Activating Endogenous Neural Precursor Cells Using Metformin Leads to Neural Repair and Functional Recovery in a Model of Childhood Brain Injury

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SUMMARY

The development of cell replacement strategies to repair the injured brain has gained considerable attention, with a particular interest in mobilizing endogenous neural stem and progenitor cells (known as neural precursor cells [NPCs]) to promote brain repair. Recent work demonstrated metformin, a drug used to manage type II diabetes, promotes neurogenesis. We sought to determine its role in neural repair following brain injury. We find that metformin administration activates endogenous NPCs, expanding the size of the NPC pool and promoting NPC migration and differentiation in the injured neonatal brain in a hypoxia-ischemia (H/I) injury model. Importantly, metformin treatment following H/I restores sensory-motor function. Lineage tracking reveals that metformin treatment following H/I causes an increase in the absolute number of subependyma-derived NPCs relative to untreated H/I controls in areas associated with sensory-motor function. Hence, activation of endogenous NPCs is a promising target for therapeutic intervention in childhood brain injury models.

INTRODUCTION

Perinatal hypoxic-ischemic (H/I) insult is an important cause of brain injury and can result in long-term neurological complications, ranging from mild behavioral deficits to severe seizure, intellectual disability, and/or cerebral palsy. Neural injury results from excitotoxicity, inflammation, and oxidative stress, leading to a loss of oligodendrocytes and neurons, which are particularly sensitive to H/I (Deng, 2010). With the exception of rehabilitation, there are no established treatments available to improve behavioral deficits in these patients to date.

The development of cell replacement strategies to repair the injured brain has gained considerable attention, with a particular interest in mobilizing endogenous neural precursor cells (NPCs). Several studies have investigated the impact of H/I on the germinal zone, a region that contains multipotent NPCs. It has been shown that neural stem cells (NSCs) remain resilient to this insult (Romanko et al., 2004) and can regenerate neurons and oligodendrocytes following H/I, albeit to a very limited extent (Felling et al., 2006; Ong et al., 2005; Plane et al., 2004; Yang and Levison, 2006; Yang et al., 2007; Zaidi et al., 2004). This injury-induced activation leads to a rare fraction of cells migrating to the injury site, which is not sufficient for self-repair (Ikedo et al., 2005). With the goal of developing new methodologies to enhance NPC activation in the neonatal brain and promote neural repair following H/I, we have built on successful NPC activation strategies that lead to tissue formation and functional recovery using small molecules to mobilize endogenous NPCs (Erlandsson et al., 2011; Kolb et al., 2007). Previous work has demonstrated that atypical PKC-dependent phosphorylation of CREB-binding protein (CBP) on serine 436 is necessary for enhancing differentiation in cortical precursor cells (Wang et al., 2010). Moreover, this pathway is activated by metformin, an AMPK agonist, shown to promote neurogenesis and enhance spatial memory in uninjured mice (Wang et al., 2012). Based on these findings, we asked whether metformin could be used to stimulate subependyma (SE)-derived NPCs and promote repair of the H/I-injured neonatal murine brain.

RESULTS

Metformin Expands the Stem Cell Pool and Promotes Differentiation via the aPKC-CBP Pathway in Neonatal Pups, In Vitro

It is currently unknown whether metformin acts on the NSCs, their downstream progeny, or both that ultimately leads to enhanced neurogenesis. To test if metformin had
direct effects on NSCs, we used an in vitro, colony-forming assay, termed the neurosphere assay on postnatal day 8 (P8) pups from C57/BL6 (control) and CBP S436A mice (CBP S436A mutants harboring a point mutation from serine to alanine in the CBP gene specifically at the aPKC phosphorylation site) (He et al., 2009). Phosphorylation at serine 436 has been shown to be essential for neural differentiation in embryonic precursors; hence, we predicted that direct effects of metformin on NPCs would not be seen in CBP S436A mutant mice. Briefly, the SE is dissected, dissociated, and plated into single cells. Over a period of 7 days, single NSCs proliferate to form a free-floating colony of cells known as a neurosphere. Each individual neurosphere is a mixed population of stem and progenitor cells (Morshead et al., 2002). In primary culture, an increase in the total numbers of neurospheres indicates an expansion of the stem cell pool. In the presence of metformin, we observed a significant (>2-fold) increase in the numbers of primary neurospheres in control mice, whereas this effect was lost in CBP S436A mutant mice (Figure 1A). Further, primary P8 SE-derived neurospheres grown and differentiated in culture in the presence of metformin gave rise to significantly more oligodendrocytes (1.9-fold increase) and neurons (3.3-fold increase) compared to controls (Figures 1B–1D), and this effect was lost for CBP S436A mutants cultured in the same conditions (data not shown). Thus, metformin expands the stem cell pool and promotes oligodendrogenesis and neurogenesis from the early postnatal brain via the aPKC-CBP pathway.

**Functional Recovery in Injured Mice following Metformin Treatment**

We asked if administration of metformin was able to enhance brain repair in a well-established model of neonatal brain injury (Vannucci and Vannucci, 2005; Vannucci et al., 1999). P8 pups received a left common carotid artery ligation followed by hypoxia for 1 hr. H/I-injured pups revealed sensory-motor deficits, with increased latencies in the righting reflex motor task and decreased latencies in the hind limb suspension task at 1 hr and 4 days post-H/I, respectively, when compared to uninjured pups (Figures 2A–2C; Figure S1). Metformin was delivered to neonatal pups starting 24 hr post-H/I via their lactating mothers, which had 7-day micro-osmotic pumps implanted subcutaneously (20 mg/kg/day), or by subcutaneous injection directly into the pups (20 mg/kg/day). We first assayed for the effects of H/I and/or metformin treatment on SE-derived NPC behavior. Uninjured pups that received metformin through the mother’s milk for 4 days (from P9–12) had a 2-fold increase in the number of neurospheres (stem cell pool expansion), similar to what was observed in culture (Figure 1A). As predicted, subcutaneous metformin injections given to CBP S436A mutant mice did not result in expansion of the stem cell pool (Figure 2D).
H/I injury alone increased the size of the NSC pool (4-fold increase) both ipsilateral (IL) and contralateral (CL) to the ischemic insult. Interestingly, H/I + metformin-treated pups gave rise to reduced numbers of SE-derived neurospheres relative to the H/I treatment (although similar to metformin-only-treated mice) (Figure 2E). One potential explanation for this relative decrease is that metformin promoted the migration of endogenous NPCs into the surrounding parenchyma. In support of this hypothesis, when we performed the neurosphere assay from cortical tissue, we observed a complete lack of neurospheres from control (uninjured) and metformin-only-treated mice, whereas neurospheres were consistently generated from the frontal cortices of H/I + metformin-treated pups (5.0 ± 2.5 neurospheres/cortices).

We then asked whether metformin treatment was able to promote functional recovery. Notably, 4 days of metformin treatment (P9–12) was not sufficient to improve the motor impairment in the hind-limb suspension task in H/I-injured pups (Figure 2C). In contrast, mice that received 7 days of metformin treatment (P9–15) displayed significant recovery in forelimb function when tested in the cylinder task 2 weeks after the lesion (P22) (Figure 2F). Mice that experienced H/I alone displayed a significant preference for use of the IL, unimpaired paw. Indeed, the performance of metformin-treated mice was not statistically different from uninjured controls. To ensure the functional recovery observed in metformin-treated mice was not due to secondary effects, H/I-injured pups received subcutaneous injections of metformin from P9–15. As was seen following exposure to metformin via the mother's milk, we observed recovery in metformin-injected H/I mice, indicating a direct effect of metformin. Next, we asked if this functional recovery was mediated by activation of the aPKC-CBP pathway. Most strikingly, CBP S436A mutants that received subcutaneous injections of metformin from P9–15 following H/I injury did not display functional recovery compared to CBP S436A
Metformin Promotes Expansion, Migration, and Differentiation of Endogenous NPCs in the Injured Brain

To determine the cellular basis for this restored function on the sensory-motor task, we performed lineage-tracking experiments using Nestin-CreERT2/R26R-YFP transgenic mice. When these mice are exposed to tamoxifen, this leads to recombination and expression of YFP in nestin-expressing NPCs and all of their progeny. To perform the lineage tracing, we provided tamoxifen to lactating mothers in their food chow, and thereby exposed their neonatal pups to tamoxifen shortly after birth. At P8, prior to H/I, the vast majority of YFP-positive cells were confined to the SE in the brains of these offspring (Figure S2A). Groups of tamoxifen-treated pups received H/I with or without metformin treatment and their brains were examined at P16 (immediately after metformin treatment, Figures S2B–S2P) and at P23 (time of observed functional recovery, Figure 3A). At both day 16 and 23, H/I + metformin-treated mice had more SE-derived YFP-positive cells in all areas examined, in both the IL and CL parenchyma. At the time of functional recovery (P23), H/I + metformin pups displayed significant increases in YFP-positive cells in the olfactory bulb and subventricular zone and an increasing trend in the hippocampus where NPCs and ongoing neurogenesis are normally observed (Figures S2G–S2K). Similarly, we observed significant increases in YFP-positive cells in regions of the brain known to play a role in motor activities (striatum, corpus callosum, and the motor cortex) compared to H/I-only- and metformin-only-treated mice (>4-fold in all areas) (Figures 3B and 3E). Indeed, H/I-only and metformin-only brains had very few YFP-positive cells in the parenchyma and were not significantly different from each other (Figure 3B).

We examined the differentiation profile of the SE-derived, YFP-positive cells by counting the numbers of differentiated cell types within the parenchyma. Further, we observed no difference in the relative percentage of differentiated cell types within the parenchyma. We found the relative distribution of SE-derived YFP+ cells to be similar between H/I-only- versus H/I + metformin-treated brains, with the majority of cells found in the olfactory bulb (Figure S2M). Taken together, these findings suggest that it is the magnitude of the contribution of newborn cells to the brain tissue that correlates with the functional recovery observed in H/I + metformin-treated pups (Figure S2N).

DISCUSSION

Our findings indicate that endogenous NPC activation following injury is able to promote self-repair and functional recovery in a rodent model of childhood brain injury. We demonstrate metformin treatment in the neonatal brain for 7 days can lead to expansion and migration into the brain parenchyma of NPCs that play a role in sensory-motor function.

Drug therapy for the H/I-injured brain has focused largely on anti-oxidants (allopurinol, N-acetylcysteine), anti-excitotoxic agents (topiramate, memantine), and anti-inflammatory agents (cromolyn) (Juul and Ferriero, 2014; Hagberg et al., 2015). These therapies have been shown to primarily mediate their effects through neuroprotection and have resulted in varying degrees of functional recovery and a reduction in infarct volume. Further, while they display similarities in cost-effectiveness, implications in different injury models, and success in adults or old age animals, metformin is the only drug treatment that has demonstrated efficacy in generating new cells in the neonatal brain. Metformin may confer additional advantages such as improving lifespan (Martin-Montalvo et al., 2013) and promoting angiogenesis (Jin et al., 2014; Liu et al., 2014; Venna et al., 2014).

The finding that metformin, an anti-diabetic drug, increases the absolute number of NPCs following injury (rather than changing the relative percentage of differentiated cells) was surprising. Several possibilities could
account for this observation. For example, metformin could have direct effects on NPCs (such as changes in cell-cycle kinetics and/or promoting cell survival) or indirect effects on non-NPCs that leads to the release of growth/trophic factors, providing immunomodulation, neuroprotection, and/or angiogenesis, which could effectively lead to an

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Figure 3. Metformin Increases the Absolute Numbers of SE-Derived NPCs in the Brain Parenchyma following Injury

(A) Experimental design for metformin administration and behavioral testing. Lineage tracing of SE-derived cells used Nestin-CreERT2/R26R-YFP transgenic mice.

(B) Quantification of SE-derived YFP+ cells found in the striatum (n ≥ 3 mice/group, two-way ANOVA), motor cortex (n ≥ 3 mice/group, two-way ANOVA), and corpus callosum (n ≥ 3 mice/group, two-way ANOVA) is shown.

(C) Quantification of YFP+ cells that were co-labeled with NeuN, indicating newborn neurons, in striatum (n ≥ 3 mice/group, two-way ANOVA) and motor cortex (n ≥ 3 mice/group, two-way ANOVA) is shown.

(D) Quantification of YFP+ cells that were co-labeled with Olig2+ or MBP+, indicating newborn immature or mature oligodendrocytes, respectively, in striatum (n ≥ 3 mice/group, two-way ANOVA), motor cortex (n ≥ 3 mice/group, two-way ANOVA), and corpus callosum (n ≥ 3 mice/group, two-way ANOVA) is shown.

(E–G) Coronal sections immunostained for YFP+ cells (green) and Hoechst (blue) (E); YFP+ (green), NeuN (red), and Hoechst (blue) (F); and YFP+ (green), Olig2+/MBP+ (red), and Hoechst (blue) (G) are shown.

(H) The relative percentages of differentiated neurons (striatum and motor cortex) and differentiated oligodendrocytes (striatum, motor cortex, and corpus callosum) were not significantly different between groups. IL and CL hemispheres were pooled together (n ≥ 4 mice/group, t test). Arrows denote labeled cells. Scale bars, 20 μm. Data are presented as mean ± SEM. TAM, tamoxifen; ST, striatum; MC, motor cortex; CC, corpus callosum. *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001.

See also Figure S2.
expansion in the size of the NPC pool. Indeed with respect to the latter, recent studies using chronic metformin treatment in adult ischemia models have demonstrated enhanced angiogenesis/neurogenesis correlating with motor function recovery (Jin et al., 2014; Liu et al., 2014). Given that the developing brain is known to exhibit a greater degree of plasticity (Kolb and Gibb, 2011) and a greater number of resident NPCs compared to the adult brain (Sachewsky et al., 2014), it is plausible that similar mechanisms of enhanced angiogenesis and neurogenesis are occurring in our H/I injury model.

Metformin is a safe, FDA-approved drug, which supports the translation of these findings to the clinic in a reasonable time frame. The therapeutic window for metformin treatment and long-term outcomes, both at the cellular and behavioral levels, will be important considerations for future studies. Further, studies to determine whether metformin treatment can ameliorate cognitive impairments that result following H/I will be important to pursue. Our findings provide compelling support for exploring endogenous NPC activation using metformin for the treatment of childhood brain injury.

**EXPERIMENTAL PROCEDURES**

**Animals**

C57/BL6, CD1 pregnant mice were purchased (Charles River Laboratories) and maintained under pathogen-free conditions. CBP S46A and Nestin-CreERT2/R26R-YFP transgenic mice were bred in laboratories and maintained under pathogen-free conditions. CBP S46A and Nestin-CreERT2/R26R-YFP transgenic mice were bred in an animal facility. Tamoxifen at a dose of 0.5 mg/kg was mixed in with rodent chow. All mice were kept on a 12-hr day/night cycle, and procedures were performed in accordance with institutional guidelines approved by the Experimental Animal Committee at the University of Toronto.

**H/I Injury Model**

Briefly, the left common carotid artery of P8 pups was ligated to give an ischemic insult. Following a 1.5-hr recovery, they were subjected to hypoxia for 1 hr at a temperature of 36°C. Pre-established endpoints were excluded from statistical analysis.

**Metformin Administration**

For the in vitro neurosphere assay, metformin stock solution was prepared by dissolving in serum-free media (SFMs) and added to culture at various concentrations. In vivo metformin was administered, through a micro-osmotic pump implanted subcutaneously into the mother, at a concentration of 20 mg/kg/day for 7 days. The pups received metformin through the mother’s milk. For injection studies, metformin at the above concentration was injected subcutaneously into the pups daily for 7 days.

**Neurosphere and Cortical Assay**

NSCs were isolated by dissecting the neonatal SE from P8 and P12 animals, as previously described (Morshead et al., 2002), and grown in the presence of mitogens for 7 days. For differentiation assays, individual neurospheres were collected from each condition and plated in the presence of 10% fetal bovine serum (FBS) and the presence and absence of metformin for 7 days. For the cortical assay, the SE was carefully dissected and discarded to ensure no contaminating SE-derived neurospheres were grown in culture. Portions of the cortex were then dissected and processed.

**Tissue Preparation and Immunohistochemistry**

Animals were sacrificed with an overdose of sodium pentobarbital, perfused with 4% paraformaldehyde at different time points (P8, P16, and P23), and cryoprotected in 20% sucrose overnight at 4°C. Coronal, cryostat sections (20 μm) were mounted on Superfrost Plus slides and stained. Briefly, brain sections were rehydrated for 5 min and permeabilized using 0.3% Triton in PBS for 20 min, rinsed, and then blocked with 1% BSA containing 0.3% Triton in PBS at room temperature. Brain sections were incubated with primary antibodies overnight at 4°C in 1% BSA with 0.3% Triton in PBS. The next day slides were washed and incubated with an appropriate secondary antibody for 2 hr at room temperature, washed, and mounted with DAPI-mounting media. A Zeiss Observer D1 inverted microscope was used to visualize immunofluorescence for differentiated neurospheres and tissue sections. Images were acquired at 20× objective using Axio Vision.

**Behavioral Testing**

The righting-reflex, hind-limb suspension test was performed as described previously (El-Khodor et al., 2008; Ten et al., 2003). The cylinder-rearing test was used evaluate functional asymmetry resulting from unilateral brain lesion and consequent hemiplegia at P22. Mice were individually placed in a Plexiglas transparent cylinder and video recorded for 4 min or until they reached ten touches per paw. The relative proportion of left (IL) forepaw contacts was calculated as follows: (left – right)/(right + left + both) × 100.

**Statistical Analysis**

Data were analyzed using Prism Software (version 6, GraphPad). An unpaired Student’s t test was used for two-group comparisons. A one-way ANOVA was used for multiple-group comparisons followed by Bonferroni’s post hoc test. For analysis comparing CL and IL hemispheres, a two-way ANOVA was performed followed by Tukey’s multiple comparisons test. A statistically significant level was defined as p < 0.05. Error bars are reported as mean ± SEM unless stated otherwise.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Supplemental Experimental Procedures and two figures and can be found with this article online at http://dx.doi.org/10.1016/j.stemcr.2015.06.011.

**AUTHOR CONTRIBUTIONS**

P.D. collected, assembled, analyzed, and interpreted data and prepared and approved the manuscript. N.M. and L.S. collected, analyzed, and interpreted data. A.A. collected data. M.F. collected...
and analyzed data. F.E.W provided CBP S436A mutant mice. F.D.M. conceived and designed the study. C.M.M. conceived and designed the study; provided financial support; analyzed, interpreted, and assembled data; and wrote and gave final approval of the manuscript.

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