Early Onset of Axonal Degeneration in Double (*plp*^{−/−}*mag*^{−/−}) and Hypomyelination in Triple (*plp*^{−/−}*mbp*^{−/−}*mag*^{−/−}) Mutant Mice

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Double (*plp*^{−/−}*mag*^{−/−}) and triple (*plp*^{−/−}*mbp*^{−/−}*mag*^{−/−}) null-allelic mouse lines deficient in proteolipid protein (PLP), myelin-associated glycoprotein (MAG), and myelin basic protein (MBP) were generated and characterized genetically, biochemically, and morphologically including their behavioral capacities. The *plp*^{−/−}*mag*^{−/−} mutant develops a rapidly progressing axon degeneration in CNS with severe cognitive and motor coordinative deficits but has a normal longevity. CNS axons of the *plp*^{−/−}*mbp*^{−/−}*mag*^{−/−} mouse are hypomyelinated and

ensheathed by “pseudomyelin” with disturbed protein and complex lipid composition. The *shiverer* trait in the *plp*^{−/−}*mbp*^{−/−}*mag*^{−/−} similar to the *plp*^{−/−}*mbp*^{−/−} mutant is significantly ameliorated, and its lifespan is considerably prolonged. The longevity of these dysmyelination mouse mutants recommends them as suitable models for the long-term evaluation of stem cell therapeutic strategies.

Key words: double mutants; triple mutant; myelin proteins; myelin lipids; ultrastructure of myelin; behavioral tests

Myelination of axons by oligodendrocytes in the CNS and Schwann cells in the PNS of vertebrates is an important event in the early development. In the CNS oligodendrocytes synthesize myelin specific proteins and lipids in a stringently regulated temporal sequence and assemble them in plasmamembrane processes that wrap axons as a highly compacted multilayer membrane to insulate internodes.

Proteolipid protein (PLP) and DM₂₀, a smaller isoform derived from alternative splicing, are the two major integral membrane proteins of CNS myelin (Stoffel et al., 1984). Their correct integration into the highly ordered membrane sheath is required for stabilizing the apposition of the adjacent extracellular membrane surfaces, ultrastructurally appearing as the interperiod dense line (IDL). Besides the naturally occurring mutants, e.g., *jimpy* (Nave et al., 1986) and *rumpshaker* (Schneider et al., 1992), two knock-out mouse models (Boison and Stoffel, 1994; Klugmann et al., 1997) have elucidated the structure–function relationship of this tetraspan membrane protein (Weimbs and Stoffel, 1992). In contrast to the point mutations (*jimpy*, *rumpshaker*), substantial myelination was observed in *plp*^{−/−} mice with a loss of the tight compaction at the IDL. The phenotype of the PLP-deficient mouse was inconspicuous, and a late onset of axon degeneration in the aging *plp*^{−/−} mouse suggests a possible function of PLP and/or DM₂₀ for axonal maintenance (Griffiths et al., 1998).

The family of myelin basic protein (MBP) isoproteins (21.5, 18.5, 17, and 14 kDa) results from alternative splicing. They comprise 30–40% of CNS and 5–15% of PNS myelin proteins (Lees and Brostoff, 1984). MBP plays a crucial role in the compaction of the opposing cytosolic surfaces of the plasma membrane processes, the main dense line (MDL) in electron microscopy (Omlin et al., 1982). The spontaneous *shiverer* mouse mutant, a “natural” knock-out, with deletion of exons 2–7 of the MBP gene (Roach et al., 1985; Molineaux et al., 1986) causes severe hypomyelination of CNS axons and leads to premature death within 3 months. No major

morphological abnormalities are observed in the PNS (Rosenbluth, 1980).

A third minor, but important constituent of the CNS myelin membrane of differentiating oligodendrocytes and in PNS, is the myelin-associated glycoprotein (MAG). MAG appears as a component of the surface (Yim et al., 1995). Its main function is thought to mediate axon–glia contact during myelin assembly (Arquint et al., 1987; Trapp, 1990). MAG expression occurs in a time-specific manner and yields the L-MAG isoform predominantly expressed at an early and S-MAG at a later stage of myelination (Lai et al., 1987; Tropak et al., 1988; Inuzuka et al., 1991; Pedraza et al., 1991). Subtle ultrastructural abnormalities in CNS myelin of *mag*^{−/−} mice were observed (Li et al., 1994; Montag et al., 1994). Axonal and myelin degeneration in the PNS but not in CNS was observed in older *mag*^{−/−} mice (Fruttiger et al., 1995; Yin et al., 1998).

The characterization of the mouse models generated in this study together with the monogenetic mutants [*plp*^{−/−}, *mbp*^{−/−} (*shiverer*) and *mag*^{−/−}] and the *plp*^{−/−}*mbp*^{−/−} double mutant provided extensive insight into the pivotal functions of the main myelin proteins for oligodendrocyte development, myelin formation, myelin structure, compaction, and maintenance of the myelin membrane, and particularly for axonal integrity.

MATERIALS AND METHODS

Animals. PLP-deficient mice (*plp*^{−/−}) (Boison and Stoffel, 1994) were crossed with MAG-deficient mice (*mag*^{−/−}) (kindly supplied by Prof. Bruce Trapp, Cleveland Clinic Foundation, Cleveland, OH) (Li et al., 1994). The resulting F1 generation was intercrossed, and mice homozygous for the PLP/MAG deficiency (*plp*^{−/−}*mag*^{−/−}) were obtained. We generated the PLP/MBP/MAG-deficient triple mutant mouse line (*mag*^{−/−}*plp*^{−/−}*mbp*^{−/−}) by crossing MAG-deficient mice (*mag*^{−/−}) with PLP/MBP-deficient animals from our transgenic breeding facility (Stoffel et al., 1997). The homozygous PLP/MBP/MAG-deficient triple mutant was obtained by intercrossing (*plp*^{Y/−}*mbp*^{−/−}*mag*^{+/−}) male with (*plp*^{−/−}*mag*^{−/−}*mbp*^{+/−}) female mice. MAG/PLP/MBP-deficient triple mutant animals are infertile. *Shiverer* mice (*shi/shi* = *mbp*^{−/−}) were selected from crossings of heterozygous animals (*mbp*^{+/−}) (The Jackson Laboratory, Bar Harbor, ME).

Genotyping. Genotyping of the wild-type and mutant PLP allele was performed by hybridizing *Nco*I-digested genomic DNA with a genomic 700 bp *Bg*/II fragment that contains sequences of exon III and intron III of the PLP gene. A 3.6 kb fragment is indicative of the wild type and a 2.7 kb signal of the mutated allele. The deletion of the *mbp* gene (exons II to VII) was verified by Southern blotting of *Bam*HI-digested genomic DNA with a 350 bp PCR fragment obtained with oligonucleotide primers hybridizing to a 5′ intron sequence upstream of the deleted exon II, 5′GAGGCCGCA-CATCAGCCCTGATTTTGTGCTAAG3′, and a corresponding 3′ primer,

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5'-CATGTATGAATGTGCATCTTGGGCAATCTATCT3', hybridizing to the sequence flanking the deletion downstream of exon VII. A 3.5 kb *Bam*HI fragment is diagnostic of the wild type and a 7 kb signal of the *shiverer* locus. The MAG genotype was examined by probing *Bam*HI/*Hind*III restricted DNA with the 270 bp *Eco*RI/*Acl*I fragment of exon VII. A 7.7 kb band indicates the wild type and a 5.5 kb signal the targeted allele.

RNA analysis. RNA was prepared from total brain of 20-d-old mice (Chomczynski and Sacchi, 1987).

Northern blot. RNA (20 µg) was separated by 1% formaldehyde-agarose gel electrophoresis and blotted to nitrocellulose (GeneScreen Plus; NEN Life Science Products, Boston, MA) and hybridized with the following genomic and cDNA fragments of PLP, MBP, MAG, and GAPDH labeled with the random priming DNA-labeling kit (Boehringer Mannheim, Mannheim, Germany): a genomic 700 bp *Bgl*II fragment containing sequences of exon III and intron III of the PLP gene, a 266 bp PCR-fragment of MAG exon IX (5' primer: 5'-GCAGTTGCCCTATATCTTCTGAGTCA3', 3' primer: 5'-CTCTGGGTGCCATAGAGGTTCTTGGAGTCA3'), a 540 bp PCR cDNA fragment of MBP (5' primer: 5'-CAAGAA-GACCCACAGCAGCTTCCGGAGGC3', 3' primer: 5'-CAGGATTCGGAAGGCTGAGCGGGGAGGGC3'), and a 550 bp *Hind*III/*Xba*I cDNA fragment of human liver glyceraldehyde 3-phosphodehydrogenase (GAPDH). Fragments were purified using the Quiaex gel extraction kit (Qiagen, Chatsworth, CA).

Quantitative RT-PCR. Five micrograms of RNA treated with RNase-free DNase were subjected to the first strand synthesis (SuperscriptII; Life Technologies GmbH, Karlsruhe, Germany). We used 1/20 of the reaction for each PCR in 30 µl in the presence of 0.1 µl of ³²P-dCTP. Samples were taken after 15, 17, 19, and 21 cycles and separated on 6% polyacrylamide gels. Gels were dried and analyzed with a PhosphorImager (Molecular Dynamics, Sunnyvale, CA) using the Imagequant software (Molecular Dynamics). The following oligonucleotide primers were used: MAG: 5': 5'-TGCTCACCAGCATCTTCACG3', 3': 5'-AGCAGCCTCTCTCAGATCC3'; PLP: 5': 5'-CAAGCTCATTCTTTGGAGCG3', 3': 5'-CAATCATGAAGGTGAGCAGG3'; MBP: 5': 5'-TACCTGGCCACAGCAAGTAC3', 3': 5'-GTCACAATGTTCTTGAAG3'; GAPDH: 5': 5'-GAGCTGAACGGGAAGCTCAC3', 3': 5'-CACCACCCTGTTGCTGTAGC3'; OMgp: 5': 5'-GCAGCAGCTGCAACTCTAAC3', 3': 5'-GAAGCATTTACTTTCCAAGCA3'; MOG: 5': 5'-CCAAGAGGAGCAGCAATGG3', 3': 5'-GTTGTAGCAGATGATCAAGG3'; NCAM120: 5': 5'-CCTGAAGAGCATCCAGTACA3', 3': 5'-ATGAATTCCAAGGACTCTG3'; NCAM140: 5': 5'-CCATCAGACACTATCTGGTC3', 3': 5'-TCAGGAAGTAGCAGGTGATG3'; CGT: 5': 5'-GAAATTCAAGGATCAACC3', 3': 5'-GTCCATTAAGTGTGCTATGC3'.

Isolation of myelin and protein analysis. Myelin of total brain of 28-d-old mice was purified by sucrose gradient centrifugation. One brain was homogenized in 10 ml of 0.32 M sucrose. The suspension was layered on top of 20 ml of 0.85 M sucrose and centrifuged at 10,000 × *g* for 20 min. The myelin-containing interphase was collected with a bent Pasteur pipette, diluted with three volumes of water, sedimented at 10,000 × *g* for 30 min, and washed twice with water (Norton and Poduslo, 1973). Aliquots were separated on 10–15% SDS-PAGE. Individual proteins were visualized by Western blot analysis (Towbin et al., 1979) with SuperSignal Substrate (Pierce, Rockford, IL), using anti-MBP antibodies (Boehringer Mannheim) and polyclonal anti-PLP (whole protein) and anti-MAG (Gln³⁷⁶-Tyr³⁹⁰) antibodies generated in our laboratory.

PNS proteins were isolated by homogenization and sonication of sciatic nerves in 200 µl of BUST buffer (50 mM Tris-HCl, pH 7.4, 8 M urea, 2% β-mercaptoethanol, and 0.5% SDS). After centrifugation at 5 × 10⁵ rpm for 20 min at 25°C the supernatant was used for protein analysis.

Lipid analysis. Lipids were extracted from mouse brain, sciatic, and pectineous nerves as described before (Bligh and Dyer, 1959). The extracted lipids were dissolved in 300 µl of chloroform/methanol (2:1), lipids from *mbp*-deficient mice in 30 µl chloroform/methanol (2:1). Lipids were separated by high-performance thin-layer chromatography (HPTLC) on silica gel-precoated plates (HPTLC 60; 10 × 10 cm plates; Merck, Darmstadt, Germany) in the solvent system chloroform/methanol/water (65:25:4). Lipid bands were visualized by charring with 50% H₂SO₄ at 120°C for 15 min and confirmed by standards. Gangliosides of purified myelin were separated in the solvent system chloroform/methanol/water/ammonia (60:35:6:2), stained with orcin-reagent at 95°C for 15 min. For semiquantitative lipid analysis, HPTLC plates were scanned, and lipid bands were integrated with a PhosphorImager. Mean values and their SEs were estimated from three independent experiments.

Light and electron microscopy. Anesthetized mice were perfused with 6% glutaraldehyde for light and electron microscopy. The optic nerve and cervical segments of the spinal cord were isolated, post-fixed in 1% phosphate-buffered OsO₄ in 0.1 M sucrose, and embedded in Epon 812. The semithin sections were stained with toluidine/pyronin. Ultrathin cross-sections of optic nerve and spinal cord were contrasted with uranylacetate and lead citrate and examined as previously described (Bussow, 1978).

Conduction velocity measurement. Nerve conduction was measured by proximal stimulation of the sciatic nerve at the knee and distal close to the hip, using a Medelec electromyograph, model MS92.

Behavioral testing (Wallace et al., 1980). The following tests were performed with eight mice of each genotype except for the *shiverer* mice with six and the PLP/MBP/MAG-deficient mice with seven mice. The behavior tests, which were performed during a 2 month period, were started with mice of the following age: controls: CD1 10 weeks; C57/Bl6 12 weeks;

Figure 1. A, Northern blot analysis; B, SDS-PAGE analysis [silver (top) and Coomassie staining (bottom)]; C, Western blot analysis of wt and six mutant mouse lines.

(*mag*^{-/-}) 20 weeks; (*plp*^{-/-}) 10 weeks; (*mbp*^{-/-}) 6–8 weeks; (*mag*^{-/-}*plp*^{-/-}) 10 weeks; (*plp*^{-/-}*mbp*^{-/-}) 10 weeks; (*mag*^{-/-}*plp*^{-/-}*mbp*^{-/-}) 8 weeks.

Morris water maze (Morris, 1984). Water mazes took place in a black circular polyethylene tub (69 cm diameter, 22 cm high) with an escape platform, a black cylinder (7.5 cm diameter), submerged 0.5 cm below the surface of the water, in a constant position. Water temperature was 21°C. The animal was set into the water at a randomly determined quadrant. A trial was finished when the mouse had reached the escape platform or at the set time of 90 sec. Animals were allowed to recover for 30 sec on the platform before another trial was started. Each mouse was tested in four subsequent trials over 5 d. Escape latency, number of squares crossed, and swimming speed were video-recorded. At the fifth day an additional 30 sec trial was performed with the escape platform being removed (probe trial). The number of crossings of the previous position of the platform was counted. The trials were registered and processed with the software MOR 005 (Tropon-Werke, Cologne, Germany) and statistically evaluated using

Table 1. Quantitative RT-PCR analysis

Genotype	PLP	MBP	MAG	MOG	OMgp	CGT	N-CAM140	N-CAM120
wt	1	1	1	1	1	1	1	1
<i>plp</i> ^{-/-}	–	1.2 ± 0.2	1.1 ± 0.4	0.8 ± 0.5	1.1 ± 0.3	1.3 ± 0.4	1.3 ± 0.2	1.1 ± 0.4
<i>mbp</i> ^{-/-}	1.4 ± 0.5	–	3.4 ± 0.7	1.3 ± 0.4	0.9 ± 0.2	1.1 ± 0.2	1.4 ± 0.4	1.1 ± 0.1
<i>mag</i> ^{-/-}	1.3 ± 0.4	1.5 ± 0.3	–	0.8 ± 0.2	0.9 ± 0.5	0.8 ± 0.5	1.6 ± 0.7	0.7 ± 0.3
<i>plp</i> ^{-/-} <i>mag</i> ^{-/-}	–	0.9 ± 0.4	–	1 ± 0.4	1.1 ± 0.2	0.8 ± 0.4	0.8 ± 0.5	1.2 ± 0.3
<i>plp</i> ^{-/-} <i>mbp</i> ^{-/-}	–	–	2.7 ± 0.4	1.4 ± 0.3	1.2 ± 0.4	1.2 ± 0.5	1.3 ± 0.5	1.2 ± 0.5
<i>plp</i> ^{-/-} <i>mbp</i> ^{-/-} <i>mag</i> ^{-/-}	–	–	–	0.9 ± 0.4	0.7 ± 0.3	1.1 ± 0.3	0.7 ± 0.3	1.5 ± 0.2

³²P-dCTP-labeled fragments were separated by PAGE, analyzed with a PhosphorImager (mean ± SEM), and normalized to the wt-expression of GAPDH and cyclophilin. The level of significance of MAG expression in *mbp*^{-/-} and in *plp*^{-/-}*mbp*^{-/-} mutants (boldface) was determined by Student's *t* test (*p* < 0.08 and *p* < 0.06, respectively).

the ANOVA software (Tropon-Werke) (Morris, 1984; Klapdor and van der Staay, 1996; Klapdor and van der Staay, 1998).

Open field. Animals were placed in the middle of a box (60 × 60 cm), the bottom of which was subdivided into 10 × 10 cm squares. Square crossings, rearing, grooming, and the location of the mouse (middle, wall, corner) were registered under video control over periods of 15 min. Mice were adapted in a quiet room under red light illumination. Data were processed with the software OBSERVE 004 (Tropon-Werke) and statistically evaluated with ANOVA software.

Rotarod test (Wallace et al., 1980). Animals were placed on a resting rod (diameter 3 cm) 20 cm above the ground for 1 min, and the rotation started at 0.4 rpm and doubled every 60 sec. The time interval during that the mice stayed on the rod was measured. The test was repeated three times a day over a period of 5 d. In the final task on day 5, the mouse was placed on the rod constantly rotating at 4 or 8 rpm, respectively. The drop off frequency within 90 sec was measured (Kuhn et al., 1995).

Descending a vertical pole (Wallace et al., 1980). A pole wrapped with cord (2 cm diameter, 80 cm length) and bordered at its upper end by a plate was held in a horizontal position. The mouse was placed with its forelimbs pointing to the platform. After turning the pole into a vertical position the time needed to turn around and required to descend and reach the floor was measured. Also the drop offs, sliding down and climbing up again, was scored. The procedure was repeated three times. A cutoff of 300 sec was chosen.

Horizontal bridge (Wallace et al., 1980). Mice were placed in the middle of a horizontal wooden bar (2 × 2 cm, 60 cm length) bridging two platforms (50 cm above the floor). The time to reach the platform, dropping off the bridge, and the time until the drop off occurs were measured.

Horizontal wire (Wallace et al., 1980). The mouse was placed with its forelimbs to the middle of a horizontal metal wire (2 mm diameter, 60 cm length) connecting two platforms (5 × 2 cm; 50 cm above the floor). The time needed to reach one of the platforms, dropping off the wire, and the time until the drop off occurred were measured. A cutoff time of 180 sec was selected.

RESULTS

We generated two mutant mouse lines, the *plp*^{-/-}*mag*^{-/-} double mutant by crossing the *plp*^{-/-} into the *mag*^{-/-} locus and a homozygous triple mutant *plp*^{-/-}*mbp*^{-/-}*mag*^{-/-} lacking PLP, MAG, and MBP by crossing the *mbp*^{+/-} genotype into the homozygous *plp*^{-/-}*mag*^{-/-} double mutant. For comparison we included the monogenic mutants, the *plp*^{-/-}*mbp*^{-/-} double mutant, and the wild type in the biochemical, morphological, and physiological analyses of six mutant genotypes: *plp*^{-/-}; *mbp*^{-/-} (*shi/shi*); *mag*^{-/-}; *plp*^{-/-}*mag*^{-/-}; *plp*^{-/-}*mbp*^{-/-}; and *plp*^{-/-}*mbp*^{-/-}*mag*^{-/-}. Their genotypes were verified by Southern blot analysis, which revealed the restriction fragments length polymorphism diagnostic for the respective targeted gene loci (data not shown). The missing mRNAs in Northern blot analysis (Fig. 1*A*), together with the absence of PLP, MBP, and MAG in SDS-PAGE and Western blot analysis proved that we had generated the two homozygous mouse lines (Fig. 1*B,C*). Sensitive protein detection methods (silver staining) and extensive Western blot analysis using polyclonal PLP and MAG antibodies and a monoclonal MBP antibody failed to detect any of the gene products in the respective mutants, including hypothetical immunoreactive PLP-related polypeptides (Fig. 1*B*) (Boison and Stoffel, 1994; Klugmann et al., 1997). Western blot analysis of sciatic nerve using the MBP-specific antibody revealed that MBP is selectively upregulated twofold in peripheral nerves of *plp*^{-/-}*mag*^{-/-} double mutant mice as previously reported for *mag*^{-/-} mice (Fig. 1*B*) (Li et al., 1994).

Figure 2. Analysis of total lipids of purified myelin and sciatic nerve. Six aliquots of brain extracts of mice carrying the *shiverer* genotype and one aliquot of the other genotypes were separated by HPTLC. *A*, Total lipids, solvent system: chloroform/methanol/water 65:25:4. *CN*, Normal fatty acid-substituted GalC; *CH*, α -hydroxy fatty acid-substituted GalC; *PE*, phosphatidylethanolamine; *SN*, normal fatty acid-substituted sGalC; *SH*, α -hydroxy fatty acid-substituted sGalC; *LPE*, lysophosphatidylethanolamine; *PC*, phosphatidylcholine; *PS*, phosphatidylserine; *PI*, phosphatidylinositol; *SPM*, sphingomyelin. *B*, Gangliosides of purified myelin, solvent system: chloroform/methanol/water/ammonia 60:35:6:2, were identified by comparison with HPLC purified standards. *C*, Semiquantitative lipid analysis, quantification with a PhosphorImager (mean ± SEM).

Table 2. Behavioral test: Morris water maze escape task

	wt C57/bl6			wt CD1			<i>plp</i> ^{−/−}			<i>mag</i> ^{−/−}			<i>plp</i> ^{−/−} <i>mag</i> ^{−/−}		
	day 1	day 3	day 5	day 1	day 3	day 5	day 1	day 3	day 5	day 1	day 3	day 5	day 1	day 3	day 5
Mean escape latency (sec)	66.1 ± 10.2	13.1 ± 2.1	8.4 ± 1.5	55.1 ± 10.0	13.8 ± 2.8	10.3 ± 1.4	53.1 ± 3.5	21.0 ± 5.9	11.1 ± 2.6	72.0 ± 7.9	30.1 ± 9.6	6.5 ± 0.6	90.0 ± 0	87.5 ± 2.5	86.3 ± 3.7
Mean swimming speed (crossed lines × sec ^{−1})	1.66 ± 0.09	1.68 ± 0.12	1.79 ± 0.15	2.12 ± 0.12	2.43 ± 0.1	1.79 ± 0.09	2.45 ± 0.1	1.66 ± 0.09	1.86 ± 0.14	1.87 ± 0.09	1.43 ± 0.15	1.66 ± 0.06	0.38 ± 0.04	0.48 ± 0.06	0.46 ± 0.06

Results in Tables 2–6 represent mean values (± SEM).

Phenotypes of myelin mutants

Plp^{−/−} and *mag*^{−/−} single null allelic mutants had no obvious phenotype. However, the double mutant *plp*^{−/−}*mag*^{−/−} developed a tremor of the hindlimbs around postnatal week 4. The initial tremor constantly progressed to dragging and jerky-like movement of the hindlimbs with increasing age. No convulsions were observed. Despite these neurological symptoms, the *plp*^{−/−}*mag*^{−/−} double mutant had a normal life span (see Table 7). The motor behavior of triple mutant mice (*plp*^{−/−}*mbp*^{−/−}*mag*^{−/−}) differed from that of the *plp*^{−/−}*mbp*^{−/−} double mutant by the combination of the *shiverer* trait with its high-frequency tremor and the PLP/MAG deficiency with its impairment described above. Seizures and convulsions did not occur until 3 months after birth but then with a high frequency. Their life span is 7–9 months, significantly longer than that of the MBP-deficient (*shiverer*) mouse of ~3 months (see Table 7) (Chernoff, 1981).

Transcriptional regulation of gene expression in the mouse mutants

We studied the complex gene expression pattern of oligodendrocytes in the single, double, and triple mutants first on the transcriptional level by Northern blot analysis (Fig. 1A). Searching for compensatory mechanisms, we expanded this study to genes possibly involved in this mechanism by quantitative RT-PCR of RNA of additional oligodendrocyte-specific proteins (MOG, OMGP, CGT) and two forms of the neural cell adhesion molecule N-CAM (140 and 120 kDa isoforms). Total RNA from p20 mouse brains and the respective oligonucleotide primers yielded PCR fragments of the expected size. Two housekeeping enzymes, GAPDH and cyclophilin, were included for comparison (Table 1).

The only significant difference in the expression pattern between the mutants and wt was a twofold to threefold overexpression of MAG in *shiverer*^{−/−} and *plp*^{−/−}*mbp*^{−/−} double mutant mice (Table 1). Further analysis of the transcripts of the MAG gene revealed that the overexpression of MAG is predominantly contributed by the smaller isoform, S-MAG, in the mutant CNS (data not shown). This overexpression was restricted to the mRNA but not to MAG protein synthesis as shown by Western blot analysis (Fig. 1C).

Deficiencies of CNS-myelin proteins are associated with alterations in myelin membrane lipid synthesis

To explore the differences in myelination the total amount of myelin, isolated from brain homogenates of 6-week-old mutant and wt mice by discontinuous sucrose gradient centrifugation, was determined (Norton and Poduslo, 1973). The yield of total myelin from brains of wt, *plp*^{−/−}, *mag*^{−/−}, and *plp*^{−/−}*mag*^{−/−} double mutant mice was approximately identical (wt: 18.1 ± 2.4 mg; *plp*^{−/−}: 18.6 ± 5.3 mg; *mag*^{−/−}: 16.7 ± 0.9 mg; *plp*^{−/−}*mag*^{−/−}: 18.3 ± 2.9 mg). However, the amount of myelin obtained from brains of the remaining mutants was strikingly low (*mbp*^{−/−}: 0.7 ± 0.3 mg; *plp*^{−/−}*mbp*^{−/−}: 0.7 ± 0.5 mg; *plp*^{−/−}*mbp*^{−/−}*mag*^{−/−}: 0.6 ± 0.4 mg). Although the amount of myelin from brains of mice carrying the *shiverer* genotype were similar, oligodendrocytes of the double mutant *plp*^{−/−}*mbp*^{−/−} and the triple mutant *plp*^{−/−}*mbp*^{−/−}*mag*^{−/−} mice wrap a significant number of larger diam-

eter axons with a “pseudomyelin” sheath (see Fig. 4A, electron microscopy).

In view of the ultrastructural finding of “pseudomyelin” membranes around larger diameter axons in the *plp*^{−/−}*mbp*^{−/−} and the triple mutant CNS (Fig. 3) and the minute amount of proteins isolated from their myelin, we studied the lipid composition of purified myelin fractions of the different genotypes, not of total brain. Previous reports focused on the lipids of whole brain of *shiverer* mouse (Norton and Poduslo, 1973; Bird et al., 1978; Cammer et al., 1984; Iwamori et al., 1985). We analyzed total lipids from purified CNS myelin, pectineus, and sciatic nerves by HPTLC. Whereas the lipid composition of CNS myelin of *plp*^{−/−}, *mag*^{−/−}, and *plp*^{−/−}*mag*^{−/−} mutants was unaffected, mutants carrying the MBP deletion showed significant differences in the amounts of total phospholipids and myelin-specific lipids. In CNS myelin of all mutants carrying the MBP deletion, the ratio of the oligodendrocyte-specific lipids, galactocerebrosides, and sulfatides to phospholipids was reduced fourfold. This ratio remained unchanged in the other mutants like in control mice (Fig. 2A,C). Cerebrosides and sulfatides were both considerably reduced in the CNS of MBP-deficient mice, although the gene expression of the key enzyme of galactocerebroside and sulfatide biosynthesis, the UDP-galactose:ceramide galactosyltransferase (CGT) (EC 2.4.1.45) (Schulte and Stoffel, 1993), was not altered (Table 1).

The lipid analysis of CNS myelin of the mutants containing the *shiverer* trait furthermore revealed that the sphingomyelin fraction consisted of only one species, stearyl-sphingomyelin. This structure was ascertained by MALDI-TOF MS (matrix-assisted laser desorption/ionization time of flight mass spectroscopy) (data not shown).

Gangliosides of CNS myelin of mice with the *mbp*^{−/−} mutation were distinctly decreased. Complex higher gangliosides (GM₁, GD_{1a}, GD_{1b}, GT_{1b}, and GQ_{1b}) were almost absent, and only GM₃ were present as main ganglioside species (Fig. 2B). Membrane lipids in the myelin sheaths of peripheral nerves were unaffected in the different genotypes.

Behavioral characteristics of the mutants

To further characterize each mutant, we studied the cognitive and motor-coordinative capacities in behavioral tests. The two double and the triple mutants differ clearly from each other by the degree of impaired neuromotor coordination. Mouse mutants carrying the *shiverer* trait (*mbp*^{−/−}; *plp*^{−/−}*mbp*^{−/−}; *plp*^{−/−}*mbp*^{−/−}*mag*^{−/−}) were excluded from the Morris water maze escape task (Morris, 1984) because they were unable to swim over the period of time required for the task. *Plp*^{−/−}*mag*^{−/−} double mutant mice were compared with their respective monogenic mutants and two wild-type strains (C57/Bl6, inbred; CD1, outbred). Whereas *plp*^{−/−} and *mag*^{−/−} mice showed a normal swimming speed and learning behavior and were indistinguishable from wt mice, none of the *plp*^{−/−}*mag*^{−/−} mutants was able to locate the platform and acquire spatial discrimination in the water maze. Because of their jerky-like movement of the hindlimbs they were not able to swim, but drifted on the water surface. That explains their distinctly reduced swimming speed compared to wt mice (Table 2). None of

Table 3. Behavioral test: open field

Genotype	Locomotion	Grooming	Rearing
wt C57/Bl6	515.0 ± 47.6	10.0 ± 2.0	92.5 ± 12.8
wt CD1	511.3 ± 45.4	9.9 ± 2.4	90.3 ± 10.4
<i>plp</i> ^{-/-}	578.5 ± 36.9	8.0 ± 2.2	57.2 ± 10.2
<i>mbp</i> ^{-/-}	43.3 ± 5.0	4.0 ± 0.8	9.3 ± 1.7
<i>mag</i> ^{-/-}	253.1 ± 79.7	2.5 ± 0.9	20.3 ± 10.0
<i>plp</i> ^{-/-} <i>mag</i> ^{-/-}	318.4 ± 78.6	7.6 ± 2.3	59.3 ± 15.5
<i>plp</i> ^{-/-} <i>mbp</i> ^{-/-}	323.6 ± 34.9	4.9 ± 1.1	43.9 ± 7.3
<i>plp</i> ^{-/-} <i>mbp</i> ^{-/-} <i>mag</i> ^{-/-}	41.6 ± 12.7	3.6 ± 1.4	10.4 ± 4.3

Table 4. Behavioral test: horizontal bridge

Genotype	% of mice reaching a platform	nr of mice dropping off	Latency to reach a platform (sec)
wt C57/Bl6	100	0	9.9 ± 9.73
wt CD1	100	0	9.4 ± 1.9
<i>plp</i> ^{-/-}	95.8	0	8.9 ± 1.2
<i>mbp</i> ^{-/-}	95.8	1	20.3 ± 2.3
<i>mag</i> ^{-/-}	45.8	2	65.3 ± 19.7
<i>plp</i> ^{-/-} <i>mag</i> ^{-/-}	62.5	7	32.0 ± 11.4
<i>plp</i> ^{-/-} <i>mbp</i> ^{-/-}	79.2	0	74.6 ± 21.1
<i>plp</i> ^{-/-} <i>mbp</i> ^{-/-} <i>mag</i> ^{-/-}	16.7	10	92.5 ± 18.3

these mice ever entered the previous training quadrant during the probe trial.

Locomotor activity of the mutants was measured in the open field test (Table 3). As expected, *shiverer*^{-/-} and triple mutant mice showed a dramatic decrease in activity compared to *plp*^{-/-} and *wt* mice. The locomotor activity of *mag*^{-/-} and the two double mutants, *plp*^{-/-}*mbp*^{-/-} and *plp*^{-/-}*mag*^{-/-}, ranged in between these two groups.

To determine the motor and posture pattern, the following tasks were applied to challenge mutant mice: clinging to a horizontal wire (Table 6), moving on a horizontal bridge (Table 4), and descending from a vertical pole (data not shown). The most striking result was that *mag*^{-/-} mice, *plp*^{-/-}*mag*^{-/-} double and triple mutants did not reach the platform when starting from the middle of the horizontal wire or bridge. In these two tasks a clear gradation between mutants became apparent: whereas the *mag*^{-/-} mutant on the average dropped off twice during the defined “cut-off” time, *plp*^{-/-}*mag*^{-/-} double mutant mice fell from the bridge seven times, and only half of tested triple mutant mice were able to remain on the bridge. In the vertical pole task the double and triple mutant mice slid down the pole instead of climbing down in a coordinate fashion like wild-type mice (data not shown).

Table 5. Behavioral test: rotarod

Genotype	Increasing speed: time remaining on rod			Constant speed: cumulative falls in 90 sec	
	day 1	day 3	day 5	4 rpm	8 rpm
wt C57/Bl6	169.4 ± 56.7	376.6 ± 34.8	425.4 ± 16.4	0	0
wt CD1	289.3 ± 41.0	305.8 ± 49.6	382 ± 31.7	0	0.4 ± 0.2
<i>plp</i> ^{-/-}	186.1 ± 59.4	316.6 ± 30.9	351 ± 19.2	0	1 ± 0.7
<i>mbp</i> ^{-/-}	17.8 ± 11.7	81.3 ± 26.0	185 ± 46.9	7.7 ± 1.4	10.2 ± 0.9
<i>mag</i> ^{-/-}	178.2 ± 33.1	270.8 ± 35.4	277.7 ± 23.8	1 ± 0.6	4.4 ± 1.9
<i>plp</i> ^{-/-} <i>mag</i> ^{-/-}	22.5 ± 9.8	92 ± 29.9	146.8 ± 39.4	3.9 ± 1.1	12.9 ± 1.4
<i>plp</i> ^{-/-} <i>mbp</i> ^{-/-}	11.3 ± 4.6	88.4 ± 30.3	173.8 ± 36.5	7.8 ± 1.7	15.3 ± 2.6
<i>plp</i> ^{-/-} <i>mbp</i> ^{-/-} <i>mag</i> ^{-/-}	40 ± 13.5	43.2 ± 15.8	61.7 ± 24.9	12.1 ± 2.3	18.4 ± 0.8

Finally, the learning behavior and skill of the mutants was scored on a rotating rod on 5 subsequent days (Kuhn et al., 1995). All mice, including the heavily affected animals with the *shiverer* trait, learned to stay on the rod rotating with increasing speed and improved their skill. The time mice remained on the rod depended significantly on the degree of impairment of their motor coordination. This is illustrated by the record of cumulative falls from the rotating rod (Table 5). The performance of *plp*^{-/-} mice was comparable to *wt* mice in this task, whereas *mag*^{-/-} mice exhibited small but significant differences compared to *wt* mice. Triple mutant mice are the most affected genotype in this task. The performance of *shiverer* and double mutant mice *plp*^{-/-}*mbp*^{-/-} and *plp*^{-/-}*mag*^{-/-} can be classified as intermediate between these genotypes, *mag*^{-/-} and the *plp*^{-/-}*mbp*^{-/-}*mag*^{-/-}.

Conduction velocity of peripheral nerves of the mutant mice is unchanged

The impact of the protein deficiency on the conduction velocity of peripheral nerves was measured by electromyography. Proximal stimulation of the sciatic nerve at the knee and distal close to the hip was applied. No significant differences in conduction velocities of peripheral nerves of *wt* and mutant mice, 4–5 months of age, were observed. They range between 26.5 and 32 m/sec.

Morphology of mutant CNS myelin structures

The genetic, biochemical, physiological, and behavioral traits of each phenotype were complemented by extensive morphological studies of CNS myelin of the different mutants by electron microscopy, which also depicts schematically these structural changes of the myelin membrane (Fig. 3). The CNS myelin sheath of *mag*^{-/-} mice showed regular MDLs and IDLs with some subtle morphological abnormalities, e.g., doubled myelin sheaths, shortened periaxonal collar, and the appearance of longitudinal channels within compacted myelin (Li et al., 1994; Montag et al., 1994). In agreement with a previous report, we observed no reduced periaxonal spacing (Montag et al., 1994).

Another compaction type of the myelin membrane is dictated by the lack of the integral membrane protein PLP. Myelin membrane stacks of *plp*^{-/-} and *plp*^{-/-}*mag*^{-/-} mice are decompacted in the absence of PLP and DM₂₀ at the IDL (Boison and Stoffel, 1994; Montag et al., 1994; Boison et al., 1995). In addition, condensed areas with main and intraperiod dense lines that are indistinguishable in electron density were observed (Fig. 3). In *plp*^{-/-}*mag*^{-/-} double mutant mice the morphological abnormalities of the *mag*^{-/-} and *plp*^{-/-} single mutants are accumulated. Most strikingly we observed in the *plp*^{-/-}*mag*^{-/-} double mutant an early onset of axon degeneration around postnatal day 40 (P40), which starts in the *plp*^{-/-} mouse only at later stages of life (Griffiths et al., 1998). An increased number of degenerated axons of the optic nerve and the ventral funiculus of the spinal cord is seen in light microscopy of optic nerves of 40-d-old mice, and a massive axonal degeneration is apparent in P120 mice (Fig. 4).

The three genotypes that we included in this study, the *shiverer*^{-/-}, the *plp*^{-/-}*mbp*^{-/-} double, and *plp*^{-/-}*mbp*^{-/-}*mag*^{-/-} triple

Table 6. Behavioral test: horizontal wire

Genotype	% of mice clinging the wire	nr of mice reaching a platform
wt C57/Bl6	87.5	3
wt CD1	66.7	4
<i>plp</i> ^{−/−}	70.8	0
<i>mbp</i> ^{−/−}	91.7	5
<i>mag</i> ^{−/−}	76.2	0
<i>plp</i> ^{−/−} <i>mag</i> ^{−/−}	20.8	0
<i>plp</i> ^{−/−} <i>mbp</i> ^{−/−}	66.7	3
<i>plp</i> ^{−/−} <i>mbp</i> ^{−/−} <i>mag</i> ^{−/−}	66.7	0

mutant mice show another type of dysmorphic myelin structure. Hypomyelination of the CNS is the dominant leading symptom in the *shiverer* mouse (Chernoff, 1981) and is combined with a remarkable increase of oligodendrocytes in the spinal cord (Nagara et al., 1983) accompanied by the appearance of numerous Schmidt-Lantermann incisures in the PNS (Gould et al., 1995). The *shiverer* genotype confers this phenotype to the two other mutants. However, unlike oligodendrocytes of the *shiverer* mutant that are nearly unable to enwrap CNS axons, oligodendrocytes of the *plp*^{−/−}*mbp*^{−/−} double and the *plp*^{−/−}*mbp*^{−/−}*mag*^{−/−} triple mutants spirally surround large-diameter axons as loosely compacted myelin consisting of only few lamellae (Fig. 4A). They adhere tightly at their extracytosolic surfaces forming a distinct electron-dense line of a myelin-like structure with a compacted IDL. The complete surrounding of an axon by alternately compacted myelin in the *plp*^{−/−}*mbp*^{−/−} double and triple mutants is obviously sufficient to facilitate a normal life span in the *plp*^{−/−}*mbp*^{−/−} and an extended lifespan in the *plp*^{−/−}*mbp*^{−/−}*mag*^{−/−} mutant under laboratory conditions. Addition of the *mag*^{−/−} genotype to the double mutant causes the early onset of axonal degenerations in the triple mutant mouse, which shortens its life span (Table 7).

DISCUSSION

Gene targeting by homologous recombination of mutant mouse genes that replace the wt genes has proven to be a useful tool in the functional analysis of structural proteins of CNS and PNS myelin. Here we describe the generation and characterization of two new mutant mouse lines, a *plp*^{−/−}*mag*^{−/−} double mutant and a homozygous triple mutant lacking PLP, MBP, and MAG (*plp*^{−/−}*mbp*^{−/−}*mag*^{−/−}). Surprisingly, *plp*^{−/−}*mag*^{−/−} double mu-

tants, in contrast to their respective single mutants, showed severe neurological symptoms with increasing age.

The triple mutant, like the other homozygous *plp*^{−/−}*mbp*^{−/−} double mutant, bears a strong resemblance to *shiverer* mice. However, the tremor contributed by the *shiverer* trait is mild, the first seizures of convulsions occur only infrequently and not before the age of 3 months, whereas the monogenic *shiverer* mutant mice developed this phenotype 1 month after birth (Bird et al., 1978; Chernoff, 1981). The shortened life expectancy of triple mutant mice (7–9 months) compared to the *plp*^{−/−}*mbp*^{−/−} double mutant (life span, 2 years) obviously derives from the additional deprivation of MAG in the myelin of these mutants. In view of the twofold to threefold overexpression of S-MAG in *shiverer* and *plp*^{−/−}*mbp*^{−/−} double mutants, it is tempting to speculate on a similar role of this MAG isoform for CNS myelin maintenance because it has been shown for PNS myelin (Fujita et al., 1998). N-CAM on the other hand, which like MAG is a member of the Ig superfamily, appears not to be involved in this process, although it has been suggested to substitute for the function of MAG in the PNS of MAG-deficient mice (Montag et al., 1994). The double mutant deficient in MAG and N-CAM (Carenini et al., 1997) supported this with regard to axonal and myelin maintenance, but not with regard to myelin formation in the PNS. However, in our study neither the upregulation of the expression of N-CAM120 nor of N-CAM140 was observed in the mutant CNS.

The results of the lipid analysis and the morphological data suggest that the loss of MBP expression causes an impairment of oligodendrocyte development. GM₃ besides GD₃ and GD_{1a} and GT_{1b} as minor components, appear to play an important role during the dichotomy of the differentiation of oligodendrocyte precursors. Addition of exogenous GM₃ to oligodendrocyte precursors in culture increased the number and the thickness of the oligodendrocyte processes (Yim et al., 1994, 1995). Also the degree of phosphorylation of MBP and MAG is dependent on GM₃. GM₃ increases the phosphorylation of MAG and simultaneously reduces MBP phosphorylation (Yim et al., 1994). This observation is of special interest in view of our finding that the gangliosides in CNS myelin of mice with the *mbp*^{−/−} mutation in general were distinctly decreased, and only GM₃ was synthesized by oligodendrocytes of these mutants. Ganglioside GM₁ on the other hand is characteristic for isolated myelin of wild type (Suzuki et al., 1967) and is also present in the myelin fraction of all mutants without the *shiverer* deletion. Mouse myelin isolated from mutants carrying the *shiverer* allele is deficient in GM₁ (Raff et al., 1978; Iwamori et al., 1985).

Gangliosides are markers in oligodendrocyte maturation (Raff et

Figure 3. Electron micrographs of cross-sections through the optic nerves of *mag*^{−/−}, double mutant (*mag*^{−/−}*plp*^{−/−}), and triple mutant (*mag*^{−/−}*mbp*^{−/−}*plp*^{−/−}) mice. The comparison of the three sections demonstrates the different types of compaction. The axon of the MAG-deficient mouse is surrounded by two distinct myelin sheaths with a normal periodicity. Myelin formation in *plp*^{−/−}*mag*^{−/−} double mutant is characterized by a decompaction of the IDL as well as by condensed areas with indistinguishable main and intraperiod dense lines. Triple mutants show areas of pseudocompacted myelin around axons with large diameter. No MDL but a pseudo-IDL formed by the compaction of outer surfaces is visible. Scale bar, 200 nm.

Figure 4. *A*, Semithin cross-sections of the spinal cord (anterior median fissure) of the indicated genotypes. Whereas *mag*^{-/-} mice are indistinguishable from *wt* controls, *plp*^{-/-}*mag*^{-/-} double mutants have comparable amounts of slightly disordered myelin; in *shiverer* mice almost no myelin is detectable, whereas in the triple mutant axons with a larger diameter show myelin sheaths (asterisks). Note the increased number of oligodendrocytes in the MBP-deficient animals (*mbp*^{-/-} and *plp*^{-/-}*mbp*^{-/-}*mag*^{-/-}) and axonal degeneration in *plp*^{-/-}*mag*^{-/-} double mutant mice (black arrows). Scale bar, 10 μ m. *B*, *Top*, Semithin cross-sections of the optic nerve of 40- and 120-d-old *plp*^{-/-}*mag*^{-/-} double mutant mice. Note the enlarged number of axonal degeneration with increasing age. Scale bar, 10 μ m. *Bottom*, Electron micrographs of cross-sections of optic nerve of *plp*^{-/-}*mag*^{-/-} double mutant and *plp*^{-/-}*mbp*^{-/-}*mag*^{-/-} triple mutant mice (black arrows, axonal degeneration). Scale bar, 2 μ m.

al., 1978; Yim et al., 1994). The ganglioside pattern of myelin of the mutants with the *shiverer* trait might express a state in the maturation of oligodendrocyte precursor cells. The *plp*^{-/-}*mbp*^{-/-} and *plp*^{-/-}*mbp*^{-/-}*mag*^{-/-} triple mutants might be useful models for studies of oligodendrocyte development and maturation.

The array and the results of behavioral tests performed in this study contribute significantly to the characterization of the different phenotypes of the mutants. This becomes most obvious by the analysis of the rather inconspicuous behavior of the mutants containing the *mag*^{-/-} allele. In the behavioral tests selected in this study, these mice showed a surprisingly poor performance. Although behavioral tests that require motor and reflex responses are known to significantly decline with age (Wallace et al., 1980), and

therefore age differences might explain the differences in locomotor activity between our findings and a previous report (Li et al., 1994), this effect obviously is caused by the MAG depletion. This suggestion is supported by other behavioral tests because mutants with the *mag*^{-/-} genotype are affected when they crossed the horizontal bridge during the defined "cutoff" time. *Plp*^{-/-}*mag*^{-/-} mice were hardly able to cling to the horizontal wire because of their motor deficits.

Axonal degeneration is missing in the CNS of *mag*^{-/-} single mutants. The early onset and rapidly progressing axonal degeneration in CNS of *plp*^{-/-}*mag*^{-/-} double mutant is demonstrated in electron microscopy of p40 and p120 optic nerves. It parallels the first signs and the progression of the neurological symptoms of this

Table 7. Genotype and phenotype of myelin mutant mice

Genotype	Phenotype					Lifespan (months)
	RNA	Protein	Lipid (CNS)	Morphology: myelin	Morphology: axons	
<i>plp</i> ^{−/−} ^a	NA	NA	NA	Dissociation of outer surfaces, IDL (CNS)	Late onset of degeneration (CNS)	>24
<i>mag</i> ^{−/−} ^b	NA	MBP ↑ (PNS)	NA	Reduced cytoplasmic collar, multiple myelin sheaths (CNS)	Late onset of degeneration (PNS)	>24
<i>plp</i> ^{−/−} <i>mag</i> ^{−/−}	NA	MBP ↑ (PNS)	NA	Phenotype of 1. and 2.	Early onset of degeneration (CNS)	>24
<i>mbp</i> ^{−/−} ^c	MAG ↑ (CNS)	NA	Gangliosides ↓, GM ₃ ↑, GalC+sGalC ↓	Hypomyelination (CNS), Schmidt-Lantermann-incisures (PNS) ↑, number of oligodendrocytes ↑	NA	3–4
<i>plp</i> ^{−/−} <i>mbp</i> ^{−/−} ^d	MAG ↑ (CNS)	NA	Gangliosides ↓, GM ₃ ↑, GalC+sGalC ↓	Hypomyelination (CNS), pseudo-myelin (CNS), Schmidt-Lantermann-incisures (PNS) ↑, number of oligodendrocytes ↑	Late onset of degeneration (CNS)	>24
<i>plp</i> ^{−/−} <i>mbp</i> ^{−/−} <i>mag</i> ^{−/−}	NA	NA	Gangliosides ↓, GM ₃ ↑, GalC+sGalC ↓	Hypomyelination (CNS), pseudo-myelin (CNS), Schmidt-Lantermann-incisures (PNS) ↑, number of oligodendrocytes ↑	Early onset of degeneration (CNS)	7–9

Summary of the phenotypic characteristics of mice of the different genotypes described in this paper. Some data have been published before: ^aBoison and Stoffel, 1994; Griffiths et al., 1998; ^bLi et al., 1994; Montag et al., 1994; ^cBird et al., 1978; Chernoff, 1981; Gould et al., 1995; Nagara et al., 1983; ^dStoffel et al., 1997.

NA, Not altered compared to wildtype.

double mutant. This again suggests an important role of MAG in maintaining the integrity of CNS axons similar to the function of MAG in PNS myelin (Fruttiger et al., 1995; Yin et al., 1998). The detrimental effect of MAG deficiency on axons occurs only in the *plp*^{−/−}*mag*^{−/−} double mutant.

A recent report discusses the function of a neuron expressed sr (soma-restricted) PLP in the maintenance of axons (Bongarzone et al., 1999). SrPLP is an alternative splice product of PLP. Exon 1 of srPLP codes for a signal sequence of 11 amino acid residues that resemble strong homology with ER retention signal sequences. The lack of the minor PLP splice product (2%) of PLP/DM₂₀ is thought to cause the late onset of axonal degeneration in the CNS of the *plp*^{−/−} mouse. However, we regard the contribution of srPLP to axonal degeneration as minute compared with the MAG deficiency in the *plp*^{−/−}*mag*^{−/−} double mutant. The additional load of the *mag*^{−/−} genotype induces dramatically axonal degeneration.

The results reported here strongly suggest that an imbalance of PLP and MBP in favor of PLP in the CNS may downregulate myelin synthesis in oligodendrocytes, because the *shiverer* phenotype is ameliorated in *plp*^{−/−}*mbp*^{−/−} and *plp*^{−/−}*mbp*^{−/−}*mag*^{−/−} triple mutant mice. This idea is supported by the phenotype of transgenic mice showing different PLP gene dosage effects, e.g., PLP or DM₂₀ overexpression (Kagawa et al., 1994; Readhead et al., 1994; Johnson et al., 1995).

Plp^{−/−}*mag*^{−/−} double mutant animals and triple mutant mice lacking two and three important constituents of CNS myelin, respectively, might be valuable models for further studies on how oligodendrocytes function during the assembly of this highly ordered membrane structure. The generation and analysis of these two new mutants strongly indicates the cooperative functions of MAG and PLP in maintaining axon integrity in the CNS.

As a perspective, the double and triple mutants are promising models for stem cell biology and therapy. The recent development in the field of totipotent embryonic and stem cell-derived glial precursor cells and their prospective use as source of cell type-specific somatic precursors for neural implantation has promising perspectives also for efficient myelination and remyelination of axons in CNS. The md rat characterized by a point mutation in exon III of the PLP gene (Thr75 to Pro) (Boison and Stoffel, 1989)

and the shaking pup (His36 to Pro) are preferred dysmyelination models in transplantation studies (Archer et al., 1997; Brustle et al., 1999). The two models suffer from the short viability of these mutants. The mouse models described and characterized here genetically, biochemically, morphologically, and behaviorally have a considerably longer life span, which recommends them as valuable tools for the long-term evaluation of stem cell therapeutic strategies.

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