INTRODUCTION

Neonatal hypoxic-ischemic brain injury (HI) remains one of the most common causes of death and severe neurologic morbidity in infants and children, resulting in chronic cognitive, motor, and sensory deficits.1,2 However, currently, there is no therapy to repair the HI except therapeutic hypothermia, which offers very limited functional recovery.3,4 Consequently, the development of supplementary or alternative therapies, that act synergistically with hypothermia or are effective when used alone, are needed to reduce the mortality and morbidity in HI.
newborn infants with HI. It is conventionally accepted that neuronal generation does not take place beyond fetal life, and therefore, neuronal regeneration does not occur in the vast majority of the post-developmental “non-neurogenic” central nervous system (CNS) after injury or disease. However, neurogenesis from neural stem/progenitor cells (NSPCs) in two restricted neurogenic niches—ventricular-subventricular zone (V-SVZ) near the lateral ventricles and hippocampal subgranular zone—of the mammalian CNS has great implications for the growing field of research on endogenous NSPCs in the mature nervous system. Manipulation of these actively dividing or relatively quiescent NSPCs could supply cell population with immense neuroregenerative potential. Stimulated endogenous NSPCs could be a source of new neurons or glial cells that replace lost neural cells in CNS diseases. Notably, the V-SVZ peaks in size during the first week of postnatal development in rodents and at approximately the 35th week of gestation in humans as a result of extensive cell proliferation. The human neonatal V-SVZ has a remarkable neurogenic capacity, suggesting that NSPCs in V-SVZ may provide an endogenous cell population for neural regeneration after neonatal brain injury.

As a model for HI, we employed the Rice-Vannucci model which has been most well-accepted as rendering one of the experimentally destructive injuries: permanently ligating unilateral common carotid artery on postnatal day 7 (P7) followed by hypoxia (8–10% O2). This procedure results in extensive HI lateral common carotid artery on postnatal day 7 (P7) followed experimentally destructive injuries: permanently ligating uni-

MATERIALS AND METHODS

Induction of unilateral HI brain injury

Through a ventral midline neck incision, the right common carotid artery of anesthetized ICR (CD1) mice on postnatal day 7 (P7) was permanently ligated with the surgical suture silk. After the incision was closed, the animals were kept warm until awake and returned to their dams for 3 h. The pups were then placed in an acrylic chamber with hypoxic atmosphere of 92% N2 and 8% O2 for 2.2 h. The animals’ body temperature was maintained at 37°C.

BrdU labeling and detection

Two hours after the induction of unilateral HI, the mice were intraperitoneally administered 50 mg/kg bromodeoxyuridine (BrdU; Sigma, St. Louis, MO, USA) every 4 h for the subsequent 12 h (four administrations total). The animals were sacrificed 24 h (n=3), 48 h (n=3), 1 wk (n=7), 2 wk (n=5), and 3 wk (n=5) after the final BrdU “pulse” and processed for BrdU detection. In some experiments designed to assess the long-term in vivo survival and permanence of BrdU-intercalated cells over months, daily “pulses” of BrdU for a week post-HI were performed. At the pre-determined timepoints for analysis, the animals were deeply anesthetized and perfused transcardially with 0.1 M PIPES buffer (pH 6.9; Sigma) containing 4% paraformaldehyde. The brain was subsequently post-fixed in the perfusing solution overnight at 4°C, cryoprotected overnight in phosphate-buffered saline (PBS) containing 30% sucrose at 4°C, and then embedded in O.C.T compound medium (Sakura Finetek, Torrance, CA, USA). Serial brain coronal cryostat sections (15-μm thick) were treated initially with 2N HCl for 30 min at 37°C to denature cellular DNA and then reacted with a Fluorescein isothiocyanate (FITC)-conjugated mouse anti-BrdU antibody and/or a mouse anti-BrdU antibody (Boehringer Mannheim, Indianapolis, IN, USA) followed by a biotinylated-horse anti-mouse secondary antibody (Vector Laboratories, Burlingame, CA, USA), in order to identify BrdU-labeled cells.

Injection of retrovirus and detection

Retrovirus vector particles encoding lacZ were collected from the supernatants of confluent ψ2 BAG helper virus-free packaging cells for 48 h at 37°C, filtered through 0.45 μm pores, aliquoted, and stored at -80°C. The BAG virus stock (6×10⁶ G418-resistant CFU/mL) was mixed with 8 μg/mL polybrene and trypan blue prior to gently expelling 2 μL through a finely drawn glass micropipette into each lateral ventricle of mice 2 h before the induction of HI. The brains were processed via standard methods described below and elsewhere for immunohistochemistry. Expression of the lacZ gene product, E. coli β-galactosidase (βgal), was detected by rabbit anti-βgal antibody (1:1000; Cappel Laboratories, Cochranville, PA, USA) followed by Texas red (TR)-conjugated goat anti-rabbit IgG (1:200; Vector Laboratories) antibodies.

The pCLPIT GFP plasmid was used to produce the murine leukemia virus (MLV)-based retroviral vector expressing GFP. The pCAG-NeuroD1-IRES-GFP sequence was subcloned from CAG-NeuroD1-IRES-GFP plasmid (Addgene, Watertown, MA, USA) and replaced the PuroR-IRES-tTA/TRE-GFP sequence in the pCLPIT GFP plasmid to generate the mouse NeuroD1/GFP-expressing retroviral vector. Retrovirus vector particles encod-
ing GFP or NeuroD1/GFP were collected from the supernatants of human embryonic kidney 293T cells and concentrated to about 10^9 TU/mL by ultracentrifugation. The virus stock solution was injected into each lateral ventricle of mice being subjected to HI, and the brains were processed as described above. The expressions of GFP and NeuroD1 gene products were detected by rabbit or mouse anti-GFP (1:100; Thermo Fisher Scientific, Waltham, MA, USA) and mouse anti-NeuroD1 (1:500; Abcam, Cambridge, MA, USA) antibodies, respectively. Species-specific secondary antibodies, conjugated with FITC (1:200; Vector Laboratories) or TR (1:200; Vector Laboratories), were used to detect the binding of primary antibodies.

**Immunohistochemistry**

Immunohistochemistry (IHC) analysis employed the anti-BrdU, anti-β-gal, anti-GFP, and/or anti-NeuroD1 antibodies described above. Dual staining was performed with primary antibodies against the following immature, undifferentiated NSPCs and established differentiated neural cell type-specific antigens: anti-NeuN (1:100; Chemicon, Billerica, MA, USA), anti-neurofilament (NF) (1:500; Sternberger, San Diego, CA, USA), anti-Doubledcortin (DCX) (1:50; Santa Cruz, CA, USA), anti-Olig2 (1:200; Millipore, Billerica, MA, USA), anti-CNPase (1:500, Sternberger), anti-glial fibrillary acidic protein (GFAP) (1:1000; Dako, Glostrup, Denmark), antinestin (1:500; PharMingen, San Diego, CA, USA). Nuclear c-fos and synaptic connection with neighboring neurons were detected by a rabbit polyclonal anti-c-fos (1:10000; Santa Cruz) and anti-synapsin 1 (1:500; Thermo Fisher Scientific) antibodies, respectively. Species-specific secondary biotinylated (Vector Laboratories), streptavidin TR-conjugated (1:200; Jackson, Cambridge, UK), FITC-conjugated, or TR-conjugated antibodies were used to detect the binding of primary antibodies. Specimens were mounted using the Vectashield mounting medium with 4,6-diamino-2-phenylindole (DAPI; Vector Laboratories), and were analyzed by an immunofluorescence microscope (BX51; Olympus, Center Valley, PA, USA) or a confocal laser scanning microscopy (LSM 700; Carl Zeiss, Oberkochen, Germany).

**Quantification**

Structures were sampled by selecting predetermined areas on each coronal section as described. Every 6th section was selected from a coronal series through the striatum by the optical dissector stereological method. BrdU+ cells were counted in three predetermined ventral, lateral, and dorsal squares (50×50 μm) of the lateral ventricle wall on all selected sections. A rectangular area of the striatum (300×600 μm) was selected at a 50-μm distance from the lateral ventricle wall and analyzed on each section. All BrdU+ nuclei in these areas were counted. Every 12th section from a coronal series [spanning from the olfactory bulb (OB) and frontal cortex rostrally and progressing caudally through the entire telencephalon] was selected and processed for BrdU IHC. As depicted, four predetermined areas (50×50 μm) in the rostral migratory stream (RMS) and granule cell layer of the OB were analyzed on each section. All BrdU+ nuclei in these areas were counted.

**Statistical analysis**

All statistical analyses were performed using SPSS version 23 (IBM Corp., Armonk, NY, USA). For statistical comparison of newly-born BrdU+ cell numbers, we performed two-way ANOVA to account for the effect of HI (non-injured vs. injured mice) and/or hemispheres (contralateral vs. ipsilateral hemisphere). Bonferroni post hoc analysis was sequentially performed for pairwise comparisons of groups at each timepoints where appropriate. Sample sizes were calculated using power analyses with an α-level of 0.05 and a power of 0.8. Data are presented as the means±SEM. p<0.05 was considered statistically significant.

**RESULTS**

**Proliferation and migration patterns of BrdU-labeled endogenous NSPCs in response to unilateral HI**

Following unilateral HI, the proliferation and migration patterns of endogenous NSPCs in the periventricular regions of both hemispheres was assessed using the serial BrdU injection method. BrdU was selectively and permanently incorporated into the nuclear genomic material of all proliferating cells entering S-phase. HI induced a significantly increased proliferation of BrdU-labeled V-SVZ NSPC population ipsilateral to the lesioned right side (Figs. 1 and 2). Furthermore, HI appeared to induce a significant redistribution of proliferative NSPCs ipsilateral to the lesioned right side (Fig. 1B, D, F, and H) compared to the grossly intact contralateral left side (Fig. 1A, C, E, and G) and uninjured control group (n=5) (Fig. 2). The accumulation of BrdU+ cells was actually most pronounced in the dorsolateral wall of the lateral ventricles, particularly the anterior one-third of the V-SVZ (Fig. 1B, D, F, and H), adjacent to the infarction cavity (IC) (Fig. 1B, D, and F); and a relatively dense “stream” of newly-born cells (arrowhead in Fig. 1B) oriented towards and into the extensively injured cortex was now apparent (arrows in Fig. 1B, D, F, H, and F’). The actual number of newly-born cells was, indeed, increased at the site of HI and its penumbra in the neocortex as well as in adjacent parenchymal areas, particularly in the striatum (Fig. 1D, F, F’, and Fig. 2). About 2–4 times as many BrdU+ cells were seen in the ipsilateral cortex and striatum 24–48 h after BrdU pulse compared to uninjured controls [e.g., cortex: 1964.2±246.5 (mean±SEM) vs. 495.6±78.8, p<0.05; striatum: 2950.8±331.8 vs. 657.6±49.4, p<0.05].

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As the migratory route from V-SVZ to OB via the RMS is so well-characterized developmentally programmed destination of migratory V-SVZ NSPCs in the intact brain, examining it in the context of unilateral HI provides an ideal opportunity to assess the role HI might play in “shifting” migratory patterns. The normal fate for most of the cells born in V-SVZ, particularly the anterior portion (“c-d” in the schematic in Fig. 1K and displayed in Fig. 1A-D), is to migrate rostrally along the RMS into the OB (“a-b” in the schematic in Fig. 1K and displayed in Fig. 1I and J), where they differentiate into the neurons. Such typical developmental program is evident in the intact left hemisphere [e.g., few BrdU+ cells surrounding the left lateral ventricle (LV) in Fig. 1A and C; many BrdU+ cells in left OB (LOB) in Fig. 1I and J].

However, although more cells were actually born in the right V-SVZ ipsilateral to the lesion in response to HI (Fig. 1B and Fig. 2), significantly fewer BrdU+ cells (about 53% of uninjured controls) were present in the right RMS (Fig. 2) 1 wk after BrdU injection. In addition, the number of newly-born cells that actually reached the right OB (ROB) was reduced significantly (about 28% of uninjured controls and 51% of that observed contralaterally).
tralateral to the lesion, e.g., ipsilateral ROB compared to contralateral LOB in Fig. 1I, J, and Fig. 2), although the migration route remained anatomically intact. Interestingly, the number of newly-born cells that reached the RMS and OB from the V-SVZ contralateral to the lesion, though certainly much greater than that ipsilateral to the lesion, was also significantly reduced compared to the uninjured control mice (Fig. 2). This suggested that injury may draw cells from even distant regions, affecting the migratory pattern of V-SVZ-derived cells even in the grossly intact contralateral hemisphere. Indeed, the greater number of BrdU+ cells in the injured ipsilateral cortex contrasted with, and may also come at the expense of, about 76% fewer BrdU+ cells noted in the contralateral cortex compared to the age-matched uninjured controls (843.8±77.7 vs. 1117.0±121.9, p<0.05) (Fig. 2), suggesting that NSPCs which normally would have taken up residence in their contiguous cortex31 may be diverted towards the lesion as the SVZ→RMS→OB axis was effected.

HI promotes the maintenance of cerebral nestin expression
Consistent with the view that HI may have a broader effect throughout the brain was the pattern of cerebral nestin expression at various times—1 day (n=5), 1 wk (n=7), and 3 wk (n=5)—following unilateral HI, both at the site of injury and in the contralateral intact hemisphere. We analyzed the expression of an intermediate filament protein nestin via IHC, which is known as NSPC and radial glia (embryonic NSPC during CNS development) markers.32-34 In contrast to the restricted nestin-immunoreactivity (IR) bordering both lateral ventricles in the intact brain of age-matched control mice on P8 (Fig. 3F), robust nestin-IR was evident not only in the periventricular regions of both lateral ventricles, but also in the septum, striatum, corpus callosum, external capsule, and cerebral cortices of both hemispheres, particularly at the right cortical infarction site, 24 h after HI (Fig. 3A and B). In addition, the density of nestin+ radial glia fibers with long processes appeared to

![Fig. 2. Density of BrdU-positive newly-born cells following unilateral hypoxic-ischemic brain injury. BrdU+ cells of V-SVZ, RMS, OB, striatum, and cortex in non-injured control and injured animal at 24 h (A), 48 h (B), and 1 week after the final BrdU pulse (C). The number of BrdU+ cells was not available to count at the cortex at 1 week after BrdU injection. Data represent the means±SEM. *p<0.05 vs. the corresponding regions in non-injured controls, †p<0.05 vs. the contralateral side to the lesion in an injured animal. V-SVZ, ventricular-subventricular zone; RMS, rostral migratory stream; OB, olfactory bulb.](https://doi.org/10.3349/ymj.2020.61.6.492)
be highly increased in both hemispheres (arrowheads in Fig. 3A and B). The robust nestin-IR was maintained in HI site and cortical penumbral area of ipsilateral hemispheres 1 wk (Fig. 3D) and even 3 wk (data not shown) after HI. However, the nestin-IR in the contralateral hemisphere decreased, except the external capsule, 1 wk after HI (Fig. 3C) and almost disappeared, except the dorsolateral SVZ adjacent to the LV, 3 wk after HI (Fig. 3E). These results suggest that neonatal HI promotes the maintenance of cerebral nestin expression and has the potential to maintain radial glia fibers after injury.

The hypothesis that such appearance of nestin+ NSPCs or radial glia fibers (even in regions distant from the lesion) might reflect the rapid mobilization or recruitment of NSPCs to regions of cell loss was supported by a further observation of BrdU+/nestin+ cells around both SVZs (Fig. 4A-C) and within both cortices (Fig. 4D-F). In addition, robust nestin-IR with
many BrdU+/nestin+ cells in the corpus callosum bearing elongated leading processes oriented towards the affected hemisphere suggested the migration of NSPCs in response to HI (Fig. 3G–I). These findings suggest that nestin+ newly-born V-SVZ cells migrate towards adjacent parenchyma and HI site, and that retained nestin+ radial glia fibers in response to HI may provide a scaffold for the migration of V-SVZ-derived neurons and glia toward non-neurogenic brain regions in the postnatal brain.35

**Differentiation pattern of BrdU-labeled NSPCs in response to HI**

To help determine the differentiation fate in vivo of HI-generated BrdU-labeled NSPCs, particularly in non-neurogenic regions, they were analyzed for their co-expression of neural cell type-specific antigens (Figs. 4 and 5). Over the 3-week period following the final BrdU pulse, many of the cells were induced to proliferate yielding new oligodendrocytes [10.8% (mean), 6.2%, and 7.4% at 1, 2, and 3 wk, respectively] (Fig. 4P-R and Fig. 5), astrocytes (28.3%, 15.6%, and 16.4%, respectively) (Fig. 4S-U and Fig. 5), and neurons (4.0%, 1.2%, and 1.2%, respectively; Fig. 4G–O and Fig. 5). These new neurons (likely an under-representation given the time course of BrdU pulses) were evident not only in the compromised hemisphere, but in the contralateral hemisphere as well, suggesting the widespread “ripple effect” of signals emanating from even a unilateral lesion. In some experiments designed to assess even longer term survival of BrdU+ cells, daily ip BrdU pulses for a full week of post-HI were performed. At 2 months after injury, about 1% of BrdU+ cells were NF+—not significantly diminished from this value at 3 weeks—suggesting permanence of the new neu-

![Fig. 4. Differentiation pattern of BrdU-labeled NSPCs in response to HI. A day after unilateral HI and the final BrdU pulse, many cells around the V-SVZ (A–C) and in the cortex (D–F) of both hemispheres showed co-localization of both anti-nestin (A, D) (red) and anti-BrdU (B, E) (green) immunoreactivity (arrows), appreciated in merged images (C, F, respectively). One-to-three weeks after HI, many of the BrdU+ cells [arrows in (H, K, N, Q, T)] had differentiated into new neurons (G–O), oligodendrocytes (P–R), and astrocytes (S–U) predominantly in the injured hemisphere, supplementing non-BrdU+ [arrowheads in (G, I, J, L, M, O, P, R, S, U)] of the respective cell type with which they seamlessly intermixed, and were often directly juxtaposed. The new neurons, recognized here as NF+ (red) cells [arrows in (G, J, M) and under the dual-wavelength filter in (I, L, O)], were present in the neocortex—predominantly ipsilateral [arrows in (G–I)] but even contralateral to the lesion [arrows in (J–L)]—and in the ipsilateral hippocampal CA1 area [arrows in (M–O)]—all classically regarded as non-neurogenic regions. The oligodendrocyte marker CNPase and the astrocyte marker GFAP were employed in (P) [and under the dual filter in (R)] and (S) [and under the dual filter in (U)], respectively. Arrowheads in (A, C) indicate cellular processes of Nestin+/BrdU+ immature cells. Confocal images showed that a BrdU+ cell (green) (arrows) colocalized with the mature neuronal marker NeuN (red) under the dual-wavelength filter in (A’–N’). Scale bars: 10 μm in (A, D, H, K, M, A’); 5 μm in (Q, T). NSPCs, neural stem/progenitor cells; HI, hypoxic-ischemic brain injury; V-SVZ, ventricular-subventricular zone.
rons. Notably, these analyses were performed caudal to the OBs and rostral to the dentate gyrus of hippocampus to prevent confusion with neurons routinely generated locally in these neurogenic regions.29,30 Following HI, neurons are the neural cell type most susceptible to elimination, yet they are the very cells deemed as incapable of being renewed in non-neurogenic regions. Given such expectations, the actual numbers were somewhat less consequential than the fact of observing any new neurons in areas that were heretofore regarded as harboring no new neurons, and were considered representative of most of the postnatal brain. The proportion of endogenous NSPCs differentiating into various neural cell types—including the proportion remaining immature and undifferentiated—was consistent with prior studies quantifying the phenotype of proliferative cells following exogenous mitogen/growth factor infusions.36-38

Tracking of LacZ-expressing V-SVZ NSPCs in response to HI
As a complement to BrdU labeling and to more rigorously track the fate of these newly proliferative HI-responsive V-SVZ NSPCs (as well as to selectively label periventricular cells independent of local cortical parenchymal progenitors), mitotic cells were identified by their incorporation of a retroviral provirus encoding lacZ following infection by a retrovirus vector21 injected into the ventricles. A cell that was in S-phase at the time of exposure to the retrovirus indicated LacZ expression; in response to HI, a subpopulation of the newly proliferative and migratory βgal+ cells now expressed mature neuronal marker (e.g., NeuN) in all of these non-neurogenic regions (Fig. 6C, E, I, L, O, and Q), suggesting de novo neurogenesis. The NeuN+ neurons were suggested to be active, integrated, and activatable in concert with other regional neurons, based on their immunohistochemically detectable upregulation and expression of c-fos and synapsin in the living, behaving animal (Fig. 6R-Z), e.g., in cortex including the infarcted area (Fig. 6R-T, X-Z) and hippocampus (CA3) where 59.6±3.2% and 25.2±4.2% of LacZ+ cells were dual immunoreactive with NeuN and nestin, respectively (Fig. 6U-W).

Tracking of NeuroD1/GFP-expressing V-SVZ NSPCs in response to HI
NeuroD1 is a basic helix-loop-helix (bHLH) transcriptional factor that is highly expressed in the developing neurons of peripherial and CNS, supporting its role as a neuronal differentiation factor. NeuroD1 may contribute to multiple levels of neural development; nevertheless, it is believed that NeuroD1 mainly plays important roles in the terminal differentiation of postmitotic neuronal cells.39 Therefore, we asked if NeuroD1-transduced V-SVZ NSPCs could enhance neuronal differentiation of internal pool of NSPCs and provide greater numbers of new neurons in non-neurogenic brain regions in response to HI. Retroviral vectors encoding NeuroD1/GFP and GFP were injected into both lateral ventricles of mice being subjected to unilateral HI (n=5 for each group) and analyzed at least 1–2 wk later (Fig. 7). In response to HI, NeuroD1+/GFP+ mitotic V-SVZ NSPCs migrated into the adjacent stratum (Fig. 6A) and hippocampus, into the cortex ipsilateral to the lesion (Fig. 6D), and into the cortical pnumbra (Fig. 6I). In addition, some LacZ-labeled V-SVZ (often in groups36,38) also migrated into the cortex (Fig. 6G) and overlying hippocampal CA1 area (Fig. 6M). Providing a broader overview under low magnification, LacZ-labeled V-SVZ NSPCs following severe HI showed a robust radiating migratory pattern towards the overlying hippocampus and neocortex (Fig. 6P and Q). Confirming the observation noted in Fig. 6G-O, a subpopulation of the newly proliferative and migratory βgal+ cells now expressed mature neuronal marker (e.g., NeuN) in all of these non-neurogenic regions (Fig. 6C, E, I, L, O, and Q), suggesting de novo neurogenesis. The NeuN+ neurons were suggested to be active, integrated, and activatable in concert with other regional neurons, based on their immunohistochemically detectable upregulation and expression of c-fos and synapsin in the living, behaving animal (Fig. 6R-Z), e.g., in cortex including the infarcted area (Fig. 6R-T, X-Z) and hippocampus (CA3) where 59.6±3.2% and 25.2±4.2% of LacZ+ cells were dual immunoreactive with NeuN and nestin, respectively (Fig. 6U-W).

Fig. 5. Cellular differentiation patterns of HI-generated BrdU+ newly-born cells in non-neurogenic regions. Counts represent the means and SEM of the sums of every 6th 15-μm coronal section extending caudally from the rostral tip of the lateral ventricle for the entire expanse of both cerebral cortices (ipsilateral and contralateral to HI). HI, hypoxic-ischemic brain injury.
cells transduced with a retrovirus encoding GFP expressed astrocyte marker GFAP (65.1±5.6%) (Fig. 7C-E") and the oligodendroglial progenitor marker Olig2 (22.4±3.8%) (Fig. 7F-H’), whereas relatively few expressed early neuronal marker DCX (11.3±2.5%) in the striatum and cortical penumbral area (Fig. 7I-K’). In contrast, most of the newly-born and migratory GFP+ cells transduced with a retrovirus encoding NeuroD1/GFP did not express GFAP or Olig2, but expressed early neuronal marker DCX or mature neuronal marker NeuN (95.1±7.4%) in all of these non-neurogenic brain regions (Fig. 7L-X), suggesting enhanced neurogenesis. In addition, very few of GFP+ cells (1.3±0.7%) expressed nestin (data not shown).

**DISCUSSION**

This study sought to elucidate the brain’s intrinsic response to HI, particularly as it involves an endogenous V-SVZ NSPC pool and specifically in areas that are designated "non-neurogenic" and hence non-regenerative. Our findings suggest that neona
tal HI alters developmental programs and does so towards the goal of repopulation and self-repair, allocating new cells to the regions of greatest need; during acute phases of neurodegeneration, many factors, such as morphogens, growth factors, neurotrophic factors, erythropoietin, and microRNAs, are elaborated to which endogenous NSPCs may respond to promote the establishment of new neurons, even within non-neurogenic regions of the postnatal brain. Some of these newly-born neurons, while generated from V-SVZ NSPCs, appear to be targeted specifically to damaged, neuron-deficient regions by redirecting otherwise stereotypical migratory patterns. Such observations may attest to an under-appreciated degree of plasticity and resilience in the mammalian CNS, even beyond the classical periods and regions of developmental malleability. The CNS may, indeed, spontaneously attempt to repair itself with its own endogenous pool of NSPCs, but that supply may simply be insufficient either in number or in factors regulating proliferation, migration, and differentiation in the context of the most destructive injuries. Therefore, the net impact of the production of new nerve cells may be limited. It is plau-
sible that this type of homeostatic response occurs routinely at a marginally detectable level, and is effective under most circumstances (where CNS insults are modest—e.g., from infections, toxins, minor trauma) in preventing such events from coming to clinical attention. However, when the damage is massive, the system—invoked but inadequate—is “overwhelmed.” Consequently, there is no apparent recovery, although endogenous NSPCs have responded in a “reparative” fashion to signals directing them towards repopulation of the damaged area. Nevertheless, this does not negate the inherent plasticity of CNS nor the potential for augmenting this response. Therefore, the relationship between the severity of HI and NSPC response re-

Fig. 7. Immunostaining of retrovirally-infected endogenous mitotic V-SVZ NSPCs with NeuroD1/GFP and GFP following HI. In response to HI, NeuroD1/GFP-expressing GFP+ cells (green) migrated into the adjacent striatum, subcortex, and cortical penumbra ipsilateral to the lesion (A, A', B, B'), and along the EC and CC (A, A”, B). The boxed areas in A and B are shown at high magnification in A', A” and B’. The asterisk in A’ and arrowheads in A” indicates the infarction cavity and LV, respectively (C-K’). Many of the newly-born and migratory GFP+ cells (C, C’) transduced with a retrovirus encoding GFP expressed astrocyte marker GFAP (D, D’, respectively) (red) with co-localization of both antibodies visualized under a dual filter (E, E’, respectively) (yellow or orange) in the cortical penumbral area. Boxed regions in (C-E) are magnified in (C’-E’), respectively. Arrows in (C’-E’) indicate dual-labeled cells (F-H). Some of newly-born GFP+ cells (F) expressed the oligodendrocyte progenitor marker Olig2 (G) (red) with co-localization of both antibodies visualized under a dual filter (H). Arrows in (F-H) indicate dual-labeled cells. (I-K) A few of newly-born GFP+ cells (I) expressed early neuronal marker DCX (J) (red) with co-localization of both antibodies visualized under a dual filter (K). Confocal images showed that GFP+ cells (arrows and arrowheads in (T) and (U), respectively) transduced with a retrovirus encoding NeuroD1/GFP expressed early neuronal marker DCX (arrows in (T)) (red) with co-localization of both antibodies visualized under a dual filter (T) (yellow or orange) in the striatum and cortical penumbral area. DAPI+ cells (arrows and arrowheads in (T’)) (blue) indicate dual-labeled cells. Confocal images showed that GFP+ cells (arrows and arrowheads in (T) and (X), respectively) colocalized with neuronal markers under a dual filter (yellow or orange). However, GFP+ cells (L, O) did not express GFAP or Olig2 (M, N, P, Q). Scale bars: 500 μm in (A); 20 μm in (H’, N) 10 μm in (A’, B’, E’, F-H). V-SVZ, ventricular-subventricular zone; NSPCs, neural stem/progenitor cells; HI, hypoxic-ischemic brain injury; EC, external capsule; CC, corpus callosum; LV, left lateral ventricle; RV, right lateral ventricle.
mains to be explored. The prediction of NSPC response based on the level of severity of HI is an important issue that should be further investigated.

While it is clear that newly-born brain cells are being produced in response to HI, data compiled to date suggest that the long-term survival of migrated neuroblasts or newly-born neurons in the injured area is not supported.41-43 The reasons for the death of newly-born neurons remain elusive and may be due to the inhospitable post-HI environment. HI initiates an inflammatory cascade, such as microglial activation, immune cells infiltration, and the release of toxic proinflammatory molecules.44 Microglial activation, proinflammatory factors, and circulating monocytes associated with inflammation not only lead to neuron death but also affect the neurogenesis and survival of newly-born neurons.45 In addition, some studies have indicated that SVZ-derived migratory NSPCs differentiate into reactive astrocytes and contribute to astrocyte scar formation after stroke.46,47 While the role of astrogliosis after stroke or HI is still elusive and requires further investigation, the modulation of pathologic environment after HI by anti-inflammatory or neurogenic factors results in substantial repair of neural tissue.47,48

Radial glial (RG) cells are embryonic neural stem cells expressing nestin, GFAP, and astrocyte-specific glutamate transporter that extend apical long processes to the pial surface from their soma, which is located in the ventricular zone.33 RG cells are not only stem cells and progenitors, but during brain development they also provide a scaffold along which newly generated immature neurons migrate from the periventricular zone into the neocortex and to subcortical structures. In the late fetal period in the human brain, and in the first postnatal week in the rodent brain, the RG fibers collapse, transform into astrocytes or ependymal cells, and normally disappear soon after birth.48 However, this study showed that signals generated in HI elicited robust cerebral nestin expression, and the density of nestin+ radial glia fibers with long processes appeared to be highly increased in both hemispheres. Moreover, the robust nestin-IR was maintained in the lesioned and cortical penumbral areas of ipsilateral hemispheres, even more than 3 weeks after HI. These findings suggest that retained nestin+ radial glia fibers in response to HI may provide a scaffold for the widespread migration of V-SVZ-derived neurons toward non-neurogenic brain regions in the postnatal brain. Similarly, earlier studies reported an increase in cells expressing RG markers after chronic hypoxia in immature rats, and demonstrated the role of RG cells in both neurogenesis and migration.50,51 A recent study also showed that RG fiber persisted in neonatal mouse brain after cryogenic injury and acted as a scaffold for the migration of V-SVZ-derived neuroblasts to the lesion site, thereby enhancing neuronal regeneration and functional recovery from neonatal brain injuries.35

Whereas there has been a great increase of studies observing neuronal replacement after neonatal HI, there have been fewer studies assessing the extent of oligodendroglial replacement. This study, along with other reports,52-54 documented an increase in the number of newly generated oligodendrocytes in the damaged hemispheres and suggested that these newly-born oligodendrocytes were produced by V-SVZ NSPCs. However, none of these studies clearly established whether these newly-born oligodendrocytes proceeded to differentiate into mature myelin-producing cells. In this study, significant astrogliosis was observed after HI, along with an increase in BrdU/GFAP cells. It is well-known that reactive astrocytes produce extracellular matrix components, such as hyaluronic acid and chondroitin sulfate proteoglycan, that inhibit oligodendrocyte differentiation.55,56 In addition, astrocytes produce numerous cytokines, including fibroblast growth factors, WNT ligands, endothelin-1, and Notch ligands, that inhibit oligodendrocyte progenitor cell maturation.57 There is also an alternative explanation for the lack of myelination depending on SVZ cell specification.57,58 However, further research is critically needed to facilitate the differentiation into mature, myelin-producing cells from V-SVZ NSPC-derived oligodendrocytes in HI.

Identifying the cues that promote the movement and differentiation of this endogenous pool of NSPCs during their normal, developmentally-appropriate migratory course28-30,59,60 and, as we observed here, during their confrontation with neurodegeneration, may allow those factors to be provided therapeutically at even more optimal, sustained levels, over broader periods and terrain. In fact, multiple cytokines, such as Delta/Notch-1, erythropoietin, interleukin-6, leukemia inhibitory factors (LIF), transforming growth factor, and vascular endothelial growth factor (VEGF), were reported to be elaborated to which endogenous NSPCs respond to change in their proliferation, migration, and differentiation patterns after neonatal HI. In addition, the treatment of erythropoietin, brain-derived neurotrophic factor, epidermal growth factor, VEGF, LIF, hyperbaric oxygen, or hypothermia has been shown to attenuate HI by modulating the behavior of endogenous NSPCs.35,40

Lastly, we have demonstrated here that neurogenic transcriptional factor NeuroD1 can efficiently convert V-SN2 NSPCs to neurons, providing a new target for cellular conversion strategies aimed at enhancing neurogenesis. Therefore, augmenting the endogenous NSPC population with exogenous neural stem cells,56 factors, and/or providing adequate synaptic neighbors and guidance cues for connectivity and trophic support may enable more significant recovery.

BrdU labeling for mitotic cells is still the most commonly used of the thymidine analogues, and much work on this subject has been done in mice. However, BrdU inhibited the expansion of NSPCs, increased cell death, and repressed neuronal and oligodendroglial differentiation.61 Although BrdU does obviously label dividing cells, BrdU positivity may be due to attempts of cells to repair themselves or a sign of cell death (apoptosis). In addition, BrdU has been well-known to have detrimental effects on chromosomes and DNA stability, as well as the cell cycle, cell differentiation, and survival.62 Besides the
totoxicity of BrdU and caveats concerning its utility, BrdU labeling, particularly at the epicenter of HI, can be hard to interpret due to the reliance on immunodetection in the injury setting, where significant labeling issues can arise. In addition, BrdU is subject to dilution effects, where the absence of labeling or decreased labeling over time can be misinterpreted. Therefore, we also utilized a retroviral vector to label mitotic NSPCs and their progeny, a technique that is not affected by the number of cell divisions. Another advantage of using a retroviral approach is the ability to direct the location of injection into V-SVZ NSPCs. Gene transfer mediated by replication-incompetent retroviral vectors has played an important role in developmental studies, particularly in revealing lineage relationships among different cell types. Replication-incompetent retroviral vectors have also been used for gain-of-function studies to explore the molecular mechanisms underlying cell differentiation in neural tissues of mice. The greatest advantage of this approach is that it can reveal cell-autonomous effects of a transgene, such as NeuroD1, on cell differentiation without disrupting the overall context of development. However, gamma retroviral vectors have a propensity to integrate near regulatory regions of active genes. The activity of promoter/enhancer elements within the vector long terminal repeats (LTRs) or aberrant splicing events can lead to the activation of genes flanking the integration site. This is commonly referred to as insertion mutagenesis. If the genes near the integration sites are proto-oncogenes, it may lead to increased proliferation of clonal populations of cells. In addition, a number of studies have shown that the promoter regions of gamma retroviral vectors may be subject to epigenetic silencing, which has been associated with DNA methylation of CpG sequences. Such silencing events gradually reduce the expression of transgenes, such as LacZ, GFP, or NeuroD1, under the control of both viral LTR and internal promoters. Taken together, it is difficult to find a single method to accurately and dynamically track the behaviors of endogenous NSPCs in response to HI. Therefore, more sophisticated measures including the use of genetic mouse models or live molecular imaging would be required to avoid the common mistake of demonstrating the behaviors of endogenous NSPCs in HI.

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