymetrix, Santa Clara, CA) between the two cell fractions at the time of isolation indicated that the A2B5+ cells had increased expression of mRNA for platelet-derived growth factor receptor (PDGFR)α (mean fold increase for the 36 probes of the PDGFRα gene = 2.14; P = 0.6 × 10−9, paired t-test) and for protein tyrosine phosphatase, receptor-type, Z1 (PTPRZ1) (mean fold increase = 6.64; P = 0.19 × 10−5) (data not shown), markers of OPC,29 supporting that this fraction is enriched in progenitor cells.

Human OPC or OL and Rat Dorsal Root Ganglion Neuron Co-Cultures

Purified dorsal root ganglion neuron (DRGN) cultures were prepared from Sprague-Dawley rat embryos as described previously.30 The cells were maintained with 12.5 ng/mL nerve growth factors (NGF) in serum-free N1-supplemented medium for 3 weeks before the addition of human cells to allow differentiation of neurons and extension of a network of neurites competent for myelination. OPCs (A2B5+) or OLs (A2B5−) at a density of 0.7 × 105 cells/cm2 suspended in defined medium were added to DRGN cultures. Defined medium consisted of Dulbecco’s modified essential medium—F12 supplemented with N1 (Sigma-Aldrich, Oakville, ON, Canada), 0.01% bovine serum albumin, 1% penicillin-streptomycin, and B27 supplement (Invitrogen, Burlington, ON, Canada), 10 ng/mL platelet-derived growth factor (PDGF)-AA, 10 ng/mL basic fibroblast factor (bFGF), and 2 nmol/L triiodothyronine (T3) (Sigma-Aldrich) (referred to as DM+GF). The OPCs and OLs were used immediately after fractionation. The extent of cell survival, differentiation, and axonal ensheathment was analyzed after 5 weeks of co-culture.

Isolated OPC and OL Cell Cultures

To assess the relative susceptibility of OPCs and OLs to injury (cell loss, TUNEL–labeled nuclei, and process retraction)
under metabolically stressful conditions, OPCs or OLs were plated (2.5 × 10⁶ cells per mL) on polystyrene-coated chamber slides and cultured for 4 to 10 days under conditions that ranged from optimal to metabolically deprived; these involved defined media supplemented with growth factors (DM+GF); minimum essential medium (MEM) with 5% fetal calf serum (FCS) (MEM+GF); DM+GF without antioxidants (DM+GF-AO), and MEM depleted of glucose and serum (MEM alone), respectively. In addition, experiments, 100 ng/mL TNF-α was added to both cell types for 48 hours under the culture conditions indicated in the Results section.

Immunocytochemistry

For assessment of cell-surface markers, cells in either isolated or co-cultures were incubated with primary antibodies (O4 and O1) (Table 1) for 30 minutes at 4°C then fixed in 4% paraformaldehyde for 10 minutes at 4°C, followed by blocking with HHG (1 mmol/L HEPES, 2% horse serum, 10% goat serum, Hanks’ balanced salt solution) for 10 minutes. Cultures were incubated with secondary antibodies conjugated with fluorescent dyes (Table 1) for 30 minutes at 4°C. Antibody isotype controls showed low nonspecific staining (data not shown). Cell nuclei were stained with 10 μg/mL Hoechst 33258 (Invitrogen) for 10 minutes at room temperature. For isolated cultures, TUNEL staining (data not shown). Cell nuclei were stained with 10 μg/mL Hoechst 33258 (Invitrogen) for 10 minutes at room temperature. For isolated cultures, TUNEL staining was determined by two-way analysis of variance, followed by either Dunnett’s or Bonferroni’s multiple comparison tests. Probability values <0.05 were considered statistically significant.

Results

Evaluation of OPC and Mature OL Numbers in MS Lesions in Situ

In an earlier publication, we reported the presence of OPCs in most early and chronic MS lesions with a significant increase of OPCs in the periplaque white matter (PPWM). Now, we assess the relative loss of OPC as compared to mature OLs in individual actively demyelinating lesions; the

### Table 1 Primary and Secondary Antibodies Used in This Study

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<th>Clonality</th>
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RT² Profiler Apoptosis PCR Array

The expression of 84 apoptosis-related genes was examined using the 84-gene human apoptosis PCR array kit (PAHS-012; SABiosciences/Qiagen, Mississauga, ON, Canada). Specific genes included are detailed in Table 2. RNA was extracted from the A2B5+ selected and A2B5− oligodendroglial lineage cells immediately after fractionation (day 0) or after 6 days in culture. Total RNA was reverse-transcribed into cDNA using the RT² First Strand Kit (SABiosciences), mixed with RT² qPCR Master Mix containing SYBR Green (SABiosciences), and aliquoted in equal volumes to each well of the real-time PCR arrays. The threshold cycle (Ct) of each gene was determined and normalized using multiple housekeeping genes and subsequently analyzed by RT² Profiler PCR Array Data Analysis software version 3.5. Data from three separate cell preparations were analyzed.

Data Analysis

Results are presented either as means ± SEM or as fold/percent difference over control for corresponding time points. The number of individual experiments for each study is indicated in the Results section. Statistical significance was determined by two-way analysis of variance, followed by either Dunnett’s or Bonferroni’s multiple comparison tests. Probability values <0.05 were considered statistically significant.
results are expressed as a percentage of mean values for the corresponding cell type found in nondemyelinating control white matter sections stained at the same time. As shown in Figure 1, A and B, the number of mature OLs (NogoAstr) was increased compared to control white matter sections. The relative number of mature OLs was reduced in the PPWM compared to control white matter (mean 55 ± 5% of control value). Examples of immunostaining for OPCs (Olig2str) and mature OLs (NogoA) in active lesions and PPWM are illustrated in Figure 1D.

Comparison of Axonal Ensheathing Capacity of A2B5+ OPCs and Mature OLs Co-Cultured with DRGNs

Figure 2, A and B, illustrates the extent of cell survival, differentiation, and axonal ensheathment of OPCs and OLs after 5 weeks in co-culture with DRGNs. OPCs establish contacts with axons and begin the ensheathment process (Figure 2A). Clustering of the paranodal protein contactin-associated protein (Caspr) indicates the initial formation of

<p>| Table 2 Categorization of PCR Array Genes |</p>
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*Member of TNF receptor superfamily.

were reduced in both lesions and PPWM, with reduction being greater in the active lesions (Figure 1C). In the remaining three cases, OPC numbers were relatively increased in both lesions and PPWM compared to control tissue (Figure 1C). The relative number of mature OLs was reduced in the PPWM compared to control white matter (mean 55 ± 5% of control value). Examples of immunostaining for OPCs (Olig2str) and mature OLs (NogoAstr) in active lesions and PPWM are illustrated in Figure 1D.
axon domains; however, sodium channel clustering (Figure 2, C, D, G, and H), which is fundamental to the nodes of Ranvier formation, is not evident. This pattern indicates that the process of myelination has begun but remains incomplete. By contrast, for the mature OLs, there was limited process formation, axonal contact, or ensheathment (Figure 2B). The few sites of Caspr aggregation occurred between axons and OL cell bodies rather than between cellular processes (Figure 2, E, F, I, and J), in contrast to OPC/axon interactions. Differences in axon ensheathment by OPCs and OLs and in Caspr redistribution along axons are quantified in Figure 2, K and L.

Comparison of OPC and Mature OL Responses to Metabolic Injury in Vitro

Comparison of Injury Response of A2B5+OPCs and A2B5−OLs in Isolated Cell Cultures

The relative survival and process outgrowth of OPCs or OLs after 6 days in isolated cell culture under different conditions are quantified in Figure 3, A–C. Figure 3, D–K, illustrates these findings. Under optimal conditions (DM+GF), both OPCs and OLs survive well (<5% TUNEL+ cells) and extend cellular processes. Relative process area per cell (Figure 3C) is greater for OPCs (10.5 ± 0.4 μmol/L²/cell) compared to OLs (8.6 ± 0.3 μmol/L²/cell; P < 0.05, n = 4 separate cultures). Under the most metabolically deficient conditions, MEM without glucose and serum (MEM alone), there was a significant increase in the percentage of TUNEL-positive OPCs as compared to OLs (25 ± 2.7% versus 15 ± 1.2%, P < 0.001) (Figure 3A) at day 6. The total number of OPCs decreased from 212 ± 2 per field under optimal conditions (DM+GF) to 95 ± 5 in MEM alone, accounting for a 56% reduction (Figure 3B). For mature OLs, the mean number of surviving cells in MEM was 142 ± 8 compared to 176 ± 6 in DF+GM, accounting for only 19% reduction. The relative process area per cell in MEM was significantly reduced for both cell types (71 ± 1.5% for OPCs versus 60 ± 2% for OLs) as compared to DM+GF (Figure 3C). The relative reduction in OPC process area was significantly greater than for OLs (P < 0.01).

Under the less severe conditions of MEM+FCS, neither OPCs nor OLs showed enhanced cell death as measured by the percentage of TUNEL+ cells or the total number of surviving cells compared to optimal conditions at day 6 (Figure 3, A and B). However, by 3 weeks, the numbers of surviving OPCs decreased by 56 ± 1.7%, whereas surviving OLs decreased by 34 ± 3.5% (P < 0.01) (data not shown). At day 6, the relative process area per cell was significantly reduced for both cell types (45 ± 2.1% and 38 ± 1.8% of values compared to that observed under optimal conditions) (Figure 3C). Under culture conditions in which antioxidants were removed from DM+GF (DM+GF-AO), OPCs, but not the OLs, showed an increase in the percentage of TUNEL+ cells (Figure 3A). The relative process area per cell for both OPCs and OLs was reduced as compared to DM+GF (Figure 3C).

Figure 2  Comparison of A2B5+ OPCs and mature OLs in co-culture with DRGN. Human adult OPCs (A, C, D, G, and H) or OLs (B, E, F, I, and J) were seeded on established rat DRGNs maintained in DFM with NGF alone or NGF+GFs (BDNF/IGF-1/PDGF-AA/bFGF) for 5 weeks. Co-cultures were fixed and immunostained either for neurofilaments (NF) (green) and MBP (red) in A and B, or 01 (green, C and E) and Caspr (red, D and F), or sodium channel (blue, G and I). Merged images are presented in H and J. In the co-cultures established with OPCs (A2B5+ cells), MBP+ cells ensheathing multiple DRGN axons are indicated by asterisks in A and B; fewer ensheathments were observed in co-cultures with the A2B5− fraction (OLs) in B. Quantitative data are presented in K. Caspr redistribution on axons in presence of ensheathing cells in OPC co-cultures is indicated by arrows in D and H; nonredistributed Caspr is observed as specks in the background. More limited Caspr (red) redistribution is seen in OL co-cultures in F and J. Quantitative data are provided in L. There was no significant redistribution of sodium channels in G and I. Data represent three independent experiments performed in duplicate for each condition (approximately 200 cells were counted per coverslip). Statistical significance was determined by one-way analysis of variance, followed by Dunnett’s test. ***P < 0.001. Images were acquired by confocal microscopy. Scale bars: 10 μm.

Comparative Changes in Apoptosis-Related Gene Expression in OPCs and OLs under Different Culture Conditions

In initial studies, we compared expression of cell death–related genes between A2B5 antibody—selected (OPCs) and nonselected (OLs) cells in an Affymetrix Human Exon gene screen; three separate central nervous system (CNS) donor samples were used. Of genes designated as being cell death relevant [Ingenuity Pathway Analysis (IPA) program version 8.6; Ingenuity Systems, Redwood City, CA] that
were significantly differentially expressed between the two populations using a paired sample analysis, all were up-regulated in the OPC fraction. Supplemental Figure S2 provides a heat map representation of the data.

When the microarray results were fitted to the Kyoto Encyclopedia of Genes and Genomes (KEGG) apoptosis pathway using Partek GS software version 6.6, differentially up-regulated proapoptotic genes were found in the OPC samples (Supplemental Figure S3).

To compare apoptosis pathway—related gene expression from both ex vivo and varying in vitro culture conditions, we used an 84-gene PCR apoptotic pathway array. Data from the 84 apoptosis-related gene PCR array analyses are presented under overall categories of anti- and proapoptotic genes; the latter are further subdivided into intrinsic, extrinsic, and final effecter apoptotic pathway genes (Table 2).

As presented in Figure 4, there was no significant variance in expression of pro- (extrinsic, intrinsic, and effecter) or antiapoptotic genes between the OPC (A2B5+) and OL (A2B5−) fractions at time of initial cell subset fractionation (day 0). The mean number of antiapoptotic and effecter pathway genes, expressed at >2-fold level by OPCs compared to OLs, did not reach statistical significance under any culture condition (Figure 4A and B; heat map analysis provided in Supplemental Figure S4). Similarly, the intrinsic pathway was not significantly altered (Figure 4C). However, under the most extreme culture condition (MEM alone), there was a net increase (>2 fold) in expression of intrinsic proapoptotic genes for both cell types compared to optimal conditions (DM+GF) (for OPCs, n = 5.3 ± 1.2 versus 1.0 ± 1.0; P < 0.05) (Table 3 and Supplemental Figure S5); (for OLs, n = 4.0 ± 2.1 versus 0.3 ± 0.4; P = 0.11) (Table 3 and Supplemental Figure S6). We did not detect any significant differences under the other culture conditions (MEM+FCS or DM+GF-AO versus DM+GF).

The most significant change in gene expression was observed for the extrinsic proapoptotic gene category that was up-regulated under all culture conditions in the OPC population compared to OLs (approximately 8 to 10 of 23 genes evaluated) (Figure 4D). These included receptors and ligands.

Figure 3 Comparative phenotypic responses of OPCs and OLs under optimal and suboptimal dissociated cell conditions. Percentage TUNEL+ cells under each condition for either cell type are summarized in A. Total cell numbers per 25× field are summarized in B. The relative process area per cell calculated as a percentage of that defined in the DM+GF condition for OPCs is shown in C. OPCs (D to G) and OLs (H to K) cultured in DM+GF (D and H), MEM alone (E and I), MEM+FCS (F and J), or DM+GF-AO (G and K) for 6 days before O4 antibody (red) and TUNEL (green), and Hoechst (blue) staining. Arrows in E, G, I, and K indicate TUNEL+ cells. Data represent three independent experiments performed in duplicate for each condition. Scale bar = 20 μm. Statistical significance was determined by two-way analysis of variance, followed by Dunnett’s test or Student’s t-test. *P < 0.05, **P < 0.01, and ***P < 0.001 versus corresponding controls; 1P < 0.05, 11P < 0.01, and 111P < 0.001 OPCs versus OLs.
alone) conditions, to 100 ng/mL TNF-α
and the percentage of TUNEL
(Figure 5). In MEM, both OPC and OLs showed an increase in
the 84 genes included in the PCR apoptosis array between cell types when
the 0.01 up-regulated versus down-regulated genes.

To assess whether the increased expression of the extrinsic
apoptotic pathway in OPCs could result in enhanced susceptibility to TNF-α-mediated injury, we exposed cultures, maintained under optimal (DM+GF) or suboptimal (MEM alone) conditions, to 100 ng/mL TNF-α for 48 hours (Figure 5). In MEM, both OPC and OLs showed an increase in the percentage of TUNEL+ cells after treatment with TNF-α. Under both optimal and suboptimal conditions, TNF-α produced a greater increase in the percentage of TUNEL+ cells for OPCs compared to OLs (P < 0.05).

**Discussion**

Our *in situ* analysis implicates OPCs as a component of the tissue injury that occurs in active demyelinating lesions in MS. The cell markers used to identify OPCs and OLs in white matter are the same as used previously, having been optimized for use in fixed human tissue sections. The percentage of mature OLs was reduced in all of the actively demyelinating lesions, as well as in the PPWM, compared to control white matter, suggesting that restoration of mature OLs remains insufficient over the time course we have evaluated. OPCs (strongly Olig2+ cells) were reduced in 11 of 20 actively demyelinating lesions when compared to non-MS control tissue. In eight of these cases, the relative reduction was greater for OPCs than for mature OLs in the corresponding sample, suggesting that OPCs were more vulnerable to injury mediators in the same microenvironment.

In the other nine cases, OPC numbers were increased in the lesions when compared to non-MS control tissue, suggesting that OPCs have migrated or proliferated to initiate tissue repair. These findings are consistent with experimental studies showing migration or proliferation of OPCs during initiation of remyelination. OPC numbers were also on average increased in PPWM, even in cases where OPC numbers in lesions remained reduced. The low numbers of OPCs in some lesions, despite a marked increase of OPCs in PPWM, may hint also to a disturbed migration of OPCs into MS lesions as described in an earlier study.

Our co-culture studies demonstrate that the cells designated as OPCs (A2B5 selected), rather than the mature OLs, have greater capacity to ensheath axons, the initial step in myelin formation. OPCs selected with the A2B5 antibody from the adult human CNS are able to myelinate following adoptive transfer into the CNS of the dysmyelinated shiverer mouse mutant. Earlier observations showing that mature OLs were present in demyelinated MS lesions led to the postulate that these cells contributed to remyelination. However, when mature OLs were transplanted into experimentally demyelinated lesions, they failed to remyelinate. Conversely, depletion of OPCs prevented successful remyelination. The mature OLs used in our study are a post-mitotic cell population that expresses myelin basic protein (MBP) and MOG. Such cells comprise approximately 30% of the total numbers recovered from adult human brain specimens, supporting the conclusion that these are previously myelinating OLs and not derived over time *in vitro* from a relatively small progenitor population. Our microarray analysis further demonstrated that the A2B5-selected cells expressed higher levels of the OPC-associated markers PDGFRα and the protein tyrosine phosphatase, receptor-type, Z1 (PTPRZ1), further strengthening the notion that these cells are indeed OPCs and that the nonselected fraction consists of mature OLs.

In the OPC-DRGN co-cultures, we observed clustering of Caspr, an indication of axonal response to contact with OPCs. However, the extent of sodium channel redistribution was limited and less than observed previously using fetal human CNS-derived OPCs. We cannot determine whether this limited response reflects the experimental paradigm, for example, due to a species difference, or the intrinsic properties of the adult CNS-derived OPCs. We note that rodent OPCs produce extensive myelination of DRGN axons in this system. Our isolated cell culture observations regarding process outgrowth under optimal conditions by mature OLs indicate that the limited capacity of mature OLs to ensheath axons cannot be attributed to an intrinsic inability to extend cellular processes.

Comparative *In Vitro* Injury Responses of OPCs and OLs to Metabolic Insults

Our comparative cell survival studies in isolated culture suggest that under all growth conditions, OPCs are more vulnerable to metabolic insults than mature OLs. The
OPCs or OLs maintained in defined culture conditions for 6 days before RNA extraction. Real-time PCR microarray was performed using a human apoptosis PCR array kit as indicated in methods. The 84 genes were classified into four groups as listed in Table 2. Data are expressed as mean number of genes ±SEM with >2 fold increased (†) or decreased expression (‡) under suboptimal conditions (MEM+FCS, MEM, DM+GF-AO) compared to optimal condition (DM+GF) in OPCs or OLs. Analyses of all gene changes are presented in Supplemental Figures S3 and S4 as heat maps.

*P < 0.05.

greatest differences in cell death were observed under the extreme growth deprivation conditions when both growth factors and antioxidants were removed. We note that under such conditions, A2B5 cells selected from the fetal human CNS do not survive at all (data not shown). However, significant differences were also observed under less extreme conditions, such as prolonged culture (3 weeks) in MEM+FCS or when antioxidants alone were withdrawn. With regard to mechanisms of injury, we found a modest increase in the number of intrinsic cell-death pathway genes up-regulated in OPCs versus OLs after 6 days under the most severe culture conditions. We cannot exclude the possibility that intrinsic or final effector signaling pathway activation would be more apparent if cultures were analyzed at earlier time points or that cell necrosis could also contribute to cell death. Both in vitro and in situ studies demonstrate that ischemia can produce cytotoxic as well as active cell death, including the death of OLs (reviewed by Dewar et al40). Our observations of a maturation-dependent susceptibility of oligodendroglial injury are consistent with earlier studies demonstrating similar results using rodent cells and animal models for periventricular leucomalacia.41,42

Under all culture conditions, OPCs up-regulated extrinsic apoptotic pathway genes to an extent greater than OLs. Such responses could be triggered either directly by the selective sensitivity of these cells to the stress of culture conditions compared to OLs or be triggered by molecules of the intrinsic pathway or molecules released by injured or dying cells. Studies on PC1243 and Jurkat cell lines,44 as well as primary dopaminergic neurons,45 showed that withdrawal of trophic support, as in our experiments, can induce expression of genes in the extrinsic pathway resulting in cell death. Overexpression of extrinsic pathway signaling receptors and ligands may have specific relevance for continuing injury in MS lesions.46,47 We previously showed that inducing sublethal increases in expression of p53, an intrinsic pathway cell death—promoting molecule in OLs by adenovirus transduction resulted in up-regulated expression of members of the TNF-R superfamily, making the OLs more vulnerable to fas- and TNF-related apoptosis-inducing ligand (TRAIL)-mediated cell death.48 Our experiments further show that the OPCs are more vulnerable to TNF-z-mediated injury than are the mature OLs when assessed under optimal and suboptimal growth conditions.

**Conclusions**

We suggest that vulnerability of human OPCs to conditions that induce injury of mature OLs in MS lesions contribute to the limited remyelination observed in both acute and chronic stages of MS. Studies based on animal models indicate that the severity and duration of an injury is an important determinant of the extent to which remyelination will be limited. Studies using the OL lineage toxin cuprizone indicate that remyelination can occur after repeated

![Figure 5](Image 87x163 to 242x295)
rounds of injury, but if the injury is sufficiently severe (eg, prolonged exposure to high-dose toxin), recovery will fail to occur. In MS, repeated or chronic insults to OPCs would be expected. Multiple additional variables are also expected to contribute to failure of remyelination, including the presence of inhibitory molecules, absence of signals needed to recruit and induce OPC differentiation, and compromised integrity of demyelinated axons.

Supplemental Data

Supplemental material for this article can be found at http://dx.doi.org/10.1016/j.ajpath.2013.04.016.

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