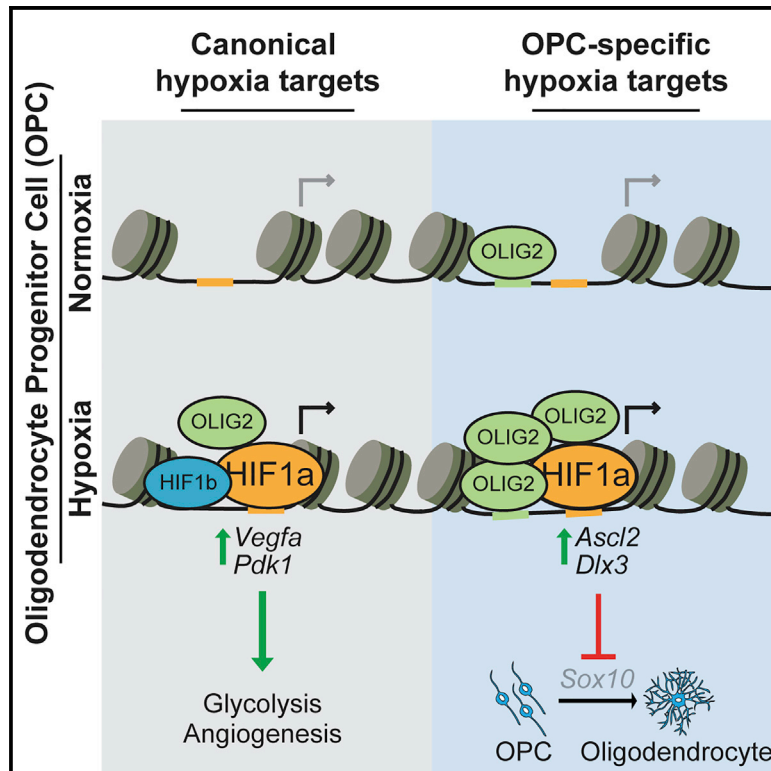# Non-canonical Targets of HIF1a Impair Oligodendrocyte Progenitor Cell Function

Graphical Abstract

Authors

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In Brief

Hypoxia impairs the generation of oligodendrocytes and myelin in various neurological disorders. Allan et al. map the genome-wide binding profile of HIF1a in oligodendrocyte progenitor cells (OPCs). HIF1a activated non-canonical, cell-type-specific target genes in OPCs that converge to suppress *Sox10* expression and block oligodendrocyte formation.

Highlights

- HIF1a activates both canonical hypoxia response and unique OPC-specific genes

- OLIG2 physically interacts with HIF1a and is enriched at OPC-specific HIF1a targets

- OPC-specific HIF1a targets suppress *Sox10* to impair oligodendrocyte formation

- MEK inhibition increases *Sox10* expression to rescue OPC function from hypoxia

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Non-canonical Targets of HIF1a Impair Oligodendrocyte Progenitor Cell Function

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SUMMARY

Mammalian cells respond to insufficient oxygen through transcriptional regulators called hypoxia-inducible factors (HIFs). Although transiently protective, prolonged HIF activity drives distinct pathological responses in different tissues. Using a model of chronic HIF1a accumulation in pluripotent-stem-cell-derived oligodendrocyte progenitors (OPCs), we demonstrate that HIF1a activates non-canonical targets to impair generation of oligodendrocytes from OPCs. HIF1a activated a unique set of genes in OPCs through interaction with the OPC-specific transcription factor OLIG2. Non-canonical targets, including *Ascl2* and *Dlx3*, were sufficient to block differentiation through suppression of the oligodendrocyte regulator *Sox10*. Chemical screening revealed that inhibition of MEK/ERK signaling overcame the HIF1a-mediated block in oligodendrocyte generation by restoring *Sox10* expression without affecting canonical HIF1a activity. MEK/ERK inhibition also drove oligodendrocyte formation in hypoxic regions of human oligocortical spheroids. This work defines mechanisms by which HIF1a impairs oligodendrocyte formation and establishes that cell-type-specific HIF1a targets perturb cell function in response to low oxygen.

INTRODUCTION

The ability to sense and respond to fluctuations in oxygen levels is required to maintain homeostasis in every cell in the body (Kaelin and Ratcliffe, 2008; Semenza, 2012). Insufficient concentrations of molecular oxygen rapidly trigger an evolutionary conserved transcriptional response that enables cell survival in low oxygen by promoting anaerobic metabolism for energy production as well as angiogenesis and erythropoiesis to increase access to local oxygen. Although this is initially protective, prolonged activation of this response leads to cellular dysfunction and disease in many tissues. For example, the response to chronic hypoxia blocks white matter formation in premature birth (Scafidi et al., 2014; Volpe, 2009; Volpe et al., 2011), promotes inflammation and insulin resistance in obesity (Lee et al., 2014), and impairs hematopoietic stem cell transplantation capacity (Takubo et al., 2010). This cellular dysfunction has been attributed largely to prolonged activation of the canonical response to low oxygen shared across all cell types; however, it is difficult to explain how activation of a conserved set of hypoxia signature genes can lead to such diverse cellular phenotypes. An alternative unexplored possibility is that cell-type-specific differences in chromatin landscape enable access to unique non-canonical targets, which could account for tissue-specific pathologies.

The response to low oxygen is mediated by hypoxia-inducible factors (HIFs), a family of transcription factors that are stabilized under hypoxic conditions in all mammalian cells and are thought to primarily upregulate multiple pathways to adapt cells to low oxygen (Cassavaugh and Lounsbury, 2011; Choudhry and Harris, 2018; Kupferschmidt, 2019). HIFs are heterodimeric complexes consisting of alpha and beta subunits. In the presence of oxygen, alpha subunits are hydroxylated by prolyl-hydroxylases, allowing recognition and ubiquitination by von Hippel Lindau (VHL) (Ivan et al., 2001; Jaakkola et al., 2001), and rapid degradation by the proteasome. In low oxygen conditions, alpha subunits escape hydroxylation, avoid degradation, and translocate to the nucleus to pair with constitutive beta subunits and regulate gene expression (Cassavaugh and Lounsbury, 2011; Choudhry and Harris, 2018; Semenza, 2007). The HIF1a motif is present more than 1 million times in the genome; however, HIF1a binds to a small fraction of these sites, suggesting that HIF1a binding is heavily regulated (Schödel et al., 2011; Smythies et al., 2019). Still, the determinants of HIF1a binding in each cell type and whether cell-type-specific targets are functional remain unknown.

The central nervous system (CNS) consumes 20% of the total body oxygen, and white matter of the CNS is highly susceptible to hypoxic insults, as seen in stroke, vascular dementia, respiratory distress syndromes, premature birth, and subsets of cerebral palsy (Hankey, 2017; Salmaso et al., 2014; Shindo et al., 2016; Volpe, 2009). In fact, chronic HIF1a activity is sufficient to block white matter development (Yuen et al., 2014). White
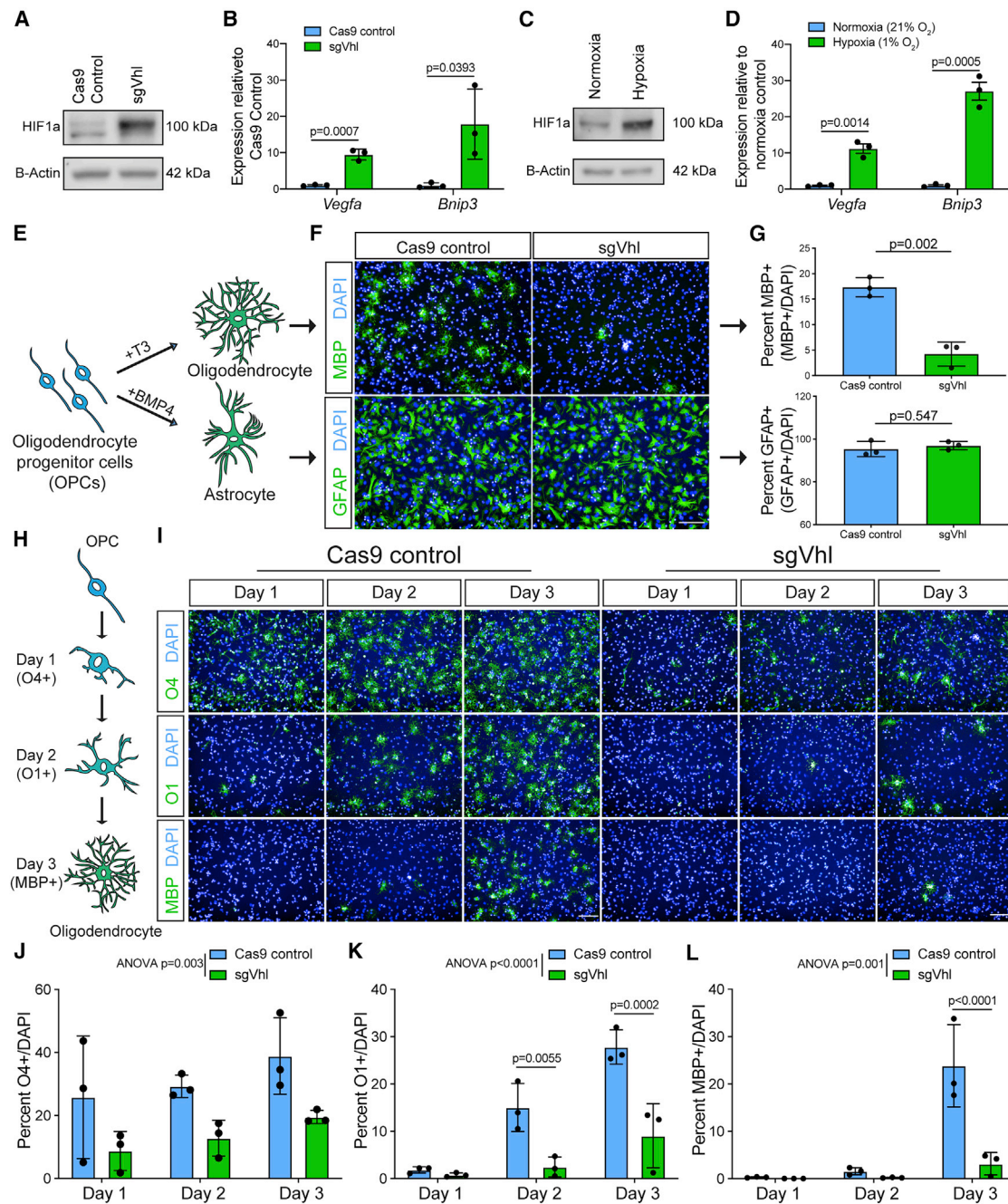
Figure 1. HIF1a Accumulation Impairs the Induction of Oligodendrocytes from OPCs

(A) Western blot of HIF1a in sgVhl and Cas9 control OPCs.

(B) qRT-PCR of *Vegfa* and *Bnip3* in Cas9 control (in blue) and sgVhl (in green) OPCs. Data are presented as mean ± SEM from three biological replicates.

(C) Western blot of HIF1a in hypoxic (1% O_2) and normoxic OPCs.

(D) qRT-PCR of *Vegfa* and *Bnip3* in hypoxic (in green) and normoxic (in blue) OPCs. Data are presented as mean ± SEM from three biological replicates.

(E) Schematic of the two *in vitro* differentiation schemes directing OPCs to either oligodendrocytes or astrocytes.

(F) Representative images of Cas9 control and sgVhl oligodendrocytes (MBP+ in green) and astrocytes (GFAP+ in green). Nuclei are marked by DAPI (in blue). Scale bar, 100 μm.

(G) Quantification of the percentage of oligodendrocytes (MBP+ cells/DAPI) and astrocytes (GFAP+ cells/DAPI) formed from sgVhl (in green) and Cas9 control OPCs (in blue). Data are presented as mean ± SD from three independent biological replicates. p values were calculated using Student's two-tailed t test.

(H) Schematic illustrating acquisition of early (O4), intermediate (O1), and late (MBP) oligodendrocyte markers during *in vitro* oligodendrocyte differentiation.

(I) Representative images of early (O4+, in green), intermediate (O1+, in green), and late (MBP+, in green) oligodendrocytes during days 1, 2, and 3 of differentiation of Cas9 control and sgVhl OPCs. Nuclei are marked by DAPI (in blue). Scale bars, 100 μm.

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matter of the CNS is formed by oligodendrocytes, which wrap neuronal axons in a lipid-rich protective sheath called myelin, allowing rapid transmission of action potentials and maintenance of axonal integrity (Chang et al., 2016; Emery, 2010; Nave, 2010). Oligodendrocytes arise from oligodendrocyte progenitor cells (OPCs), which are prevalent in the developing and adult CNS, and HIF1a accumulation has been shown to be sufficient to impair oligodendrocyte formation from OPCs (Jablonska et al., 2016; van Tilborg et al., 2018; Yuen et al., 2014). However, the mechanism of the HIF1a-mediated block in oligodendrocyte formation from OPCs remains unclear. In this study, we use OPCs as an archetypal hypoxia disease-relevant cell type to define the mechanism by which chronic HIF1a drives cell dysfunction.

RESULTS

Knockout of *Vhl* Models Chronic HIF1a Accumulation in OPCs

Defining the mechanisms by which HIF activity perturbs cell function is notoriously challenging, as HIFs are rapidly degraded, in minutes, when cells are restored to normoxia. Because of this instability and the low abundance of HIFs, biochemical studies often require extraordinary numbers of cells, which is challenging for hypoxia disease-relevant cell types outside of cancer cell lines. To explore the mechanisms underlying the HIF-mediated block in oligodendrocyte development from OPCs, we generated a cellular model of chronic HIF1a accumulation in mouse pluripotent stem cell-derived OPCs, which are uniquely scalable and amenable to genetic manipulation (Hubler et al., 2018; Lager et al., 2018; Najm et al., 2015; Najm et al., 2011). CRISPR-Cas9-mediated knockout of *Vhl*, a central component of the ubiquitin-proteasome system that degrades HIFs (Choudhry and Harris, 2018; Haase, 2009; Rechsteiner et al., 2011), in OPC cultures resulted in stable HIF1a protein accumulation and significant 9-fold and 18-fold activation of downstream hallmark HIF1a targets *Vegfa* and *Bnip3*, respectively, without affecting viability (Figures 1A, 1B, and S1A–S1C). The response of OPCs to *Vhl* knockout mirrored that of OPCs cultured in hypoxia at both the protein and RNA levels (Figures 1C, 1D, and S1D). *Vhl*-knockout OPCs were generated with two independent single guide RNAs (sgRNAs) targeting *Vhl* (sgVhl and sgVhl.2), each of which caused significant decreases in *Vhl* transcript and protein levels through Cas9-mediated insertion-deletions (in-dels) at the respective target sites compared with control (Cas9 expressing OPCs with no sgRNA) (Figures S1E–S1G). It is possible that loss of VHL stabilizes pathways independent of HIF1a; therefore we confirmed key findings from *Vhl*-knockout OPCs in wild-type OPCs treated with hypoxia (1% O$_2$), primary mouse OPCs exposed to hypoxia (1% O$_2$), and human OPCs in pluripotent stem cell-derived oligocortical spheroids (Madhavan et al., 2018).

HIF1a Accumulation Specifically Delays OPC Differentiation into Oligodendrocytes

HIF1a accumulation in OPCs is sufficient to impair oligodendrocyte formation (Yuen et al., 2014). To test whether HIF1a accumulation is a general or specific inhibitor of OPC differentiation, sgVhl OPCs were stimulated to form either astrocytes (Grinspan et al., 2000) or oligodendrocytes (Baas et al., 1997; Barres et al., 1994; Gao et al., 1998; Najm et al., 2011; Figure 1E). This revealed a significant 4-fold reduction in oligodendrocyte formation by staining for myelin basic protein (MBP), a marker of mature oligodendrocytes, with no change in astrocyte formation by staining for glial fibrillary acidic protein (GFAP) in sgVhl OPCs compared with control (Figures 1E–1G and S1H), which shows that the effects of HIF1a are cell type specific. Staining for early (O4+), intermediate (O1+), and late (MBP+) oligodendrocyte markers throughout the differentiation process demonstrated a significant and delayed acquisition of all oligodendrocyte markers in sgVhl and sgVhl.2 OPCs compared with control (Figures 1H–1L and S1I–S1K). These data suggest that HIF1a accumulation specifically impairs early OPC differentiation, thereby delaying the formation of oligodendrocytes and ultimately myelin.

HIF1a Upregulates Unique Target Genes in OPCs

To delineate the gene targets of HIF1a in OPCs responsible for blocking oligodendrocyte development, chromatin-linked immunoprecipitation sequencing (ChIP-seq) was used to map its genome-wide chromatin binding profile. Using 100 million control and sgVhl OPCs, we identified 503 high-stringent HIF1a peaks (false discovery rate [FDR] < 0.001) in sgVhl OPCs with clear enrichment proximal to the annotated transcription start site (TSS), which agrees with HIF1a as a promoter centric transcription factor (Schödel et al., 2011; Smythies et al., 2019; Figures 2A and S2A). HIF1a was enriched at canonical target genes *Vegfa* and *Bnip3*, and globally peaks were enriched for HIF motifs and motifs for ubiquitously expressed transcription factors that have been shown to interact with HIF1a, including Sp1, c-Myc, and Bmal1 (Huang, 2008; Kaluz et al., 2003; Wu et al., 2017; Figures S2A and S2B; Tables S1, S2, and S3). HIF1a was not found proximal to *Sirt1*, *Wnt7a*, or *Wnt7b* (Figures S2C–S2E), each of which has previously been suggested as a putative HIF target that impairs oligodendrocyte formation in hypoxia-treated OPCs (Jablonska et al., 2016; Yuen et al., 2014). Moreover, sgVhl OPCs, wild-type OPCs, and primary OPCs exposed to hypoxia exhibited no increase in expression of *Sirt1*, *Wnt7a*, or *Wnt7b* transcripts, suggesting that other targets are likely functioning to block oligodendrocyte formation (Figures S2F–S2I; Zhang et al., 2020). Without HIF1a ChIP-seq data in OPCs, these previous studies focused on potential HIF1a target genes identified in other cell types, which overlooked that HIF1a might bind to unique cell-type-specific targets in OPCs that suppress oligodendrocyte formation.

(J–L) Quantification of the percentage of early O4+ (J), intermediate O1+ (K), and late MBP+ (L) oligodendrocytes in sgVhl (in green) and Cas9 control OPCs (in blue) at days 1, 2, and 3 of differentiation. Data are presented as mean ± SD from three independent biological replicates. p values were calculated using two-way ANOVA (reported as ANOVA p =) for overall group differences and Sidak's multiple-comparisons test for individual time point differences. p values for qRT-PCRs were calculated using Student's two tailed t test. See also Figure S1.
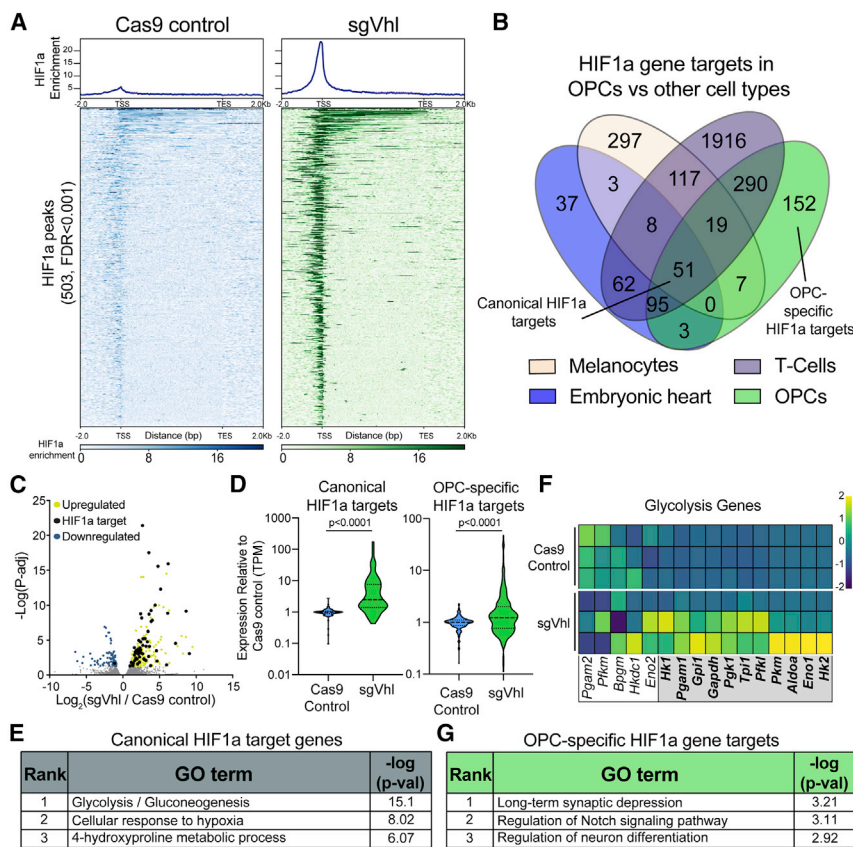
Figure 2. HIF1a Directly Upregulates Unique Targets in OPCs

(A) Aggregate binding profile and heatmap of 503 HIF1a peaks (FDR < 0.001) in sgVhl OPCs within 2 kb of the transcription start site (TSS) and transcription end site (TES) of the closest expressed gene in Cas9 control and sgVhl OPCs. See Table S1 for full list of peaks.

(B) Venn diagram of direct HIF1a target genes in diverse mouse tissues overlapped with HIF1a targets in OPCs. See Table S2 for full list of peaks.

(C) Volcano plot of genes that significantly increase and decrease, in yellow and blue, respectively (p-adj < 0.05), in sgVhl OPCs compared with Cas9 control OPCs with direct targets of HIF1a, in black. Gray dots are genes not significantly different between conditions. Data are from three biological replicates.

(D) Violin plots of expression data (TPM) normalized to Cas9 control OPCs for canonical and OPC-specific HIF1a targets in Cas9 control (in blue) and sgVhl (in green) OPCs. Bold dashed line represents the median, and thin dashed lines represent the upper and lower quartiles. p values were calculated using the Mann-Whitney test.

(E) Gene Ontology (GO) analysis of shared canonical targets of HIF1a. Table shows the rank of the GO term along with −log(p value).

(F) Heatmap of row-normalized expression of glycolysis genes (TPM) between Cas9 control and sgVhl OPCs. Genes in the gray box are direct targets of HIF1a. Each row represents a biological replicate.

(G) Gene Ontology (GO) analysis of OPC-specific HIF1a target genes. Table shows the rank of the GO term along with −log(p value).

See also Figure S2.

To define the cell-type-specific targets of HIF1a in OPCs, we overlapped HIF1a targets in OPCs with HIF1a targets from the limited number of publicly available datasets derived from other mouse cell types, including melanocytes (Loftus et al., 2017), T cells (Ciofani et al., 2012), and embryonic heart (Guimarães-Camboa et al., 2015). This analysis identified 51 genes that were HIF1a targets across all four cell types ("canonical HIF1a targets"), 152 genes that were HIF1a targets only in OPCs on the basis of these datasets ("OPC-specific HIF1a targets"), and 2,250 genes that were specific to heart, T cells, or melanocytes ("other tissue-specific HIF1a targets") (Figure 2B).

To determine functional targets of HIF1a in OPCs, we performed RNA sequencing (RNA-seq) of sgVhl and control OPCs. Overlapping transcripts that significantly changed between sgVhl and control OPCs (p-adjusted [adj] < 0.05) with direct targets of HIF1a revealed that HIF1a directly bound to 61 significantly increased genes and only 1 significantly decreased gene (Figures 2C and S2J; Table S1), consistent with the role of HIF1a as a transcriptional activator (Dengler et al., 2014; Guimarães-Camboa et al., 2015). Genes that were both significantly upregulated (p-adj < 0.05) and not called as top targets of HIF1a (at a stringent threshold FDR < 0.001) were enriched for immune and hypoxia pathways, suggesting that these genes are weaker direct targets of HIF1a or upregulated indirectly in the context of constitutive HIF accumulation

(Figure S2J). Both shared canonical and OPC-specific HIF1a target genes collectively increased in expression in sgVhl OPCs compared with control, with canonical targets induced to a greater extent than non-canonical targets (Figures 2D, S2K, and S2L). However, both of these gene sets showed a significantly greater increase in expression compared with the "other tissue-specific" HIF1a targets in sgVhl OPCs (Figure S2K).

Gene Ontology (GO) analysis of canonical HIF1a target genes showed enrichment for metabolic and hypoxia pathways, which agrees with HIF1a's role to promote glycolysis in a majority of cell types (Figure 2E; Choudhry and Harris, 2018; Majmundar et al., 2010; Miska et al., 2019; Nagao et al., 2019). In fact, more than half of the enzymes in the glycolysis pathway were direct HIF1a targets in OPCs, and both sgVhl and sgVhl.2 OPCs exhibited 2-fold increases in levels of the glycolysis byproduct lactate compared with control OPCs (Figures 2F and S2M).

Interestingly, GO analysis for cell-type-specific HIF1a targets demonstrated enrichment for pathways separate from the canonical hypoxic response and related to the tissue of origin (Figure S2N). In particular, GO analysis for OPC-specific HIF1a targets showed enrichment for neural development and differentiation pathways (Figure 2G). This demonstrates that HIF1a regulates genes critical for the canonical response to low oxygen as well as separate cell-type-specific targets.

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Chromatin Accessibility and Cell-Type-Specific Transcription Factors Define Non-canonical HIF1a Targets

How HIF1a is able to bind uniquely in a cell-type-specific manner remains unknown. To better understand the determinants of the cell type specificity of HIF1a binding profiles, we compared the open and active chromatin landscape in OPCs at canonical, non-canonical OPC-specific, and "other tissue-specific" HIF1a peaks (Figure S3A; Table S2). ChIP-seq for the active chromatin mark H3K27Ac (Creyghton et al., 2010) demonstrated a significant gain of H3K27Ac in sgVhl OPCs compared with control (FDR < 0.1), and these regions were enriched for HIF motifs (Figures S3B and S3C). Shared canonical and OPC-specific HIF1a peaks exhibited greater enrichment for HIF1a, H3K27Ac, and open chromatin (defined by ATAC-seq [assay for transposase-accessible chromatin using sequencing] regions in non-transduced OPCs) compared with "other tissue-specific" HIF1a sites (Figures 3A and S3D). This agrees with previous findings that HIF1a preferentially binds open and active chromatin and that the level of HIF1a binding correlates with the magnitude of expression of gene targets in hypoxia (Smythies et al., 2019; Xia and Kung, 2009). However, there was a subset of "other tissue-specific" HIF1a peaks enriched for open and active chromatin that lacked HIF1a binding in OPCs, suggesting that chromatin accessibility and activity were not the sole predictors of HIF1a binding. HIF1b, a common binding partner with HIF1a, was more strongly enriched at canonical HIF1a targets compared with OPC-specific targets, which further suggests that other interacting transcription factors could define cell-type-specific targets of HIF1a (Figure S3D).

Motif enrichment analysis under cell-type-specific HIF1a peaks demonstrated enrichment for lineage-defining transcription factors such as the Mitf family in melanocytes (Levy et al., 2006), Nkx2 family in embryonic heart (Bartlett et al., 2010), and basic-helix-loop-helix (bHLH) motifs in OPCs (Figure S3E; Table S3). Specifically, the motif for Olig2, a lineage-defining bHLH transcription factor in OPCs (Yu et al., 2013), was highly enriched (in the top 5% of motifs) under OPC-specific HIF1a peaks, whereas the Olig2 motif was not enriched under any other tissue-specific HIF1a peaks (Figures 3B and S3E; Table S3). ChIP-seq for OLIG2 in control and sgVhl OPCs validated the enrichment for OLIG2 at OPC-specific HIF1a peak loci compared with canonical and "other tissue-specific" peaks (Figures 3C, S3F, and S3G; Table S4). Furthermore, both canonical and non-canonical HIF1a peaks demonstrated an increase in OLIG2 binding in sgVhl OPCs compared with control, suggesting that HIF1a and OLIG2 physically interact (Figures 3C and S3G). Co-immunoprecipitation (coIP) of OLIG2 and HIF1a validated this physical interaction specifically in sgVhl OPCs compared with Cas9 control OPCs (Figures 3D, S3H, and S3I). Given this physical interaction between HIF1a and OLIG2, we observe an appearance of OLIG2 at canonical HIF1a target genes in sgVhl OPCs, such as Bnip3 (Figures 3E, S3J, and S3K; Table S4). Non-canonical HIF1a gene targets, such as Ascl2, are already enriched for OLIG2 in control OPCs and display increased OLIG2 binding in sgVhl OPCs, suggesting that OLIG2 also interacts with HIF1a at these loci (Figure 3E). Collectively, these data demonstrate that outside of canonical HIF1a targets, lineage-defining transcription factors, such as OLIG2, generate a permis-

sible chromatin landscape to drive HIF1a to unique non-canonical targets in diverse cell types.

OPC-Specific Targets of HIF1a Suppress *Sox10* Expression and Oligodendrocyte Formation

To understand whether OPC-specific targets are sufficient to impair oligodendrocyte formation, we first examined whether HIF1a induction affects the expression of critical lineage transcription factors and lineage markers in OPCs. This revealed a selective reduction in *Sox10*, a transcription factor required for formation of oligodendrocytes from OPCs (Stolt et al., 2002, 2004), and downstream SOX10 target genes *Plp1* and *Pdgfra*, whereas other lineage transcription factors, such as *Olig2* and *Nkx2-2*, remained unchanged (Figures 4A, S4A, and S4B). Of note, HIF1a did not bind proximal to any master regulators of oligodendrocyte development; however, the proximal promoter region of *Sox10* showed a significant reduction of H3K27Ac in sgVhl OPCs (Figure 4B). The decrease in expression of *Sox10* in sgVhl OPCs was validated at the mRNA and protein levels as well as by reduction in expression of downstream SOX10 target genes *Plp1* and *Pdgfra* in sgVhl OPCs and primary OPCs treated with hypoxia (1% O_2) (Figures 4C, 4D, and S4C–S4E). These data suggest that HIF1a indirectly represses *Sox10* through activation of other gene targets to block oligodendrocyte development.

To define the top functional targets of HIF1a in OPCs, we overlapped direct HIF1a targets (FDR < 0.001) with genes that exhibited both increased transcription (p-adj < 0.05) and increased H3K27Ac (FDR < 0.1) in sgVhl OPCs compared with control (Figures 2C and 4E; Table S1). Hits were independently validated in OPCs treated with hypoxia (1% O_2), which showed that all of our top 10 HIF1a targets were significantly upregulated compared with normoxia (Figures 1D and 4F). Of the top ten targets of HIF1a in OPCs, *Ascl2* and *Dlx3* were the only non-canonical, OPC-specific HIF1a targets (Figure 4E; Table S2). Both are transcription factors that regulate differentiation of somatic stem cells in the periphery and are not normally expressed by any cell type in the mouse CNS (Tabula Muris Consortium et al., 2018; Zhang et al., 2014), which is reflected by the lack of active chromatin at *Ascl2* and *Dlx3* in control OPCs (Figures 4G and S4F). However, OLIG2 binds to both of these loci in Cas9 control OPCs, which agrees with our finding that OLIG2 is enriched at non-canonical, OPC-specific HIF1a target genes (Figure 4G). Both targets display robust HIF1a peaks proximal to their promoters specifically in sgVhl OPCs compared with heart, melanocytes, and T cells along with gained H3K27Ac and OLIG2 binding in sgVhl OPCs compared with control OPCs (Figure 4G). Ascl2 and Dlx3 both accumulate at the protein level in sgVhl OPCs and are induced in primary OPCs treated with hypoxia compared to normoxic controls (Figures S4G–S4I). In fact, ASCL2 was induced *in vivo* in the brains of mouse pups reared in chronic hypoxia (10% O_2), and this correlated with a decrease in SOX10 and white matter proteins MBP and MAG compared with normoxia-reared controls (Figures S4J and S4K).

Ectopic expression of *Dlx3* and *Ascl2* in OPCs using CRISPR activation (CRISPRA) technology was sufficient to impair the acquisition of early (O4+), intermediate (O1+), and late (MBP+) oligodendrocyte markers across the course of differentiation (Figure 4H). Moreover, ectopic expression of *Dlx3* and *Ascl2*
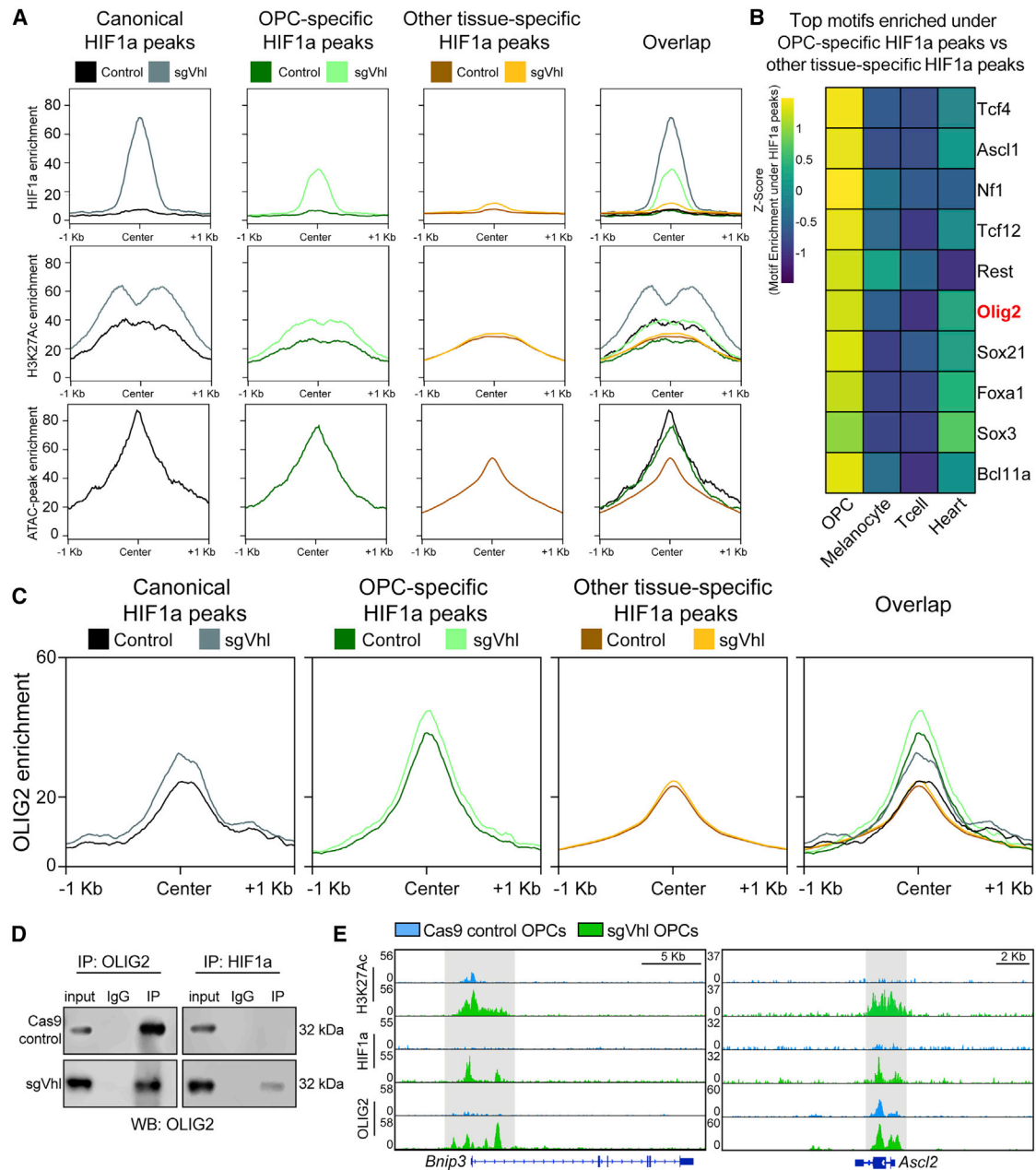
Figure 3. Chromatin Accessibility and Cell-Type-Specific Transcription Factors Define Non-canonical HIF1a Targets

(A) Aggregate plots of HIF1a and H3K27ac enrichment in Cas9 control and sgVhl OPCs and open chromatin enrichment in non-transduced OPCs at shared canonical HIF1a peaks, OPC-specific HIF1a peaks, and "other tissue-specific" HIF1a peaks.

(B) Heatmap of the enrichment of the top 10 transcription factor motifs under non-canonical HIF1a peaks in OPCs compared with other tissue types. Olig2 (highlighted in red) is a dominant transcription factor of the oligodendrocyte lineage.

(C) Aggregate plots of OLIG2 enrichment in Cas9 control and sgVhl OPCs at canonical HIF1a peaks, OPC-specific HIF1a peaks, and "other tissue-specific" HIF1a peaks.

(D) Western blot of OLIG2 from total cell lysate (input), IgG control lysate, and lysate from pull-down (IP) of either OLIG2 (on left) or HIF1a (on right) in Cas9 control (top) and sgVhl OPCs (bottom). See also Figures S3H and S3I for reciprocal coIP and replicate coIP in sgVhl OPCs.

(E) Genome Browser view of H3K27Ac, HIF1a, and OLIG2 ChIP-seq in Cas9 control (blue) and sgVhl (green) OPCs at *Bnip3* and *Ascl2*. Scale bars, 5 and 2 kb, respectively.

See also Figure S3.
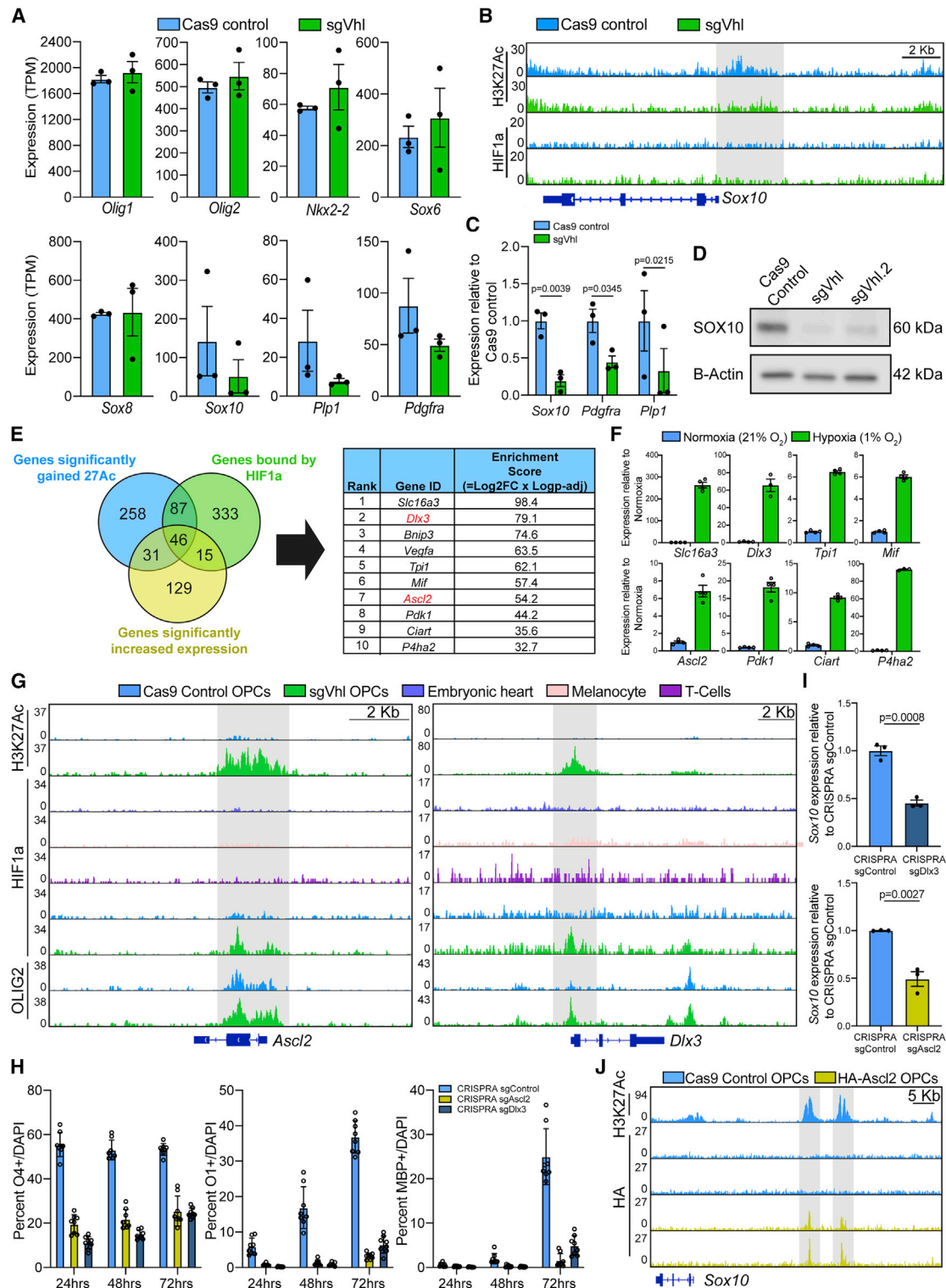
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Figure 4. OPC-Specific HIF1a Targets *Ascl2* and *Dlx3* Suppress *Sox10* and Impair Oligodendrocyte Formation

(A) Quantification of the normalized expression (TPM) of oligodendrocyte lineage markers in Cas9 control (in blue) and sgVhl (in green) OPCs. Data represent mean ± SEM from three independent RNA-seq replicates.

(B) Genome Browser view of H3K27ac and HIF1a ChIP-seq in Cas9 control (in blue) and sgVhl (in green) OPCs at the locus for *Sox10*. The gray bar highlights significant H3K27Ac enrichment at the *Sox10* promoter in control cells ($-\log_{10}$[q value] = -5.6) that is absent in Vhl-knockout OPCs. Scale bar, 2 kb.

led to a significant reduction in *Sox10* expression (Figures 4I and S4L), whereas activation of canonical HIF1a targets *Slc16a3*, *Bnip3*, and *Vegfa* did not (Figures S4L and S4M). ASCL2 is known to impair somatic stem cell differentiation and can act as a repressor at critical differentiation genes (Murata et al., 2020; Schuijers et al., 2015; Wang et al., 2017), so we hypothesized that ASCL2 could directly bind to upstream regulatory regions of *Sox10* (Yu et al., 2013). ChIP-seq for HA-tagged ASCL2 in OPCs revealed that ASCL2 is able to directly bind upstream enhancers of *Sox10* and reduce expression of *Sox10* compared with control OPCs (Figures 4J and S4N–S4P).

Although Ascl2, Dlx3, and possibly other non-canonical OPC-specific HIF1a targets are each sufficient to decrease *Sox10* expression and impair oligodendrocyte formation, it was not clear if these targets were individually necessary for the functional effect. RNAi-mediated knockdown of *Ascl2* or *Dlx3* in sgVhl OPCs did not rescue oligodendrocyte formation (Figures S4Q and S4R) suggesting that non-canonical HIF1a targets collectively or redundantly converge to suppress *Sox10* expression and impair oligodendrocyte formation from OPCs.

Chemical Inhibition of MEK/ERK Increases Oligodendrocyte Formation from sgVhl OPCs

To identify potential pathways that could overcome the differentiation block imposed by HIF1a, we screened a library of 1,753 bioactive compounds for the ability to increase the formation of MBP+ oligodendrocytes relative to DMSO-treated sgVhl OPCs (Figures 5A and S5A–S5D; Table S5). Compounds that were non-toxic (fold change in total cell number >0.7 relative to DMSO-treated sgVhl OPCs at the primary screening dose of 3 μM) and enhanced the number and percentage of MBP+ oligodendrocytes (fold change >3 relative to DMSO-treated sgVhl OPCs) were considered primary hits (Figure S5E). MEK inhibitors were enriched among the primary hits and, as a class, demonstrated a significant increase in oligodendrocyte formation compared with all other non-toxic compounds tested in the primary screen (Figures 5B, 5C, and S5F; Table S5). Interestingly, drugs previously identified to enhance oligodendrocyte formation, such as miconazole, clemastine, and tamoxifen (Gonzalez et al., 2016; Mei et al., 2014; Najm et al., 2015), were not enriched as hits in this screen, highlighting the ability of this screen to identify context-specific modulators of the differentiation block imposed by HIF1a (Figures 5A and S5B).

To identify compounds that were effective across a range of doses, we performed an eight-point dose curve from 78 nM to 10 μM of the top 14 hits. Performing unbiased hierarchical clustering of the results revealed that all five MEK inhibitors clustered together and led to a pronounced increase in oligodendrocyte formation relative to DMSO-treated sgVhl OPCs, with AZD8330 and PD0325091 significantly outperforming PD318088 and selumetinib, while PD184352 was not significantly different from the other four MEK inhibitors (Figures 5D, 5E, and S5G; Table S6). The ability of the top two MEK inhibitors to affect oligodendrocyte formation correlated with their on-target half maximal inhibitory concentration (IC_{50}) values for MEK1 and MEK2 as well as their on-target ability to reduce ERK1/2 phosphorylation in sgVhl OPCs (Figures 5F, 5G, and S5H). Performance of an eight-point dose curve consisting of 12 drugs that each inhibit a potential downstream target of MEK revealed that ERK1/2 inhibitors led to the greatest increase in oligodendrocyte formation compared with the other classes of drugs tested (Shaul and Seger, 2007; Yohe et al., 2018; Figures S5I and S5J). Compounds targeting the closely related MEK5 pathway did not increase oligodendrocyte formation from sgVhl OPCs, and AZD8330, the most potent MEK inhibitor, did not impair ERK5 phosphorylation in sgVhl OPCs (Figures S5K and S5L). Collectively, these results demonstrate that chemical inhibition of MEK/ERK acts as a node of intervention to restore the formation of oligodendrocytes in the context of HIF1a accumulation.

MEK/ERK Inhibition Drives *Sox10* Expression without Changing HIF1a Activity

We next asked whether MEK inhibition increased differentiation of sgVhl OPCs by modulating HIF1a or its direct downstream targets. Treating OPCs with 300 nM AZD8330, our most potent MEK inhibitor (Figures 5F and 5G), for 14 h did not alter nuclear accumulation of HIF1a (Figures 6A–6C). Furthermore, downstream non-canonical HIF1a targets *Ascl2* and *Dlx3* did not increase activation of MEK/ERK signaling (Figure S6A). These data suggest that MEK inhibitor treatment does not directly counter HIF1a or downstream HIF1a target activity but rather circumvents the effect of HIF1a accumulation by increasing oligodendrocyte differentiation despite persistent HIF signaling.

(C) qRT-PCR of *Sox10* and downstream SOX10 target genes *Pdgfra* and *Plp1* in sgVhl (in green) and Cas9 control (in blue) OPCs. Data are presented as mean ± SEM from three biological replicates.

(D) Western blot for SOX10 in sgVhl, Vhl.2 OPCs, and Cas9 control OPCs. Data represent results from a single biological replicate.

(E) Venn diagram overlapping genes with proximal HIF1a binding (in green), significant gains in H3K27ac (in blue), and significantly increased expression (in purple) in OPCs. The top ten genes were then ranked and listed by their enrichment score, which is the product of the log_2(fold change in gene expression) and $-log$(p-adj). See Table S1 for the full target list.

(F) qRT-PCR of eight top HIF1a target genes (see Figure 1D for *Vegfa* and *Bnip3*) in hypoxic (1% O_2, in green) and normoxic (21% O_2, in blue) OPCs. Data are presented as mean ± SEM from three or four technical replicates.

(G) Genome Browser view of HIF1a ChIP-seq at *Ascl2* and *Dlx3* in sgVhl OPCs (in green), Cas9 control OPCs (in blue), and other diverse cell types. H3K27ac and OLIG2 enrichment are also shown in sgVhl and Cas9 control OPCs. The gray bars highlight HIF1a accumulation in sgVhl OPCs. Scale bars, 2 kb.

(H) Quantification of the percentage of early O4+, intermediate O1+, and late MBP+ oligodendrocytes from sgAscl2 (in yellow), sgDlx3 (in dark blue), and sgControl (in light blue) CRISPRA OPCs at days 1, 2, and 3 of differentiation. Data are presented as mean ± SD of six to eight technical replicates (individual wells) per condition.

(I) qRT-PCR of *Sox10* in sgAscl2 (in yellow), sgDlx3 (in dark blue), and sgControl (in light blue) CRISPRA OPCs. Data are presented as mean ± SEM from three biological replicates.

(J) Genome Browser view of HA ChIP-seq upstream of *Sox10* in Cas9 control (in blue) and HA-Ascl2 (in yellow) OPCs. The gray bars highlight enhancers for *Sox10* defined by H3K27Ac ChIP-seq in Cas9 Control OPCs. Scale bar, 5 kb.

p values for qRT-PCRs were calculated using Student's two-tailed t test. See also Figure S4.
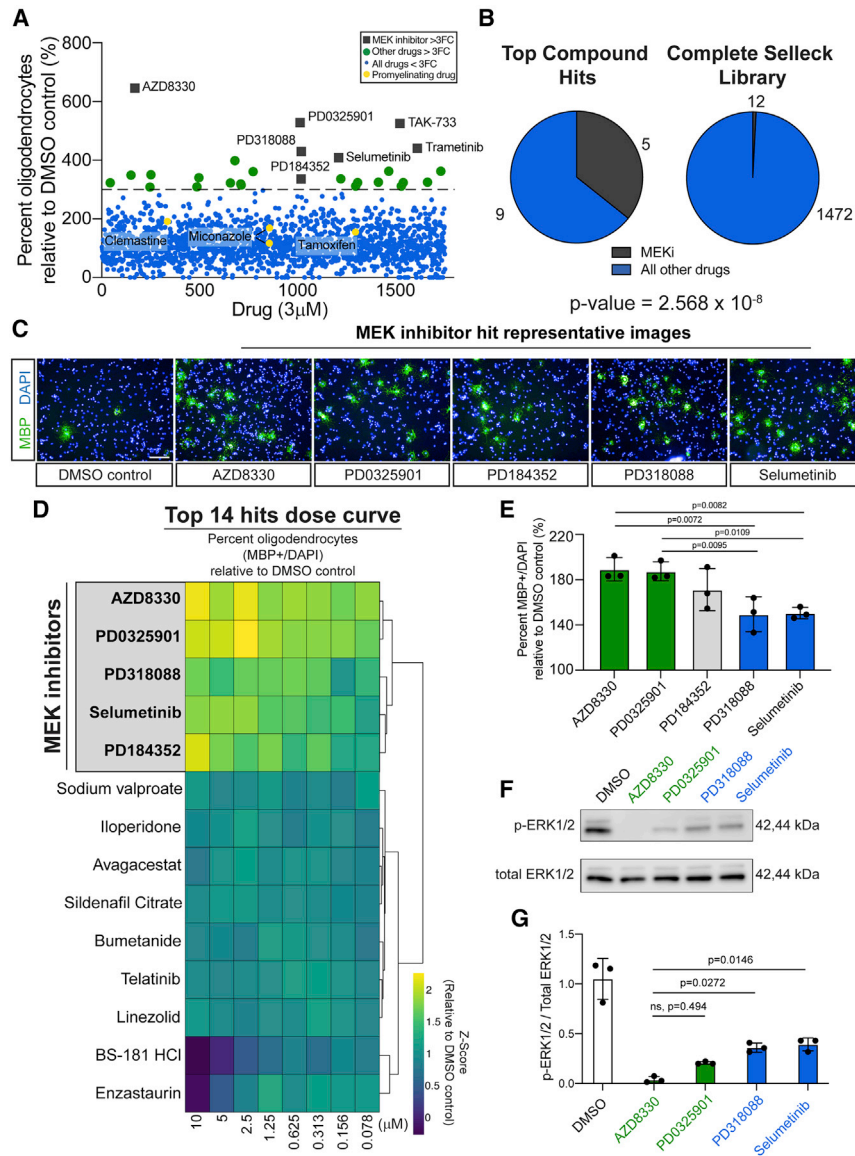
Figure 5. Chemical Inhibition of MEK/ERK Increases Oligodendrocyte Formation from sgVhl OPCs

(A) Primary bioactives library screen showing the effect of 1,753 molecules on percentage of oligodendrocytes (MBP+ cells/total DAPI) formed by sgVhl OPCs relative to DMSO-treated sgVhl OPCs. The dotted line represents a 3-fold increase in oligodendrocyte formation, and compounds that clear this threshold are indicated as green dots. MEK inhibitors are highlighted as gray boxes and pro-myelinating drugs are highlighted as yellow dots. See also Table S5.

(B) Pie charts of the number of MEK inhibitors (in dark gray) and other compounds classes (in blue) within top hits compared with their prevalence in the non-toxic compounds of the Selleck library. p value was calculated using hypergeometric analysis.

(C) Representative immunocytochemistry images of oligodendrocytes (MBP+, in green) from the primary drug screen of the top 5 MEK inhibitor hits along with the DMSO negative control. Nuclei are marked by DAPI (in blue). Scale bars, 100 μm.

(D) Heatmap showing the row-normalized fold change in the percentage (MBP+/DAPI) of oligodendrocytes relative to DMSO-treated sgVhl OPCs of an eight-point dose curve of the primary hits. The heatmap rows are sorted by unsupervised hierarchical clustering with columns in order from high (10 μM) to low dose (78 nM). MEK inhibitors are highlighted in gray and boldface type. Data are presented as the mean from three separate dose-curve plates. See also Table S6.

(E) Averaging all eight doses shows the ability of each MEK inhibitor to increase the formation of oligodendrocytes (MBP+/DAPI) relative to DMSO-treated sgVhl OPCs. Green and blue columns represent the most and least effective compounds respectively. Data are presented as the mean ± SD from three separate dose-curve plates. p values were calculated using a one-way ANOVA with Tukey's multiple-comparisons test.

(F) Representative western blot for phosphorylated ERK1/2 (p-ERK1/2) relative to total ERK1/2 of sgVhl OPCs incubated with 100 nM of indicated MEK inhibitors for 30 min.

(G) Quantification of the ratio of p-ERK1/2 to total ERK1/2 for the most (in green) and least (in blue) effective MEK inhibitors relative to DMSO-treated controls. Data are presented as mean ± SD from three biological replicates. p values were calculated using one-way ANOVA with Dunnett's multiple-comparisons test. See also Figure S5.

Gene set enrichment analysis (GSEA) revealed that AZD8330 treatment of sgVhl OPCs led to an enrichment for "oligodendrocyte differentiation" and "oligodendrocyte development" pathways, which were normally depleted in sgVhl OPCs compared with control (Figures 6D and 6E). Further supporting this, AZD8330 treatment of sgVhl OPCs led to a significant increase in the subset of genes within the GO term "oligodendrocyte development, GO:0014003" that were normally decreased in sgVhl OPCs compared with control (fold change of sgVhl to control OPCs < 0.75), such as *Sox10* and *Myrf* (Figures 6F and 6G). We validated the AZD8330-mediated increase in expression of both of these transcription factors, which are critical for oligodendrocyte differentiation, using qPCR (Figure 6G). To confirm that this was a function of impaired MEK/ERK signaling, treat-ment of sgVhl OPCs with ERK1/2 inhibitors AZD0364 and SCH772984 led to a similar increase in *Sox10* expression compared with DMSO (Figure S6B).

Given that MEK inhibitors do not change HIF1a translocation to the nucleus or downstream HIF1a target gene expression, it is unlikely that the MEK inhibitor-mediated increase in *Sox10* expression is specific to sgVhl OPCs. In agreement with this point, *Sox10* expression increases in Cas9 control OPCs treated with 300 nM AZD8330 for 14 h (Figures S6C and S6D). However, MEK inhibitors lead to a pronounced increase oligodendrocyte formation in sgVhl OPCs, with minimal impact on the oligodendrocyte formation from control OPCs, highlighting the context-specific block imposed by HIF1a (Figures S6E and S6F). Collectively, these data suggest that the reduction of *Sox10* expression
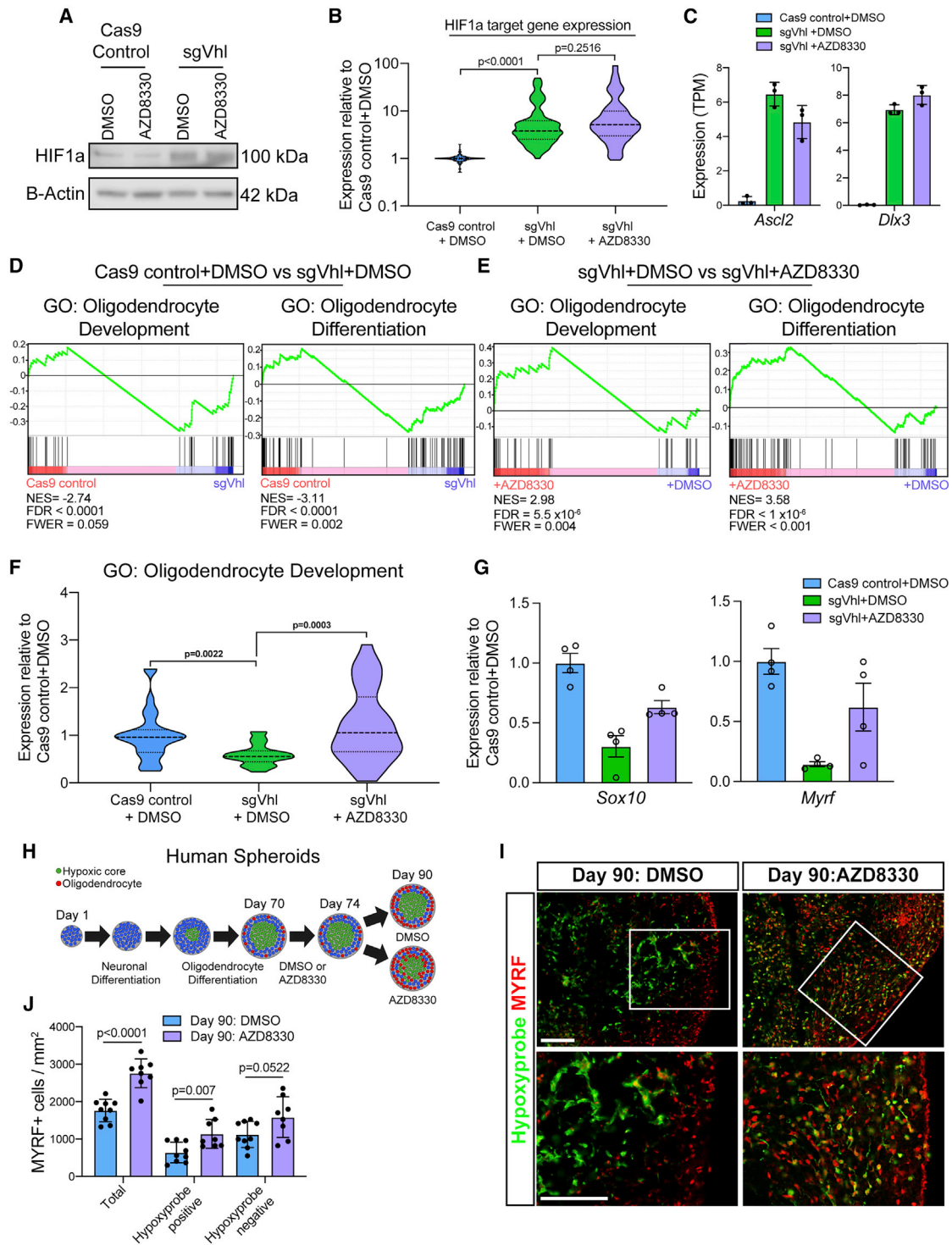
Figure 6. MEK Inhibitors Restore *Sox10* Expression Independent of HIF1a Activity in OPCs

(A) Western blot for HIF1a in sgVhl OPCs treated with AZD8330 or DMSO. Data represent results from a single biological replicate.

(B) Violin plot of normalized expression (TPM values normalized to Cas9 control + DMSO) of direct functional targets of HIF1a (see Figure 2B) in Cas9 control + DMSO (in blue), sgVhl + DMSO (in green), and sgVhl + AZD8330 (in purple) OPCs.

(C) Quantification of normalized expression (TPM) for both *Ascl2* and *Dlx3* in Cas9 control + DMSO (in blue), sgVhl + DMSO (in green), and sgVhl+AZD8330 (in purple) OPCs. Data represent mean ± SD from three independent RNA-seq replicates.

(D) Gene set enrichment analysis (GSEA) of gene program changes in sgVhl compared with Cas9 control OPCs demonstrates a significant reduction (FDR < 0.001) in GO terms for oligodendrocyte development (normalized enrichment score [NES] = −2.74) and oligodendrocyte differentiation (NES = −3.11).

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by cell-type-specific HIF1a targets is critical for the HIF-mediated block in oligodendrocyte differentiation, such that restoration of *Sox10* expression without altering canonical HIF function in sgVhl OPCs restores oligodendrocyte formation.

MEK/ERK Inhibition Drives Oligodendrocyte Formation in Hypoxic Regions of Human Oligocortical Spheroids

To evaluate the effect of MEK inhibition in hypoxic human OPCs, we leveraged a previously established method of generating human myelinating cortical spheroids from pluripotent stem cells (Madhavan et al., 2018). The interior of human brain spheroids is hypoxic (Brawner et al., 2017; Giandomenico and Lancaster, 2017), and we hypothesized that these hypoxic regions would inhibit SOX10 expression and oligodendrocyte formation. Immunohistochemistry using hypoxyprobe to define hypoxic regions of the spheroid (Pogue et al., 2001) demonstrated that SOX10 expression is significantly impaired in hypoxic regions of the spheroid (Figures S6G and S6H). We next tested whether hypoxic regions of the oligocortical spheroid were inhibitory to oligodendrocyte formation and whether this could be overcome using a MEK inhibitor. To test this, we treated oligocortical spheroids with either DMSO or 300 nM AZD8330 for 4 days starting at 70 days *in vitro*, immediately following induction of oligodendrocytes. At day 90, spheroids were treated with hypoxyprobe and harvested for analysis (Figure 6H). Immunohistochemistry for oligodendrocytes (MYRF+ cells) and hypoxic regions (defined by hypoxyprobe staining) demonstrated a significant 2.8-fold reduction in the number of MYRF+ oligodendrocytes in hypoxic regions compared with normoxic regions within the spheroids (Figures 6I, 6J, and S6H). MEK/ERK inhibition with AZD8330 treatment led to a significant 1.8-fold increase in the number of oligodendrocytes within hypoxic regions of the spheroid (Figures 6I, 6J, and S6I). Collectively these results show that oxygen tensions shape oligodendrocyte development and that MEK inhibition circumvents the hypoxia-mediated inhibition of oligodendrocyte formation in three-dimensional (3D) models of human brain development.

DISCUSSION

Cells are equipped to translate external cues from the environment into internal signals that ultimately alter transcriptional programs. Molecular oxygen is crucial to support energy production of the cell, and low oxygen upregulates a rapid and conserved transcriptional response mediated largely by HIF transcription factors in all mammalian cells. HIF1a promotes an adaptive response by upregulating oxygen-independent metabolism and increasing blood vessel formation; however, chronic accumulation of HIF1a negatively affects the function of almost every organ system (Kullmann et al., 2020; Lee et al., 2019; Menendez-Montes et al., 2016; Takubo et al., 2010).

Here, we profiled the genome-wide functional targets of HIF1a in OPCs and found that HIF1a not only binds to canonical hypoxia response genes that are shared across multiple cell types but also activates a unique set of non-canonical genes in a cell-type-specific manner. In the context of the brain, these non-canonical HIF1a targets that impair oligodendrocyte formation could have implications in the numerous hypoxia-driven pathologies of white matter such as diffuse white matter injury of prematurity (Salmaso et al., 2014; van Tilborg et al., 2018), white matter stroke in adults (Hankey, 2017; Marin and Carmichael, 2019), and multiple sclerosis (Ding et al., 2020; Graumann et al., 2003; Zeis et al., 2008). More broadly, we suggest that non-canonical HIF1a targets in diverse cell types affect a variety of cell-type-specific functions, such as oligodendrocyte differentiation, heart morphogenesis, and T cell activation. This concept has previously been overlooked, as many studies have focused on canonical targets of HIF1a or those that were discovered in immortalized cell lines, which may behave differently in response to HIF1a accumulation. *Wnt7a* and *Wnt7b*, for example, were identified as direct HIF1a targets in mouse fibroblasts and extrapolated to be direct targets in mouse OPCs (Yuen et al., 2014). However, we demonstrate that Wnt7 ligands are not direct HIF1a targets in OPCs, a finding corroborated by a recent study demonstrating that Wnt7 ligands are targets of HIF1a in astrocytes but not in OPCs (Zhang et al., 2020). This highlights that the response to hypoxia is not universal across all cell types and that the cell-type-specific response to HIF1a induction, such as Wnt signaling in astrocytes or downregulation of *Sox10* in OPCs, have critical phenotypic consequences (Zhang et al., 2020).

The same transcription factor can bind to different gene targets in different cell types through interaction with transcriptional machinery unique to each cell type (Mullen et al., 2011; Trompouki et al., 2011). HIF1a binding has been shown to depend on the openness and activation status of chromatin; however, we and others show that chromatin accessibility is not the sole predictor of HIF1a binding (Schödel et al., 2011; Smythies

(E) GSEA of gene program changes in sgVhl + AZD8330 compared with sgVhl + DMSO OPCs demonstrates a significant enrichment (FDR < 0.001) in GO terms for oligodendrocyte development (normalized enrichment score [NES] = 2.98) and oligodendrocyte differentiation (NES = 3.58).

(F) Violin plot showing normalized expression (TPM values normalized to Cas9 control + DMSO OPCs) of genes in the GO term oligodendrocyte development (GO:0014003) that decrease (fold change [FC] < 0.75) in sgVhl + DMSO OPCs (in green) relative to Cas9 control + DMSO OPCs (in blue) as well as sgVhl OPCs following treatment with AZD8330.

(G) qRT-PCR of *Sox10* and *Myrf* in Cas9 control + DMSO (in blue), sgVhl + DMSO (in green), and sgVhl + AZD8330 (in purple) OPCs. Data are presented as mean ± SEM from four technical replicates (individual wells).

(H) Schematic of human brain oligocortical spheroids treated at days *in vitro* (DIV) 70 with either DMSO or AZD8330 for 4 days. At day 90, organoids were incubated with hypoxyprobe, fixed, and sectioned for immunohistochemistry.

(I) Representative immunohistochemistry images for oligodendrocytes (MYRF+, in red) and hypoxic regions (hypoxyprobe, in green) of DIV 90 oligocortical spheroids at low and high magnification that were treated from DIV 70–74 with DMSO or AZD8330. Scale bars, 100 μM.

(J) Quantification of oligodendrocytes (MYRF+ [mm²]) in the whole oligocortical spheroid (total), hypoxic region of the spheroid (hypoxyprobe positive), and normoxic region of the spheroid (hypoxyprobe negative) in DIV 90 spheroids that had been treated with DMSO or AZD8330 from DIV 70–74. Data represent mean ± SD from eight or nine individual spheroids. p values were calculated using Student's two-tailed t test.

AZD8330 was used at a dose of 300 nM for all experiments, and p values for violin plots were calculated using the Kruskal Wallis one-way ANOVA with Dunn's multiple-comparisons test. See also Figure S6.

et al., 2019; Xia and Kung, 2009). Our results highlight that HIF1a binds more strongly and induces a greater increase in canonical hypoxia response genes compared with non-canonical cell-type-specific genes, which are enriched for open chromatin as well as motifs for lineage-defining transcription factors. This implies that chronic HIF1a upregulation could allow sufficient accumulation of non-canonical HIF1a targets that then impair development and function in a cell-type-specific manner. These non-canonical targets could represent either a pathological "off-target" effect of sustained HIF1a accumulation or a normal cell-type-specific response to molecular oxygen levels that is coopted in the context of hypoxia disease.

To cast a wide net to uncover pathways that circumvent the disruption of oligodendrocyte formation imposed by HIF, we performed a chemical genetics screen revealing that MEK/ERK inhibition enhanced oligodendrocyte programs. This adds to the ongoing debate of the functionality of MEK/ERK signaling in the oligodendrocyte lineage. Previous studies have shown that MEK/ERK signaling can promote myelination (Ishii et al., 2012; Yang et al., 2016), while others demonstrate that inhibition of MEK/ERK signaling enhances OPC differentiation to oligodendrocytes (Newbern et al., 2011; Suo et al., 2019). We believe that this discrepancy owes to a potential stage-specific function of MEK/ERK such that this pathway impairs the transition from OPC to immature oligodendrocyte but then promotes mature oligodendrocyte formation and myelination. However, the mechanism of how inhibition of MEK/ERK promotes early oligodendrocyte programs remains elusive. One possibility could be that impairing ERK2 phosphorylation leads to transcriptional disinhibition at *Sox10* and other key development genes, as has been shown in cancer and mouse embryonic stem cells (Tee et al., 2014; Yohe et al., 2018). MEK inhibitors have also been shown to directly upregulate SOX10 translocation to the nucleus, which would then drive downstream oligodendrocyte gene programs (Fufa et al., 2019). Continued dissection of the roles of MEK/ERK signaling during the process of differentiation would provide critical insight to how this pathway plays a bidirectional role regulating oligodendrocyte formation from OPCs.

Overall, this work advances our conceptual understanding of the tissue-specific response to chronic HIF accumulation and how oxygen tensions regulate tissue physiology and pathology.

Limitations of Study

Although we show that *Ascl2* and *Dlx3* individually are not necessary for the HIF1a-mediated block in differentiation, it is possible that inactivation of combinations of *Ascl2*, *Dlx3*, and other non-canonical targets might elucidate a core set of non-canonical HIF1a targets that are necessary to impair oligodendrocyte formation. The mechanism by which MEK/ERK signaling regulates *Sox10* expression remains unknown and will benefit from further study with a particular focus on developmental stage-specific effects. Because of the number of cells required in this study, most experiments were limited to *in vitro* cultured mouse OPCs. Given that hypoxia also impairs oligodendrocyte formation *in vivo* in rodents and *in vitro* in human oligocortical spheroid tissue, it will be important for future studies to validate the effects of non-canonical HIF1a targets in these more therapeutically relevant contexts. Finally, it will be exciting to understand if cell-type specific

non-canonical HIF1a targets are the key drivers of hypoxia-mediated dysfunction in other cell types outside of OPCs.

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

- KEY RESOURCES TABLE
- RESOURCE AVAILABILITY
  - Lead Contact
  - Materials Availability
  - Data and Code Availability
- EXPERIMENTAL MODEL AND SUBJECT DETAILS
  - Culture of 293T Cells
  - Pluripotent stem cell-derived OPC culture
  - Primary mouse OPC culture
  - Mouse OPC differentiation to oligodendrocytes and astrocytes
  - Human oligocortical spheroid production
  - Tissue from mild chronic hypoxia (MCH) mice
- METHOD DETAILS
  - *In vitro* hypoxia experiments
  - Immunocytochemistry
  - High content imaging and quantification
  - Generation of CRISPR and CRISPRA OPCs
  - Generation of HA-Ascl2 OPCs
  - Validating CRISPR knockout in OPCs
  - Kinetics of OPC differentiation to oligodendrocytes
  - Compound screening and assessment
  - Western blot
  - Co-immunoprecipitation
  - qRT-PCR
  - Cell viability assay
  - siRNA knockdown in OPCs
  - RNA-seq sample preparation and analysis
  - Violin plots
  - Gene ontology analysis and heatmaps
  - Gene set enrichment analysis
  - ChIP-seq and analysis
  - Omni ATAC-seq
  - Diffbind analysis
  - HIF1a ChIP-seq overlap analysis
  - Motif enrichment analysis
  - Mass spectrometry for lactate and pyruvate
  - Oligocortical spheroid immunohistochemistry
- QUANTIFICATION AND STATISTICAL ANALYSIS

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AUTHOR CONTRIBUTIONS

DECLARATION OF INTERESTS

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KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
CD140a-APC	eBioscience	Cat#17-1401-81; RRID:AB_529482
NG2-AF488	Millipore	Cat#AB5320A4; RRID:AB_11203143
OLIG2	Proteintech	Cat#13999-1-AP; RRID:AB_2157541
OLIG2	Millipore	Cat#AB9610; RRID:AB_570666
OLIG2	Millipore	Cat#MABN50; RRID:AB_10807410
MBP	Abcam	Cat#ab7349; RRID:AB_305869
O1	CCF Hybridoma Core Facility	N/A
O4	CCF Hybridoma Core Facility	N/A
ASCL2	Millipore	Cat#MAB4417; RRID:AB_10562654
GFAP	Agilent/Dako	Cat#Z033401-2
Donkey anti-rat AF488	Thermo Fisher Scientific	Cat#A-21208; RRID:AB_2535794
Donkey anti-mouse AF488	Thermo Fisher Scientific	Cat# A-21202; RRID:AB_141607
Donkey anti-mouse AF647	Thermo Fisher Scientific	Cat#A31571; RRID:AB_162542
Donkey anti-rabbit AF647	Thermo Fisher Scientific	Cat#A31573; RRID:AB_2536183
HIF1a	Abcam	Cat#Ab2185; RRID:AB_302883
Phospho-p44/42. (phosphorylated-ERK1/2)	Cell Signaling Technologies	Cat#9101S; RRID:AB_331646
P44/42 (total ERK1/2)	Cell Signaling Technologies	Cat#9102S; RRID:AB_330744
Phospho-ERK5 (Thr218/Tyr220)	Cell Signaling Technologies	Cat#3371S; RRID:AB_2140424
ERK5 (total)	Cell Signaling Technologies	Cat#3372S; RRID:AB_330491
DLX3	Abcam	Cat#Ab178428
SOX10	R&D Systems	Cat#AF2864; RRID:AB_442208
B-Actin peroxidase	Sigma Aldrich	Cat#A3854; RRID:AB_262011
VHL	BD Biosciences	Cat#556347; RRID:AB_396376
MBP	Abcam	Cat#ab7349; RRID:AB_305869
MBP	BioLegend	Cat#808401; RRID:AB_2564741
MAG	Thermo Fisher Scientific	Cat#346200; RRID:AB_2533179
H3K27Ac	Abcam	Cat#Ab4729; RRID:AB_2118291
MyRF	Gift from Michael Wegner	N/A
HA	Cell Signaling Technologies	Cat#3724S; RRID:AB_1549585
Normal Rabbit IgG	Cell Signaling Technologies	Cat#2729S; RRID:AB_1031062
SOX6	abcam	Cat#30455; RRID:AB_1143033
A2B5	Millipore	Cat#MAB312; RRID:AB_94709
NKX2-2	DSHB	Cat#74.5A5; RRID:AB_531794
Bacterial and Virus Strains		
Stbl3 E.Coli	ThermoFisher	Cat#C737303
Lenti-X shots for lenti-virus generation	Clonetech	Cat# 631276
Biological Samples		
MCH forebrain mouse protein lysate samples	Gift from Brian Popko	N/A
Chemicals, Peptides, and Recombinant Proteins		
N2 Max	R&D Systems	Cat#AR009
B27	Thermo Fisher Scientific	Cat#12587010
bFGF	R&D Systems	Cat#23-3FB-01M
PDGFA	Thermo Fisher Scientific	Cat#221-AA-050
DMEM/F12	Thermo Fisher	Cat#11320-033

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
DMEM	Thermo Fisher	Cat# 11960-044
Fetal Bovine Serum (FBS)	Fisher	Cat#A3160402
MEM Non-essential amino acids	Thermo Fisher	Cat#11140-050
Glutamax	Thermo Fisher	Cat#35050061
2-Mercaptoethanol	Sigma	Cat#M3148
Poly-L-ornithine	Sigma Aldrich	Cat#P3655-50MG
Laminin	Sigma Aldrich	Cat#L2020-1MG
Noggin	R&D Systems	Cat#3344NG050
IGF-1	R&D Systems	Cat#291G1200
cyclic AMP	Sigma Aldrich	Cat#D0260-100MG
NT3	Thermo Fisher Scientific	Cat#267N3025
T3 (Thyroid hormone)	Sigma Aldrich	Cat#T-6397
Hb-EGF	R&D Systems	Cat#259-HE-050
CNTF	R&D Systems	Cat#557-NT-010
BMP4	R&D Systems	Cat#314-BP-050
CloneR	Stem Cell Technologies	Cat#5889
LDN193189	Sigma	Cat#SML0559
SB-43152	Sigma	Cat#S4317
Normal donkey serum	Jackson ImmunoResearch	Cat#017-000-121
Paraformaldehyde	Electron microscopy sciences	Cat#15710
Triton X-100	Sigma Aldrich	Cat#1086431000
DAPI	Sigma Aldrich	Cat#D8417
Puromycin	Thermo Fisher Scientific	Cat#A1113802
Blasticidin	Thermo Fisher Scientific	Cat#A1113903
Hygromycin	Thermo Fisher Scientific	Cat#10687010
SCH772984	Selleck Chemicals	Cat#S7101
AZD0364	Selleck Chemicals	Cat#S8708
VX-11e	Selleck Chemicals	Cat#S7709
BIX02188	Selleck Chemicals	Cat#S1530
BIX02189	Selleck Chemicals	Cat#S1531
RIPA Buffer	Sigma	Cat# R0278
Protease and Phosphatase Inhibitor	Thermo Fisher Scientific	Cat#78441
NuPAGE Bis-Tris gels	Thermo Fisher Scientific	Cat#NP0335BOX
PVDF membranes	Thermo Fisher Scientific	Cat#LC2002
TRIzol	Ambion	Cat#15596-026
Protein G magnetic DynaBeads	Thermo Fisher Scientific	Cat#10004D
Proteinase K	Thermo Fisher	Cat#AM2546
Ampure XP beads	Aline Biosciences	Cat#C-1003-5
Hypoxyprobe-1	Hypoxyprobe Inc	Cat#HP1-100Kit
Penicillin-streptomycin	ThermoFisher	Cat#15070063
Bioactive Compound Library-I	Selleck Chemicals	Cat#1700
Critical Commercial Assays		
QIAquick Gel Extraction Kit	QIAGEN	Cat#28115
QIAGEN Plasmid Maxi Kit (25)	QIAGEN	Cat#12163
KAPA HiFi HotStart ReadyMix	KAPA Biosystems	Cat#kk2602
KAPA Sybr Fast qPCR kit	KAPA Biosystems	Cat# KK4605
Taqman gene expression master mix	Thermo Fisher	Cat#4369016
NEBNext® Library Quant Kit for Illumina®	New England 641 Biolabs	Cat#E7630
Protein Assay Kit	Bio-Rad	Cat#5000001

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
RNeasy Mini Kit	QIAGEN	Cat#74104
iScript cDNA synthesis kit	Bio-Rad	Cat#1708891
NEBNext Poly(A) mRNA Magnetic Isolation Module	NEB	Cat#E7490L
NEBNext Ultra RNA Library Prep Kit for Illumina	NEB	Cat#E7530L
truChIP Chromatin Shearing Kit with Formaldehyde	Covaris	Cat#520154
Hypoxyprobe Kit	Hypoxyprobe Inc	Cat#HP1-100Kit
ON-TARGETplus Mouse siRNA smartpool: Ascl2	Horizon/Dharmacon	Cat#L-047262-01-0005
ON-TARGETplus Mouse siRNA smartpool: Dlx3	Horizon/Dharmacon	Cat#L-041957-01-0005
siGENOME Mouse siRNA pool: Non-Targeting Control #2	Horizon/Dharmacon	Cat#D-001206-14-05 5 nmol
Basic NucleofectorTM Kit for Primary Mammalian Glial Cells	Lonza	Cat# VPI-1006
QIAGEN MinElute PCR purification Kit	QIAGEN	Cat#28004
MTS assay kit	abcam	Cat# ab197010
Deposited Data		
RNA-seq datasets	This paper	GSE143474; GSE143473 (subseries)
ChIP-seq datasets	This paper	GSE143474; GSE143472 (subseries)
ATAC-seq datasets	This paper	GSE143474; GSE155366 (subseries)
Experimental Models: Cell Lines		
Lenti-X 293T Cell line	Takara Biosciences	Cat#632180
Epiblast stem cell (EpiSC) derived OPCs	Najm et al., 2011	N/A
Human iPSCs (H7) to derive Oligocortical spheroids	NIH hESC registry	NIHhESC-10-0061; RRID:CV-CL_9772
Experimental Models: Organisms/Strains		
Mouse, Male: 129S/SvEv E3.5 for EpiSC isolation	Jackson Laboratory	RRID:MGI:3050593
Mouse, mixed sex, postnatal day 2 and 7: C57BL/6J	Jackson Laboratory	RRID:IMSR_JAX:000664
Oligonucleotides		
Primer: sgVHL Forward: TCCCTACACGACGCTCTTCCGATCT CTCTCAGGTCATCTTCTGCAACC	This manuscript	IDT
Primer: sgVHL Reverse: AGTTCAGA CGTGTGCTCTTCCGATCTGACAAG ATGCTCGGGGTCGG	This manuscript	IDT
Primer: sgVHL.2 Forward: TCC CTACACGACGCTCTTCCGATC TAATAAACAGGTGCCATGCCC	This manuscript	IDT
Primer: sgVHL.2 Reverse: AGTTCA GACGTGTGCTCTTCCGATCTAGAT TGACTATTAACCTGGCAATG	This manuscript	IDT
sgRNAs	See STAR Methods	IDT
Taqman probes	See STAR Methods	IDT
Recombinant DNA		
CRISPRv2 constructs	Sanjana et al., 2014	Addgene plasmid #52961; RRID:Addgene_52961
lentiSAMv2 constructs	Joung et al., 2017	Addgene plasmid #75112; RRID:Addgene_75112

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Activation helper	Joung et al., 2017	Addgene plasmid #89308; RRID:Addgene_89308
N terminus tagged 3x HA Ascl2 construct	Genecopoeia	Cat# EX-Mm03774-Lv117
Software and Algorithms		
Columbus Image Data Storage and Analysis System	Perkin Elmer	https://www.perkinelmer.com/Product/columbus
Outknocker	Schmid-Burgk et al., 2014	http://www.outknocker.org/outknocker2.htm
Graphpad Prism	Graphpad Software	https://www.graphpad.com/scientific-software/prism/; RRID: SCR_002798
Image Studio Acquisition software	LI-COR	LI-COR Image Studio Software; RRID:SCR_015795
Salmon 0.14.1	Patro et al., 2017	https://github.com/COMBINE-lab/salmon
DESEQ2	Love et al., 2014	https://bioconductor.org/packages/release/bioc/html/DESeq2.html; RRID:SCR_015687
Metascape	Zhou et al., 2019	http://metascape.org/gp/index.html#/main/step1; RRID:SCR_016620
GSEA	Subramanian et al., 2005	https://www.gsea-msigdb.org/gsea/index.jsp; RRID:SCR_003199
Trim Galore! Version 0.4.1	Krueger, 2012	http://www.bioinformatics.babraham.ac.uk/projects/trim_galore/; RRID:SCR_011847
Bowtie2 Version 2.3.2	Langmead and Salzberg, 2012	http://bowtie-bio.sourceforge.net/bowtie2/index.shtml; RRID:SCR_005476
MACs Version 2.1.1	Feng et al., 2012	https://github.com/macs3-project/MACS; RRID:SCR_013291
Integrative Genomics Viewer	Broad Institute	https://www.broadinstitute.org/igv/; RRID:SCR_011793
Galaxy	Afgan et al., 2018	https://galaxyproject.org/; RRID:SCR_006281
Diffbind software	Ross-Innes et al., 2012	https://bioconductor.org/packages/release/bioc/html/DiffBind.html; RRID:SCR_012918
HOMERv4.11.1	Heinz et al., 2010	http://homer.ucsd.edu/; RRID:SCR_010881
BEDTools Version 2.25.0	Quinlan and Hall, 2010	https://github.com/arq5x/bedtools2; RRID:SCR_006646
Deeptools Version 3.3.1	Ramírez et al., 2016	https://deeptools.readthedocs.io/en/develop/; RRID:SCR_016366
Microsoft Excel	Microsoft	2016
Illustrator CC	Adobe	2020
Other		
C-Chamber	BioSpherix	Cat#C-274
96-well plate	ThermoFisher	Cat#167008
384-well PDL-coated plate	PerkinElmer	Cat#6057500
Operetta High Content Imaging and Analysis System	PerkinElmer	N/A
EL406 Microplate Washer Dispenser	BioTek	N/A
5μl dispense cassette	BioTek	N/A
Janus automated workstation	PerkinElmer	N/A
Odyssey Fc Imaging System	LI-COR	N/A
7300 real-time PCR system	Applied Biosystems	N/A
5973 Mass Spectrometer	Agilent	N/A
Hamamatsu Nanozoomer S60 Slide Scanner	Hamamatsu	C13210-01
SuperSignal West Pico PLUS Chemiluminescent Substrate	ThermoFisher	Cat#34580

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Halt Protease and Phosphatase Inhibitor Cocktail, EDTA-free (100x)	ThermoFisher	Cat#78441
Nextera Tn5 Transposase	Illumina	Cat#FC-121-1030
Covaris S2 Focused Ultrasonicator	Covaris	Cat#17015
Amaxa Nucleofector 2b	Lonza	Cat#AAB-1001
E12.5 heart HIF1a ChIP-Seq	Guimarães-Camboa et al., 2015	GSM1500750, GSM1500751, GSM1500749
Melanocyte HIF1a ChIP-Seq	Loftus et al., 2017	GSM2305570, GSM2305571, GSM2305572
Th17 T cell HIF1a ChIP-Seq	Ciofani et al., 2012	GSM1004819, GSM1004991, GSM1004820, GSM1004993
Mus musculus mm10 genome build	NCBI; Mouse Genome Sequencing Consotium	https://www.ncbi.nlm.nih.gov/assembly/GCF_000001635.20/

RESOURCE AVAILABILITY

Lead Contact
Further information and requests for materials, datasets, and protocols should be directed to and will be fulfilled by the Lead Contact, Paul Tesar (paul.tesar@case.edu).

Materials Availability
This study did not generate new unique reagents.

Data and Code Availability
All datasets generated in this study have been deposited in Gene Expression Omnibus (https://www.ncbi.nlm.nih.gov/geo/) under SuperSeries accession code GEO: GSE143474 with subseries for RNA-seq (GEO: GSE143473), ChIP-seq (GEO: GSE143472), and ATAC-seq (GEO: GSE155366).

Software used to analyze the datasets are freely or commercially available and detailed in the Key Resources Table.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Culture of 293T Cells
293T cells (Takara Bioscience, 632180) were used for lentiviral production according to the vendor's instructions and were cultured in the following medium: DMEM (Thermo Fisher, 11960-044) supplemented with 10% FBS (Fisher, A3160402), 1x MEM Non-essential amino acids (Thermo Fisher, 11140-050), 1x Glutamax (Thermo Fisher, 35050061), and 0.1mM 2-Mercaptoethanol (Sigma, M3148).

Pluripotent stem cell-derived OPC culture
Mouse OPCs were generated from epiblast stem cells (EpiSCs) as previously described except SHH was not used for maintenance or differentiation of OPCs (Najm et al., 2011). Mouse protocols were approved by Case Western Reserve University School of Medicine's Institutional Animal Care and Use Committee (IACUC). In brief, epiblast stem cells (EpiSCs) were isolated from 129S/SvEv male embryos (E3.5; The Jackson Laboratory), pushed to form neural rosettes, and then rosettes were passaged into nunclon plates coated with poly-L-ornithine (Sigma, P3655-50MG) and laminin (Sigma, L2020-1MG) in OPC growth media consisting of DMEM/F12 supplemented with N2 Max (R&D Systems, AR009), B27 (Thermo Fisher, 12587010), 20ng/mL bFGF (R&D Systems, 23-3FB-01M), and 20ng/mL PDGFA (R&D Systems, 221-AA). Media was changed every 48 hours and cultures were maintained in 37°C with 5% CO₂. After 4 passages these EpiSC derived OPCs were sorted to purity by fluorescence activated cell sorting using conjugated CD140a-APC (eBioscience, 17-1401-81; 1:80) and NG2-AF488 (Millipore, AB5320A4; 1:100) antibodies.

Primary mouse OPC culture
Primary mouse OPCs were derived using two methods. In the first method, cerebral cortices were harvested from postnatal day 2 (P2) C57BL/6J pups and dissociated using a Tumor Dissociation Kit (Miltenyi). Cells were then filtered through a 70 μm filter, washed in DMEM/F12, and plated on poly-ornithine (Sigma, P3655-50MG) and laminin (Sigma, L2020-1MG) coated plates to be expanded, passaged, and used in experiments. The second method follows the immunopanning protocol from P7 C57BL/6J mice (Barres et al., 1992). Primary *in vivo* derived OPCs using either method were pooled from multiple pups such that they are a combination of male and female cells. Similar to EpiSC-derived OPCs, primary OPCs were grown on Poly-L-ornithine and laminin coated flasks in OPC growth media. Media was changed every 48 hours.

Mouse OPC differentiation to oligodendrocytes and astrocytes

For oligodendrocyte generation, OPCs were seeded at either 40,000 cells per well (96-well plate, Fisher, 167008) or 15,000 cells per well (384-well PDL-coated cell carrier plates, PerkinElmer, 6057500) on plates coated with PO and laminin. Oligodendrocyte differentiation media consisted of DMEM/F12 supplemented with N2 Max, B27, 100ng/mL noggin (R&D, 3344NG050), 100ng/mL IGF-1 (R&D, 291G1200), 10uM cyclic AMP (Sigma, D0260-100MG), 10ng/mL NT3 (R&D, 267N3025) and 40ng/mL T3 (Thyroid hormone, Sigma, T-6397). Cells were analyzed after 3 days unless otherwise noted.

For OPC differentiation to astrocytes, 30,000 OPCs were plated per well in a 96 well plate (Fisher, 167008) containing astrocyte differentiation media described in Liddelow et al. (2017). This media consisted of a 1:1 (v/v) mixture of neurobasal media and high glucose DMEM supplemented with sodium pyruvate, glutamax, N2 Max, and N-acetyl-cysteine and with growth factors including 20ng/mL bFGF, 5ng/mL Hb-EGF (R&D, 259-HE-050), 10ng/mL CNTF (R&D, 557-NT-010), and 10ng/mL BMP4 (R&D, 314-BP-050) for 3 days.

Human oligocortical spheroid production

Human embryonic stem cells (line H7, WiCell) were grown in mTesR1 media and oligocortical spheroids were generated with minor modifications to the protocol previously described (Madhavan et al., 2018). Briefly, in the first step of generating oligocortical spheroids, CloneR (Stem Cell Technologies, 5889) was used instead of Y-27632 and Dorsomorphin was replaced with 150nM LDN193189 (Sigma, SML0559). Spheroids were treated with 150nM LDN193189 and 10μM SB-43152 (Sigma, S4317) for the first 6 days followed 20ng/ml FGF-2 (R&D Systems, 233-FB-25/CF) and 20ng/ml EGF (R&D Systems, 236-EG-200) from day 7 to 25. This was followed by 10ng/ml BDNF (R&D Systems, 248-BD) and 20ng/ml NT-3 (R&D Systems, 267-N3) treatment every other day between days 27 and 40. For OPC development and oligodendrocyte differentiation cultures 10ng/ml PDGF-AA (R&D Systems, 221-AA) and 10ng/ml IGF (R&D Systems, 291-GF-200) were added to cultures every other day between days 51 and 60 an 40n/ml T3 (Sigma, T6397) every other day between days 61 and 70. Spheroids were treated every other day with vehicle DMSO or 300nM MEKi between days 70 and 74 and harvested on day 90. Spheroids were treated with 200μM Hypoxyprobe-1 two hours prior to harvesting for IHC (pimonidazole, Hypoxyprobe Inc, Burlington MA, HP1-100Kit).

Tissue from mild chronic hypoxia (MCH) mice

MCH is a well described model of Diffuse White Matter Injury (DWMI) (Clayton et al., 2017a, 2017b; Fancy et al., 2011; Scafidi et al., 2014; Yuen et al., 2014) and protein samples were generously provided by Brian Popko. In brief, postnatal day 3 (P3) C57Bl6 pups were placed into a BioSpherix chamber maintained at 10+/− 0.5% O2 by displacement of nitrogen until P11. Animals were then quickly sacrificed by CO$_2$ asphyxiation followed by decapitation and the frontal cortex was isolated as this region has been shown to contain subcortical white matter that is susceptible to hypoxia-induced DWMI (Clayton et al., 2017b; Jablonska et al., 2012; Sanjana et al., 2014; Yuen et al., 2014).

METHOD DETAILS

In vitro hypoxia experiments

OPCs were plated in OPC growth media and then placed into a 2 shelf C-Chamber from BioSpherix (C-274). Oxygen tension was controlled using the ProOx 110 from BioSpherix such that nitrogen gas would flush out oxygen to maintain the chamber at the desired oxygen level. The subchamber was set at 1% O$_2$ and cells were cultured for 48 hours unless otherwise noted. Hypoxia treated cells were then rapidly lysed for RNA or protein to minimize degradation of HIFs upon exposure to room air. The normoxic controls were cultured concurrently in the same cell culture incubator containing the BioSpherix C-chamber.

Immunocytochemistry

For antigens requiring live staining (O1 and O4), antibodies were diluted in N2B27 base media supplemented with 10% Donkey Serum (v/v) (Jackson ImmunoResearch, 017-000-121) and then added to cells for 18 minutes at 37°C. Cells were then fixed in cold 4% PFA (Electron microscopy sciences) for 18 minutes at room temperature, washed with PBS, and permeabilized and blocked in blocking solution containing 0.1% Triton X-100 in PBS supplemented with 10% normal donkey serum (v/v) for 30 minutes at room temperature. Primary antibodies were diluted in blocking solution and incubated with samples overnight at 4°C. Primary antibodies used included anti-OLIG2 (1.2μg/mL, Proteintech, 12999-1-AP), anti-MBP (1:100, Abcam, ab7349), anti-O1 (1:50, CCF Hybridoma Core), anti-O4 (1:100, CCF Hybridoma Core), anti-ASCL2 (1:10, EMD Millipore, MAB4417), anti-GFAP (1:5000, Dako, Z033401-2), anti-SOX6 (1:2000, abcam, 30455), anti-A2B5 (2μg/mL, Millipore, MAB312), and anti-NKX2-2 (1:200, DSHB, 74.5A5). The next day, cells were rinsed with PBS and incubated in blocking solution for one hour with the appropriate secondary antibody conjugated to an Alexa Fluor (4μg/mL, Thermo Fisher) along with the nuclear stain DAPI (Sigma, 1ug/mL).

High content imaging and quantification

Both 96-well and 384-well plates were imaged using the Operetta High Content imaging and analysis system (PerkinElmer). For 96-well and 384-well plates, 8 fields and 5-fields were captured from each well at 20x magnification respectively. Images were analyzed with PerkinElmer Harmony and Columbus software as described previously (Hubler et al., 2018; Najm et al., 2015). In brief, nuclei of live cells were identified using a threshold for area of DAPI staining to exclude pyknotic nuclei or debris. To identify oligodendrocytes, each DAPI

positive nucleus was expanded by 50% to determine potential intersection with staining of an oligodendrocyte marker (O4/O1/MBP) in a separate channel. Expanded nuclei that intersected O4/O1/MBP staining were scored as oligodendrocytes. Percentage of oligodendrocytes was then calculated by dividing the number of oligodendrocytes by total number of DAPI positive cells per image.

Generation of CRISPR and CRISPRA OPCs

Guide sequences were curated from the Brie library (Doench et al., 2016) and cloned into the CRISPRv2 backbone (Addgene 52961) (Sanjana et al., 2014) for generating CRISPR-mediated knockout OPCs. Guide sequences were curated from the CRISPRav2 library (Horlbeck et al., 2016) and cloned into the lenti-SAMv2 backbone (Addgene 75112) (Joung et al., 2017) for CRISPR activation. The activation helper plasmid (Addgene 89308) (Joung et al., 2017) was co-transduced for all CRISPR activation OPCs. Plasmids containing cloning sites for the sgRNA were digested and annealed oligomers containing the sgRNA were then ligated to their respective backbones. sgRNAs used in this study include: Cas9 control (empty- no sgRNA), sgVhl (GCCCGGTGGTAAGATCGGGT), sgVhl.2 (TGTCCATCGACATTGAGGGA), CRISPRA-NTC (GATCGGTTATGTTTAGGGTT), CRISPRA-Ascl2 (CCAAGTTTACCAGCTTTACG), CRISPRA-Dlx3 (GTTAGGGGTAACAACAAAGA), CRISPRA-Vegfa (GAGGCCGTGGACCCTGGTAA), CRISPRA-Slx16a3 (GCCTGGC CGCTGTTCCCCGA), and CRISPRA-Bnip3 (GGGGGAGCGTGTGGCACGTG).

Positive stbl3 bacterial clones (Thermo Fisher, C737303) were confirmed to have the correct inserted oligomer using Sanger sequencing. HEK293T cells were then transfected using lenti-X shots following the manufacturer's protocol (Clonetech, 631276). The next day, transfection media was switched to OPC growth media without growth factors for virus collection. After 2 days, the media from transfected HEK293T cells was collected, filtered, and supplemented with OPC growth factors PDGF and FGF. This lentivirus-containing media was added to OPCs at a ratio of 1:2 (v/v) with fresh OPC growth media. After 24 hours of incubation with virus, transduced cells were switched to fresh OPC growth media and allowed to recover for 48 hours. OPCs were then selected for 96 hours in OPC growth media supplemented with a lethal dose of puromycin (500ng/mL, Thermo Fisher, A1113802) for CRISPR knockout cells or a lethal dose of blasticidin (10μg/mL, Thermo Fisher, A1113903) and hygromycin (100μg/mL, Thermo Fisher, 10687010) for CRISPR activation cells. OPCs were then allowed to recover for at least 24 hours following removal of selection and frozen down in aliquots for future use. For all experiments, the lentivirally transduced CRISPR/CRISPRA targeting and non-targeting control OPCs were derived from the same batch of OPCs and infected and selected simultaneously. qPCR was performed to validate a reduction or overexpression of gene targets of interest for each batch of CRISPR/CRISPRA OPCs generated.

Generation of HA-Ascl2 OPCs

The lentiviral HA-ASCL2 overexpression construct was obtained from Genecopoeia (EX-Mm03774-Lv117) and transformed into stbl3 bacteria (Thermo Fisher, C737303). Positive clones were selected on ampicillin agar plates and amplified for maxi-prep (QIAGEN, 12163) to obtain sufficient quantities of plasmid. This construct was used to generate lentivirus and transduce OPCs following the same protocol used to generate CRISPR knockout OPCs using puromycin (500ng/mL, Thermo Fisher, A1113802) to select for infected OPCs.

Validating CRISPR knockout in OPCs

Primers were identified surrounding the target cut site for the two sgVhl constructs (see previous section for sgRNAs) that generate 200-250 base pair amplicons.

For the sgVhl cut site the primers were:

F 5′ TCCCTACACGACGCTCTTCCGATCTCTCTCAGGTCATCTTCTGCAACC 3′
R 5′ AGTTCAGACGTGTGCTCTTCCGATCTGACAAGATGCTCGGGGTCGG 3′

For the sgVhl.2 cut site the primers were:

F 5′ TCCCTACACGACGCTCTTCCGATCTAATAAACAGGTGCCATGCCC 3′
R 5′ AGTTCAGACGTGTGCTCTTCCGATCTAGATTGACTATTAACCTGGCAATG 3′

PCR products were run on an agarose gel, excised by gel extraction (28115, QIAGEN), and submitted to the Case Western Reserve University Genomics core for library preparation and sequencing. Libraries were prepared by adding unique indices by PCR using KAPA HiFi HotStart ReadyMix. Samples were then pooled evenly, quantified using NEBNext® Library Quant Kit for Illumina® (New England 641 Biolabs, E7630), and denatured and diluted per Illumina's MiSeq instructions. These finished libraries were then sequenced using an Illumina MiSeq (250bp paired-end). Results were analyzed using Outknocker software (Schmid-Burgk et al., 2014) (http://www.outknocker.org/outknocker2.htm) to calculate the percentage of reads with insertions or deletions at the sgRNA target site.

Kinetics of OPC differentiation to oligodendrocytes

This protocol was adapted from a previous study from our lab (Elitt et al., 2018). Control and CRISPR/CRISRPA OPCs were seeded at 40,000 cells per well in 96-well poly-ornithine and laminin coated plates into oligodendrocyte differentiation medium. At day 1, 2, and 3 post-plating, cells were live stained with mouse anti-O4 (CCF Hybridoma Core Facility) or mouse anti-O1 (CCF Hybridoma Core Facility) followed by fixation with 4% PFA. Cells were then stained using rat anti-MBP (1:100, Abcam, ab7349), followed by counter-

staining with DAPI (1 μg/ml, Sigma, D8417). 6-8 fields were captured per well and the percentage of MBP+ cells were quantified relative to total cell number (DAPI) to control for any possible variations in plating density.

Compound screening and assessment

Compound screening was carried out as described in Lager et al. (2018). Poly-D-lysine-coated 384-well CellCarrier Ultra plates (PerkinElmer) were coated with laminin diluted in N2B27 base media using an EL406 Microplate Washer Dispenser (BioTek) equipped with a 5μl dispense cassette (BioTek) and were incubated at 37°C for at least 1 hour. A 3mM stock of the Selleck bioactive library in dimethylsulfoxide (DMSO) was then added to the plates using a 50nL solid pin tool attached to a Janus automated workstation (Perkin Elmer) at a 1:1000 dilution such that each well received a single compound at a final concentration of 3μM. Compounds for dose response validation were sourced from the Selleck library, except for ERK1/2 inhibitors SCH772984 (Selleck, S7101), AZD0364 (Selleck, S8708), and VX-11e (Selleck, S7709) and MEK5 inhibitors BIX02188 (Selleck, S1530) and BIX02189 (Selleck, S1531) which were purchased separately. OPCs were dispensed in oligodendrocyte differentiation media at 12,500 cells per well into the laminin-coated 384 well plates using the BioTek EL406 Microplate Washer Dispenser and differentiated at 37°C for 3 days. At this point, cells were fixed, washed and stained using the BioTek EL406 Microplate Washer Dispenser. Cells were stained with anti-MBP (1:100, Abcam, ab7349) along with DAPI (1 μg/ml, Sigma, D8417) and imaged using the Operetta High Content Imaging and Analysis system (PerkinElmer).

Western blot

For cell culture derived protein samples, at least 1 million OPCs were collected and lysed in RIPA buffer (Sigma, R0278) supplemented with protease and phosphatase inhibitor (Thermo Fisher, 78441) for at least 15 minutes and cleared by centrifugation at 13,000 g at 4°C for 15 minutes. Protein concentrations were determined using the Bradford assay (Bio-Rad Laboratories). Protein was then diluted and added to Laemmli loading buffer, boiled at 95°C for 5 minutes, run using NuPAGE Bis-Tris gels (Thermo Fisher, NP0335BOX), and then transferred to PVDF membranes (Thermo Fisher, LC2002). Blocking and primary/secondary antibody solutions were performed for at least 30 min with 5% nonfat drug milk (Nestle Carnation) in TBS plus 0.1% Tween 20 (TBST). Primary antibodies used included anti-HIF1a (1:500, Abcam, ab2185), anti-phospho-p44/42 (ERK1/2) (100ng/mL, CST, 9101), anti-p44/42 (total ERK1/2) (100ng/mL, CST, 4696), anti-DLX3 (2μg/mL, abcam, ab178428), anti-SOX10 (1:100, R&D, AF2864), anti-B-Actin peroxidase (1:50,000, Sigma, A3854), anti-VHL (5μg/mL, BD Biosciences, 556347), anti-ASCL2 (1:10, EMD Millipore, MAB4417), anti-MBP (1μg/mL, Biolegend, 808401), anti-OLIG2 (100ng/mL, Millipore, MABN50), anti-HA (200ng/mL, Cell Signaling Technologies, 3724S), and anti-MAG (1μg/mL, Thermo Fisher, 346200). Membranes were then imaged using the Odyssey Fc Imaging System (LI-COR) and analyzed using Image Studio™ software that is integrated into the Odyssey imaging suite. Westerns were normalized to loading control Beta-Actin unless otherwise noted.

Co-immunoprecipitation

For immunoprecipitation, 15 million OPCs were lysed in co-IP buffer (50mM Tris-HCL, pH 7.4, 120mM NaCl, 1mM EDTA, 1% Nonidet P-40 with Complete EDTA-free protease inhibitor tablet) for 15 minutes with intermittent vortexing. Lysate was cleared by centrifugation at 16,000 g at 4°C for 15 minutes. The supernatant was divided into three groups for input, immunoprecipitation, and IgG control. Samples were rotated at 4°C for 2 hours with 10uL of Protein G magnetic Dynabeads (Thermo Fisher, 10004D) to remove off target binding. Beads were removed by magnet. The supernatant of the input sample was stored at −80°C while the supernatant of the immunoprecipitation and IgG samples were combined with 15μg of HIF1a antibody (abcam, 2185), 10μg of OLIG2 antibody (Millipore, ab9610), or 2μg of normal rabbit IgG antibody (Cell signaling technologies, 2729S) and rotated overnight at 4°C. The following day, 20μL of protein G beads were added to the mixture and rotated overnight at 4°C. Samples were placed on a magnetic bar to remove supernatant, and beads were washed with 1mL of co-IP buffer 6 times. On the final wash, all supernatant was removed and 50uL of Laemmeli buffer was added. Sample was heated at 95°C for 5 minutes and collected to run for western blot analysis.

qRT-PCR

At least 500,000 OPCs were lysed in TRIzol (Ambion) followed by phenol-chloroform extraction and processing with the RNeasy Mini Kit (QIAGEN, 74104). RNA quality and quantity was determined using a NanoDrop spectrophotometer. cDNA was generated using the iSCRIPT kit following the manufacturer's instructions (Biorad, 1708891). qRT-PCR was performed using pre-designed TaqMan gene expression assays (Thermo Fisher, 4369016). qPCR was performed using the Applied Biosystems 7300 real-time PCR system and probes were normalized to *Rpl13a* endogenous control. Taqman probe IDs and cat numbers used in the study include: *Sox10* (Mm00569909_m1), *Rpl13a* (Mm05910660_g1), *Vhl* (Mm00494137_m1), *Ascl2* (Mm01268891_g1), *Tpi1* (Mm00833691_g1), *Ciart* (Mm01255905_g1), *P4ha2* (Mm00477940_m1), *Pdk1* (Mm00554300_m1), *Pfkl* (Mm00435587_m1), *Slc16a3* (Mm00446102_m1), *Dlx3* (Mm00438428_m1), *Vegfa* (Mm00437306_m1), *Bnip3* (Mm01275600_g1), *Pdgfra* (Mm00440701_m1), *Plp1* (Mm01297210_m1), and *Myrf* (Mm01194959_m1). As the taqman probe for *Ascl2* bind regions outside of the *Ascl2* cDNA sequence, SYBR qPCR (Kappa Biosystems, KK4605) was performed to verify the increase in *Ascl2* expression in HA-ASCL2 OPCs using *Rpl13a* as a loading control with the following primers: *Ascl2* (F: ATGGAAGCACACCTTGACTG, R: AGTGGACGTTTGCACCTTCA) *Rpl13a* (F: TTCTCCTCCAGAG TGGCTGT, R: GGCTGAAGCCTACCAGAAAG).

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Cell viability assay

OPCs were seeded into 96 well plates at 25,000 cells/well (96-well plate, Fisher, 167008) and recovered for 24 hours. Cell viability was assessed using an MTS assay kit (abcam, ab197010) following the manufacturer's protocol. Colorimetric measurements were recorded 9 hours after the addition of the MTS dye solution to the OPCs.

siRNA knockdown in OPCs

siRNAs were purchased from Horizon/Dharmacon (Ascl2: L-047262-01-0005, Dlx3: L-041957-01-0005, Non-Targeting Control: D-001206-14-05 5 nmol) and nucleofected into control and sgVhl OPCs using the Basic Nucleofector Kit for Primary Mammalian Glial Cells (Lonza, VPI-1006) following manufacturer's instructions. In brief, 5 million OPCs were resuspended in nucleofection solution with 0.6μl of the desired 50μM siRNA solution, nucleofected using the Amaxa Nucleofector 2b (Lonza, AAB-1001) using the A033 setting, and then plated in differentiation media supplemented with penicillin-streptomycin (Thermo Fisher, 15070063) for 72 hours. Leftover OPCs were plated in growth media supplemented with penicillin-streptomycin (Thermo Fisher, 15070063) for 48 hours and then lysed with TRIzol (Ambion, 15596026) for RNA extraction and qPCR to assess knockdown efficiency.

RNA-seq sample preparation and analysis

At least 1 million OPCs were lysed in TRIzol and RNA was isolated as described for qPCR. Libraries were prepared following protocols from NEBNext Poly(A) mRNA Magnetic Isolation Module (NEB, E7490L) and NEBNext Ultra RNA Library Prep Kit for Illumina (NEB, E7530L). In brief, samples were enriched for mRNA using oligo(dT) beads, which were then fragmented randomly and used for cDNA generation and subsequent second-strand synthesis using a custom second-strand synthesis buffer (Illumina), dNTPs, RNase H and DNA polymerase I. cDNA libraries then went through terminal repair, A-base ligation, adaptor ligation, size selection and PCR enrichment. Final libraries were pooled evenly and sequenced on the Illumina NovaSeq with paired-end 150bp reads with a read-depth of at least 20 million reads per sample.

For gene expression analysis, reads were aligned to the mm10 genome build using salmon 0.14.1 (https://github.com/COMBINE-lab/salmon) (Patro et al., 2017) to quantify transcript abundance in transcripts per million (TPM) values. Transcripts were summarized as gene-level TPM abundances with tximport. A gene with TPM > 1 was considered expressed. Differential expression analysis was then performed using DESEQ2 (https://bioconductor.org/packages/release/bioc/html/DESeq2.html) (Love et al., 2014). Significant genes were called based on p-adj and fold change values as described in the results section.

Violin plots

For violin plots in Figures 2 and 6 and S2, RNaseq replicates were divided by the average of the respective control (TPM) on a per gene basis in each category and then all individual replicates were plotted together. For Figure S2, the average expression for each gene in each category in sgVhl OPCs was divided by the average expression in control OPCs. In all cases the genes in each category were limited to those that were expressed in sgVhl OPCs (TPM > 1) and all TPM values were floored to 1.

Gene ontology analysis and heatmaps

Metascape (http://metascape.org/) (Zhou et al., 2019) was used to identify significant pathways from RNaseq data. Glycolysis genes used to construct the heatmap in Figure 2F were taken from Panther ("Glycolysis: P00024"). In Figure 6F, RNA-seq data was fit to the genes within "Oligodendrocyte Development" (GO: 0014003) that were downregulated in sgVhl OPCs compared to Cas9 control OPCs (fold change less than 0.75).

Gene set enrichment analysis

Gene set enrichment analysis (GSEA) scores were generated for gene sets in C5.bp datasets using classic scoring, 1000 gene-set permutations, and signal-to-noise metrics. Normalized enrichment scores, false discovery rate and FWER o-values were all calculated by GSEA software (https://www.gsea-msigdb.org/gsea/index.jsp) (Subramanian et al., 2005).

ChIP-seq and analysis

Nuclei isolation and chromatin shearing were performed using the Covaris TruChIP protocol following manufacturer's instructions for the "high-cell" format. In brief, 5-20 million cells (H3K27Ac) or 100 million cells (HIF1a) were crosslinked in "Fixing buffer A" supplemented with 1% fresh formaldehyde for 10 minutes at room temperature with oscillation and quenched for 5 minutes with "Quench buffer E." These cells were then washed with PBS and either snap frozen and stored at −80°C, or immediately used for nuclei extraction and shearing per the manufacturer protocol. The samples were sonicated with the Covaris S2 using the following settings: 5% Duty factor 4 intensity for four 60 s cycles. Sheared chromatin was cleared and incubated overnight at 4 degrees with primary antibodies that were pre-incubated with protein G magnetic DynaBeads (Thermo Fisher, 10004D). Primary antibodies used included anti-H3K27Ac (9μg/sample, Abcam, ab4729), anti-HIF1a (25μg/sample, Abcam, ab2185), anti-Olig2 (15μg/samples, Millipore, ab9610), and anti-HA (15μg/sample, cell signaling technologies, 3724S). These beads were then washed, eluted, reverse crosslinked and treated with RNase A followed by proteinase K. ChIP DNA was purified using Ampure XP beads (Aline Biosciences, C-1003-5) and then used to prepare Illumina sequencing libraries as described previously (Schmidt et al., 2009). Libraries were sequenced on the Illumina HiSeq2500 with single-end 50bp reads with a read-depth of at least 20 million reads per sample.

For peak calling, reads were quality and adaptor trimmed using Trim Galore! Version 0.4.1. Trimmed reads were aligned to mm10 with Bowtie2 version 2.3.2 (Langmead and Salzberg, 2012). Duplicate reads (potential artifacts of PCR in library preparation) were removed using Picard MarkDuplicates. Peaks were called with MACS version 2.1.1 (Feng et al., 2012) to define broad peaks for histone marks (H3K27Ac) and narrow peaks for transcription factors (HIF1a), OLIG2, and HA-ASCL2 and normalized to background genomic DNA with matched inputs. Thresholding was set at FDR < 0.001 for calling both H3K27Ac, HIF1a, and OLIG2 peaks and was set at FDR < 0.1 for calling HA-ASCL2 peaks. Peaks were visualized with the Integrative Genomics Viewer (IGV, Broad Institute). Peaks were assigned to the nearest expressed gene (TPM > 1 in Cas9 control or sgVhl OPCs) using bedtools available in Galaxy (https://usegalaxy.org) (Afgan et al., 2018).

Omni ATAC-seq

Omni ATAC-Seq was performed on 50,000 OPCs in two independent biological replicates following the protocol outlined in Corces et al. (2017). In brief, nuclei were extracted from cells and treated with transposition mixture containing Nextera Tn5 Transposase for (Illumina, FC-121-1030). Transposed fragments were then purified using QIAGEN MinElute columns (QIAGEN, 28004), PCR amplified, and libraries were purified with Agencourt AMPure XP magnetic beads (Aline Biosciences, C-1003-5) with a sample to bead ratio of 1:1.2. Final libraries were sequenced on the illumine HiSeq2500 with single-end 50bp reads with nearly 100 million reads per sample. Reads were aligned to the mm10 mouse genome following the same pipeline used for ChIP-seq data (see ChIP-seq and analysis) and peaks were called using the "narrowPeaks" function of MACS version 2.1.1 as outlined for HIF1a ChIP-seq (see ChIP-seq and analysis).

Diffbind analysis

H3K27Ac and Olig2 ChIP-seqs were performed in duplicate from two independent batches of Cas9 control and sgVhl OPCs. Differential H3K27Ac and Olig2 analysis between sgVhl and Cas9 control OPCs was performed using "Diffbind software" (https://bioconductor.org/packages/release/bioc/html/DiffBind.html) (Ross-Innes et al., 2012). A false discovery rate of 0.1 was used to call significantly enriched and depleted regions of H3K27Ac and 0.001 for Olig2.

HIF1a ChIP-seq overlap analysis

HIF1a ChIP-seq raw data were re-analyzed for: two HIF1a replicates ChIP-seqs and 1 input of E12.5 heart (GSM1500750, GSM150075, GSM1500749) (Guimarães-Camboa et al., 2015), two HIF1a replicates ChIP-seqs and 1 input of melanocyte (GSM2305570, GSM2305571, GSM2305572) (Loftus et al., 2017), and two HIF1a ChIP-seqs and 2 inputs of Th17 cells (GSM1004819, GSM1004991, GSM1004820, GSM1004993) (Ciofani et al., 2012). For each of the HIF1a ChIP-seqs, the peaks were called as described above (MACS2, narrow peak, FDR < 0.001). Peaks unique to OPCs were identified by finding peaks not found in the union of all the replicates of every other tissue sample using the bedtools (v 2.25.0) (Quinlan and Hall, 2010) "intersect" command. Common peaks across tissues were also identified using the bedtools "intersect." RPCG-normed bigwigs were used to create aggregate plots for HIF1a, ATAC-seq, and H3K27ac using the deeptools (version 3.3.1) (Ramírez et al., 2016) "computeMatrix" and "plotHeatmap" commands, centering on the HIF1a peak locations in the sgVhl OPCs. Genes within 5Kb of these HIF1a peaks were called as HIF1a target genes. For melanocytes and T cells, HIF1a target genes were called as only those at the intersection of both of the available replicates. For embryonic heart and OPCs, there were fewer peaks called compared to the other tissues, so HIF1a gene targets were called as the union of the two available replicates. These gene and peak lists were used to generate venn diagrams using the following webtool: (http://bioinformatics.psb.ugent.be/webtools/Venn/). Cell type specific gene targets of HIF1a were called if the gene was only a target of HIF1a in that specific cell type (see Table S2).

Motif enrichment analysis

Motifs were called under significant HIF1a peaks (FDR < 0.001), OLIG2 peaks (FDR < 0.001), and HA-ASCL2 peaks (FDR < 0.1) or regions of significantly gained H3K27Ac (FDR < 0.1) or OLIG2 (FDR < 0.05) using HOMERv4.11.1 (Heinz et al., 2010). The FindMotifsGenome.pl tool was used with mm10 as the reference genome. Known motifs were compiled into Tables S3 and S4 and ranked based on p value. To calculate enrichment of motifs under tissue-specific HIF1a peaks, the percentile of the rank for each motif from HOMER was calculated in all four tissue types (OPCs, heart, melanocytes, and T cells). The average percentile for a motif in the three other tissues (heart, melanocyte, and T cells) was then subtracted from the percentile of the same motif in OPCs. The magnitude of the difference was then ranked and filtered for transcription factors expressed in OPCs to obtain the top motifs enriched under OPC-specific HIF1a peaks compared to other tissue types.

Mass spectrometry for lactate and pyruvate

OPCs were cultured in OPC growth media and after growth media was removed, cells were washed with ice-cold saline (3 times) and collected with 1 mL of 80% ethanol pre-chilled on dry ice. Cells were frozen at −80°C until analyses. This cell extract was then vortexed and sonicated for 30 s on, 30 s off, alternating for 10 minutes. Next, cells were pelleted by centrifugation at 4°C for 10 min at 14,000 rpm. Supernatant was transferred to GC/MS vials and evaporated to dryness under gentle stream of nitrogen. Keto- and aldehyde groups were reduced by addition of 10 μl of 1 N NaOH plus 15 μl NaB^{2}H$_4$ (prepared as 10 mg/ml in 50 mM NaOH). After mixing, samples were incubated at room temperature for 1 hour and then acidified by 55 μl of 1 N HCl by dropping the acid slowly. Next, samples were evaporated to dryness. Next, 50 μl of methanol were added to precipitate boric acid. Internal standard was added

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(10 μl of 17:0 FA, 0.1 mg/ml). Samples were evaporated to dryness and reacted with 40 μl of pyridine and 60 μl of tert-butylbis(dimethylsilyl) trifluoroacetamide with 10% trimethylchlorosilane (Regisil) TBDMS at 60°C for 1 hour. Resulting TBDMS derivatives were injected into GC/MS. Analyses were then carried out on an Agilent 5973 mass spectrometer equipped with 6890 Gas Chromatograph. A HP-5MS capillary column (60 m × 0.25 mm × 0.25 μm) was used in all assays with a helium flow of 1 ml/min. Samples were analyzed in Selected Ion Monitoring (SIM) mode using electron impact ionization (EI). Ion dwell time was set to 10 msec. Lactate and pyruvate were both measured for all samples.

Oligocortical spheroid immunohistochemistry

Oligocortical spheroids were treated every other day with DMSO or 300nM AZD8330 between days 70 and 74 and harvested on day 90. 2 hours prior to harvesting, spheroids were treated with 200uM Hypoxyprobe-1 (pimonidazole, Hypoxyprobe Inc, Burlington MA, HP1-100Kit). Spheroids were washed in PBS and fixed overnight in ice cold 4% Paraformaldehyde (Electron Microscopy Sciences, 15710) and then washed in PBS and cryoprotected in a 30% sucrose solution. Spheroids were frozen in OCT and sectioned at thickness of 15 μm. Slides were washed in PBS and incubated overnight with anti-MyRF (1:1000, gift from Michael Wegner) and anti-pimonidazole antibodies (1:250, Hypoxyprobe Inc, Burlington MA, HP1-100Kit) followed by labeling with appropriate Alexa Fluor labeled secondary antibodies (2μg/mL, Thermo Fisher). Images were captured using a Hamamatsu Nanozoomer S60 Slide scanner with NDP 2.0 software. Images spanning the edge to the central region of each oligocortical spheroid were used for analysis. SOX10+ cells and MyRF+ oligodendrocytes in Hypoxyprobe positive and Hypoxyprobe negative areas within each image were quantified using ImageJ software.

QUANTIFICATION AND STATISTICAL ANALYSIS

GraphPad Prism was used to perform statistical analyses unless otherwise noted. Statistical tests and replicate descriptions are detailed in each figure legend. In brief, black filled-in circles for bar graphs indicate biological replicates whereas open circles represent technical replicates. Statistics were only performed on samples with biological replicates. Data is typically graphed as mean ± standard deviation (SD) or ± standard error of the mean (SEM) as detailed in the figure legend. A p value less than 0.05 was considered significant unless otherwise noted.