One-step Reprogramming of Human Fibroblasts into Oligodendrocyte-like Cells by SOX10, OLIG2, and NKX6.2

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https://doi.org/10.1016/j.stemcr.2021.03.001

SUMMARY

Limited access to human oligodendrocytes impairs better understanding of oligodendrocyte pathology in myelin diseases. Here, we describe a method to robustly convert human fibroblasts directly into oligodendrocyte-like cells (dc-hiOLs), which allows evaluation of remyelination-promoting compounds and disease modeling. Ectopic expression of SOX10, OLIG2, and NKX6.2 in human fibroblasts results in rapid generation of O4+ cells, which further differentiate into MBP+ mature oligodendrocyte-like cells within 16 days. dc-hiOLs undergo chromatin remodeling to express oligodendrocyte markers, ensheath axons, and nanofibers in vitro, respond to promyelination compound treatment, and recapitulate in vitro oligodendrogial pathologies associated with Pelizaeus-Merzbacher leukodystrophy related to PLP1 mutations. Furthermore, DNA methylome analysis provides evidence that the CpG methylation pattern significantly differs between dc-hiOLs derived from fibroblasts of young and old donors, indicating the maintenance of the source cells’ “age.” In summary, dc-hiOLs represent a reproducible technology that could contribute to personalized medicine in the field of myelin diseases.

INTRODUCTION

Oligodendrocytes (OLs), the myelinating cells of the central nervous system (CNS), form and maintain the myelin sheath. Contribution of OLs to axon potential propagation, and trophic and metabolic support of neurons underlines their multifunctional role in CNS and implicates OL dysfunctionality and myelin loss in a wide range of CNS diseases including leukodystrophies, inflammatory demyelinating, neurodegenerative, and psychiatric diseases as well as ischemic or traumatic insults (Franklin and Ffrench-Constant, 2017; Micu et al., 2018; Stangel et al., 2017).

In dys- and demyelinating diseases the myelin sheath is either not properly formed or destroyed (Van Der Knaap and Bugiani, 2017). Promotion of (re-)myelination is a promising treatment strategy; however, the development of new therapeutic methods is hampered by the limited availability of primary human oligodendrocytes. In the past few years, a number of protocols have been developed to generate myelinating oligodendrocytes from human induced pluripotent stem cells (iPSC) in two- and three-dimensional (3D) culture systems (Douvaras and Fossati, 2015; Garcia-Leon et al., 2018; Madhavan et al., 2018). We have recently shown that forced expression of the transcription factors SOX10, OLIG2, and NKX6.2 in human fibroblasts results in rapid generation of O4+ cells, which further differentiate into MBP+ mature oligodendrocyte-like cells within 16 days. dc-hiOLs undergo chromatin remodeling to express oligodendrocyte markers, ensheath axons, and nanofibers in vitro, respond to promyelination compound treatment, and recapitulate in vitro oligodendrogial pathologies associated with Pelizaeus-Merzbacher leukodystrophy related to PLP1 mutations. Furthermore, DNA methylome analysis provides evidence that the CpG methylation pattern significantly differs between dc-hiOLs derived from fibroblasts of young and old donors, indicating the maintenance of the source cells’ “age.” In summary, dc-hiOLs represent a reproducible technology that could contribute to personalized medicine in the field of myelin diseases.
Figure 1. Characterization of dc-hiOLs
(A) Schematic representation of dc-hiOL generation. Fibroblasts are infected with SON lentivirus at day -1 in fibroblast medium (FM). Cells are cultured in glia induction medium (GIM) from day 0 to day 9, after which the medium is switched to glia differentiation medium (GDM) to promote terminal differentiation of oligodendrocyte-like cells.

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experimental models. A limited number of studies have presented transdifferentiation of murine embryonic fibroblasts into oligodendrocytes using different combinations of transcription factors (Matjusaitis et al., 2019; Najm et al., 2013; Yang et al., 2013). Very recently, Pouya et al. (2020) reported direct conversion of human oligodendrocytes but with limited characterization of the induced cells.

Here, we present a robust and highly reproducible protocol for direct conversion of human fibroblasts into oligodendrocyte-like cells. We found that overexpression of SOX10, OLIG2, and NKX6.2, previously shown adequately to differentiate iPSC-derived neural progenitor cells (NPCs) to OLs, is also sufficient to directly reprogram young, adult, and old-aged fibroblasts to directly converted human induced oligodendrocyte-like cells (dc-hiOLs) within 16 days. dc-hiOLs undergo chromatin remodeling to induce key oligodendrocyte markers, display ensheathment capacity in vitro, respond to promyelination compound treatment, and recapitulate oligodendrogial pathologies associated with Pelizaeus-Merzbacher leukodystrophy due to PLP1 mutation. Additionally, DNA methylation profiling indicates that dc-hiOLs retain the epigenetic age signature of donor cells. The technique described here significantly facilitates and advances compound screening and disease-modeling studies as well as the possibility of personalized medicine in the field of dys- and demyelinating diseases.

**RESULTS**

**Overexpression of SOX10, OLIG2, and NKX6.2 Directly Reprograms Human Fibroblasts into Oligodendrocyte-like Cells**

To develop a time- and cost-efficient method to generate human oligodendrocytes, we aimed to establish a transdifferentiation approach. Based on our previous observation that ectopic expression of SON in human iPSC-derived NPCs promotes their differentiation into oligodendrocytes, we reasoned that the same factors may directly convert fibroblasts into oligodendrocytes (Ehrlich et al., 2017). Overexpression of SON using a polycistronic lentiviral vector in three different human fetal/newborn fibroblast lines of diverse tissue origin (skin and lung) (Fib A, Fib B, Fib C) resulted in morphological changes as early as 9 days post infection. The cells lost their typical fibroblast spindle-like morphology and acquired a rounder shape followed by progressive extension of branches upon medium enrichment with differentiation factors including NT3, insulin-like growth factor 1 (IGF-1), and dibutyryl cyclic AMP (dbcAMP) for 7 days (Figures 1A and S1A). To further characterize the identity of the cells, we performed flow-cytometry and immunocytochemistry (ICC) analyses using the immature oligodendrogial marker O4. O4+ dc-hiOLs were already detected by post-infection day 8 (Figure 1B). Their percentage increased progressively until day 16, reaching an average conversion efficiency of

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(B) Quantification of O4+ cells using flow-cytometry analysis at different time points during the transdifferentiation process for three fetal/neonatal fibroblast cell lines (Fib A, Fib B, and Fib C). Data are shown as the mean ± SEM of n = 4 different transdifferentiation experiments for each cell line.

(C) Comparison of donor cell age effect on transdifferentiation efficiency. Quantification of O4+ cells by fluorescence-activated cell-sorting analysis at day 16. Data are presented as mean of n = 3 different cell lines for fibroblasts from newborn to 5 months of age, adult (25–32 years of age), and older (66–71 years of age) donors.

(D) Immunostaining of dc-hiOLs for O4 (red)/Ki67 (green) and O4 (red)/MBP (green) at days 12 and 16 post infection. Scale bar, 50 μm.

(E) Quantification of Ki67+ over O4+ cells at days 12 and 16. Data are presented as the mean ± SEM of n = 4 different transdifferentiation experiments of one cell line. Unpaired Student’s t test was performed for statistical analysis.

(F) Quantification of MBP+ over O4+ cells at days 12 and 16. Data are presented as the mean ± SEM of n = 4 different transdifferentiation experiments of one cell line. Unpaired Student’s t test was performed for statistical analysis.

(G) Migration assay using live cell imaging. Graph showing the velocity of purified O4+ cells at days 12 and 14 of direct conversion. Data are presented as the mean ± SEM of n = 3 different transdifferentiation experiments of one cell line. Unpaired Student’s t test was performed for statistical analysis.

(H) Immunostaining of purified O4+ dc-hiOLs for GalC, CNPase, MBP, and PLP1 (green) at day 16. Scale bar, 50 μm.

(I) Low- and high-magnification confocal images of dc-hiOLs cultured on nanofibers. O4+ cells were purified at day 11 and cultured on FITC-labeled nanofibers for 10 days. Immunostaining of dc-hiOLs for MBP (magenta) shows the alignment of oligodendrocyte processes along the nanofibers (green). Scale bar, 100 μm.

(J) Quantification of MBP+ dc-hiOLs aligned to nanofibers generated from three different cell lines. Three inserts were counted per cell line.

(K) Three-dimensional reconstruction of confocal image of dc-hiOLs stained for MBP (magenta) on FITC-labeled nanofibers illustrates the ensheathment of nanofibers by dc-hiOLs. The clipped view on the rendered surfaces enables one to follow individual nanofibers covered by dc-hiOLs (arrows). Scale bar, 5 μm.

(L) Confocal image of dc-hiOLs co-cultured with iPSC-derived neurons for 6 days. The image illustrates the colocalization of MBP+ (magenta) oligodendrocyte process with neuronal process visualized by TUJ1 (green). Scale bar, 25 μm.

*p < 0.05, ***p < 0.001. See also Figures S1 and S2.
Figure 2. SON Induce Chromatin Remodeling to Activate the Expression of Oligodendrocyte Genes
(A) Global transcriptome comparison using RNA-seq. MA plot comparing gene expression values between dc-hiOLs and fibroblasts. Differentially expressed genes accounting for donor heterogeneity with FDR < 0.01 are color coded. Light-blue dots, dc-hiOL-upregulated genes (1,410); light-green dots, fibroblast-upregulated genes (1,418); dark-blue dots, representative oligodendrocyte marker genes; dark-green dots, representative fibroblast marker genes.

(B) Normalized gene expression values of typical oligodendrocyte progenitor cell (OPC), oligodendrocyte (OL), neuronal, astrocyte, microglia, and fibroblast genes in comparison with single-nuclei RNA-seq data from Jäkel et al. (2019). Fibroblasts (FibA–FibC), induced oligodendrocyte-like cells (dc-hiOLa–dc-hiOLc), and primary human oligodendrocytes (pOl1–pOL3) indicate samples of the present studies. Data from Jäkel et al. (2019) include different subclusters of neurons (Neuron1–Neuron5) and oligodendrocytes (Oligo1–Oligo6), and additional clusters of OPCs, committed OL precursors (COPs), immune oligodendroglia (imOLG), astrocytes (Astrocytes), pericytes (Pericytes), and immune cells (Macrophages and Microglia_Macrophages). Unsupervised hierarchical clustering shows that dc-hiOL1–3 cluster with the other oligodendroglial clusters, whereas fibroblasts (Fib1–Fib3) form a separate cluster. Unexpectedly, in the dataset published by Jäkel et al. (2019), MBP was expressed in comparable levels in oligodendrocytes and astrocytes. This finding might be an RNA-seq artifact.

(C) Global chromatin accessibility comparison using ATAC-seq. MA plot comparing accessibility at ATAC-seq peaks between dc-hiOLs and fibroblasts. Blue dots, peaks preferentially accessible in dc-hiOLs (18,052); light-green dots, fibroblast-accessible peaks (4,075).

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20.4% ± 11.3% and yield of 39.1% ± 13.6% (Figures 1B and S1B–S1D). Control fibroblast cultures infected with the backbone lentiviral vector without SON showed no morphological changes and did not generate any O4+ cells (Figure S1C). Noteworthy, we assessed the reproducibility of the protocol using additional fibroblast samples of various age groups and found no significant effect of donor cell age on the transdifferentiation efficiency (Figure 1C). We followed the kinetics of transdifferentiation by using double-staining for O4 and the cell-cycle marker Ki67 as well as myelin basic protein (MBP), a marker specific for mature oligodendrocytes. Between days 12 and 16 the population of proliferating O4+ cells reduced by 3-fold, whereas the number of MBP+/O4+ double-positive cells significantly increased, indicating downregulation of proliferation and rapid maturation of dc-hiOLs (Figures 1D–1F). In agreement, live imaging revealed decreased migratory velocity of O4+ cells over time (Figure 1G; Videos S1 and S2). By day 16, besides MBP, dc-hiOLs expressed additional mature oligodendrocyte markers including CNP, GALC, and PLP1 (Figure 1H). Almost no NPCs, neurons, or astrocytes were found using ICC for Nestin, TUJ1, and GFAP, demonstrating the specific transdifferentiation into the oligodendroglial lineage (Figure S1E).

dc-hiOLs Are Stable after Transgene Silencing and Exhibit Ensheathment Capacity In Vitro
To test the dependence of dc-hiOLs on transgene expression, we used an SON-internal ribosomal entry site (IRES)-red fluorescent protein (RFP) plasmid for lentiviral transduction of fibroblasts. By day 16, 28.6% of the cells expressed O4 but no longer RFP, demonstrating the maintenance of the oligodendroglial identity after transgene silencing (Figure S1F). Moreover, in support of oligodendroglial fate stability, we observed an upregulated expression of endogenous SOX10 and OLIG2 transcripts in cells with silenced transgenes (Figure S1G). Of note, when comparing the expression levels of endogenous SOX10 and OLIG2 in fibroblast derived dc-hiOLs and NPC-derived hiOLs, we observed higher expression of both genes in hiOLs. This finding may be attributed to significant expression levels of SOX10 and OLIG2 already in NPCs (Figure S1H).

We next analyzed the functional characteristics of the cells by performing in vitro myelination assays. O4+ dc-hiOLs were sorted at day 11 post transduction and replated on fluorescein isothiocyanate (FITC)-labeled nanofibers that mimic the structure of neuronal axons or directly on iPSC-derived neurons. After 10 days of culture with nanofibers, dc-hiOLs proceeded to terminal differentiation and exhibited marked morphological change by forming complex MBP+ processes aligning along (Figures 1I and 1J) and wrapping around nanofibers, as showcased by 3D reconstruction analysis (Figures 1K and S2A). Comparison of dc-hiOLs with iPSC-derived oligodendrocytes co-cultured with nanofibers revealed comparable morphology although with fewer but longer processes (Figures S2B–S2E). Consistently, co-culture of purified O4+ dc-hiOLs with human iPSC-derived neurons already revealed dc-hiOL capacity to recognize and ensheath neuronal processes after 6 days of co-culture (Figure 1L).

Fibroblasts Undergo Changes in Chromatin Accessibility to Induce the Oligodendrocyte-Related Transcriptional Program
To further characterize the induced cells, we analyzed the global gene expression profile of O4+ dc-hiOLs purified at day 16. The transcriptome of dc-hiOLs was analyzed by RNA sequencing (RNA-seq) and compared with that of the original fibroblasts and adult primary human oligodendrocytes from three non-related individuals. Pairwise differential gene expression analysis showed a clear upregulation of the oligodendrocyte transcriptional program as showcased by the high expression of key oligodendrocyte marker genes such as MOG, MAG, PLP1, and OLIG1 (Figure 2A). Importantly, this is comparable with that of primary human oligodendrocytes (Figure S3A). Unsupervised hierarchical clustering (Figure S3B) confirmed a high similarity of biological replicates within the same cell type (minimum replicate correlation coefficient $\rho = 0.89$) and showed an increased similarity between the signature of dc-hiOLs and primary oligodendrocytes compared with fibroblasts. The expression signature of dc-hiOL clusters together with that of its progenitor fibroblasts suggested a partially preserved fibroblast identity. However, gene

(D) Example of oligodendroglial markers genes upregulated upon SON overexpression. Upper half: normalized RNA-seq coverage across donors (reads per kilobase per million [RPKM]). Lower half: normalized ATAC-seq accessible fraction coverage across donors (RPKM). Blue tracks, dc-hiOL signal; green tracks, fibroblast signal. Coordinates: ‘chr19:35,289,212-35,305,038’ (MAG), ‘chr6:29,642,061-29,674,746’ (MOG), ‘chr18:76,970,725-77,144,324’ (MBP), ‘chrX:103,772,927-103,797,741’ (PLP1).

(E) Ranked TF motif enrichment within top ATAC-seq peaks as assessed by HOMER (Heinz et al., 2010). Motifs with FDR < 0.05 are color coded. Circle color denotes TF family and circle size the percentage of peaks containing the motif. Inset trees quantify the similarity between obtained motifs (hierarchical clustering using Euclidean distance; bar indicates Euclidean distance of 1). Similar motifs (Pearson correlation coefficient >0.75) were merged into ensemble motifs. Original motifs are presented in Figure S3H. See also Figure S3.
samples and were clearly separated from fibroblasts dendrocyte progenitor cells isolated from human brain with primary pOls as well as oligodendrocytes and oligo-neuronal, and fibroblast genes, dc-hiOLs clustered together comparing the expression levels of representative glial, hiOLs (Figures 2C and S3F). Higher incidence of ATAC-seq ATAC-seq peaks between fibroblasts and the derived dc-
differentiation the cells undergo marked chromatin rearrange-
ment as denoted by the analysis of differentially accessible

data was confirmed by qPCR analysis (Figures S3D and S3E). Importantly, we compared our RNA-seq data of dc-
hiOLs and primary oligodendrocytes (pOls) with a previ-
sely published single-nuclei RNA-seq dataset from white
matter areas of human brain (Jäkel et al., 2019). When

We next utilized an assay for transposable-accessible chromatin using sequencing (ATAC-seq) to interrogate changes in chromatin accessibility induced by SON. During transdiff-

genomes. Among the top enriched TF motifs in dc-hiOLs were several mem-
bers of the SOX family that are crucial for oligodendroglial
cell fate and/or regulation of oligodendroglial differen-
tiation, such as SOX2 and SOX3 (Hoffmann et al., 2014) (Figures

Distribution of peaks of open chromatin regions were identified within the regulatory regions in core oligo-
dendrocyte markers concomitantly with their increased expression in dc-hiOLs (Figure 2D). To gain more insights into the mechanism of reprogramming, we analyzed transcrip-
tion factor (TF) binding sites (false discovery rate [FDR] < 0.05) within the open chromatin regions. Among

Test of Promyelination Compounds on dc-hiOLs
Promotion of endogenous remyelination is a key target for therapeutic intervention in multiple sclerosis, dysmyeli-

Starting at day 5 of the protocol, cultures were treated with 1 μM of the aforementioned compounds for 11 days (Figure 3A). Vehicle (DMSO) and T3 treatment were used as negative and positive control, respectively, and assay validation was performed by immunostaining for O4 and MBP. None of the compounds had a positive effect on the number of immature O4+ cells (Figures 3B and 3C). Benz-
trope and clemastine enhanced the terminal differentiation and maturation of dc-hiOLs to MBP+ OLs to a similar extent as T3, while miconazole showed a milder but still significant effect (Figures 3B and 3D). These findings suggest that dc-hiOLs can serve as a suitable system for rapid testing of promyelination candidates.

dc-hiOLs Derived from PMD Patients Exhibit Disease-Related Features
We then sought to test whether directly converted OLs from patients with leukodystrophies display pathogenic pheno-
types enabling in vitro disease modeling. We focused on Pel-
izeus-Merzbacher disease (PMD), a rare monogenic disor-
der caused by distinct types of PLP1 mutations that is characterized by dysmyelination and broad clinical severity. To assess whether dc-hiOLs can recapitulate the phenotypic heterogeneity of PMD, we analyzed the pathol-
ogy of OLs derived from a patient with a severe PMD (form 0) related to a PLP1 missense mutation (cPLP1 643C>T, p.Pro215Ser) as well as from three patients with intermediate forms of PMD (form 2) related to PLP1 duplication (PLP1 dupl.). Diseased cells were compared with sex- and age-
matched controls. As demonstrated by the quantification of O4+ and MBP+ cells, PLP1 643C>T dc-hiOLs, but not PLP1 dupl., was associated with impaired terminal differen-
tiation to MBP-expressing cells (Figures 4A–4D, S4A, and S4B). Previous works associated defective myelination of OLs with point mutations in PLP1 with ER stress and vulner-
ability to cell death (Nevin et al., 2017; Numasawa-Kuroiwa et al., 2014). Accordingly, immunostaining for PLP1 and apoptosis assay at day 13 showed strong perinuclear accumu-
lation of PLP1, indicative of ER stress, and significantly higher number of PLP1 643C>T dc-hiOLs with active caspase-3/7 compared with control cells, respectively (Figures 4E–4G). On the contrary, PLP1 dupl. led to a milder pheno-
type with preserved PLP1 expression throughout the cyto-

Related Features
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dc-hiOLs Maintain the Epigenetic Age of Donor Cells

A fundamental challenge in using reprogrammed cells to study age-related traits is the preservation of donor cells’ “age.” While iPSC reprogramming is accompanied by cell rejuvenation and erasure of aging signature (Lapasset et al., 2011; Meissner et al., 2008), directly converted cells maintain many features of aging (Huh et al., 2016; Mertens et al., 2015). The strong correlation between cellular age and DNA methylation (Fraga and Esteller, 2007) prompted us to study whether dc-hiOLs retain the epigenetic age signature of their cells of origin.

DNA methylation patterns of dc-hiOLs and corresponding fibroblasts were analyzed across three donor age groups: neonatal (between newborn and 5 months of age), adult (between 25 and 32 years of age), and old age (between 66 and 71 years of age). We first analyzed differentially methylated CpG sites in key oligodendrocyte genes. Comparing methylation β values between dc-hiOLs and fibroblasts shows significant hypomethylation of CpGs mapping to MBP, whereas there were no differences in methylation levels of MAG and MOG (Figure 5A) although they are highly expressed in dc-hiOLs, suggesting additional transcriptional regulatory mechanisms. Unsupervised analysis of differentially methylated regions between dc-hiOL and fibroblast samples confirmed a significantly hypomethylated region of 17 CpG sites mapping to exon 1 of MBP transcripts typically expressed in the brain (Figure 5B, HMM-Fisher, p = 1 × 10^-19). We observed a high epigenetic divergence of adult/old samples as compared with neonatal samples (adult/old versus neonatal: 57,106 differentially methylated CpG sites, adjusted p < 0.05). Analysis of the top 10,000 most variable CpG sites of all 57,106 age-associated CpGs showed that the age signature of dc-hiOLs is maintained throughout transdifferentiation and reflects the age of donor cells (Figure 5C). Cells from adult and old-aged donors rather display global DNA methylation patterns similar to those shown on unsupervised t-distributed stochastic neighbor embedding (t-SNE) analysis (Figure 4D), indicating that age-related epigenetic changes take already place in early adulthood. Gene ontology term analysis of the differentially methylated CpGs between neonatal and adult/old-aged samples revealed many developmental annotations as well as other processes associated with transcriptional regulation and post-translational modifications (Table S1). In line with t-SNE dimensionality reduction, unsupervised hierarchical clustering of the top 5,000 most variable CpGs clearly separated neonatal samples from dc-hiOLs derived from adult and old-aged donors (Figure 5SA). We applied Horvath’s epigenetic clock (Horvath, 2013) as age predictor to our
Figure 4. dc-hiOLs Derived from PMD Patients Recapitulate Disease-Related Pathologies

Characterization of dc-hiOLs derived from PMD patients. dc-hiOLs derived from one PMD patient with a point mutation (PLP1 643C>T) and three PMD patients with PLP1 duplication (PLP1 dupl.) were compared with sex- and aged-matched healthy controls, respectively.

(A and B) Representative immunofluorescence images of O4+ (red), MBP+ (green), and PLP1+ (green) dc-hiOLs at day 16 of transdifferentiation. Scale bars, 50 μm.

(C–H) Comparison of dc-hiOLs derived from one PMD patient with point mutation PLP1 643C>T with one control cell line. Percentage of O4+ (C) and MBP+ (D) cells at day 16. Data are shown as the mean ± SEM of n = 3 different transdifferentiation experiments of one cell line for each group. Two-tailed unpaired t test was used for statistical analysis. (E) Representative plot profiles comparing the distribution of PLP1 in control (black) and PLP1 643C>T (green) dc-hiOLs. Two-dimensional graph of pixel intensity along cell diameter. (F) Apoptosis assay for PLP1 643C>T dc-hiOLs. MR-(DEVD)2 reagent is cleaved by active caspase-3/7 and produces a fluorescent product (magenta). ICC for O4+

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methylated dataset and confirmed the distinction of fibroblasts and the derived dc-hiOls in separate age groups (Figure S5B). Overall, our results show that dc-hiOls preserve the epigenetic age of donor cells, at least with respect to DNA methylation profile, and represent a promising model to study age-associated myelin deficits.

**DISCUSSION**

Limited access to human oligodendrocytes has been an obstacle in biomedical research of myelin disorders. Differentiation of human iPSCs into oligodendrocytes enables large-scale production of cells and provides a valuable tool in clinical research. However, this technology faces the challenges of high-cost, complex, and time-consuming cell-culture techniques and genetic instability (Popp et al., 2018). Our results suggest that direct conversion of fibroblasts into OLs is a rapid and inexpensive technique, which represents a complementary tool to study myelin diseases and identify remyelination-promoting compounds.

Ectopic expression of SON directly transdifferentiates human fibroblasts within 16 days into O4+ dc-hiOls, which exhibit oligodendroglial properties. Although the efficiency varies between fibroblast lines of different tissue origin, in the vast majority of the lines the efficiency is higher than in earlier protocols using mouse cells (Najm et al., 2013; Yang et al., 2013), which may be explained by the polycistronic construct used here and the thereby increased transduction efficiency as shown recently for directly converted neurons (Herdy et al., 2019). Notably, the fibroblast line that showed the lowest transdifferentiation efficiency was derived from lung tissue (Fib C, Figure 1C), in contrast to all other lines which originated from skin, suggesting that the source tissue might affect conversion efficiency; however, studies examining systematically the influence of tissue origin of fibroblasts on the transdifferentiation efficiency are currently lacking.

Gene expression and chromatin analysis revealed a chromatin switch that coincides with activation of the oligodendrocyte transcriptional program during reprogramming. Several members of the SOX TF family are among the top enriched TF motifs in induced oligodendrocytes. SOX10 is critical for the determination of oligodendroglial fate (Garcia-Leon et al., 2018), whereas SOX2 and SOX3 have been shown to play a redundant role in promoting oligodendrocyte differentiation partially through downregulation of miR-145 (Hoffmann et al., 2014). Furthermore, in spinal cord precursor cells SOX3 suppresses astrocytic genes, whereas SOX9 and SOX10 promote the differentiation into oligodendrocytes (Klum et al., 2018). We observed remaining activity of fibroblast TFs including the AP1 family, which may explain the incomplete silencing of the fibroblast program. However, we showed that the acquired oligodendroglial fate is stable and that cells do not revert back to fibroblast fate even upon transgenic silencing. Inhibition of AP1 activity and use of small molecules that regulate the epigenetic landscape of the donor cells might further increase the efficiency of oligodendroglial transdifferentiation, as has been recently shown for directly converted neurons (Herdy et al., 2019).

O4+ dc-hiOls can be easily purified and used for drug screens and testing of promyelinating compounds. We demonstrated that clemastine, benzotropine, and miconazole increase significantly the terminal differentiation of dc-hiOls into MBP+ cells. Since fibroblasts differentiate rapidly into O4+ dc-hiOls, dc-hiOls might be less suitable to identify compounds that have a positive effect on early oligodendroglial differentiation stages.

Reprogramming technology provides a unique opportunity to generate scarcely accessible cell types and study related disorders. As demonstrated here, dc-hiOls can serve as a system for disease modeling of rare demyelinating diseases. PMD is a monogenic X-linked leukodystrophy related to PLP1 mutations. PLP1-related disorders are characterized by genetic heterogeneity reflected in a wide spectrum of pathology, ranging from absence of motor acquisition with early-age lethality (PMD form 0) to walking capacities with normal life expectancy (form 4 or SPG2) (Cailloux et al., 2000; Osorio and Goldman, 2018). We found that dc-hiOls bearing the PLP1 643C>T mutation suffer from ER stress and exhibit impaired terminal differentiation to myelinogenic MBP+ cells, whereas PLP1 duplication develops a milder phenotype. Our data confirm the results of prior publications studying iPSC-derived OLs from patients with PMD and provide evidence that dc-hiOls are suitable for disease modeling and for the identification of correlations between genotypes and phenotypes (Madhavan et al., 2018; Nevin et al., 2017; Numasawa-Kuroiwa et al., 2014). The rapidity of the described protocol enables

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* indicates p < 0.05, ** indicates p < 0.01. See also Figure S4.
generation of dc-hiOLs from numerous patients at low cost, providing a platform for studying additional myelin
or CNS diseases and the identification of clinical sub-
groups. dc-hiOLs might have the potential to facilitate
the development of personalized medicine by bypassing
the relative expensive and time-consuming generation
of iPSCs. A key feature of dc-hiOLs, as revealed by CpG
methylation analysis, is that cells circumvent rejuvena-
tion and retain the epigenetic age of donor cells. Aging
is a major risk for many CNS diseases, and our data
suggest the applicability of dc-hiOLs in studies of age-
related myelin alterations. However, assessment of addi-
tional age-related features such as mitochondrial and
telomerase activity needs to be performed for future use
of dc-hiOLs in aging studies. A limitation of the available
transdifferentiation methods, including ours, is the
limited yield of produced cells due to the lack of an inter-
mediate stem cell type, restricting the conduction of large-
scale assays. An alternative strategy for the generation of
human OLs is represented by the reprogramming of

Figure 5. Epigenetic Characterization of Fibroblast and dc-hiOL Samples
(A) Focused analysis of methylation β values of CpG sites mapping to MBP, MAG, and MOG genes showing that only MBP is differentially
methylated on the global gene level between dciOL and fibroblast samples (p < 0.01, Wilcoxon’s test).
(B) Unsupervised analysis of differentially methylated regions between dciOL and fibroblast samples revealed a region of 17 CpG sites
mapping to exon 1 (hg19 coordinates: chr18:74,728,834-74,729,551) of MBP transcripts typically expressed in the brain being hypo-
methylated in dc-hiOL samples as compared with fibroblast samples (HMM-Fisher, p = 1 × 10^-19).
(C) Heatmap showing methylation β values of the top 1,000 differentially methylated probes between aged samples and neonatal samples.
Note the shared aging signature between fibroblast and dc-hiOL samples across neonatal and aged samples.
(D) Unsupervised t-distributed stochastic neighbor embedding (t-SNE) analysis of all dc-hiOL samples showing that aged and old samples
group together and are clearly separated from neonatal samples.
See also Figure S5.
fibroblasts into induced neural progenitor cells (iNPCs) and subsequent differentiation into oligodendrocytes (Meyer et al., 2014). Additional studies comparing side-by-side oligodendrocytes generated from fibroblasts, iNPCs, or iPSCs will help to further evaluate the advantages and disadvantages of the different techniques. Of note, although the in vitro data provided here support the oligodendroglial identity of dc-hiOLs, further analysis of their capacity to form compact myelin sheaths in vivo is needed.

In summary, direct conversion of fibroblasts into dc-hiOLs represents a rapid, efficient, and inexpensive alternative to iPSC-derived human oligodendrocytes for disease modeling and the identification of remyelination-promoting compounds, and advances significantly the possibility of personalized medicine in the field of de- and dysmyelinating diseases.

**EXPERIMENTAL PROCEDURES**

**Plasmids and Lentiviral Preparation**

The construction of the polycistronic lentiviral SF-SON-puro and SF-SON-RFP vectors has been previously described (Ehrlich et al., 2017). Detailed description of the plasmids and the lentiviral production can be found in supplemental experimental procedures.

**Cell Lines and Culture**

A summary of the cell lines used in this study is provided in Table S2. Detailed description can be found in supplemental experimental procedures.

**Direct Conversion of Fibroblasts to dc-hiOLs**

Fibroblasts were seeded at a density of 12 × 10³ cells/cm² on a poly-L-lysine (Sigma)/mouse laminin (Sigma) coated plate in fibroblast medium (FM). The next day cells were transduced with the SF-SON lentivirus in fresh FM, and 16 h later the medium was changed to glial induction medium (GIM), which consisted of DMEM-F12 (Gibco), 1:200 N2 supplement (Gibco), 1:100 B27 supplement lacking vitamin A (Gibco), and 1% penicillin/streptomycin/glutamine enriched by 0.5 mM Smoothened agonist (SAG) (Cayman Chemical), 10 ng/mL platelet-derived growth factor (PDGF) (Peprotech), 200 μM ascorbic acid (AA) (Sigma), and 20 ng/mL T3 (Sigma). GIM was changed every other day and the concentration of T3 was increased to 60 ng/mL at day 4. Puromycin (1 μg/mL, Gibco) selection of the transduced cells took place for the first 5 days. At day 9 the GIM was replaced by glial differentiation medium (GDM) consisting of DMEM-F12, 1:200 N2 supplement, 1:100 B27 supplement lacking vitamin A, 1% penicillin/streptomycin/glutamine enriched by 0.5 μM SAG, 60 ng/mL T3, 10 ng/mL NT3 (Peprotech), 10 ng/mL IGF-1 (Peprotech), 200 μM AA, and 10 ng/mL PDGF. PDGF was replaced by 100 μM dbcAMP (Sigma) at day 12. Medium change was performed every other day. Cells of low passage (fewer than 10) were used for all direct reprogramming experiments. Conversion efficiency is defined as the number of O4⁺ cells divided by the total number of 4’,6-diamidino-2-phenylindole (DAPI)-positive cells at day 16. Yield is defined as the number of O4⁺ cells at day 16 divided by the number of plated fibroblasts at day 0.

**Immunocytochemistry**

Cells were fixed using 4% paraformaldehyde (PFA) for 20 min and permeabilized with 0.5% Triton X-100 solution for 10 min. For O4 no cell permeabilization was performed. After three PBS washes, cells were blocked for 30 min with 5% normal goat serum and 5% fetal bovine serum in PBS. Primary antibodies were added in blocking buffer overnight at 4°C. Upon three washes, the secondary antibodies were added in PBS at room temperature for 2 h. Nuclei were stained with 1 μg/mL DAPI (Sigma-Aldrich). Cells on glass coverslips were visualized on a Zeiss LSM700 confocal microscope, while cells cultured on a plastic surface were visualized on a Leica DMi6000 B inverted microscope. The primary antibodies are listed in Table S3. Anti-PLP1 antibody was generously provided by Prof. Bruce Trapp.

**Flow Cytometry**

Details of flow cytometry are provided in supplemental experimental procedures.

**Migration Assay**

The methodology of migration assay is described in supplemental experimental procedures.

**Isolation of Primary Human Oligodendrocytes**

Brain tissue was obtained from adults undergoing surgical resections as treatment for non-tumor-related intractable epilepsy in accordance with the guidelines set by the Biomedical Ethics Unit of McGill University. Details are provided in supplemental experimental procedures.

**Quantitative RT-PCR**

Total RNA was extracted from cells using the RNeasy Mini Kit (Qiagen). DNase (Qiagen) treatment was used to avoid genomic DNA contamination, and RNA was reversely transcribed to cDNA using the High Capacity cDNA Reverse Transcription Kit (Applied Biosystems). qRT-PCR was carried out using SYBR Green-based detection and gene expression was normalized to GAPDH. The primers can be found in Table S3.

**RNA-seq, ATAC-seq, and DNA Methylation Array Analysis**

The methodology of RNA-seq, ATAC-seq, and DNA methylation analysis is described in supplemental experimental procedures.

**In Vitro Myelination Assays**

Nanofiber chamber slides (PCL NanoAligned) were coated with FITC-conjugated poly-L-lysine (Sigma) for 2 h at 37°C followed by two washes with sterile H2O and overnight incubation with mouse laminin (Sigma) at 4°C. Twenty-four hours before cell plating, nanofibers were incubated with GDM at 37°C. O4-expressing dc-hiOLs were purified at day 11 of the transdifferentiation process and replated on the nanofiber slides (60,000 cells/slide).
in GDM. Half-medium change was performed every other day and cells were analyzed with ICC after 10 days. A published heuristics algorithm was used to analyze the morphology of dc-hiOLs and iPSC-derived hiOLs on nanofibers (Xu et al., 2019). Three-dimensional reconstruction of confocal image stacks was performed with IMARIS analysis software (Oxford Instruments). 3D surface rendering was applied.

For the neuronal/dc-hiOL co-culture experiments, neurons were generated from human iPSC-derived NPC as previously described (Ehrlich et al., 2017) and replated at densities of 2 × 10^3 per well in 24-well plates containing Matrigel-coated (BD Biosciences) glass coverslips. After 22 days of differentiation, neurons were used for co-culture assays. dc-hiOLs purified for O4 at day 11 of transdifferentiation were added to 22-day-old neurons at densities of 9 × 10^3 cells per well in 24-well plates. Co-cultures were maintained in GDM supplemented with 2 ng/mL brain-derived neurotrophic factor (Peprotech) and 2 ng/mL glial cell line-derived neurotrophic factor (Peprotech). Following 6 days of co-culture, cells were fixed in 4% PFA for ICC analysis.

**Compound Treatment**
Methodological details are provided in supplemental experimental procedures.

**Apoptosis Assay**
A Magic Red Caspase3/7 Assay Kit (Bio-Rad) was used for measurement of caspase activity. Further details are provided in supplemental experimental procedures.

**Statistics**
Data of at least three biological replicates (n) are presented as the mean ± SEM. Statistical significance was determined by Student's t test. Significance was set at *p < 0.05, **p < 0.01, and ***p < 0.001.

**Data and Code availability**
The accession number for the RNA-seq data reported in this paper is ArrayExpress: E-MTAB-7838. The RNA-seq data and will be made publicly available upon publication. The R script for the comparison of our RNA-seq data with the published dataset by Jaékel et al. is ArrayExpress: E-MTAB-7838. The RNA-seq data and will be made publicly available upon publication. The R script for the comparison of our RNA-seq data with the published dataset by Jaékel et al. is available on github (https://github.com/ctho1/compareRNAseq).

**SUPPLEMENTAL INFORMATION**
Supplemental Information can be found online at https://doi.org/10.1016/j.stemcr.2021.03.001.

**AUTHOR CONTRIBUTIONS**
T.K., K.C., and M.E. conceived the study; B.H.-R. and J.M.V. designed and analyzed the RNA-seq and ATAC-seq experiments; C.T. performed biostatistical analyses; K.C., F.W., S.A., M.S., S.V., Q.L.C., and M.E. performed the in vitro experiments; K.C., S.M., A.B.-V.E., and Y.K.T.X. analyzed data; O.B.-T. provided the PMD duplicated fibroblasts and clinical expertise; J.A., G.M., L.O., J.W., and K.-P.K. provided human cells/cell lines; K.C. and T.K. drafted the manuscript; B.H.-R., L.O., H.R.S., A.B.-V.E., O.B.-T., J.M.V., and M.E. were involved in editing and discussion.

**CONFLICTS OF INTEREST**
T.K. and M.E. have filed a patent for the generation of human oligodendrocytes. T.K. received compensation for serving on scientific advisory boards (Frequency Therapeutics, Inc.) and speaker honoraria and research funding from Novartis.

**ACKNOWLEDGMENTS**
We thank Dr. Thomas Zobel from the Imaging Network, Münster for helping with the 3D reconstruction analysis and Dr. Laura Starost for providing hiOLs. This work was supported by the German Research Foundation (DFG CRC-TR-128B07 to T.K.), Interdisciplinary Center for Clinical Research (KiTZ/007/20 to T.K.), Progressive MS Alliance (collaborative research network PA-1604-08492 (BRAVEinMS) to G.M., J.A., A.B.-V.E., and T.K.), AFM-TELETHON 20270 (to A.B.-V.E.), the National MS Society (RG-1801-30020 to T.K.), and the EC FP7 project LEUKOTREAT and ‘les amis de Ianis’ association (to O.B.-T.). Work in the Vaquerizas laboratory was funded by the Max Planck Society, the Deutsche Forschungsgemeinschaft (DFG) Priority Program SPP 2202 ‘Spatial Genome Architecture in Development and Disease’ (project no. 422857230 to J.M.V.), the DFG Clinical Research Unit CRU326 ‘Male Germ Cells: from Genes to Function’ (project no. 329621271 to J.M.V.), the European Union’s Horizon 2020 research and innovation program under the Marie Skłodowska-Curie grant agreement no. 643062—ZENCODE-ITN to J.M.V., and the Medical Research Council, UK.

Received: March 22, 2020
Revised: March 1, 2021
Accepted: March 1, 2021
Published: March 25, 2021

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