Overexpression of EphB4 promotes neurogenesis, but inhibits neuroinflammation in mice with acute ischemic stroke

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Abstract. Ischemic stroke is one of the most common diseases that has a high rate of mortality, and has become a burden to the healthcare system. Previous research has shown that EPH receptor B4 (EphB4) promotes neural stem cell proliferation and differentiation in vitro. However, little is known regarding its role in the neurogenesis of ischemic stroke in vivo. Thus, the present study aimed to verify whether EphB4 was a key regulator of neurogenesis in ischemic stroke in vivo. Cerebral ischemia was induced in C57BL/6J mice via middle cerebral artery occlusion (MCAO), followed by reperfusion. Immunofluorescence staining was performed to evaluate the effect of EphB4 on the neurogenesis in cerebral cortex. The levels of inflammatory cytokines were determined using an ELISA kit. The expression levels of ABL proto-oncogene 1, non-receptor tyrosine kinase (ABL1)/Cyclin D1 signaling pathway-related proteins were detected via western blotting. The current findings indicated that EphB4 expression was significantly increased in the cerebral cortex of MCAO model mice in comparison with sham-operated mice. Moreover, EphB4 appeared to be expressed in neural stem cells (Nestin+), and persisted as these cells became neuronal progenitors (Sox2+), neuroblasts [doublecortin (DCX)+], and eventually mature neurons [neuronal nuclei (NeuN)+]. Overexpression of EphB4 elevated the number of proliferating (bromodeoxyuridine+, Ki67+) and differentiated cells (Nestin+, Sox2+, DCX+ and NeuN+), indicating the promoting effect of EphB4 on the neurogenesis of ischemic stroke. Furthermore, EphB4 overexpression alleviated the inflammation injury in MCAO model mice. The expression levels of proteins-related to the ABL1/Cyclin D1 signaling pathway were significantly increased by the overexpression of EphB4, which suggested that restoration of EphB4 promoted the activation of the ABL1/Cyclin D1 signaling pathway. In conclusion, this study contributes to the current understanding of the mechanisms of EphB4 in exerting neurorestorative effects and may recommend a potential new strategy for ischemic stroke treatment.

Introduction

Stroke remains the second major cause of mortality worldwide and is the leading cause of death in China (1). According to a Global Burden of Disease study, there were 5.5 million deaths and 116.4 million disability adjusted life years due to stroke in 2017 (2). Although rapid progress has been achieved in the understanding of the pathogenesis of ischemic stroke, there is currently no effective therapeutic approach for nerve repair (3). Accumulating evidence has shown that cerebral ischemia can stimulate the activation of quiescent neural stem cells (NSCs) in the subventricular zone (SVZ) into transient amplifying progenitors, which become neuroblasts after several divisions (4,5). The neuroblasts migrate into the damaged cortex and differentiate into mature neurons to promote the recovery of the neurological function (6). The growth peak of NSCs in the adult brain after ischemia is 7-10 days (7). However, this internal response cannot functionally compensate for the ischemic damage, and the migrating cells do not differentiate into mature neurons in the cortex (8,9). Thus, it is critical to search for therapeutic approaches and specific targets to augment the differentiation of endogenous NSCs into neurons.

Neuroinflammation participates in the pathophysiological progress of secondary brain injury after ischemic stroke by increasing pro-inflammatory cytokines and neuronal apoptosis (10). A previous study has shown that the levels of inflammatory cytokines gradually increase within 0-24 h of cerebral ischemia reperfusion in rats, and reach a peak at 24 h (11). The interaction between neuroinflammation and subsequent neurogenesis remains unknown, and thus, has gained significant interest in recent years. Moreover, further understanding of neuroinflammation and its association with neurogenesis could provide a novel approach for brain repair.

Previous studies have reported that ephrin (Eph) receptors can regulate the proliferation of stem cells and progenitor cells in the central nervous system (CNS) (12). The Eph receptor family is the largest known receptor tyrosine kinase family. The members of this family bind their ligand ephrins to initiate bidirectional signaling and regulate different...
physiological activities, which have become popular research subjects internationally (13). Previous studies have shown that ephrinB/EphB signaling served an early role in the regulation of stem cell behavior. For instance, ephrinB3/EPH receptor B3 (EphB3) signaling exhibited an inhibitory effect on NSC proliferation in the developing SVZ (14). In the adult CNS, overexpression of ephrinB2 or EphB2 promotes NSC proliferation and represses neuroblast migration (15). However, research regarding EphB4 focuses on tumor cells. Restoration of EphB4 promotes tumor cell proliferation, migration and angiogenesis (16,17). Moreover, a previous study revealed that EphB4 regulated the self-renewal, proliferation and neuronal differentiation of human embryonic NSCs in vitro (18).

The present study aimed to investigate the role of EphB4 in the neurogenesis and neuroinflammation of ischemic stroke in vivo.

Materials and methods

Animals and middle cerebral artery occlusion (MCAO) model.

The animal experiment was approved by the Animal Care and Use Committee of Inner Mongolia Baogang Hospital (approval no. BG201802034YY). A total of 120 male C57BL/6J and EphB4+/− C57BL/6J mice (age, 6-8 weeks; weight, 21-23 g), bred by Cyagen Biosciences, Inc., were maintained on a 12-h light/dark cycle under specific pathogen-free conditions (25°C; 50-70% humidity) with free access to food and water.

Transient MCAO was performed as previously reported (19), with a slight modification. The mice were randomly divided into four groups (n=5/group): Sham, EphB4+/− Sham group, MCAO group and EphB4+/− MCAO group. Mice were anaesthetized with isoflurane (3% induction; 1-1.5% maintenance) in a mixture of 75% N₂O and 25% O₂ and the rectal temperature was maintained at 37.0±0.5°C. Then, the right common carotid artery and the right external carotid artery (ECA) were exposed. The ECA was then dissected distally, ligated and coagulated. The right MCA was occluded with a heparinized intraluminal filament (0.26 mm; rounded tip). After 30 min of ischemia, the intraluminal filament was withdrawn to accomplish reperfusion. The sham group underwent the same procedure without insertion of the intraluminal filament. After reperfusion for 1, 3, 7 and 14 days, the mice were sacrificed with 120 mg/kg pentobarbital sodium via intraperitoneal injection. Death was ascertained based on pupil dilation and an inability to palpate the carotid pulse. Brain tissues were collected after the mice were sacrificed. Due to severe injuries, eight mice were lost to control and analyzed using the 2−ΔΔCq method (20). The primers used were as follows: EphB4 forward (F), 5′-CCC ATTTGAGCCCTGTCAATGTC-3′ and reverse (R), 5′-TCA AGCTGCTGAGGTGAGA-3′; ephrinB2 F, 5′-TATGCAGA TGCGATTITCCAA-3′ and R, 5′-TGGGTATAGTTACG ATCTTTGTC-3′; and GAPDH F, 5′-CATGGAATGATGAC AACAGCCT-3′ and R, 5′-AGTCCTTTCACGATACCA AA GT-3′; GAPDH was used as an internal reference.

Immunofluorescence analysis. Immunofluorescence staining was used to visualize nerve cells. Bromodeoxyuridine (BrdU; 50 mg/kg; Sigma-Aldrich; Merck KGaA) was injected intraperitoneally into all mice twice daily for 4 consecutive days after MCAO. After being fixed with 4% paraformaldehyde at 37°C for 24 h, brain tissue was embedded in paraffin and sectioned at a thickness of 5 μm. For BrdU staining, the sections were pretreated with 50% formamide, 280 mM NaCl and 30 mM sodium citrate at 65°C for 2 h, incubated in 2 M HCl at 37°C for 30 min to denature the DNA and rinsed in boric acid (0.1 M; pH 8.5) at room temperature for 10 min. After blocking the sections with 5% donkey serum (Jackson ImmunoResearch Europe) for 2 h at room temperature, they were incubated for 48 h at 4°C with the following primary antibodies: Anti-EphB4 (Santa Cruz Biotechnology, Inc.; cat. no. sc-365510; 1:50), anti-BrdU (Abcam; cat. no. ab152095; 1:100), anti-Ki67 (Abcam; cat. no. ab15580; 1:200), anti-NeuN (Abcam; cat. no. ab104224; 1:200). Subsequently, the sections were incubated with Alexa Fluor 488- or 594-conjugated secondary antibodies (Thermo Fisher Scientific, Inc.; cat. nos. A48296 and A10106; both 1:2,000) for 2 h at room temperature. The fluorescent images of cells were captured with a fluorescence microscope system (Nikon Eclipse 80i; Nikon Corporation; magnification, x200). In total, three areas of interest were selected around the margin of infarct area, and an average of the number of cells in the peri-infarct cortex for each section was obtained (21).

Western blotting. Total proteins were extracted from the infarct side of cerebral cortex with RIPA lysis buffer (Beyotime Institute of Biotechnology). BCA Protein Assay reagent (Beyotime Institute of Biotechnology) was applied to analyze proteins concentration. Proteins (50 μg/lane) were separated by 10% SDS-PAGE and transferred onto a nitrocellulose membrane. Subsequently, the membranes were blocked with 5% non-fat milk powder for 2 h at room temperature.
and incubated with primary antibodies, including anti-EphB4 (Abcam; cat. no. ab254300; 1:1,000), anti-ephrinB2 (Abcam; cat. no. ab75868; 1:500), anti-phosphorylated (p)-ABL proto-oncogene 1, non-receptor tyrosine kinase (ABL1; Cell Signaling Technology, Inc.; cat. no. 2864; 1:1,000), anti-ABL1 (Cell Signaling Technology, Inc.; cat. no. 2862; 1:1,000), anti-Cyclin D1 (Abcam; cat. no. ab134175; 1:2,000), anti-CDK4 (Abcam; cat. no. ab108355; 1:1,000) and β-actin (Abcam; cat. no. ab8227; 1:1,000) at 4°C overnight, followed by the horse-radish peroxidase-labeled secondary antibodies (Abcam; cat. nos. ab205718 and ab205719; both 1:2,000) at 37°C for 1 h. The immunoreactive proteins were visualized using an ECL kit (MilliporeSigma) and the data were analyzed using ImageJ software (version 1.8.0; National Institutes of Health). The expression values were normalized against β-actin.

ELISA. The productions of IL-6, IL-1β, TNF-α, IL-10 and monocyte chemoattractant protein (MCP)-1 in the tissues and plasma of mice were measured using the corresponding mouse ELISA kits (cat. nos. RAB0314, RAB0274, RAB0477 and
EphB4 expression in the cerebral cortex of MCAO model mice. To evaluate the association of EphB4 with ischemic stroke, the expression levels of EphB4 and ephrinB2 in the cortex of MCAO model mice, followed by a reperfusion for 1, 3, 7 and 14 days, were detected. The TTC results confirmed the successful ischemia induction (Fig. 1A). The results from RT-qPCR assay demonstrated that the expression levels of EphB4 and ephrinB2 were increased after MCAO and peaked at 7 days (Fig. 1B and D), which was also confirmed by the results of western blotting assay (Fig. 1C and E).

EphB4 expression in nerve cells of the cerebral ischemic cortex.

As reported in previous studies, in the cerebral cortex, NSCs express Nestin, the generated neuronal progenitors express Sox2, neuroblasts express DCX and the mature neurons express NeuN (22, 23). In total, three areas of interest were selected around the margin of infarct area, and an average of the number of cells in the peri-infarct cortex for each section was calculated (Fig. 2A). The current immunofluorescence results identified that EphB4 was expressed by Nestin+, Sox2+, DCX+ and NeuN+ cells in the peri-infarct cortex of MCAO model mice (Fig. 2B-E). Moreover, EphB4 appeared to be expressed in NSCs (Nestin+) and persisted as these cells became neuronal...
progenitors (Sox2$^+$), neuroblasts (DCX$^+$) and eventually mature neurons (NeuN$^+$). These results indicated the potential relationship between the EphB4 and neurogenesis.

Overexpression of EphB4 promotes NSC proliferation in the cerebral cortex of MCAO model mice. To confirm whether EphB4 modulated the proliferation of nerve cells, NSCs

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Figure 3. Overexpression of EphB4 enhances neuronal stem cell proliferation in the cortex at 7 days following MCAO. (A) Reverse transcription-quantitative PCR and (B) western blot analysis of EphB4 expression in EphB4$^{+/+}$ mice and wild-type mice. (C) Images showing the immunostaining for Nestin$^+$, and the bar graph presented the quantification of the number of Nestin$^+$ cells in the peri-infarct cortex. (D) Images showed the immunostaining for BrdU$^+$ and the bar graph showed the quantification of the number of BrdU$^+$ cells in the peri-infarct cortex. (E) Images showed the immunostaining for Ki67$^+$ and the bar graph showed the quantification of the number of Ki67$^+$ cells in the peri-infarct cortex. Scale bar, 25 µm. n=5 for each group. Data are presented as mean ± SD. *P<0.05 vs. Sham; **P<0.01 vs. control; #P<0.05 vs. MCAO. EphB4, EPH receptor B4; MCAO, middle cerebral artery occlusion; BrdU, bromodeoxyuridine.
labelled with Nestin⁺, and newborn cells labeled with Brdu⁺ or Ki67⁺ were observed in the peri-infarct cortex 7 days after MCAO using an immunofluorescence assay. EphB4 expression was measured via RT-qPCR and western blotting in EphB4⁺/⁺ mice prior to MCAO. The results demonstrated that EphB4 was significantly elevated in EphB4⁺/⁺ mice compared with wild-type mice (Fig. 3A and B). Immunofluorescence results revealed that there were no differences in the number of positive cells (Nestin⁺, Brdu⁺ and Ki67⁺) between sham mice and the EphB4⁺/⁺ sham mice, while the number of these positive cells was significantly increased in MCAO mice vs. sham-operated mice. Overexpression of EphB4 further enhanced the number of these positive cells in MCAO model mice (Fig. 3C-E). Overall, these data indicated that overexpression of EphB4 promoted NSC proliferation after focal cerebral ischemia in vivo.

Overexpression of EphB4 promotes NSC differentiation in the cerebral cortex of MCAO model mice. Sox2 is an important molecular marker of adult NSCs (24). DCX is
a microtubule-associated protein expressed in immature
neurons, which is essential for neuronal migration and
differentiation (25). Thus, Sox2+, DCX+ and NeuN+ expression
in the peri-infarct cortex 7 days after ischemia reperfusion
was detected. As shown in Fig. 4A and B, Sox2+ and DCX+
cells were significantly increased at 7 days in MCAO model mice
compared with sham-operated mice. EphB4 overexpression
further increased the number of Sox2+ and DCX+ cells
compared with the MCAO model mice. However, NeuN+ cells exhibited no significant difference between MCAO
and EphB4+ MCAO groups at 7 days (Fig. 4C). However,
at 14 days, the number of NeuN+ cells was significantly
increased in the EphB4+ MCAO model mice compared with
the MCAO model mice (Fig. 4D), suggesting that restoration
of EphB4 promoted NSC differentiation after focal cerebral
ischemia in vivo.

Overexpression of EphB4 attenuates the inflammatory
response in MCAO model mice. To investigate the effect of
EphB4 on the inflammation injury in ischemic stroke model
mice, a series of inflammatory cytokines, including IL-1β, IL-6, TNF-α, IL-10 and MCP-1, in the tissues of cerebral cortex and the plasma of mice were evaluated. The results demonstrated that the levels of IL-1β, IL-6, TNF-α and MCP-1 were quickly increased in the tissues of MCAO group compared with sham operation group. However, the stroke-evoked enhancement of cytokine levels was markedly inhibited in EphB4+/+ mice after MCAO (Fig. 5A-D). Interestingly, the level of IL-10, generally regarded as an anti-inflammatory cytokine (26), was significantly decreased after stroke; however, the low level of IL-10 was significantly increased in the EphB4+/+ MCAO group (Fig. 5E). Consistent with the aforementioned results, the levels of IL-1β, IL-6, TNF-α and MCP-1 were quickly inhibited in the plasma of EphB4+/+ MCAO model mice, which were enhanced by MCAO operation (Fig. 5F-I). However, the level of IL-10 was significantly decreased in MCAO mice vs. sham-operated mice, and EphB4+/+ increased the low level of IL-10 in MCAO model mice (Fig. 5J). These results suggest that overexpression of EphB4 attenuated the MCAO-induced inflammatory response.

**EphB4 activates the ABL1/Cyclin D1 signaling pathway in the cerebral cortex of MCAO model mice.** To assess the potential mechanism of EphB4 in neurogenesis, the related proteins of the ABL1/Cyclin D1 signaling pathway were examined. As shown in Fig. 6A-D, the expression levels of p-ABL1, Cyclin D1 and CDK4 showed a significant increase in the MCAO-treated ischemia mice compared with the sham-operated group. Moreover, compared with the MCAO-treated ischemic mice, the abovementioned protein expression levels were significantly increased in the EphB4+/+ MCAO model mice at 7 days after ischemia (Fig. 6B-D). These results indicated that overexpression of EphB4 activated the ABL1/Cyclin D1 signaling pathway, thereby promoting cell proliferation and differentiation in mice after ischemia stroke.

**Discussion**

Ischemic strokes result in numerous deaths and leave survivors with significant disability, despite various reasonable treatments (27). Neurogenesis has been proven to be a primary neurovascular response during stroke recovery (28,29). In the present study, it was found that overexpression of EphB4 promoted the proliferation and differentiation of NSCs in the cerebral cortex by modulating the ABL1/Cyclin D1 signaling pathway after cerebral ischemia, suggesting that EphB4 may be an effective strategic target for improving neurological dysfunction in ischemic stroke.

EphB4 tyrosine kinase receptor and its ephrinB2 ligand have been identified as important regulators of tumor cell proliferation, migration and differentiation (30-32). For instance, EphB4 was highly expressed in hepatocellular carcinoma and knockdown of EphB4 inhibited cell proliferation and migration and tumor growth (33). Kadife et al (34) discovered that a high expression level of EphB4 facilitated colorectal cancer cell migration and tumor growth. Interestingly, EphB4 was reported to be upregulated after stroke, and was suggested to be involved in neoangiogenesis and neuronal survival (35,36), which was consistent with the present findings that EphB4 was significantly increased in mice after MCAO. A previous study also revealed that EphB4 was expressed in NSCs and instructed NSC neuronal differentiation in the subgranular zone (37). The present study demonstrated that EphB4 was expressed in NSCs and may participate in the differentiation of NSC in the cortex following MCAO.
BrdU is usually employed to label cells in the cell cycle of S phase (38). Ki67 is a nuclear protein associated with ribosomal RNA transcription that acts as a marker for cell proliferation (39). A previous study showed that Nestin+ cells can be observed extensively in restricted regions where they may serve as a niche of stem/progenitor cells with the capacity for proliferation and differentiation in the adult mouse brain (40). Therefore, in the present study, the increased levels of Nestin+, Ki67+ and BrdU+ cells in the peri-infarct cortex of the ischemia-treated mice indicated the endogenous NSC response following MCAO. In the EphB4 overexpression group, the increase of Nestin+, Ki67+ and BrdU+ cells was further enhanced. These results indicated that EphB4 overexpression promotes cell proliferation post-ischemia.

Sox2, a member of the Sox family of transcription factors, is highly expressed during the development of embryonic stem cells and adult NSCs, and is considered to be the key to the proliferation and differentiation of NSCs (41). DCX, a protein facilitating microtubule polymerization, is expressed in migrating neuroblasts and immature neurons, and can be classified as a marker for adult neurogenesis (42). NeuN is a neuronal protein and a neuron marker utilized to specifically label mature neurons (43). Hence, in the current study, the increased number of Sox2+; DCX+ and NeuN+ cells in the cortex of the EphB4+/- MCAO model mice indicated that EphB4 may participate in promoting NSC differentiation into neurons post-ischemia. The upregulation of these neural cells is considered to contribute to functional recovery of brain.

The downstream factor of the ephrinB2/EphB4 signaling pathway, ABL1, binds to EphB4 in an activity-dependent manner (16). Previous findings have shown that ABL1 served an important role in the stimulation of neurogenesis (44). Moreover, ABL1 could mediate the signaling between EphB4 and Cyclin D1/CDK4 to regulate proliferation in NSCs (18). The current results revealed that overexpression of EphB4 increased the protein expression levels of p-ABL1, Cyclin D1 and CDK4, suggesting that EphB4 may promote neurogenesis after ischemic stroke via the ABL1/Cyclin D1 signaling pathway in vivo.

There were certain limitations in the present study. Firstly, the effect of EphB4 on neurogenesis was not assessed in vitro. Secondly, the level of EphB4 in patients with ischemic stroke was not examined and the association between EphB4 levels and lesion volume requires further study. The underlying mechanism of EphB4 on neurogenesis and neuroinflammation should also be investigated both in vitro and in vivo.

In conclusion, the present study demonstrated that restoration of EphB4 promoted cell proliferation and differentiation at 7 days after MCAO, and facilitated NSC directional differentiation into neurons at 14 days after MCAO. However, EphB4 upregulation inhibited neuroinflammation at 1 day after MCAO. Moreover, it was found that overexpression of EphB4 promoted the activation of the ABL1/Cyclin D1 signaling pathway. These findings suggested that EphB4 may be a valuable therapeutic target for ischemic stroke.

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Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors' contributions

JW guaranteed the integrity of the entire study, performed the experiments and wrote the manuscript. ZZ, SF and XiaL constructed the MCAO model and collected brain tissues. ZZ, XinL and SW performed the data analysis and statistical analysis. LY designed the study, supervised the research and participated in reviewing the manuscript. All authors read and approved the final manuscript. JW and LY confirmed the authenticity of all the raw data.

Ethics approval and consent to participate

The study was approved by the Ethics Committee of Inner Mongolia Baogang Hospital (approval no. BG201802034YY).

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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