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Familial t(1;11) translocation is associated with disruption of white matter structural integrity and oligodendrocyte–myelin dysfunction

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Abstract
Although the underlying neurobiology of major mental illness (MMI) remains unknown, emerging evidence implicates a role for oligodendrocyte–myelin abnormalities. Here, we took advantage of a large family carrying a balanced t(1;11) translocation, which substantially increases risk of MMI, to undertake both diffusion tensor imaging and cellular studies to evaluate the consequences of the t(1;11) translocation on white matter structural integrity and oligodendrocyte–myelin biology. This translocation disrupts among others the DISC1 gene which plays a crucial role in brain development. We show that translocation-carrying patients display significant disruption of white matter integrity compared with familial controls. At a cellular level, we observe dysregulation of key pathways controlling oligodendrocyte development and morphogenesis in induced pluripotent stem cell (iPSC) derived case oligodendrocytes. This is associated with reduced proliferation and a stunted morphology in vitro. Further, myelin internodes in a humanized mouse model that recapitulates the human translocation as well as after transplantation of t(1;11) oligodendrocyte progenitors were significantly reduced when compared with controls. Thus we provide evidence that the t(1;11) translocation has biological effects at both the systems and cellular level that together suggest oligodendrocyte–myelin dysfunction.

Introduction

Schizophrenia (SZ) and other major mental illnesses (MMI) such as bipolar disorder and major depression show high heritability. Accumulating evidence from GWAS studies points to a multifactorial polygenic inheritance, with...
individual genes conferring a modest increased susceptibility, as well as pleiotropy [1]. In contrast, rare genetic variants, such as a balanced chromosomal translocation in a large Scottish family, that co-segregate with MMI show high penetrance [2–6]. The range of psychiatric phenotypes observed in people carrying the balanced t(1;11) translocation suggests its study will be of considerable value for an improved understanding of biological processes underlying MMI. This translocation disrupts the DISC1 gene and segregates with SZ and affective disorders in this large family.

Despite multiple lines of evidence from pathological, gene expression, and radiological studies, the role of glia is understudied [2, 7–15]. As oligodendrocytes enable rapid impulse propagation and provide trophic and metabolic support to axons [16–19], their dysfunction is likely to result in altered neuronal homeostasis. Further, the sole protein-coding gene disrupted by the t(1;11) translocation; Disc1, is known to affect the specification and differentiation of oligodendrocytes [20–22] in animal models. Hence a direct study of the impact of the t(1;11) translocation on oligodendrocytes is important. Furthermore, given the limitations of animal models of neuropsychiatric disorders, complementary models of human glia in the context of MMI are particularly necessary. Indeed, recent studies using patient derived iPSC cells have shown impaired glial maturation suggesting a causal link with SZ [23, 24].

A powerful approach to interrogate the structural and cellular white matter consequences of the t(1;11) translocation is to undertake combined water diffusion MRI (dMRI) and biological studies of iPSC-derived oligodendrocytes from patients carrying the t(1;11) translocation. Previous studies using iPSCs from individuals with DISC1 mutations and MMI have predominantly studied neural precursor (NPCs) and/or neuronal processes [2, 25, 26]. Here, we demonstrate abnormalities of white matter integrity using brain dMRI in t(1;11) translocation-carrying cases compared with familial controls. In addition, case iPSC-derived oligodendrocytes display cellular and structural abnormalities in vitro as well as upon transplantation into hypomyelinated mice. This study elucidates a cell biological basis of oligodendrocyte–myelin deficits in MMI. In addition, we establish a human platform for future mechanistic studies.

**Results**

**Global changes in white matter structure and connectivity due to the t(1;11) translocation**

To begin to understand the effect of t(1;11) translocation at a whole brain as well as cellular level, we used a multi-tiered approach integrating patient and control whole brain imaging, and in vitro and in vivo stem cell experiments along with transgenic studies (Fig. 1a).

To study the impact of t(1;11) translocation on global white matter structure and structural connectivity, we undertook whole-brain probabilistic tractography using dMRI on 21 individuals from the previously reported Scottish family known to carry the t(1;11) translocation of whom eight were carriers of the t(1;11) translocation [27]. All eight carriers had a psychiatric diagnosis; one with SZ, four with major depression (MDD), three with cyclothymia while only 1 of the 13 family members without the t(1;11) translocation, had a psychiatric diagnosis [27]. The affected noncarrier however is described to carry modifier loci on chr11q2 and chr5q that might contribute to the development of MMI (described in [28]). High-resolution T1-weighted structural and dMRI data were combined to create structural connectivity matrices of each participant’s brain. In these matrices (Fig. 1b–e), the 85 gray matter regions, parcellated from the structural MRI data, are the nodes of the brain structural network while the connecting white matter pathways, identified from whole brain dMRI tractography, are its edges. The edge connection strength between nodes is obtained by recording the mean fractional anisotropy (FA), a measure of white matter microstructure, along tractography streamlines connecting all ROI (network node) pairs (Fig. 1b–e). Global graph theory measures of brain structural connectivity, such as network degree (number of connections one node has to other nodes), network strength (the average sum of edge connections per node), and global efficiency (the average of the inverse shortest path length between nodes), can then be calculated for each subject and compared across populations to assess differences in brain structure. In our case, we hypothesized that carriers have reduced connectivity (e.g., lower degree, strength, and global efficiency) than noncarriers. We estimated the effect of translocation status (carrier versus non-carrier) and age on global connectivity measures using a Markov chain Monte Carlo (MCMC) approach [29]. Significant differences between carriers and non-carriers were found for global network degree (posterior mean = $-1.81$; 95% CI = $[-3.43, -0.34]$; pMCMC = 0.02) and global network strength (posterior mean = $-1.64$; 95% CI = $[-3.09, -0.22]$; pMCMC = 0.03) (Fig. 1f, g). Global network efficiency showed a tendency towards significance (posterior mean = $-1.19$; 95% CI = $[-2.65, 0.26]$; pMCMC = 0.09) while clustering coefficient was not significantly different between groups (posterior mean = 0.93; 95% CI = $[-2.64, 0.59]$; pMCMC > 0.05) (Data not shown). Together these findings show that carriers have reduced structural connectivity compared with controls and are consistent with t(1;11) translocation causing widespread changes in white matter structural integrity.
Fig. 1 Global changes in structural networks and connectivity in t(1;11) carriers. a Overview of the approach used in this study to investigate the effect of t(11) on oligodendrocyte development and myelination capability. Brain imaging of control and affected t(11) carriers was undertaken to identify differences in connectivity. Separately, patient derived iPSCs were differentiated towards oligodendroglial fate to study developmental differences. A combinatorial approach of cellular phenotyping along with transcriptomic analysis was employed. Finally, the myelination capability was studied in transgenic mice (Der1/+) recapitulating the human mutation as well as by transplanting hiPS derived OPCs into a hypomyelinated mouse model (MBPShi/Shi, Rag2−/−). b–e Average FA-weighted structural networks (a, b) and connectivity matrices (c, d) for controls (b, d) and carriers (c, e) obtained using diffusion and structural MRI. Network nodes were identified from high-resolution T1-weighted volume scans using Freesurfer (http://freesurfer.net), with connecting edges created using probabilistic tractography (bedpostx/probtrackx; http://fsl.fmrib.ox.ac.uk/fsl) from high angular resolution diffusion MRI data. In b, c network nodes are colored to indicate gray matter in different lobar structures, while in d, e stronger connections between different cortical regions are indicated by lighter yellow colors. Careful examination of the connectivity matrices show that a number of connections have higher FA in d than e suggesting that brain connectivity is increased in controls compared with carriers. (f, g) Quantification of Global network degree (f) and Global network strength (g) from d, e shows statistically significant changes in carriers (n = 13 controls and eight translocation carriers, Markov chain Monte Carlo (MCMC) test). Median with upper and lower quartiles is shown. The whisks depict the range. Cases studied using iPS cells in this particular study have been highlighted.
Altered differentiation and gene expression in t (1;11) derived oligodendroglia

In order to begin to determine the cellular and molecular basis underlying the observed abnormalities in white matter integrity, we next studied oligodendrocytes from affected and unaffected individuals—see Suppl. Table 1. iPSC lines were established, using an episomal non-integrating method, from four individuals carrying the mutation (Case 1—cyclotymia, Case 2—MDD, Case 3—MDD, Case 4—SZ; see Supplementary table 1) and three unaffected family controls (Supplementary Fig. 1). H3K27 trimethylation staining indicated typical X-chromosome inactivation in the female iPSC lines as seen by distinct foci in the nucleus while the male lines showed diffuse staining (Supplementary Fig. 1). Quantitative analysis showed no difference in progenitor cell numbers between control and case lines (Supplementary Fig. 4d).

We found that full-length DISC1 mRNA expression was significantly reduced in case lines compared with control samples (OPCs: Controls: 0.72 ± 0.05, Cases: 0.22 ± 0.05, n = 3 independent conversions, p < 0.05 One-way ANOVA with Holm–Sidak’s correction) (Supplementary Fig. 4d).

Methods.

To begin to understand the molecular consequences of the translocation on oligodendrocytes, we next undertook RNA-seq analysis on differentiated week 3 oligodendrocyte cultures. We compared three cases and two control lines by...
paired-end RNA sequencing on an Illumina platform. By setting an adjusted p-value of <0.05, we identified 228 genes (164 upregulated and 64 downregulated) (Fig. 3a).

Gene ontology (GO) analysis identified novel candidate pathways dysregulated in MMI (Fig. 3b). The differentially regulated genes included those relating to nervous system development, activation of GTPase activity, galactosylceramide biosynthesis, myelination, and calcium ion binding (Fig. 3b, c). Genes involved in oligodendrocyte differentiation, actin binding, and ion-transmembrane transport such as NKX2-2, ZNF804A, SLC8A3, and CTTNBP2 despite not being enriched in GO analysis were also found to be dysregulated in t(1;11) carrying lines (NKX2.2 Controls: 1.0 ± 0.46 Cases: 7.16 ± 4.41; ZNF804A Controls: 2.13 ± 1.27 Cases: 25.31 ± 24.39; SLC8A3 Controls: 1.22 ± 0.44 Cases: 9.56 ± 5.71; CTTNBP2 Controls: 1.04 ± 0.37 Cases: 5.09 ± 3.28, n = 3 independent conversions, p < 0.05 F-test for comparing variances).

In vitro morphology as well as in vivo internodal length is severely affected in t(1;11) carrying oligodendrocytes

In support of the dysregulation of actin-related genes in case lines, DISC1 has previously been shown to affect microtubule organization and neurite outgrowth [32, 33].
oligodendrocytes, this is likely to affect morphological development and myelin formation. Case-derived O4+ oligodendrocytes were found to be severely stunted and dysmorphic in comparison to familial controls (Fig. 4a), a finding confirmed by quantification of surface area covered by individual oligodendrocytes (Fig. 4b) (Controls: 2170 μm² ± 179.3, Cases: 1495 μm² ± 173.6, p < 0.05 unpaired t-test, n ≥ 30 cells per line from three independent conversions). The reduction in surface area was prominent across the case lines with the exception of Case 1 which did not show a significant change (Supplementary Fig. 5a, b). Sholl analysis also showed a severe reduction in oligodendrocyte
Familial t(1;11) translocation is associated with disruption of white matter structural integrity and...

Fig. 4 Case-derived oligodendrocytes are morphologically impaired in vitro and, in a mouse-model recapitulating the translocation. a O4 staining of week 3 oligodendrocytes shows decreased cellular size and complexity of case-derived cells. b Quantification of cellular area showing the effect of t(1;11) translocation among different case-derived oligodendrocytes. n = 95 for control (three lines) and 120 for case (4 lines) oligodendrocytes. Mean ± SEM shown with paired t-test used for assessing statistical significance (p-value < 0.01). c Case oligodendrocytes have simpler morphology with fewer branch points away from the cell soma as assessed by Sholl analysis. Mean values shown and unpaired t-test used to assess statistical significance (p-value < 0.001). d Low magnification image of myelin basic protein (MBP) staining in the P21 Der1/+ cortex shows absence of gross cortical abnormalities. (Ctx cerebral cortex, Str striatum, CC corpus callosum). e Reconstructions of myelinated segments of upper cortical layer neurons from boxed region in a of wild-type (+/+) and Der1 heterozygous (Der1+) brains stained for MBP and paranodal protein Caspr. Arrowheads demarcate myelin segments used for quantification. f Graph showing internodal length measurements from b. Each data point represents one myelin segment and the graph contains pooled data from n ≥ 50 segments each from three animals per genotype. p < 0.05 Unpaired t-test with Welch’s correction. g-i Electron microscopy analysis of the corpus callosum of wild-type and Der1/+ mice (g) shows that thickness of myelin (h) and the relative frequency of myelinated fibers (i) are not affected due to the translocation. More than 250 axons measured in each group from 4 animals per genotype. j-l Single-oligodendrocyte analysis using by CNPase staining j at P42 supports findings in e, f and shows an increase in number of processes per oligodendrocytes (k) but a decrease in internodal length (l). Each data points represents one cell and the graph contains pooled data from n = 28 cells each from 4 animals per genotype. p < 0.05 Unpaired t-test. Scale bars: a 20 µm, d 500 µm, e 10 µm, g 1 µm, j 20 µm

process complexity (Fig. 4c). To evaluate whether this in vitro phenotype might be replicated in an in vivo context we next examined a novel mouse model engineered to model the effects of the t(1;11) translocation on DISC1 expression [34]. This mouse mimics the translocation-induced gene fusion that abolishes full-length DISC1 expression from one allele and produces chimeric transcripts encoding deleterious truncated/chimeric forms of DISC1 from the derived chromosome 1 [34, 35]. Here, upon co-staining for MBP, a myelin marker, and CASPR, a paranodal marker, in mouse cortices at postnatal day 21 (Fig. 4d, e), we found a significant reduction in internode lengths in the cortical layers of Der1/+ mutant mice (WT: 33.99 ± 0.93 µm, Der1/+: 21.72 ± 0.37 µm, n = 3 mice each, >190 internodes per genotype; p < 0.05 t-test with Welch’s correction) (Fig. 4f). Analysis of individual myelinated axons using electron microscopy however did not show any difference in g-ratio or in the frequency of myelination (Fig. 4g–i) (n = 4 mice each). To assess the number of myelin sheaths produced by a single oligodendrocyte as well as their respective lengths, we performed immunofluorescence for CNPase to stain entire oligodendrocytes and hence allow for tracing processes from the cell body to myelin sheaths. Employing this technique, we analyzed oligodendrocytes in the sparsely myelinated layers II/III of the medial prefrontal cortex of 6 weeks old mice (Fig. 4j). Here we found that the number of myelin sheaths produced by a single cell was significantly increased in heterozygous Der1 mice when compared with wild-type littermates (WT: 42.57 ± 2.46, Der1+/+: 49.50 ± 1.51, n = 4 mice each, 28 cells per genotype; p < 0.05 t-test) (Fig. 4k). In addition, in agreement with our in vitro analysis the average length of internodes produced by single oligodendrocytes was significantly reduced in heterozygous Der1 mice (WT: 66.94 ± 1.80 µm, Der1+/+: 60.63 ± 2.71 µm, n = 4 mice each, 28 cells per genotype; p < 0.05 t-test) (Fig. 4l).

To next evaluate the in vivo consequences of t(1;11) mutation on human oligodendrocytes we established human glial chimeras by transplantation of t(1;11) and control glial cells into neonatal hypomyelinated immune-deficient mice (MBPshih/+, Rag2−/−). Neonatal animals (P1–P2) were unilaterally injected with case (Case 2 and Case 4) or control (Control 1 and Control 2) derived OPCs (200,000 cells per injection) into the developing neocortex and analysed at 8- and 13-weeks post transplantation—the latest time-point allowed by animal regulations—by staining for MBP, GFAP, and human nuclei (hNu). Human cells were found in the corpus callosum around the site of injection where they expressed either MBP or GFAP (Fig. 5a, b). In addition, cells had also crossed the midline and differentiated. Myelin sheath lengths were greatly reduced in case-derived oligodendrocytes (Fig. 5c–f, Supplementary Fig. 5c, d) with a majority of myelin sheaths (56% for Case 2 and 48% for Case 4) between 5 and 10 µm. In comparison, only 10% of control oligodendrocytes fell in that range (Supplementary Fig. 5e). Indeed, mean sheath length showed a significant difference both at 8- and 13-weeks post transplantation (8 weeks Controls: 23.20 ± 0.84 µm Cases: 9.48 ± 0.26 µm, n = 6 mice each and 281, 320 segments measured, respectively p < 0.001 unpaired t-test with Welch’s correction; 13 weeks Controls: 29.44 ± 1.39 µm Cases: 20.24 ± 0.76 µm, n = 3–6 mice and 269, 145 segments measured, respectively, p < 0.05 unpaired t-test with Welch’s correction.) (Fig. 5g, h). Together the mutant mice and human transplantation studies reveal reduced myelin internode formation associated with the t(1;11) translocation.

Discussion

In this study we provide converging lines of evidence from human imaging, molecular, and cellular analysis including of chimeric mice that suggests that the t(1;11) translocation causes oligodendrocyte–myelin deficits.

The results from structural connectivity, measured using graph theory analysis are consistent with t(1;11) translocation dependent widespread changes to white matter connectivity. It has previously been shown that family members
who carry the balanced translocation have a pattern of cortical thinning similar to that observed in patients with SZ [36]. Here, we provide evidence of disruption of white matter topology and organization building on earlier voxel-based findings from ourselves showing a link between FA values and psychotic symptoms in those with the t(1;11) translocation [37] and between white matter organization, genetic risk markers, and cognition in the patients with “idiopathic” SZ scanned as part of the same project [38]. The recent large ENIGMA SZ diffusion tensor imaging study of over 4000 individuals has recently shown white matter microstructural differences in SZ [39] and it is likely that the t(1;11) translocation causes an altered regulation of SZ candidate genes and the DISC1 core interactome [40].

There has been much debate about the importance of DISC1 as a genetic risk factor for SZ [41, 42] and it is more appropriate to instead classify DISC1 as a genetic risk factor for MMI with a broad clinical phenotype. As such, DISC1 interacts with many protein partners and mechanistically affects several neurodevelopmental pathways despite not being identified as having a clear association with SZ from genome-wide studies [1]. This is exemplified from the clinical phenotypes of the four affected carriers in this study, one with SZ and three with affective disorders. Interestingly, OPCs and oligodendrocytes derived from a juvenile-onset, treatment resistant individual showed severe developmental, structural, and functional changes across tests (Case 4; described in Supplementary Table 1). In contrast, cells derived from individuals presenting a less severe symptomatology (as in Case 1) also showed comparatively milder changes in vitro. This both highlights the need for multiple patient lines as well as the promise of iPS-based systems to model aspects of disease variation. The use of several iPS lines allows stratification based on cellular or gene expression changes as well as identification of common phenotypes to aid drug-screening approaches. In line with this, we uncovered shared gene expression changes across patient lines.

Robust and comparable differentiation of case lines compared with familial controls showed that the translocation does not interfere with early specification and patterning of cells to the oligodendroglial lineage. Case lines however, showed premature cell cycle exit of OPCs, a finding consistent with reports for cortical neurons derived from DISC1 exon 8 interrupted iPSC lines [26] and supported also by the observed increased expression of NEX2.2, ZNF488, and SLC8A3 genes. Of these, NEX2.2 and ZNF488 are described to be specific to the oligodendroglial lineage as well as having a pro-differentiation effect on OPCs [43–45]. Thus, their increased expression in our dataset is consistent with the observed phenotype of
Familial t(1;11) translocation is associated with disruption of white matter structural integrity and... precocious differentiation. Our analysis also highlights genes identified via GWAS studies and implicated in psychosis (ZNF804A) [46] and suicidal behavior (SKA2) [47]. In addition, several genes (UGT8, GAL3ST1, ZNF488, CTTNBP2, SLC8A3, GRIA4, and KCND2) identified in our analysis agree with a recent study of glial progenitors from childhood onset SZ patient-derived iPSC lines [23]. This study of humanized glial chimeric mice suggests a potential causal role for impaired glial progenitor cell differentiation and found premature migration of GPCs, reduced white matter expansion, and hypomyelination relative to controls as well as deficits in astrocyte differentiation and morphology [23]. This points towards convergent mechanisms for oligodendrocyte dysfunction in MMI despite differences in genetic basis. Future studies employing functional genetics approaches such as CRISPR and siRNA mediated knockdown could clarify the role of these genes in the pathophysiology.

Our finding of dysregulation of actin associated genes, is of considerable interest noting that DISC1 has been shown to play a role in microtubule reorganization and neurite outgrowth [33]. In keeping with these findings, we observed case-derived oligodendrocytes to have a morphological phenotype of shorter and less complex processes. Critically, we also observed in the Der1 mouse model, that recapitulates the effects of the human translocation upon DISC1 expression, shorter internodes that were further supported by a reduction in myelin segment length found upon transplantation of case and control-derived OPCs into hypomyelinated mice. Transgenic rodent models of Disc1 have shown a range of anatomical changes including cerebral cortical thinning, reduced neurite outgrowth as well as behavioral changes (reviewed in [48]) but finer structure deficits in myelination have thus far not been shown. A crucial role for the actin remodeling pathway in myelination has been described [49–51] and could serve as an important pharmacological target in future studies. Finally, our study also points towards a galactolipid imbalance in case-derived oligodendrocytes leading to myelin defects. In keeping with this, mice genetically altered for galactolipid pathway genes present neurological manifestations due to altered axo–glia interactions and unstable myelination [52–54]. Together these findings are consistent with t(1;11) translocation dependent oligodendrocyte–myelin dysfunction.

The future evaluation of in vivo functional consequences of shorter internodes would be of great interest since shorter internodes are associated with reduced conduction velocities and even degenerative changes. This is particularly relevant when considering that gamma oscillations between different brain regions are known to be affected in MMI. Parvalbumin expressing interneurons, the major cell-type thought to regulate gamma oscillations are extensively myelinated [55, 56] and its disruption could potentially contribute to the development of MMI.

In summary, our findings here of reduced white matter connectivity in t(1;11) translocation carriers add to the growing evidence of white matter deficits observed in MMI [57] and suggest that oligodendroglia play an important role in the underlying pathophysiology of MMI.

Material and methods

Participants

Individuals with and without the t(1;11) translocation were recruited from a previously reported extended Scottish family. The t(1;11) translocation status of family members was originally ascertained by karyotyping and these were extended in subsequent studies. Researchers within the Division of Psychiatry have been in contact with members of the family for many years and through them other members of the family were invited to participate. The study was approved by the Multicentre Research Ethics Committee for Scotland. A detailed description of the study was given and written informed consent was obtained from all individuals prior to participation. A summary of the individual participants in this study, their translocation status, diagnosis, and medication at the time of biopsy is given in Supplementary Table 1. Psychiatric diagnoses were established by consensus between two trained psychiatrists according to DSM-IV (TR) criteria. Diagnostic information was obtained by a face-to-face semi-structured interview using the Structured Clinical Interview for DSM-IV supplemented by reviews of hospital records and collateral information from hospital psychiatrists and general practitioners. The operational criteria symptom checklist was completed based on psychiatric case notes and interview data.

MRI imaging and analysis

All imaging data were collected on a Siemens Magnetom Verio 3T MRI scanner running Syngo MR B17 software (Siemens Healthcare, Erlangen, Germany). For each subject, whole-brain dMRI data were acquired using a single-shot spin-echo echo-planar imaging sequence with diffusion-encoding gradients applied in 56 directions ($b = 1000$ s/mm²) and six T2-weighted ($b = 0$ s/mm²) baseline scans. Fifty-five 2.5-mm-thick axial slices were acquired with a field-of-view of 240 mm and matrix 96×96 giving 2.5 mm isotropic voxels. In the same session, a 3D T1-weighted inversion recovery-prepared fast spoiled gradient-echo volume was acquired in the coronal plane with 160 contiguous slices and 1 mm isotropic voxel resolution.
Image processing and tractography is described in Supplementary Methods.

**Animal ethics**

All animal experiments were conducted in accordance with the UK Animals (Scientific Procedures) Act (1986). Shiverer mice (C3Fe.SWV-Mbpsh/i; 001428) were purchased from Jackson Laboratories. MbpShi/Shi; Rag2−/− homozygous males were crossed with MbpShi+/+; Rag2−/− heterozygous females to obtain 50% double homozygous pups. Der1 heterozygous animals and control littermates were obtained by crossing wild-type males with heterozygous females. The date of vaginal plug was observed as E0.5 and embryos timed accordingly. All mice were maintained on a C57BL/6J background on a reversed 12-h light cycle with ad libitum access to food and water.

Shiverer, Rag2, and Der1 mice were identified by genotyping from tail/ear-clips using the following primers (in 5′–3′ orientation):

- **Mbp_F** TCC CTG GTG GCA GCT ATG AGC AGA CAC TGA
- **Mbp_R** CCC CGT GGT AGG AAT ATT ACA TTA CCA GCT
- **Shi_F** AGG GGA TGG GGA GTC AGA AGT GAG GAA AGA
- **Shi_R** ATG TAT GTG TGT GTG TGC TTA TCT AGT GTA
- **Rag2_F** GGT CAT CCT TTG CAA CAC AG
- **Rag2_M** CAG CGC TCC TCC TGA TAC TC
- **Rag2_R** TGC ATT CCT AGA GCG TCC TT
- **Disc1_F** CCT GCA TCC ACA GAC GTG C
- **Disc1_R** CAG TAG TAA GAA AAG AGA CCA CCC CC
- **Disc1_M** ATA ACG GTC CTA AGG TAG CGA GC

**Mouse model of t(1;11) translocation**

This mouse line is described in [34]. Briefly, to generate a complementary model of the translocation that mimics its effects upon DISC1 expression, the endogenous mouse Disc1 locus was genetically engineered using Regeneron’s GEMM platform (VelociMouse®). The mouse Disc1 allele in C57BL/6 × 129/Sv ES cell clones was targeted by deleting ~98.5 kb of genomic DNA encompassing exons 9–13, thus removing the 3' half of the gene that is translocated to chromosome 11 in human t(1;11) carriers. This was replaced with ~114 kb of human chromosome 11 genomic DNA containing exons 4–7 of the DISC1FP1 gene, corresponding to the region of the gene that is translocated to chromosome 1 in t(1;11) carriers. The edited endogenous mouse Disc1 locus thus mimics the DISC1/DISC1FP1 gene fusion event on the derived chromosome 1 in t(1;11) carriers.

**CNPase single-oligodendrocyte tracing**

For CNPase immunofluorescence studies, female mice were transcardially perfused with 4% PFA at 6 weeks of age. Brains were cut into 100-µm coronal sections on a vibratome (Leica). Free-floating sections containing the medial prefrontal cortex were blocked and permeabilized with 10% normal goat serum in 0.25% Triton X-100/PBS and subjected to antigen retrieval in 0.05% Tween20/10 mM trisodium citrate (pH 6.0) at 95 °C for 20 min. Subsequently, sections were washed in PBS, blocked as above, and incubated with primary mouse anti-CNPase antibody (Atlas antibodies; 1:2000) in blocking buffer for 24 h at 4 °C. Thereafter, sections were washed in PBS, incubated with an Alexa Fluor 488-labeled secondary antibody raised in goat (Invitrogen; 1:2000) in blocking buffer for 24 h at 4 °C. Hoechst33258 (5 µg/ml) for 20 min to label nuclei. After final washes in PBS, sections were mounted on slides and z-stack images through the whole section were taken using a Leica SP8 confocal microscope. Images were analyzed blind to genotype by assigning random numbers to the animals and using the ImageJ simple neurite tracer plugin.

**Neonatal transplantation of OPCs into MbpShi/Shi, Rag2−/− mice**

Homzygous MbpShi/Shi; Rag2−/− pups between P1-P2 were anaesthetised with isoflurane and maintained on a heat-mat for the duration of the transplantation. Dissociated case and control-derived OPCs at day 51 (day 0 = iPSCs) were delivered (200,000 cells in 2 μl) using a Hamilton syringe (Hamilton Company) through the skull. Transplants were directed at the presumptive corpus callosum and surrounding cortex. Pups were transcardially perfused with 4% paraformaldehyde at 8- and 13-weeks post injection and brains collected for histology. Three to four mice were transplanted per line.

**Electron microscopy**

For electron microscopy, male and female mice at the age of 3 weeks were anaesthetised and intracardially perfused with 4% paraformaldehyde/2% glutaraldehyde in 0.1 M phosphate buffer. After overnight post fixation, brains were cut into 1-mm-thick coronal sections using a brain slicer. Sections between Bregma 1.0 and −1.0 were trimmed further.
to isolate the medial aspect of the corpus callosum in a 1 mm × 2 mm × 1 mm block (L × W × H). Tissue samples were fixed in perfusion buffer for 24 h and 1% OsO₄ in 0.1 M phosphate buffer for 30 min. Using an automated processor (Leica), samples were dehydrated in rising concentrations of Ethanol, incubated in propylene oxide and rising concentrations of resin (TAAB laboratories) in propylene oxide before being embedded in resin and hardened for 22 h at 60 °C. Ultrathin sections were produced and analysed with a TEM-1400 Plus (Joel).

**Induced pluripotent stem cell line maintenance**

All iPSC were derived from human donor dermal skin fibroblasts using integration-free episomal methods [58]. iPSC lines were maintained in Matrigel (BD Biosciences) coated plastic dishes in E8 medium (Life Technologies) at 37 °C and 5% CO₂.

**Karyotyping, pluripotency markers, and X-chromosome inactivation staining**

Standard G-banding chromosome analysis was performed to confirm chromosome number and gross genetic abnormalities over the course of this study. Pluripotency of case and control iPSC lines was confirmed by staining for SOX2, OCT3/4, and TRA1-60. X-chromosome inactivation in female iPSC lines was confirmed by staining for H3K27 trimethylation marks [59].

**OPC and oligodendrocyte generation**

Derivation of NPCs was performed as described [30, 60] (also described in Supplementary Methods). Briefly, NPCs were cultured in suspension as neurospheres in Advanced DMEM/F12 containing: 1% N-2 supplement, 1% B27 supplement, 1% Anti-anti, 1% GlutaMAX (all from Life Technologies), FGF2 20 ng/ml (PeproTech), PDGFα 20 ng/ml (PeproTech), purmorphamine 1 μM (Calbiochem), SAG 1 μM (Calbiochem), IGF-1 10 ng/ml (PeproTech), and Tri-iodothyronine (T3) 30 ng/ml (Sigma) for 6–12 weeks. Following papain dissociation of neurospheres (Worthington), cells were plated on 1/100 diluted Matrigel (BD Biosciences), 20 μg/ml Fibronectin (Sigma), and 10 μg/ml Laminin (Sigma) coated coverslips or plates at a density of 20,000–30,000 cells per 0.3 cm². Differentiation of oligodendrocytes was achieved by mitogen withdrawal (except for T3 and IGF-1).

**RNA sequencing**

Library preparation, sequencing, and analysis are described in detail in Supplementary methods. Sequencing reads will be made available via ArrayExpress or European Genome-Phenome Archive prior to publication.

**Quantitative real-time PCR (qRT-PCR)**

RNA was extracted from cell pellets using the RNeasy Mini kit (QIAGEN). Five hundred nanograms of RNA was used to prepare cDNA by random hexamer extension using MMuLV reverse transcriptase (ThermoFisher) according to the manufacturer’s instructions.

Gene specific primers were used in a 96-well plate format to quantify differences in cDNA levels using a BioRad CFX96 qPCR machine. ΔΔCt method was used to normalize and quantify relative fold changes in gene expression.

Primers used in this study were (in 5’–3’ orientation):

<table>
<thead>
<tr>
<th>Primer</th>
<th>Sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>DISC1_F</td>
<td>CCA GCC TTG CTT GAA GCC AAA A</td>
</tr>
<tr>
<td>DISC1_R</td>
<td>TGA GGA GTC CCT CCA GCC CT T C</td>
</tr>
<tr>
<td>GAL3ST1_F</td>
<td>CAA GAC CCG GAT CGT CAA TAC TAC</td>
</tr>
<tr>
<td>GAL3ST1_R</td>
<td>TGT CAT AGC CCA GGT CAA AGA</td>
</tr>
<tr>
<td>SLC8A3_F</td>
<td>CGT CAC CAT TGG TCTCAA AAAAA</td>
</tr>
<tr>
<td>SLC8A3_R</td>
<td>TGT CAT AGC CCA GGT CAA AGA</td>
</tr>
<tr>
<td>NKK2.2_F</td>
<td>GTC AGG GAC GGC AAA CCA T</td>
</tr>
<tr>
<td>NKK2.2_R</td>
<td>GCG CTG TAG GCA GAA AAG G</td>
</tr>
<tr>
<td>CTTNBP2_F</td>
<td>AAG AGC GTG GCA AGA ACA AG</td>
</tr>
<tr>
<td>CTTNBP2_R</td>
<td>TGG GCC TCC TCT ATG ACT TTG</td>
</tr>
<tr>
<td>ZNF804A_F</td>
<td>AGT GGC CCC ATG TAC AAA TCA</td>
</tr>
<tr>
<td>ZNF804A_R</td>
<td>CCA CAA CAA CTC GTT GGG AAA A T</td>
</tr>
<tr>
<td>GAPDH_F</td>
<td>GAG TCC ACT GGC GTC TTC AC</td>
</tr>
<tr>
<td>GAPDH_R</td>
<td>ATG ACG AAC ATG GGG GCA T</td>
</tr>
<tr>
<td>β-ACTIN_F</td>
<td>CAC CTC CCC TGT GTG GAC TTG GG</td>
</tr>
<tr>
<td>β-ACTIN_R</td>
<td>GTT ACA GGA AGT CCC TTG CCA TCC</td>
</tr>
</tbody>
</table>

**Immunocytochemistry**

All steps were performed at ambient temperature. Cells were fixed with 4% paraformaldehyde in PBS for 5 min, permeabilized with 0.1% Triton X-100 containing PBS (except for PDGFα and O4), blocked in 3% goat serum, and incubated with appropriate primary and secondary antibodies (Alexa Fluor conjugated; Life Technologies). Nuclei were counterstained with DAPI (Sigma) and coverslips were mounted on slides with FluorSave (Merek). Fields were selected based upon uniform DAPI staining and...
imaged in three channels. Antibodies; OLIG2 (1:200, Millipore), PDGFRα (1:500, Cell Signaling), O4 (1:500, R&D Systems), MBP (1:50, Abcam), GFAP (1:500, DAKO), GFAP (1:500, Cy3 conjugated; Sigma), CASPR (1:1000, Abcam), Sox2 (1:1000, Abcam), Oct3/4 (1:250, Santa Cruz), Nanog (1:100, R&D Systems), TRA1-60 (1:100, StemGent), hNu (1:300, Millipore), Human GFAP (1:1000, Biologic), Neurofilament-H (1:10,000, Biologic).

Proportion of mitotic cells was studied by labeling OPCs with 10 μM ethynyl deoxyuridine (EdU, Life Technologies) on day 3 for 24 h. Media was replaced completely the next day and cells cultured for an additional 48 h in the absence of EdU before fixation using 4% PFA (Sigma). For detection, cells were permeabilized using 0.1% saponin (Sigma) for 2 h and detected using the EdU detection kit (Life Technologies) according to manufacturer’s instructions.

**Image acquisition and analysis**

Images were acquired either using a Zeiss ObserverZ1 wide field or Zeiss LSM710 confocal microscope (Carl Zeiss Microimaging). Images were converted to TIFF format, corrected for brightness and contrast, and analyzed using ImageJ (NIH) or Adobe Photoshop (Adobe Inc.). Figures for preparation were assembled using Adobe Illustrator (Adobe Inc.).

**Cell morphometry**

Cellular area was measured in ImageJ using O4 stained images [61]. Images were thresholded, segmented, and individual cells outlined using the wand selection tool.

Measurement of internodal lengths was performed using Simple Neurite Tracer plugin in ImageJ. Electron microscopy measurements were made using the freehand line tool in ImageJ. Areas were calculated by the in-built area measurement function.

**Statistical analysis**

Statistical analysis of the data was performed with Prism (GraphPad software, USA). All tests were performed as two-tailed. Tractography data was analyzed by MCMC modeling [29].

For experiments involving iPSCs, each derivation was considered as n = 1 and data was collected from a minimum of three derivations for each line. Normality of data distribution was analyzed by D’Agostino and Shapiro–Wilk’s tests. Normally distributed data was analyzed using Student’s t test or by one-way ANOVA test with Holm–Sidak’s multiple comparison correction (for >2 groups). Welch’s correction was applied when standard deviations of the two groups differed from each other. Quantitative PCR data was analyzed using nonparametric tests such as Mann–Whitney or Kruskal–Wallis tests (for 2 and >2 groups, respectively).

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**Author contributions** This study was conceived by NAV, AMM, JKM, and SC. DHRB, AMM, SML, and MJ were involved in recruiting family members, analyzing medical records and obtaining clinical and parasitical data. CA, MEB, and SML analyzed MRI and tractography data. NAV, DM, DS, SKB, BTS, PAT, KB, MJ, PM, EG, MRL, DJAW undertook human stem cell culture including reprogramming, iPS propagation and oligodendrocyte lineage differentiation, characterization and functional assessment and/or contributed to design, analysis, and interpretation of data. OD and GEH analysed RNA sequencing data. KE designed the Dermo mouse model. NAV and SEM undertook animal studies and, with CFFC, data analysis, and interpretation. NAV, MJ, and SC wrote the paper. All authors approved the paper.

**Compliance with ethical standards**

**Conflict of interest** KE was a former employee at Sanofi. The other authors declare that they have no conflict of interest.

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Familial t(1;11) translocation is associated with disruption of white matter structural integrity and...