Cortical Network Dysfunction Caused by a Subtle Defect of Myelination

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Subtle white matter abnormalities have emerged as a hallmark of brain alterations in magnetic resonance imaging or upon autopsy of mentally ill subjects. However, it is unknown whether such reduction of white matter and myelin contributes to any disease-relevant phenotype or simply constitutes an epiphenomenon, possibly even treatment-related. Here, we have reanalyzed Mbp heterozygous mice, the unaffected parental strain of shiverer, a classical neurological mutant. Between 2 and 20 months of age, Mbp1/- versus Mbp1/1 littersmates were deeply phenotyped by combining extensive behavioral/cognitive testing with MRI, 1H-MR spectroscopy, electron microscopy, and molecular techniques. Surprisingly, Mbp-dependent myelination was significantly reduced in the prefrontal cortex. We also noticed a mild but progressive hypomyelination of the prefrontal corpus callosum and low-grade inflammation. While most behavioral functions were preserved, Mbp1/- mice exhibited defects of sensorimotor gating, as evidenced by reduced prepulse-inhibition, and a late-onset catatonia phenotype. Thus, subtle but primary abnormalities of CNS myelin can be the cause of a persistent cortical network dysfunction including catatonia, features typical of neuropsychiatric conditions.

Key words: myelin basic protein (Mbp) mutant, Cnp mutant, prepulse inhibition, catatonia, MRI, electron microscopy

Introduction

The expansion of white matter tracts is among the most striking features of mammalian brain evolution and long-range connectivity along myelinated tracts is thought to underlie many aspects of higher brain function (Fields et al., 2014; Nave, 2010; Zatorre et al., 2012). The role of myelin for motor-sensory functions is well recognized, and illustrated by the grave neurological symptoms in patients with severe myelin abnormalities, such as leukodystrophies. In contrast, a potential impact of mildly disturbed myelination in psychiatric diseases is poorly understood. It is unknown, whether subtle alterations in myelin thickness or the number of myelinated axons in cortex and subcortical white matter tracts compromise higher brain functions. Theoretically, such subtle alterations could perturb the millisecond precision crucial for signal propagation in higher brain networks (Uhlhaas and Singer 2012), thereby explaining the often postulated “disconnectivity” syndromes (Catani and Ffytche, 2005; Schmahmann et al., 2008). Psychiatric symptoms have been documented in white matter defects, such as metachromatic leukodystrophy (Hyde et al., 1992) and multiple sclerosis (Feinstein et al., 2014). However, the role of oligodendrocyte and myelin defects in higher brain functions and diseases such as schizophrenia is largely unclear (Takahashi et al., 2011).

Myelin is made by oligodendrocytes as a multilayered membrane sheath and enables rapid axonal impulse propagation (Nave, 2010). Myelin membranes have a unique composition of lipids and proteins, including abundant structural...
components of CNS myelin, such as myelin basic protein (MBP), proteolipid protein (PLP), and cyclic nucleotide phosphodiesterase (CNP). MBP is important for myelin membrane compaction and therefore also essential in initiating and driving the axonal wrapping process (Jahn et al., 2009).

While myelination takes place largely during early postnatal life, there is continuous myelin growth in specific cortical areas, such as prefrontal cortex (PFC), lasting well into adult life (Miller et al., 2012). Recently, it has been shown that these sparsely myelinated areas regulate myelin growth differently than robustly myelinated subcortical regions, with neuronal activity and social interactions becoming a driving force of PFC development including its underlying white matter (Benes, 1989; Gibson et al., 2014; Liu et al., 2012; Makinodan et al., 2012; Tomassy et al., 2014; Uda et al., 2015).

Numerous reports have highlighted white matter abnormalities in psychiatric patients. This comprises findings by magnetic resonance imaging (MRI) of reduced white matter size and accelerated white matter aging (Balevich et al., 2015; Kochunov et al., 2013). Moreover, diffusion tensor imaging (DTI) and magnetic transfer resonance (MTR) have revealed abnormal connectivity and loss of white matter integrity in the corpus callosum of schizophrenia and bipolar patients (Balevich et al., 2015; Kubicki et al., 2005; Kubicki et al., 2002; McIntosh et al., 2008). Increased numbers of microglia in the proximity of myelinated fibers, suggesting neuroinflammation (Uranova et al., 2011), and reduced density of oligodendrocytes in PFC were reported in schizophrenia, bipolar disorder and major depression (Uranova et al., 2004).

Human autopsy material repeatedly revealed a decrease of several structural myelin proteins, such as MBP, and respective mRNAs (Chambers and Perrone-Bizzozero, 2004; Dracheva et al., 2006; Flynn et al., 2003; Hakak et al., 2001; Honer et al., 1999; Matthews et al., 2012; Parlapani et al., 2009; Tkachev et al., 2003). These correlative observations were made decades after disease onset. We also note that long-term antipsychotic medication is suspected to cause brain atrophy (Bartzokis et al., 2007; Ho et al., 2011) which includes myelinated fiber loss, and that pharmacological models of schizophrenia can trigger secondary myelin deficits (Zhang et al., 2012). Thus, distinguishing cause and consequence of myelin abnormalities in psychiatric disease has remained difficult. Whether mild changes of CNS myelination (in absence of overt neurological disease) affect cortical network functions can theoretically be addressed in animal models. Importantly, hypomyelination in these models must be subtle and not interfere with basic behavior if a causal impact on altered higher brain function shall be established.

Here, we studied mice heterozygous for the oligodendroglial Mbp gene, i.e., the healthy-appearing parents of dysmyelinated homozygous Shiverer mice (Chernoff, 1981; Readhead et al., 1987; Roach et al., 1985; Shine et al., 1992). We report a very subtle hypomyelination phenotype in Mbp+/- mice, which is more pronounced in the PFC than in caudal white matter tracts and progresses over time. Importantly, both basic behavior and cognition were normal, but we demonstrate a persistent defect of prepulse inhibition of the startle response (PPI), a surrogate marker for gating defects in human patients with schizophrenia and their first-degree relatives (Bráff et al., 1978; Kumari et al., 2005). Moreover, we find late-onset catatonia and signs of low-grade inflammation. Our data in mice show that minor myelin abnormalities can be causal of cortical network dysfunctions, in the absence of other behavioral defects. These abnormalities alone can explain some of the pivotal phenotypes common to mental diseases.

Materials and Methods

Important Note
All experiments were performed by investigators unaware of group assignment (“fully blinded”).

Mice
Experiments were approved by the local animal care and use committee. Mice were group-housed in standard plastic cages with access to food/water ad libitum, 12 h light-dark-cycle (light-on at 7:00 am), 20–22°C. Male Mbp+/- versus Mbp+/- mice [C57BL/6N background; genotyped as previously described (Klugmann et al., 1997)], and for comparator experiments, Cup+/- versus Cup+/- mice (Hagemeyer et al., 2012), were used.

Behavioral Testing
Male Mbp+/- versus Mbp+/- mice (littermates) underwent behavioral testing at 2 age-periods: 3–6 and 17–18 months. Experiments were conducted during light phase (9:00 – 17:00) in following order: Elevated plus maze, open field, hole board, prepulse inhibition of the startle response (PPI), rotarod, acoustic startle assessment, grip-strength, hot-plate, marble burying, olfaction (buried-food-test), visual-cliff, LABORAS home-cage observation, social interaction, Y-maze, sucrose preference, Morris water maze, and novel object recognition. Inter-test interval was ≥24 h. All tests were conducted as published in detail (Bodda et al., 2013; Dere et al., 2014; El-Kordi et al., 2013; Moy et al., 2004; Radyushkin et al., 2010). Tests yielding significance are described below.

Prepulse Inhibition of the Acoustic Startle Reflex (PPI), Testing Sensorimotor Gating
Mice are put in metal cages (82x40x40mm, restricting major movements), placed in sound-attenuating cabinets. Cages are equipped with a movable platform-floor attached to a sensor, recording vertical movements (TSE-Systems). Startle reflexes are evoked by acoustic stimuli delivered by a loudspeaker above the cage, connected to an
acoustic generator. The startle reaction to an acoustic stimulus inducing a movement of the force-sensitive platform is recorded over 260 ms beginning with pulse onset. An experimental session consists of 2-min habituation to a 65dB continuous background white noise, followed by 1 min baseline recording. Then, 6 pulse-alone trials (120 dB, 40 ms) are applied to decrease within-session habituation (not included in 120 dB/40 ms PPI analysis). For PPI, the startle pulse is applied either alone or preceded by a prepulse stimulus of 20-ms duration and 70-, 75-, or 80 dB intensity. A delay of 100 ms with background noise is interposed between prepulse and pulse. Trials are presented in pseudorandom order with variable intervals ranging from 8 to 22 s. The amplitude of the startle response (arbitrary units) is defined as the difference between the maximum force detected during the recording window and that measured immediately before stimulus onset. For each animal, amplitudes are averaged separately for trials with stimulus alone or stimulus preceded by prepulse. PPI is calculated as a percentage of startle response using following formula: \[ \text{PPI} = 100 - \left[ \frac{\text{startle amplitude after prepulse}}{\text{startle amplitude after pulse only}} \right] \times 100 \].

**Startle Response to 120dB**

Response to startle stimulus of 120 dB alone, without any prepulse, and response without stimulus are recorded. The plain startle reaction to an acoustic pulse is a short-latency reflex mediated by an olivesynaptic neural circuit that includes lower brainstem, spinal and cranial motor neurons, and cerebellum (Kim et al., 2010; Takeuchi et al., 2001).

**Bar Test for Catatonia**

This test was performed as described (Hagemeyer et al., 2012; Kuschinsky and Hornykiewicz, 1972) and is illustrated in Supporting Information videos.

**MRI And 1H-MR Spectroscopy (1H-MRS)**

Mice were anesthetized with 5% isoflurane, intubated and kept under 1.75% isoflurane and 5% oxygen by active ventilation with constant respiratory frequency of 85 breaths/min (Animal-Respirator-Advanced™, TSE-Systems). MRI and 1H-MR spectroscopy (MRS) were performed at a magnetic field strength of 7T (ClinScan, Bruker-BioSpin). MRI comprised T2-weighted images (2D-FSE, TR/TE= 4000/50ms), diffusion weighted images (2D-EPI, TR/TE= 7000/28 ms, 12 directions, b = 0/1000s/mm²) and 3 differently weighted 3D FLASH based datasets (TR/TE = 28/1.9 ms, flip-angle = 25° for T1-weighting and 5° for proton-weighting, the later with/without additional magnetization-transfer weighting by Gaussian-shaped off-resonance pulses with a flip angle of 50° and an off-resonance frequency of 1200 Hz). Localized proton MR spectra (PRESS, TR/TE= 6000/10 ms) were obtained from a volume-of-interest in hippocampus (1.8x0.7x1.8mm³), cortex (3.9x0.7x3.2 mm³) and corpus callosum (3.9x0.7x1.7 mm³). Metabolite quantification involved spectral evaluation by LCModel (Version 6.3-0G (Provencher, 1993)). Results with Cramer-Rao lower bounds >20% were excluded from further analyses. Maps of magnetization transfer ratio (MTR), magnetization saturation (MTSat), and diffusion tensor imaging (DTI) readouts, fractional anisotropy (FA), apparent diffusion coefficient (ADC), axial diffusivity (AD), and radial diffusivity (RD) were calculated using in-house Matlab scripts (Mathworks).

**Electron Microscopy (EM)**

Anesthetized mice were perfused intracardially with 15 mL HBSS, followed by 50 mL fixative (2.5% glutaraldehyde, 4% paraformaldehyde in phosphate buffer with 0.5% NaCl). Brains were dissected and 200μm sagittal sections cut using Leica-VT1200S Vibratom (Leica-Microsystems). The genu of the corpus callosum was punched out of the sliced tissue, post-fixed with 2% OsO₄ (Science Services), dehydrated with ethanol/proplylene oxide and Epon-embedded (Serva). Semi-thin sections were stained with methylene blue and Azurll. Ultrathin sections were contrasted with 4% uranyl acetate and lead citrate, scanned via Zeiss EM900 (Zeiss) and digital pictures obtained by wide-angle dual speed 2K-CCD-Camera (TRS).

Myelin/axon tracings were performed on EM pictures (>15/animal) via ImageJ (http://rsb.info.nih.gov/ij/). As myelin thickness index and inner tongue ratio, g-ratio and t-ratio were calculated.

**Reverse Transcription and Real-Time PCR**

For RNA analysis (PFC/brainstem), tissues were homogenized in Qiazol, chloroform added, samples centrifuged at 12,000g, 15 min, 4°C, extractions performed with miRNeasy mini-kit (QIAGEN), RNA concentration measured via absorbance at OD260nm and purity controlled (OD260/OD280). RNA (200 ng) was mixed with 0.6 pmol oligo dT-mix, 120 pmol of Random-Hexamer (Roche) and retro-transcribed to cDNA with SuperScriptIII Reverse-Transcriptase-Kit (Life-Technologies). Derived cDNA (8 ng) was used as template for SYBR GREEN-based real-time PCR in Roche LightCycler 480. Samples were assayed in triplicate. Myelin mRNAs were normalized to 18s rRNA, microglia markers to the geometric mean of Gapdh, β-Axin, Pdgf. Following primers were used:

- **Mhp F5'-ACGGACACCTTCCAAGTT-3' R5'-GTGTGCCG TCACCGTGAAAAA-3'**
- **Cop F5'-TAACCTTCCTTAGCCCTG-3' R5'-GTCTCCTA GCATGTGCGACCT-3'**
- **Plp F5'-GGCTAGACATCCCCAAGAAG-3' R5'-GCAAACA CCAGGAGCCATA-3'**
- **18s F5'-GCTTCTAATACACACACATCCCA-3' R5'- AAATGATTTGTTCTTCGCC-3'**
- **iNos F5'-GGGCTTGTCAGCGGAGATCA-3' R5'-CCATGAT GGTCACAATTCTGC-3'**
- **Tufz F5'-GCAACATCCCTACCTCCT-3' R5 '-CCCCA GGGCCAGGAAAT-3'**
- **Arginase1 F5'-AAGGAAAAGTTCCCCAGATGCCA-3' R5'- GC AAGCCAATGTACAGCTG-3'**
- **β-Axin F5'-CTCTCTTCCTGGAGAAGAGC-3' R5'-ATGCA CAGGATTCAC-3'**
- **Gapdh F5'-CAATGGATACCGGCTACAGCAA-3' R5'- TTACTCTTGGAGGCCATGT-3'**
- **Pdgf F5'-AACAGATCTTCTGATCAGCCTACACT-3' R5'- CCTTCAGGGAGTAACCA-3'**
Western Blotting
Tissue was homogenized in lysis buffer (50 mM Tris/Cl pH 7.5, 150 mM NaCl, 1 mM EDTA, 1% Triton x-100, 0.5% sodium deoxycholate) with addition of freshly prepared protease inhibitors (50 μg/mL Leupeptin, 0.2 mM phenylmethanesulfonylfluoride, 1 mM of activated sodium orthovandate, 30 μg/mL Aprotinin), incubated on ice (30 min) and centrifuged at 9000 rpm, 4°C, 10 min. After determination of protein concentration (Lowry et al., 1951), samples were mixed with 4X Lämmli-buffer and incubated at room temperature (30 min). Comparable amounts of total protein/sample were separated on 15% SDS polyacrylamide gels and transferred onto nitrocellulose. Membranes were blocked for ≥1 h at room temperature (30 min). Comparable amounts of total protein/sample were separated on 15% SDS polyacrylamide gels and transferred onto nitrocellulose. Membranes were blocked for ≥1 h at room temperature (Tris-buffered saline, 1% Tween, 5% powder milk) and incubated overnight at 4°C with primary antibody: anti-MBP (1:2000, Dako), anti-CNPase (1:5000, Sigma-Aldrich), anti-PLP (1:500, A431, produced in-house (Jung et al., 1996) and MBP (1:2000, Dako), anti-CNPase (1:5000, Sigma-Aldrich). Incubation with secondary antibody, anti-mouse or anti-rabbit (1:5000, Rockland) followed for 2 h at room temperature. Bands were detected with Odyssey Infrared Imaging system and quantified via Image-Studio-Lite software. Intensities of Mbp, Cnp and Plp were normalized to a-Tubulin.

Immunohistochemistry
Anaesthetized mice were perfused with 4% paraformaldehyde (PFA), brains stored at −80°C. Coronal sections (30 μm) were immunolabeled for microglia marker anti-Iba1 (ionized calcium-binding adapter molecule 1; Wako, 1:1000 in 3% normal horse serum/0.5% TritonX-100), followed by donkey anti-rabbit antibody (R&D System, 594nm). Cell nuclei were counterstained with DAPI (4',6-diamidino-2-phenylindole 1:10000, Sigma-Aldrich). Brain slices were scanned with inverted epifluorescence microscope (LEICA, DMI 6000B) and analyses performed with IMARIS software (http://www.bitplane.com/) and Fiji (http://fiji.sc/Fiji).

Statistical Analysis
Between-group comparisons were performed using t-test for independent samples and 2-way analysis of variance (ANOVA or repeated-measures ANOVA). Mann-Whitney U and Wilcoxon tests were applied when normality assumption was violated (Kolmogorov-Smirnov test). All statistics were performed using Prism GraphPad software or MatLab scripts (Mathworks). Data are expressed as mean ± SEM, P values <0.05 considered significant.

Results
Heterozygous Mbp+/− mice were long-lived, of normal body size, and without obvious abnormalities, including cage behavior and litter sizes, in agreement with classical studies of Shiverer (Mbp−/−) mice (Chernoff, 1981; Readhead et al., 1987; Roach et al., 1985; Shine et al., 1992). At much lower level of Mbp expression (in partly complemented Mbp-transgenic Shiverer mice) Mbp is rate-limiting for myelination (Readhead et al., 1987). Thus, we hypothesized that also Mbp+/− mice might exhibit subtle defects of white matter formation, previously overlooked.

Spatio-Temporal Differences of Myelin Gene Expression
We quantified steady-state mRNA levels for the major myelin-specific proteins in PFC and brainstem (Fig. 1A–C). Plp1 and Cnp mRNA, taken as indicator of oligodendrocyte number and differentiation, did not reveal any appreciable difference between Mbp+/− and Mbp+/+ mice. Since oligodendrocytes do not die in Shiverer mice (Bu et al., 2004; Rosebrough, 1980), Plp1 and Cnp mRNA data confirmed that oligodendrocyte numbers are unchanged in aged Mbp+/− mice. In agreement with decreased Mbp gene dosage, Mbp mRNA was reduced by about 50% in Mbp+/+ mice throughout postnatal development. Interestingly, at 20 months, this difference vanished (significance lost), both in PFC and brainstem (Fig. 1A). Thus, at older age, Mbp gene dosage appears no longer rate-limiting for Mbp mRNA abundance.

Mbp protein was reduced in tissue lysates by 50% in 3- and 6-month-old Mbp+/− mice, confirming previous studies (Barbaresi et al., 1983). Plp/Dm20 and Cnp (normalized to a-Tubulin) were not different in Mbp+/− mice, in line with mRNA data. However, in older mice (20 months), the Mbp content began to differ between brain regions. In the PFC, Mbp+/− mice expressed <25% of Mbp compared to Mbp+/+, accompanied by a reduction of CNP by about 50%. In contrast, no further Mbp decrease was detected in brainstem (Fig. 1A–C). By EM, CNS myelin was well compacted and without obvious pathology (example in Fig. 2).

Subtle White Matter Defects Revealed by MRI And 1H-NMR Spectroscopy
The qualitative inspection of T2-weighted MR images revealed minor thinning of the corpus callosum in 6 and 18 month-old Mbp+/− compared to Mbp+/+ mice, associated with general mild reduction of white to grey matter contrast (Fig. 2A). Volume decrease together with 4% reduction of MTR indicates lower myelin content per pixel in Mbp+/− mice. This difference in MTR becomes even more evident when considering the observed 10% increase in T1 relaxation time. Due to the known dependence of MTR from T1 (Hendelman et al., 2001), this corresponds to a reduction of MTsat by 20% in the corpus callosum of Mbp+/− mice, both at 6 and 18 months (Fig. 2B, Table 1). MTsat increased with age, independent of genotype. Microstructural changes in neuronal fibers were analyzed by DTI. Upon aging, DTI of the corpus callosum revealed mild increase in FA in combination with minor decrease of AD, RD, and ADC (Table 1). However, all these changes were not Mbp genotype-dependent.

By 1H-MRS, reduced Mbp gene dosage had a small but (nominally) significant impact on brain metabolism. Most pronounced at 18 months, we noted increases in myo-inositol,
taurine and total creatine in cortex and corpus callosum of Mbp^{+/−} compared to Mbp^{+/+} mice (Table 2). Since we found no difference for the neuronal marker NAA, and an increase in taurine (a potential neuroprotective agent (El Idrissi and Trenkner 1999)), myo-inositol elevation likely reflects age-dependent microglial activation in Mbp^{+/−} mice (Badar-Goffer et al., 1992; Ross et al., 1997), and also total creatine would support glial changes (Urenjak et al., 1993).

Ultrastructure of CNS Myelin
To assess morphological correlates of reduced Mbp expression at the EM level, we analyzed the rostral corpus callosum in 18 month-old mice (Fig. 2C). Myelinated axons in cross-sections showed regularly compacted sheaths and no genotype-related pathology. Quantification revealed (as expected for EM micrographs) significant heterogeneity between subregions. This could explain why the percentage of

**FIGURE 1:** Myelin gene expression (mRNA, protein) in prefrontal cortex and brainstem of Mbp^{+/−} versus Mbp^{+/+} mice at 3 different ages. A: Mbp/Mbp, B: Cnp/Cnp, C: Plp/Plp. N = 4 per group; 2-sided t-tests, mean ± SEM presented.
myelinated axons or inner tongue size did not yield statistically significant differences between $\text{Mbp}^{+/+}$ and $\text{Mbp}^{-/-}$ mice. Whereas g-ratio measurements showed just a tendency of overall reduction of myelin sheath thickness in $\text{Mbp}^{+/+}$ mice, reduced myelin sheath thickness of small caliber axons ($<0.6 \mu m$) was significant.
| Indices                        | Brain area                | Volumetry (mm$^3$) | 6 months | 18 months | 6 months vs. 18 months (P values) |
|-------------------------------|---------------------------|-------------------|----------|-----------|----------------------------------|
|                               |                           | $Mbp^{+/+}$       | ($n = 14$) | $Mbp^{+/−}$ | ($n = 12$)                  | $P$ value | $Mbp^{+/+}$       | ($n = 15$) | $Mbp^{+/−}$ | ($n = 17$) | $P$ value | Genotype | Age   | Interaction |
| Volumetry (mm$^3$)            | Ventricles                | 10.73 ± 0.38      | 12.76 ± 0.48 | 11.96 ± 0.53 | 12.57 ± 0.65                  | 0.1       | 0.13               | 0.44       | 0.56       |
|                               | Cerebellum                | 59.9 ± 0.67       | 61.02 ± 0.64 | 59.16 ± 0.73 | 57.89 ± 1.03                  | 0.24      | 0.93               | 0.03       | 0.17       |
|                               | Olfactory Bulb            | 26.27 ± 0.44      | 26.76 ± 0.3  | 27.48 ± 0.28 | 27.07 ± 0.16                  | 0.37      | 0.78               | 0.07       | 0.09       |
|                               | Brainstem                 | 58.8 ± 1.12       | 52.99 ± 1.77 | 62.35 ± 2.35 | 61.39 ± 2.0                   | 0.01      | 0.17               | 0.002      | 0.36       |
|                               | Brain Matter (excluded the previous) | 315.2 ± 3.92    | 314.4 ± 3.1  | 323.6 ± 2.5  | 314.8 ± 3.81                  | 0.88      | 0.19               | 0.22       | 0.26       |
|                               | Fractional Anisotropy     |                  |           |           |                                 |           |                    |            |            |
|                               | Corpus Callosum           | 0.34 ± 0.006      | 0.35 ± 0.004 | 0.37 ± 0.008 | 0.36 ± 0.011                  | 0.17      | 0.87               | 0.02       | 0.17       |
|                               | Radial Diffusivity        |                  |           |           |                                 |           |                    |            |            |
|                               | ($10^{-6}$ mm$^2$/s)      | 546 ± 6.3         | 556 ± 1.1  | 506 ± 12     | 516 ± 9.7                     | 0.07      | 0.32               | $<0.0001$  | 0.52       |
|                               | Axial Diffusivity         |                  |           |           |                                 |           |                    |            |            |
|                               | ($10^{-6}$ mm$^2$/s)      | 908 ± 11          | 963 ± 12   | 905 ± 15     | 897 ± 13                      | 0.03      | 0.71               | 0.02       | 0.03       |
|                               | Apparent Diffusion Coefficient | 667 ± 7        | 700 ± 11   | 639 ± 12.2   | 619 ± 27                      | 0.01      | 0.72               | 0.005      | 0.15       |
|                               | ($10^{-6}$ mm$^2$/s)      |                  |           |           |                                 |           |                    |            |            |
|                               | Magnetization Transfer Ratio |                  |           |           |                                 |           |                    |            |            |
|                               | Corpus Callosum           | 0.67 ± 0.004      | 0.64 ± 0.003 | 0.67 ± 0.002 | 0.65 ± 0.004                  | **0.001** | 0.67 ± 0.002       | **<0.0001**| 0.03       | 0.9       |
|                               | Magnetization Transfer Saturation |            |           |           |                                 |           |                    |            |            |
|                               |                          | 0.05 ± 0.0005     | 0.04 ± 0.004 | 0.06 ± 0.002 | 0.04 ± 0.001                  | **0.001** | 0.06 ± 0.002       | **<0.0001**| 0.0002     | 0.42      |
|                               | T1 Relaxation             | 1353 ± 21         | 1509 ± 19  | 1322 ± 88    | 1450 ± 18                     | **0.001** | 1322 ± 88          | **<0.0001**| 0.005      | 0.36      | 0.78     |

Bolded $P$ values, “survived” multiple testing adjustment via Bonferroni (MT and DTI: 7 parameters evaluated in one brain region).
### TABLE 2: Metabolic Alterations in the Brain of Mbp<sup>+/−</sup> Versus Mbp<sup>+/+</sup> Mice Measured by 1H-MRS

| Metabolite (mM) | Brain area  | 6 months | 18 months | 6 months vs. 18 months (P values) |
|----------------|-------------|----------|-----------|-----------------------------------|
|                | Genotype   | Age      | Interaction |
| Myo-inositol   | Cortex     | 3.49 ± 0.12 | 3.62 ± 0.1 | 0.44  |
|                | Corpus Callosum | 3.65 ± 0.06 | 3.77 ± 0.08 | 0.09  |
|                | Hippocampus | 4.67 ± 0.16 | 4.62 ± 0.22 | 0.85  |
|                | Mbp<sup>+/−</sup> (n = 12) | 3.25 ± 0.12 | 3.59 ± 0.1 | 0.02  |
|                | Mbp<sup>+/+</sup> (n = 15) | 3.71 ± 0.11 | 4.15 ± 0.14 | 0.01  |
| Taurine        | Cortex     | 10.54 ± 0.21 | 11.17 ± 0.34 | 0.17  |
|                | Corpus Callosum | 10.18 ± 0.16 | 9.848 ± 0.2 | 0.3  |
|                | Hippocampus | 11.18 ± 0.22 | 11.62 ± 0.35 | 0.28  |
|                | Mbp<sup>+/−</sup> (n = 17) | 9.21 ± 0.35 | 10.48 ± 0.22 | 0.006 |
|                | Mbp<sup>+/+</sup> (n = 15) | 9.71 ± 0.22 | 10.46 ± 0.15 | 0.007 |
| Choline        | Cortex     | 1.24 ± 0.04  | 1.4 ± 0.13  | 0.34  |
|                | Corpus Callosum | 1.2 ± 0.02  | 1.19 ± 0.03 | 0.74  |
|                | Hippocampus | 1.14 ± 0.04  | 1.2 ± 0.03  | 0.21  |
|                | Mbp<sup>+/−</sup> (n = 12) | 1.2 ± 0.05  | 1.2 ± 0.04  | 0.49  |
|                | Mbp<sup>+/+</sup> (n = 15) | 1.31 ± 0.06 | 1.32 ± 0.05 | 0.87  |
| tCreatine      | Cortex     | 7.03 ± 0.16  | 7.44 ± 0.25 | 0.45  |
|                | Corpus Callosum | 7.72 ± 0.1  | 7.66 ± 0.11 | 0.77  |
|                | Hippocampus | 8.2 ± 0.12   | 8.33 ± 0.17 | 0.53  |
|                | Mbp<sup>+/−</sup> (n = 17) | 6.83 ± 0.22 | 7.55 ± 0.08 | 0.002 |
|                | Mbp<sup>+/+</sup> (n = 15) | 8.07 ± 0.11 | 8.4 ± 0.08 | 0.02  |
| tNAA           | Cortex     | 8.22 ± 0.18  | 8.62 ± 0.14 | 0.1  |
|                | Corpus Callosum | 8.06 ± 0.12 | 7.9 ± 0.11 | 0.48  |
|                | Hippocampus | 7.5 ± 0.1    | 7.39 ± 0.17 | 0.55  |
|                | Mbp<sup>+/−</sup> (n = 17) | 8.27 ± 0.3  | 8.69 ± 0.12 | 0.6  |
|                | Mbp<sup>+/+</sup> (n = 15) | 7.99 ± 0.12 | 8.21 ± 0.11 | 0.19  |

Bolded P values “survived” testing adjustment via Bonferroni (10 comparisons: 5 metabolites and 2 time points per brain region).
Microglial Cells in the Corpus Callosum

Since elevated myo-inositol has been associated with microgliosis (Badar-Goffer et al., 1992; Ross et al., 1997), we immunostained brain sections of 14-month-old mice for Iba1, a microglia/macrophage-specific cytosolic protein. When quantified in rostral corpus callosum, an approximately 30% increase in Iba1

$^{+}$ cells was found in Mbp

$^{-/-}$ compared to Mbp

$^{+/-}$ mice. Notably, there was no obvious change of microglial morphology from the ramified appearance. Activation of microglia/macrophages is supported by a 2-fold increase in TNF-alpha mRNA in PFC of Mbp

$^{-/-}$ (Fig. 3A–E).

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$^{+/-}$ mice. Notably, there was no obvious change of microglial morphology from the ramified appearance. Activation of microglia/macrophages is supported by a 2-fold increase in TNF-alpha mRNA in PFC of Mbp

$^{+/-}$ (Fig. 3A–E). A similar picture of low-grade inflammation, including myo-inositol increase and microgliosis as shown here for Mbp

$^{+/-}$ mice, was found in 18 month-old Cnp

$^{-/-}$ mice (Fig. 3F–G and (Hagemeyer et al., 2012)). We note that in aging Mbp

$^{+/-}$ mice, Cnp was reduced by 50% in the PFC (see above).

Behavioral Consequences of a Subtle Hypomyelination Phenotype

To clarify whether the subtle hypomyelination phenotype found in Mbp

$^{+/-}$ is by itself the cause of behavioral abnormalities in absence of overt neurological deficits, several cohorts of male mice underwent comprehensive behavioral testing. As listed in Table 3, Mbp

$^{+/-}$ mice showed normal sensory and motor functions as well as anxiety-like behavior, tested in elevated plus maze and open field. There were no signs of anhedonia, compulsive or abnormal home-cage behavior. Sociability and learning abilities were similar between genotypes. However, Mbp

$^{+/-}$ mice showed persistent impairment in PPI (Fig. 4A,B). PPI deficit is among the most reliable objective features of severe psychiatric diseases and as such one of the most relevant translational phenotypes (Braff et al., 1978; Fendt and Koch 2013; Kumari et al., 2005). The here-with documented network dysfunction likely affects the ability to adequately filter and interpret environmental stimuli, a common trait in psychiatric disorders. Another network function, the plain startle response, a measure of cerebellar reflex circuitry, tended to be lower in Mbp

$^{+/-}$ mice at 3 months, and was reduced at 17 months (Fig. 4C). Importantly, Mbp

$^{+/-}$ mice exhibited upon aging (>16 months) an increasingly catatonic phenotype (Fig. 4D,E, Supp. Info. videos), reminiscent of that described earlier in aged Cnp

$^{-/-}$ mice (Hagemeyer et al., 2012). Together, our behavioral data...
| Behavioral paradigms                                           | 3 months | 17 months |  |  |
|---------------------------------------------------------------|----------|-----------|--|---|
|                                                               | $Mbp^{+/+}$ | $Mbp^{+-}$ | $Mbp^{+/+}$ | $Mbp^{+-}$ |
| Anxiety and activity                                         |          |           | $P$ value |          |
| Elevated plus maze (time close arm [s])                      | 22       | 34        | 176 ± 9   | 166 ± 10  | 0.67      |
| Elevated plus maze (time open arm [s])                       | 22       | 34        | 31 ± 5    | 31.28 ± 5 | 0.79      |
| Elevated plus maze (time centre [s])                         | 22       | 34        | 75 ± 8    | 78 ± 5    | 0.67      |
| Elevated plus maze (total distance [m])                      | 22       | 34        | 10.1 ± 0.3| 9.46 ± 0.51| 0.6       |
| Elevated plus maze (average speed [mm/s])                    | 22       | 34        | 31.8 ± 1.5| 33.8 ± 1.2| 0.31      |
| Open field (total distance [m])                              | 22       | 34        | 48 ± 1.6  | 47.6 ± 1.4| 0.72      |
| Open field (average velocity [mm/s])                         | 22       | 34        | 116.2 ± 3.4| 115.3 ± 3.4| 0.6      |
| Open field (periphery [s])                                   | 22       | 34        | 341 ± 9   | 341 ± 7   | 0.77      |
| Open field (intermediate [s])                                | 22       | 34        | 60 ± 7    | 60 ± 5    | 0.71      |
| Open field (centre [s])                                     | 22       | 34        | 19 ± 2    | 19 ± 2    | 0.9       |
| Open field (latency to the periphery [s])                    | 22       | 34        | 16 ± 4    | 15 ± 2    | 0.21      |
| Exploratory behavior                                         |          |           |           |          |
| Hole board (holes visited [n])                               | 22       | 34        | 5 ± 1     | 6 ± 1     | 0.38      |
| Impulsivity                                                  |          |           |           |          |
| Marble burying (marble buried [n])                           | 22       | 34        | 7.23 ± 1.14| 7.1 ± 0.92| 0.9      |
| Motor learning and coordination                               |          |           |           |          |
| Rota-rod day 1 (latency to fall [s])                         | 22       | 34        | 123 ± 15  | 140 ± 17  | 0.87      |
| Rota-rod day 2 (latency to fall [s])                         | 22       | 34        | 214 ± 18  | 208 ± 15  | 0.79      |
| Muscle strength                                              |          |           |           |          |
| Grip strength [p]                                            | 22       | 34        | 142 ± 3   | 139 ± 2   | 0.41      |
| Heat/pain perception                                         |          |           |           |          |
| Hot plate (latency to lick [s])                              | 22       | 34        | 15 ± 1    | 16 ± 1    | 0.15      |
| Sight                                                        |          |           |           |          |
| Visual cliff (latency in the “air” [%])                      | 22       | 34        | 36 ± 4    | 34 ± 3    | 0.57      |
| Olfaction                                                    |          |           |           |          |
| Buried food test (digging latency [s])                       | 22       | 34        | 87 ± 21   | 74 ± 13   | 0.88      |
| Behavioral paradigms                  | 3 months | 17 months |   |   | P value | 3 months | 17 months |   |   | P value |
|--------------------------------------|----------|-----------|---|---|---------|----------|-----------|---|---|---------|
| **Hearing**                          |          |           |   |   |         |          |           |   |   |         |
| Hearing (startle amplitude [AU])     | 22       | 34        |   |   |         |          |           |   |   |         |
| Anhedonia                           |          |           |   |   |         |          |           |   |   |         |
| Sucrose preference test [%]          | 22       | 34        | 74 ± 2 | 74 ± 2 | 0.89    |          |           |   |   |         |
| **Homecage behavior**                |          |           |   |   |         |          |           |   |   |         |
| LABORAS (climbing [s])               | 10       | 16        | 2188 ± 429 | 2480 ± 419 | 0.64 |          |           |   |   |         |
| LABORAS (grooming [s])               | 10       | 16        | 5363 ± 463 | 4899 ± 457 | 0.5 |          |           |   |   |         |
| LABORAS (locomotion [s])             | 10       | 16        | 2734 ± 295 | 2859 ± 374 | 0.85 |          |           |   |   |         |
| LABORAS (drinking [s])               | 10       | 16        | 84.5 ± 23   | 260 ± 56   | 0.12 |          |           |   |   |         |
| LABORAS (eating [s])                 | 10       | 16        | 2468 ± 362  | 2486 ± 272 | 0.85 |          |           |   |   |         |
| LABORAS (scratching [s])             | 10       | 16        | 1065 ± 120  | 869 ± 217  | 0.73 |          |           |   |   |         |
| LABORAS (circular movements [s])     | 10       | 16        | 541.5 ± 59.2| 711.9 ± 107.1| 0.51 |          |           |   |   |         |
| **Sociability**                      |          |           |   |   |         |          |           |   |   |         |
| Three-chambered social test (sociability [%]) | 22 | 34 | 62 ± 2 | 63 ± 2 | 0.81 |          |           |   |   |         |
| **Learning and memory**              |          |           |   |   |         |          |           |   |   |         |
| Morris water maze, training (escape latency [s]) | 22 | 34 |          |          |          |          |          |   |   |         |
| Probe trial - time in target zone [%] | 22       | 34        | 41 ± 2 | 45 ± 2 | 0.17 | 22       | 33        | 47 ± 2 | 54 ± 2 | 0.09 |
| Escaping latency - reversal [s]      | 22       | 34        |          |          |          | 22       | 33        |          |          |         |
| Probe trial reversal - time in target zone [%] | 22 | 34 | 33 ± 2 | 32.7 ± 2 | 0.54 | 22       | 33        | 41 ± 2 | 43 ± 2 | 0.3    |
| Novel object recognition (novelty preference [%]) | 22 | 34 | 59 ± 3 | 58 ± 3 | 0.81 |          |           |   |   |         |
| Y-maze (alternation index [#])       | 22       | 34        | 0.7 ± 0.05 | 0.8 ± 0.03 | 0.63 |          |           |   |   |         |
demonstrate that the isolated Mbp/Mbp reduction of Mbp \(^{+/−}\) mice is sufficient to induce measurable network dysfunction in both PFC (PPI) and brainstem (startle), as well as age-related catatonia (frontal corpus callosum), all mental disease-typical features.

**Discussion**

In human patients with psychiatric diseases, numerous MRI and post-mortem findings have been published that point to subtle white matter abnormalities and myelin loss (Balevich et al., 2015; Chambers and Perrone-Bizzozero, 2004; Dracheva et al., 2006; Flynn et al., 2003; Hakak et al., 2001; Honer et al., 1999; Kochunov et al., 2013; Kubicki et al., 2005; Kubicki et al., 2002; Matthews et al., 2012; McIntosh et al., 2008; Parlapani et al., 2009; Tkachev et al., 2003).

However, once complex diseases such as schizophrenia become clinically manifest it is virtually impossible to distinguish cause and consequence of correlative findings. While minor defects of subcortical myelination are unlikely the sole cause of a complex psychiatric disorder, the failure to adequately myelinate cortical fibers is an obvious risk factor for long-range connectivity and may play a causal role when combined with other genetic and environmental risk factors or aging (Nave and Ehrenreich, 2014). On the other hand, clinical and autopsy findings of white matter loss could be merely secondary effects in a long-lasting disease process that primarily perturbs the neuronal circuitry. Myelination defects could finally be secondary after years of pharmacological treatment (Bartzokis et al., 2007; Ho et al., 2011). All these scenarios are not strictly alternative and might act in combination. Mechanistic insight into any of these key variables requires well-defined experimental animal models.

In the present study we addressed the potentially primary role of minor myelination defects for behavior and brain network function. We comprehensively reanalyzed the phenotype of “healthy” Mbp heterozygous Shiverer mice, a classical line of natural mouse mutants, in which only Mbp homozygous mice are dysmyelinated and exhibit severe neurological defects (Chernoff, 1981). Contrary to the early literature with its focus on the homozygous phenotype, we found that central myelination of Mbp heterozygous mice is not identical to wildtype controls. Upon gross histological analysis, and when judged by cage-behavior, Mbp \(^{+/−}\) mice cannot be distinguished from Mbp \(^{+/+}\) littermates. However, the detailed analysis at biochemical, histological, electron microscopic, MRI/MRS, and behavioral level, revealed intriguing differences. This includes a mild hypomyelination of the PFC, with signs of microglial activation, combined with prepulse inhibition
(PPI) deficits and clinical signs of catatonia upon aging in the presence of otherwise normal behavioral performance.

The principle finding (by histology and MRI) is a subtle hypomyelination of the PFC of adult mice and a thinning of the underlying corpus callosum, i.e., regions relevant for cognitive functions and catatonia. Oligodendrocytes were probably not diminished in number, as suggested by unaltered mRNA abundance of myelin protein genes (except for Mbp itself), indicating that hypomyelination is unlikely caused by a paucity of mature oligodendrocytes.

Our attempt to confirm the histologically documented hypomyelination by EM as a reduction of myelin sheath thickness yielded only borderline significance, except for small caliber fibers which had a larger g-ratio. This supports an alternative working model, in which myelination by Mbp heterozygous oligodendrocytes is less efficient at an early stage of ensheathment. For example, a small fraction of oligodendroglial processes might normally contact axons and initiate myelination, but then fail to continue wrapping and instead retract, which would lead to a reduced number of myelinated internodes. This model is compatible with current concepts of CNS myelination and the role of activity-dependent Mbp mRNA translation at the tip of an oligodendrocyte process (Waké et al., 2011). Here, Mbp is essential in stabilizing the first membrane wraps that have been deposited in order for myelination to proceed (Simons and Nave, 2016; Snaiadero et al., 2014). We further note that in wild type mice, axons of cortical projection neurons are often incompletely myelinated along their length (Tomassy et al., 2014). This suggests that myelination signals in the cortex are indeed weak in comparison to other CNS regions, such as spinal cord or optic nerve, where virtually all axons are fully myelinated at late postnatal stages. We therefore propose that translating 50% Mbp mRNA (in Mbp/+ mice) may be sufficient for oligodendrocytes in the spinal cord, which are exposed to strong axonal signals, to stabilize the initial wraps and robustly myelinate axons. Thus, in Mbp heterozygous oligodendrocytes of the PFC myelination may be inefficient because here axonal signals, including their spiking activity, are weak compared e.g. to the spinal cord. Such a working model is clearly speculative, but fully compatible with experimental observations in other labs (Lundgaard et al., 2013; Tomassy et al., 2014; Waké et al., 2011). It could explain why at higher age the amount of Mbp in the PFC of heterozygotes has dropped below the expected 50%. This differs from the brainstem, where axonal activity is less likely to decrease with age and where myelination remains robust. Our data also match prior experiments in wild type mice, which have shown that reduced social activity during postnatal development has a negative impact on myelination, specifically of the PFC and its underlying white matter (Liu et al., 2012; Makinodan et al., 2012). Collectively, these in vivo observations support the idea that cortical myelination is stimulated by neuronal activity, and that the strength of this signal differs between CNS regions and presumably at different stages of life.

By 1H-MRS, we detected elevated myo-inositol levels in white matter that corresponded to the histologically determined increase in microglial cells and elevated TNF-alpha expression. We have no evidence for underlying axonal degeneration in white matter tracts, or that low-grade microglial activation is a response to the assembly of unstable myelin, as by electron microscopy compact myelin appears fully stable with only 50% Mbp incorporated. Interestingly, neuroinflammation in white matter tracts is a major feature of mice homozygous and heterozygous null for the Cnp gene (Hagemeyer et al., 2012; Lappe-Siefke et al., 2003), and we measured at the protein level a 50% reduction of Cnp (but not Plp) in the cortex of aged Mbp+/− mice. Cnp is a structural protein of non-compacted myelin that antagonizes MBP during myelin compaction by maintaining “myelinic channels” open (Snaiadero et al., 2014). The mechanism of reduced Cnp/CNP in cortical white matter of Mbp+/− mice or in autopsy material from patients with schizophrenia (Dracheva et al., 2006; Flynn et al., 2003; Hakak et al., 2001; Prabakaran et al., 2004) is unknown. However, it is plausible that loss of Cnp/CNP-dependent metabolic support for myelinated axons triggers (unknown) stress signals that activate resident microglial cells. Above threshold or upon “secondary hits,” as shown for Cnp heterozygous mice (Wieser et al., 2013), neuroinflammation in white matter tracts can also drive axonal degeneration, which generates a vicious circle of events that is relevant for neuropsychiatric diseases.

At the behavioural level, Mbp+/− mice are still below threshold of a clear psychiatric disease phenotype, as evidenced by a comprehensive battery of tests, in which they exhibit normal basic behavior and higher brain functions including cognition. However, already at age 3 months they showed deficits of sensorimotor gating with reduced PPI, a physiological phenomenon shared between human patients with schizophrenia and rodent disease models (Swerdlov et al., 2008). The link between Mbp heterozygosity and PPI deficits is unclear, but is likely reflecting subtle myelin abnormalities and desynchronized impulse propagation. We note that also Plp1 overexpressing mice share mild PPI deficits, but in the presence of other schizophrenia-relevant defects (Tanaka et al., 2009). Another network disturbance (i.e. of a cerebellar reflex circuitry) of Mbp+/− mice is indicated by the here observed reduction in the startle response. Interestingly, a similar phenomenon has been reported in a mouse model of chronic stress (Dirks et al., 2002). Most importantly, later in life, Mbp+/− mice exhibit a catatonia-like behavior that we have previously seen in Cnp+/− mice upon aging (Hagemeyer et al., 2012). Collectively, these in vivo observations support the idea that cortical myelination is stimulated by neuronal activity, and that the strength of this signal differs between CNS regions and presumably at different stages of life.
et al., 2012), suggesting again that reduced Cnp expression and mild neuroinflammation, downstream of a slightly abnormal CNS myelin assembly, may predispose to this schizophrenia-relevant frontal cortical phenotype.

In summary, the reduction of Mbp/Mbp induces subtle hypomyelination, accompanied by low-grade inflammation, leading to cortical network dysfunction. This reveals that Mbp/Mbp decrease per se can be causative of psychiatric illness-relevant phenotypes.

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Author Contributions

Concept and design of the study: HE, KAN
Data acquisition and analysis: GP, SB, WM, TR, JB, NM, GW, IH, SG, HW
Drafting the manuscript and display items: HE, KAN, GP
All authors read and approved the final version of the manuscript.

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