

The neuroprotective effect of recombinant human erythropoietin via an antiapoptotic mechanism on hypoxic-ischemic brain injury in neonatal rats

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Purpose: The neuroprotective effects of erythropoietin (EPO) have been recently shown in many animal models of brain injury, including hypoxic-ischemic (HI) encephalopathy, trauma, and excitotoxicity; however, limited data are available for such effects during the neonatal periods. Therefore, we investigated whether recombinant human EPO (rHuEPO) can protect against perinatal HI brain injury via an antiapoptotic mechanism.

Methods: The left carotid artery was ligated in 7-day-old Sprague-Dawley (SD) rat pups (*in vivo* model). The animals were divided into 6 groups: normoxia control (NC), normoxia sham-operated (NS), hypoxia only (H), hypoxia+vehicle (HV), hypoxia+rHuEPO before a hypoxic insult (HE-B), and hypoxia+rHuEPO after a hypoxic insult (HE-A). Embryonic cortical neuronal cell culture of SD rats at 18 days gestation (*in vitro* model) was performed. The cultured cells were divided into 5 groups: normoxia (N), hypoxia (H), and 1, 10, and 100 IU/mL rHuEPO-treated groups.

Results: In the *in vivo* model, Bcl-2 expressions in the H and HV groups were lower than those in the NC and NS groups, whereas those in the HE-A and HE-B groups were greater than those of the H and HV groups. The expressions of Bax and caspase-3 and the ratio of Bax/Bcl-2 were in contrast to those of Bcl-2. In the *in vitro* model, the patterns of Bcl-2, Bax, and caspase-3 expression and Bax/Bcl-2 ratio were similar to the results obtained in the *in vivo* model.

Conclusion: rHuEPO exerts neuroprotective effect against perinatal HI brain injury via an antiapoptotic mechanism.

Key Words: Erythropoietin, Apoptosis, Hypoxic-ischemic brain injury, Bcl-2, Bax, Caspase-3

Introduction

Erythropoietin (EPO), a hematopoietic cytokine known best for its erythropoietic effects, has neuroprotective effects in many animal models of brain injury, including hypoxic-ischemic (HI) encephalopathy, trauma and excitotoxicity¹. The biological effects of EPO in the central nervous system (CNS) are mediated by binding to its specific receptor (EPO-R). The expression of EPO and EPO-R in the nervous system is modulated by hypoxia and metabolic insult². EPO-R expression is abundant in the embryonic, fetal and adult brain of rats, mice, monkeys and humans^{3,4}. EPO has been identified as a neurotrophic and neuroprotective agent in a wide variety of experimental paradigms, from neuronal cell culture to *in vivo* models of brain injury. Although the exact mechanisms of the neuroprotective action of EPO are not completely known, promotion of cell survival signaling cascades, modulation of intracellular calcium metabolism, attenuation of nitric oxide (NO) production, inhibition of glutamate release, anti-apoptosis, antioxidative and anti-inflammatory actions have been suggested as possible mechanisms⁵.

EPO is known to promote cell survival via Bcl-2 family of genes in non-neuronal tissues and regulates the expression of apoptosis-related genes from Bcl-2 family in different experimental paradigms⁶. EPO is able to shift the Bcl/Bax ratio towards a net antiapoptotic effect in cultured microglia⁷. It also upregulates Bcl-xl mRNA and protein expressions in cultured neurons⁸. EPO reduced retinal ganglion cell death induced by glutamate and NO, and reversed Bcl-2 downregulation⁹. EPO also increases Bcl-2 gene expression in rat cardiac myocytes after traumatic brain injury¹⁰.

Various studies have demonstrated a protective effect of EPO in models of brain injury *in vitro* and *in vivo*^{7, 10, 11}. Here we investigated whether recombinant human EPO (rHuEPO) can protect the developing rat brain from HI injury via an anti-apoptosis analyzing possible molecular mechanisms accounting for the neuroprotective effect of rHuEPO.

Materials and methods

1. Materials

rHuEPO was purchased from CJ Cheiljedang Corporation (Seoul, Korea). Primary antibodies included the following: Bcl-2 (1:1,000; Santa Cruz Biotechnology, Santa Cruz, CA, USA), Bax (1:1,000; Cell Signaling technology, Danvers, MA, USA) and caspase-3 (1:1,000; Cell Signaling technology). Secondary antibodies used were goat anti-mouse or rabbit IgG-HRP (1:2000, Santa Cruz Biotechnology).

2. Animal protocol and drug administration

The neonatal rat HI procedure was performed as described by Rice et al.¹². We chose 7-day-old Sprague-Dawley (SD) rat pups weighing between 12 and 16 g. The rat pups were anesthetized with ether. The left common carotid artery (CCA) was exposed, isolated and permanently doubly ligated with 5-0 surgical silk. The incision was sutured and the animal was allowed to recover in a warm environment. The whole surgical procedure was completed in less than 5 minutes. After a recovery period of 1 hour, the pups were placed in plastic chambers, which were submerged in a 37°C water bath and exposed to hypoxia (mixture of 8% O₂ and 92% N₂) for 2 hours. After this hypoxic exposure, the pups were returned to their dams for the indicated time.

rHuEPO was prepared in phosphate-buffered saline (PBS) and injected intraperitoneally at a dose of 1,000 IU/kg 30 minutes before or after a hypoxic insult.

The animals were divided into six groups. In group 1 (Normoxia control, NC, n=7), they were not exposed to hypoxia. In group 2 (Normoxia Sham-operated, NS, n=8), they were subjected only to open-close surgery without CCA ligation. In group 3 (Hypoxia only, H, n=8), they were subjected to hypoxia without rHuEPO injection. In group 4 (Hypoxia+vehicle, HV, n=7), they were injected with PBS (the same volume of rHuEPO). In group 5 (Hypoxia+rHuEPO "before a hypoxic insult", HE-B, n=8), they were injected with 1,000 IU/kg of rHuEPO 30 minutes before a hypoxic insult. In group 6 (Hypoxia+rHuEPO "after a hypoxic insult", HE-A, n=8), they were injected with the same dose of rHuEPO 30 minutes after a hypoxic insult. Pups from each litter were randomly assigned and marked to the NC, NS, H, HV, HE-B or HE-A groups.

3. Hematoxylin and eosin (H & E) staining

Histologic studies were performed 7 days after a HI insult. After intracardiac perfusion with saline, the brains were removed, fixed in 4% paraformaldehyde, embedded in paraffin, and prepared for light microscopy. Four µm sections were mounted on glass cover slides. Sections were deparaffinized in xylene for 30 minutes and serially treated with 100% (5 minutes), 96% (10 minutes), and 70% (10 minutes) ethanol. Slides were stained with hematoxylin, rinsed for a few seconds with water to remove excess stain, placed in 1% eosin, rinsed briefly in water to remove the acid and washed through the following series: 70%, 95% and 100% ethanol at 5 minutes each. Finally, the slides were transferred to xylene for clearing, mounted, and covered with cover slips.

Each slides were taken a photograph and were measured brain area with a densitometer (Multi-gauze software, Fuji Photofilm,

Tokyo, Japan).

4. Cortical neuronal cell culture of rat embryos

The cortical neuronal cell culture of rat embryos based on the Brewer¹³ method was performed. SD rats pregnant for 19 days were anesthetized with ether for 5 minutes at room temperature and the uteri were removed. The fetal pups were washed in 100% ethanol and Hanks' balanced salt solution (HBSS) (GibcoBRL, Grand Island, NY, USA). The fetal pup brains were dissected at 37°C HBSS containing 1 mM sodium pyruvate and 10 mM HEPES (pH 7.4). The dissected brain cortical tissues were then placed in 2 mL trypsin and incubated at 37°C water bath for 1 minute. Trypsin was removed by washing the tissues five times with 10 mL HBSS. The cells were moved in 1 mL HBSS, passed 6-7 times via pasteur pipette with tiny hole and dispersed. The cell suspension was centrifuged at 1,000 rpm at 25°C for 5 minutes and pellets were washed with HBSS (without phenol red).

After cell counting, they were inoculated in plating Neurobasal media (GibcoBRL) (100 mL neurobasal, 2 mL B27 supplement, 0.25 mL glutamax I, 0.1 mL 25 mM glutamate, 0.1 mL 25 mM 2-mercaptoethanol) with about 2×10^6 cells/mm² in each dish. The cell cultures were incubated in a CO₂ chamber. A fifth of the culture solutions were changed every three days and were replaced with feeding Neurobasal media (GibcoBRL).

The cultured cells were divided into five groups: normoxia group (N), hypoxia group (H) and hypoxia+1, 10 and 100 IU/mL rHuEPO groups (H+E1, H+E10 and H+E100). The N group was prepared in 5% CO₂ incubators while the H group and H+E groups in 1% O₂ incubators (94% N₂, 5% CO₂) for 6 hours. The experiments were repeated four times (n=4).

5. 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide (MTT) assay

The MTT assay was used for estimation of cell viability and growth as originally described by Mosmann¹⁷). MTT was dissolved at a concentration of 5 mg/mL. Ten μ L of the 5 mg/mL MTT stock solution was added to each well. After 4 hour of incubation at 37°C, media was removed and added 100 μ L of the lysing buffer (Dimethyl sulfoxide (DMSO) : 95% ethanol = 1 : 1). Absorbance of the samples was read at 492 nm using a microtiter plate enzyme-linked immunosorbent assay (ELISA) reader. The amount of formazan produced is proportional to the number of live and metabolically active cells.

6. Brain extraction and protein isolation

Pups were killed 7 days after a hypoxic insult. Left cerebral

hemispheres from rat brains were immediately removed, frozen in liquid nitrogen and stored at -70°C until use. Frozen tissues were homogenized in protein lysis buffer containing complete protease inhibitor cocktail tablets (Roche Applied Science, Mannheim, Germany), 1 M Tris-HCl (pH 8.0), 5 M NaCl, 10% Nonidet P-40 and 1 M 1,4-dithio-DL-threitol (DTT). After incubation for 20 minutes on ice, the samples were centrifuged at 12,000 rpm at 4°C for 30 minutes and the supernatant was transferred to a new tube. Total protein was measured with a Bio-Rad Bradford kit (Bio-rad Laboratories, Hercules, CA, USA).

7. Western blotting

Samples of equal amounts of proteins (30 μ g) were subjected to a 12% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) after denaturing in SDS gel-loading buffer (100 mM Tris-HCl (pH 6.8), 200 mM DTT, 20% glycerol, 4% SDS and 0.2% bromophenol blue) in boiling water for 10 minutes. After electrophoresis proteins were electrotransferred onto polyvinylidene difluoride (PVDF) membrane (Millipore, Billerica, MA, USA) at a constant voltage of 10 V for 30 minutes. After transfer, the membrane was washed twice with 1 x tris-buffered saline (TBS) plus 0.1% Tween-20 (TBST pH 7.4) and then pre-incubated with a blocking buffer (5% nonfat dry milk in TBST) at room temperature for 1 hour. The membrane was then incubated with the primary antibodies 1:1,000 dilutions in TBST at 4°C overnight. After washing, the blots were incubated with the secondary antibodies (1:2,000 dilution) at room temperature for 1 hour. Finally, the membrane was washed and developed with an enhanced chemiluminescence (ECL) Plus Western Blotting Detection System (Amersham Biosciences, Piscataway, NJ, USA) or SUPEX (Neuronex, Pohang, Korea).

8. Semiquantitation of the western blots

The intensity of the corresponding western blot band was measured by using a densitometer (Multi Gauge Software) and was calculated as the ratio of the signal intensity in the ischemic (left) hemisphere compared to the contralateral (right) hemisphere.

9. RNA extraction and real-time PCR

Total RNA extraction was carried out with TRIzol reagent (Invitrogen Corporation, Carlsbad, CA, USA). Briefly, a piece of tissue was homogenized in 1 mL of TRIzol reagent. Total RNA was separated from DNA and proteins by adding chloroform and was precipitated using isopropanol. The precipitate was washed twice in 100% ethanol, air-dried and re-diluted in diethylpyrocarbonate (DEPC)-treated distilled water. The amount and purity of extracted

RNA was quantitated by GeneQuant 1,300 spectrophotometer (GE Healthcare, UK). The RNA was then stored at -70°C before further processing. For real-time PCR (for reverse transcription), total RNA ($1\ \mu\text{g}$) was reversely transcribed for 1 hour at 37°C in a reaction mixture containing 20 U RNase inhibitor (Promega, Madison, WI), 1 mM dNTP (Promega, Madison, WI, USA), 0.5 ng Oligo (dT) 15 primer (Promega, Madison, WI, USA), 1 x RT buffer and 200 U M-MLV reverse transcriptase (Promega, Madison, WI, USA). The reaction mixture was then incubated at 95°C for 5 minutes to stop the reaction. The cDNA was then stored

at -20°C before further processing.

Real-time PCR was performed in 48 well PCR plates (Mini Opticon™ Real-Time PCR System) using the FINNZYMES DyNAmo SYBR green qPCR kit (Finnzymes, Beverly, MA, USA). Amplification conditions are shown in Table 1. The thermal profile was 95°C for 15 minutes followed by 40 cycles of denaturation, annealing and extension as indicated in Table 1. Real-time PCR data were analysed with LightCycler software (Bio-rad Laboratories, Hercules, CA, USA). Each sample was conducted in triplicate.

10. Statistics analysis

Data were analyzed using the SPSS version 12 statistical analysis package (SPSS Inc., Chicago, IL, USA). Examined data were assessed using the t-test, GLM (general lineal model), and ANOVA. In each test, the data were expressed as the mean \pm SD, and $P < 0.05$ was accepted as statistically significant.

Table 1. Primer Pairs and Annealing Temperature for Real-time Polymerase Chain Reaction

Name	Primer Sequence (5'-3')	Annealing
Bcl-2	F:TTGACGCTCTCCACACACATG R:GGTGGAGGAACTCTTCAGGGA	57°C
Bax	F:TGCTGATGGCAACTTCAACT R:ATGATGGTTCTGATCAGCTCG	55°C
caspase-3	F:AATTCAAGGGACGGGTCATG R:GCTTGTGCGGTACAGTTTC	56°C

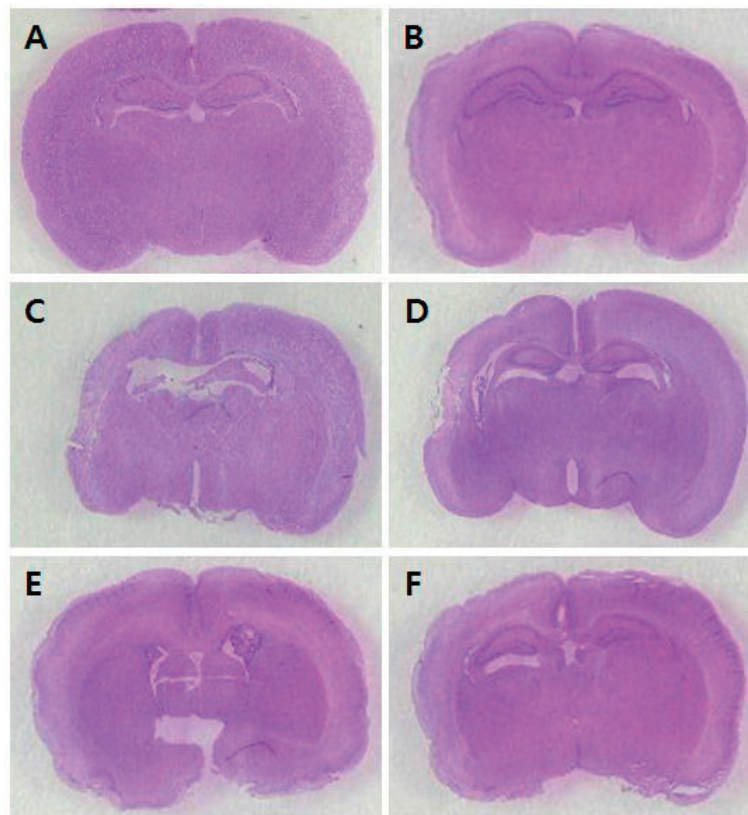


Fig. 1. Hematoxylin and eosin (H&E) staining after hypoxic insult of neonatal rat brain. A, Normoxia control group; B, Normoxia sham-operated group; C, Hypoxia group; D, Hypoxia+Vehicle group; E, Hypoxia+Recombinant human erythropoietin (rHuEPO)-treated group before hypoxic insult; F, Hypoxia+rHuEPO-treated group after hypoxic insult. Comparison of the percentage of the area of the left hemisphere area with that of the right hemisphere: A, 102.8%; B, 94.8%; C, 58.5%; D, 68.6%; E, 94.3%; and F, 81.4%.

Results

1. H & E staining after a hypoxic insult of neonatal rat brain

H & E stain of 7-day-old rat brain after a HI insult revealed that the brain areas in the H (C, 58.5%) and HV (D, 68.6%) groups were decreased compared to the NC (A, 102.8%) and NS (B, 94.8%) groups, whereas those in the HE-B (E, 94.3%) and HE-A (F, 81.4%) groups were increased compared to the H (C, 58.5%) and HV (D, 68.6%) groups (Fig. 1). Percentages were the ratios of left hemisphere area compared to right hemisphere area.

2. The expressions of Bcl-2, Bax and caspase-3 proteins by western blot analysis in the neonatal HI brain injury

The expressions of Bcl-2 in the H and HV groups were decreased compared to the NC and NS groups, whereas those in the HE-A and HE-B groups were increased compared to the H and HV groups ($P<0.05$) (Fig. 2A). The expressions of Bax and caspase-3 and the ratio of Bax/Bcl-2 in the H and HV groups were increased

compared to the NC and NS groups, whereas those in the HE-A and HE-B groups were decreased compared to the H and HV groups ($P<0.05$) (Fig. 2B, 2C and 2D).

3. The expressions of Bcl-2, Bax and caspase-3 mRNAs by real-time PCR on the HI brain injury in neonatal rats

The expressions of Bcl-2 in the H and HV groups were decreased compared to the NC and NS groups, whereas those in the HE-A and HE-B groups were increased compared to the H and HV groups ($P<0.05$) (Fig. 3A). The expressions of Bax and caspase-3 and the ratio of Bax/Bcl-2 in the H and HV groups were increased compared to the NC and NS groups, whereas those in the HE-A and HE-B groups were decreased compared to the H and HV groups ($P<0.05$) (Fig. 3B, 3C and 3D).

4. The images of cortical neuronal cell culture of rat embryos

The cortical neuronal cells were observed using light microscopy

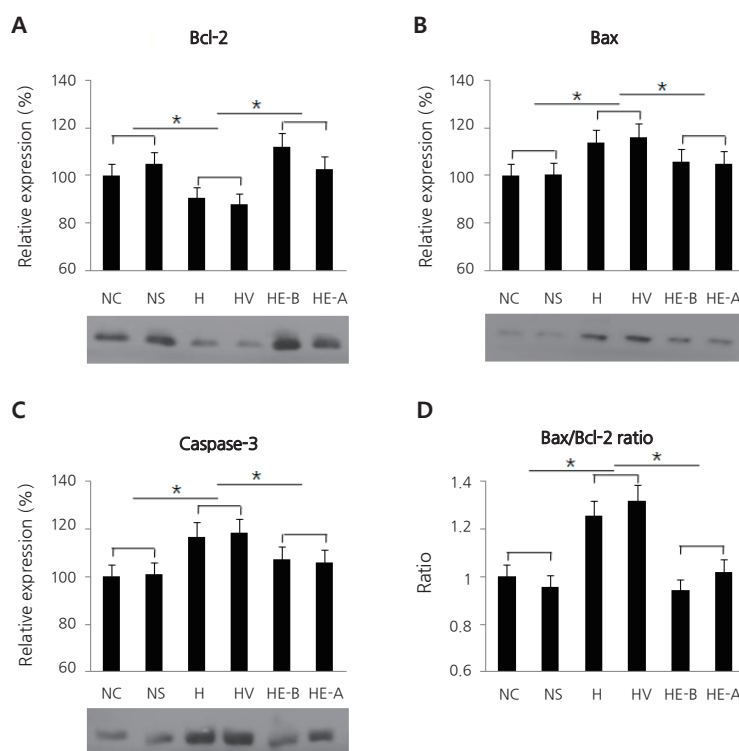


Fig. 2. Western blots of Bcl-2 (A, NC, 100±0; NS, 104.8±1.0; H, 90.4±1.14; HV, 87.9±0.56; HE-B, 112±1.12; HE-A, 102.8±0.37), Bax (B, NC, 100±0; NS, 100.3±0.73; H, 113.7±1.23; HV, 115.9±1.08; HE-B, 105.7±1.48; HE-A, 104.9±1.07), and caspase-3 (C, NC, 100±0; NS, 100.9±0.41; H, 116.9±1.57; HV, 118.5±1.76; HE-B, 107.2±1.63; HE-A (105.9±2.51) antibodies 7 days after hypoxic insult, and the ratio of Bax/Bcl-2 expression (D). Recombinant human erythropoietin (rHuEPO) was administered intraperitoneally at 1,000 IU/kg. NC, Normoxia control group (n=7); NS, Normoxia sham-operated group (n=8); H, Hypoxia group (n=8); HV, Hypoxia+Vehicle group (n=7); HE-B, Hypoxia+rHuEPO-treated group before hypoxic insult (n=8); HE-A, Hypoxia+rHuEPO-treated group after hypoxic insult (n=8). * $P<0.05$ compared with hypoxia.

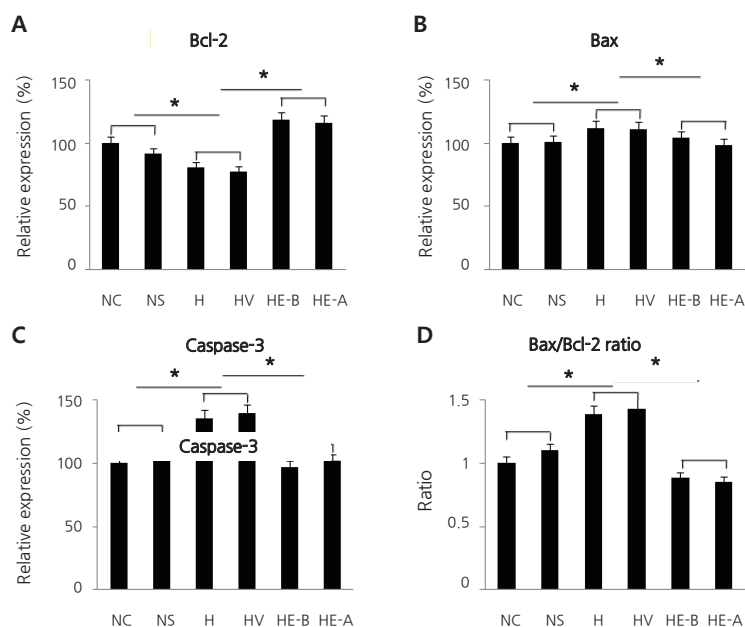


Fig. 3. Real-time PCR of Bcl-2 (A; NC, 100 ± 0 ; NS, 91.4 ± 2.63 ; H, 80.7 ± 1.1 ; HV, 77.4 ± 1.39 ; HE-B, 118.1 ± 1.63 ; HE-A, 115.7 ± 2.32), Bax (B; NC, 100 ± 0 ; NS, 100.7 ± 0.51 ; H, 111.7 ± 0.6 ; HV, 110.9 ± 1.24 ; HE-B, 104.2 ± 0.97 ; HE-A, 98.6 ± 1.35), and caspase-3 (C; NC, 100 ± 0 ; NS, 116.5 ± 1.23 ; H, 135.7 ± 2.8 ; HV, 139.5 ± 1.11 ; HE-B, 97.3 ± 0.68 ; HE-A, 102.1 ± 0.74) primers 7 days after hypoxic insult, and the ratio of Bax/Bcl-2 expression (D). Recombinant human erythropoietin (rHuEPO) was administered intraperitoneally at 1,000 IU/kg. NC, Normoxia control group ($n=7$); NS, Normoxia sham-operated group ($n=8$); H, Hypoxia group ($n=8$); HV, Hypoxia+Vehicle group ($n=7$); HE-B, Hypoxia+rHuEPO-treated group before hypoxic insult ($n=8$); HE-A, Hypoxia+rHuEPO-treated group after hypoxic insult ($n=8$). * $P < 0.05$ compared with hypoxia.

under high magnification ($\times 200$). The cells in the N group (Fig. 4A) appeared normal, whereas those in the H group (Fig. 4B) showed cellular damages. The cellular patterns of the H+E1 (Fig. 4C) and H+E10 group (Fig. 4D) were similar to those in the N group, whereas the cells in the H+E100 group (Fig. 4E) appeared similar to those in the H group.

5. Cell viabilities of cultured cortical neuronal cells determined by the MTT assay

The relative cell viabilities in the H group were decreased compared to the N group. The relative cell viabilities in the H+E1 and H+E10 group were similar to those in the N group, whereas those in the H+E100 group appeared similar to those in the H group (Fig. 5).

6. The expressions of Bcl-2, Bax and caspase-3 by proteins western blot analysis in the cortical neuronal cell culture

The expressions of Bcl-2 in the H group was decreased compared to the N group, whereas those in the H+E10 group were increased compared to the H group ($P < 0.05$) (Fig. 6A). The expressions

of Bax and the ratio of Bax/Bcl-2 in the H group were increased compared to the N group, whereas those in the H+E10 group were decreased gradually depending to the concentration of rHuEPO ($P < 0.05$) (Fig. 6B and 6D). The expressions of caspase-3 in the H group were increased compared to the N group, whereas those in the H+E1 and H+E10 groups were decreased compared to the H group ($P < 0.05$) (Fig. 6C).

7. The expressions of Bcl-2, Bax and caspase-3 mRNA by real-time PCR in cortical neuronal cell culture

The expressions of Bcl-2 in the H group were decreased compared to the N group, whereas those in the H+E1 group was increased compared to the H group ($P < 0.05$) (Fig. 7A). The expressions of Bax and the ratio of Bax/Bcl-2 in the H group were increased compared to the N group, whereas those in the H+E1, H+E10 and H+E100 groups were decreased compared to the H group ($P < 0.05$) (Fig. 7B and 7D). The expressions of caspase-3 in the H group were increased compared to the N group, whereas those in the H+E1 and H+E10 groups were decreased compared to the H group ($P < 0.05$) (Fig. 7C).

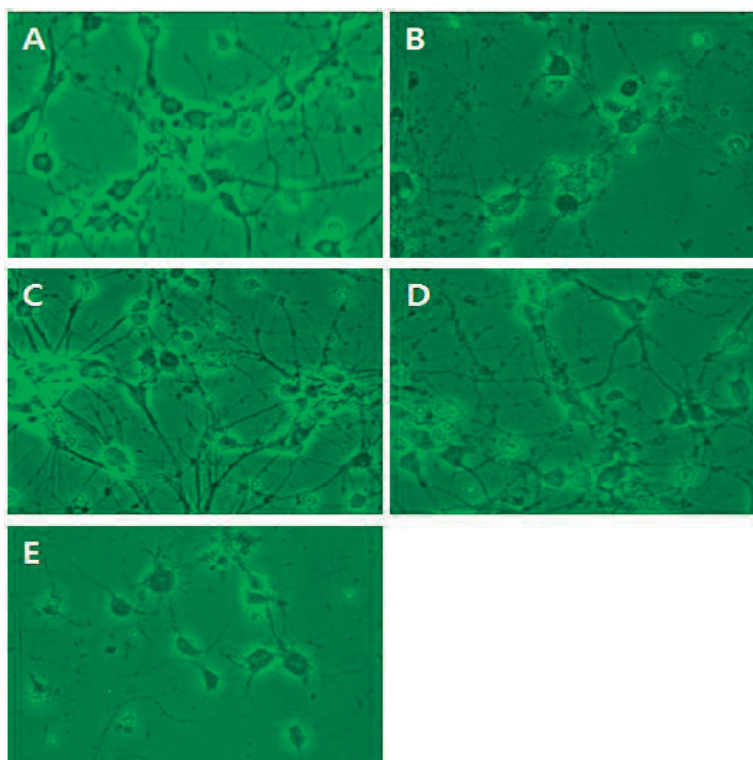


Fig. 4. High-magnification ($\times 200$) photomicrograph of brain cortical cell culture from a 19-day-pregnant Sprague-Dawley rat. Recombinant human erythropoietin (rHuEPO) was administered at 1, 10, and 100 IU/mL. A, Normoxia group (N); B, Hypoxia group (H); C, Hypoxia+1 IU/mL rHuEPO-treated group (H+E1); D, Hypoxia+10 IU/mL rHuEPO-treated group (H+E10); E, Hypoxia+100 IU/mL rHuEPO-treated group (H+E100).

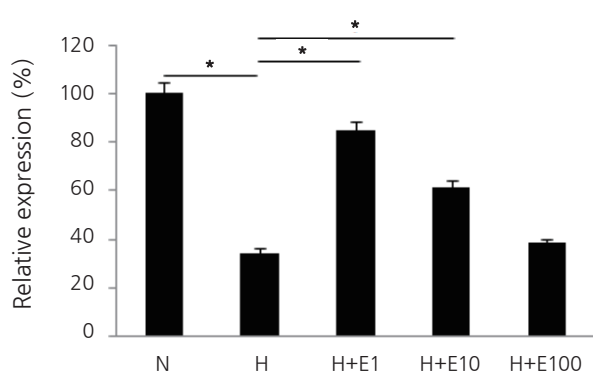


Fig. 5. Cell viabilities were measured by the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide (MTT) assay. N (100 ± 0), normoxia group; H (34.6 ± 0.7), hypoxia group; H+E1 (84.6 ± 1.72), hypoxia+1 IU/mL recombinant human erythropoietin (rHuEPO)-treated group; H+E10 (61.5 ± 1.14), hypoxia+10 IU/mL rHuEPO-treated group; H+E100 (38.5 ± 0.89), hypoxia+100 IU/mL rHuEPO-treated group. $*P < 0.05$ compared with hypoxia.

Discussion

HI brain injury in the human perinatal period can result from several pathological processes such as premature placental detachment or other complications of pregnancy and delivery. It is an important cause of neonatal mortality and subsequent sequelae

such as cerebral palsy, mental retardation, learning disability and epilepsy¹⁵. Hypoxia, ischemia, and more recently, intrauterine and neonatal infections, have been implicated in the pathogenesis of injury to the developing brain¹⁵. Neonates who were exposed to HI brain injury sustain both acute and chronic morbidities that can affect the sensory, motor, cognitive and behavioral systems. The full extent of this injury may not be apparent until the child reaches school age or later⁵. Premature neonates are particularly vulnerable to HI exposure and may develop a variety of deficits that will reveal themselves when tested at school entry¹⁶.

A variety of therapeutic strategies are employed to protect against the pathologic mechanisms of HI injury. These include induction of hypothermia, antagonism of excitotoxic glutamate receptors, anti-inflammatory agents, neurotropic factors and inhibitors of the caspase pathway and of stress kinases⁵. Pharmacological agents which mediate one or more of these processes may provide neuroprotection for patients against this condition. Furthermore, study of neuroprotective agents may provide important information about treatment for the perinatal HI brain injury. The *in vivo* neonatal rat HI model and the *in vitro* cortical cell culture model of rat embryos have been well characterized and used extensively to search for neuroprotective agents¹⁷.

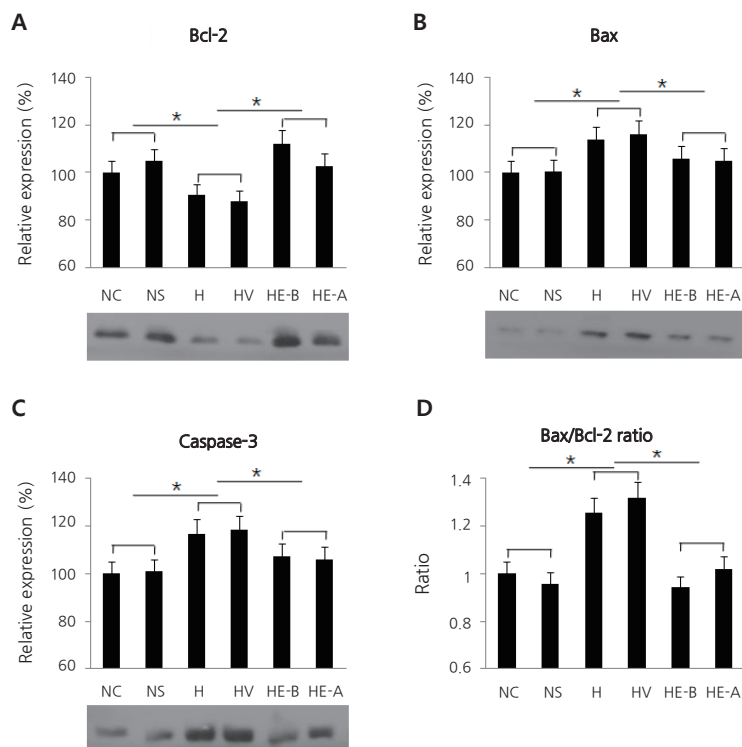


Fig. 6. Western blots of Bcl-2 (A; N, 100 ± 0 ; H, 97.6 ± 0.96 ; H+E1, 95.2 ± 0.45 ; H+E10, 102.9 ± 0.75 ; H+E100, 98.7 ± 0.85), Bax (B; N, 100 ± 0 ; H, 104.3 ± 0.57 ; H+E1, 97.7 ± 0.7 ; H+E10, 91 ± 0.87 ; H+E100, 84.3 ± 0.54), and caspase-3 (C, N, 100 ± 0 ; H, 107.8 ± 0.71 ; H+E1, 99.2 ± 0.63 ; H+E10, 104.6 ± 0.89 ; H+E100, 126.9 ± 0.61) in cultured cortical neuronal cells from 19-day-old rat embryos, and the ratio of Bax/Bcl-2 expression (D). Recombinant human erythropoietin (rHuEPO) was administered at 1, 10, and 100 IU/mL. A, Normoxia group (N, n=4); B, Hypoxia group (H, n=4); C, Hypoxia+1 IU/mL rHuEPO-treated group (H+E1, n=4); D, Hypoxia+10 IU/mL rHuEPO-treated group (H+E10, n=4); E, Hypoxia+100 IU/mL rHuEPO-treated group (H+E100, n=4). * $P < 0.05$ compared with hypoxia.

EPO, a 34-kDa glycoprotein hormone, was first characterized as a hematopoietic growth factor. It is now widely used for the treatment of anemia associated with renal failure, cancer, prematurity, chronic inflammatory disease and human immunodeficiency virus infection. EPO is initially produced in the liver but shortly after birth, its production is shifted to the kidney¹⁸. EPO has been originally characterized as the principal regulator of erythropoiesis by inhibiting apoptosis and by stimulating the proliferation and differentiation of erythroid precursor cells¹⁹. EPO exerts remarkable neuroprotection in both cell cultures and in animal models of nervous system disorders and has been implicated in the development of the CNS. Neuroprotective effect of EPO has also been demonstrated on various animal models of cerebral ischemia²⁰.

Recent studies have shown that EPO protects neonatal brains as well from HI injury by reducing apoptotic cell death *in vivo*²¹ and protects cultured neurons against oxygen glucose deprivation injury by a paracrine protective pathway through astrocytes expressing

EPO²². EPO is also found to protect the developing rat brain against NMDA receptor antagonist neurotoxicity and protects the mouse retina against light-induced degeneration by inhibiting photoreceptor cell apoptosis²³. Our *in vivo* experimental study revealed that the brain areas in hypoxia were decreased compared to normoxia, whereas those in EPO-treated groups were increased compared to hypoxia.

In *in vitro* experiments, EPO has been shown to protect cultured hippocampal and cortical neurons against glutamate toxicity²⁴, hypoxia and glucose deprivation²⁵ and serum deprivation-induced and kainic acid-induced apoptosis²⁴. In addition, systemic administrations of rHuEPO as a pretreatment or posttreatment have been shown to effectively reduce brain injury after focal ischemia²⁶, subarachnoid hemorrhage²⁷ and HI²⁸. Our *in vitro* experimental study showed the cells in normoxia appeared normal, whereas those in hypoxia showed cellular damages. And the cellular patterns in the low dose EPO-treated groups were similar to those in normoxia, whereas the cells in the high dose EPO-treated groups

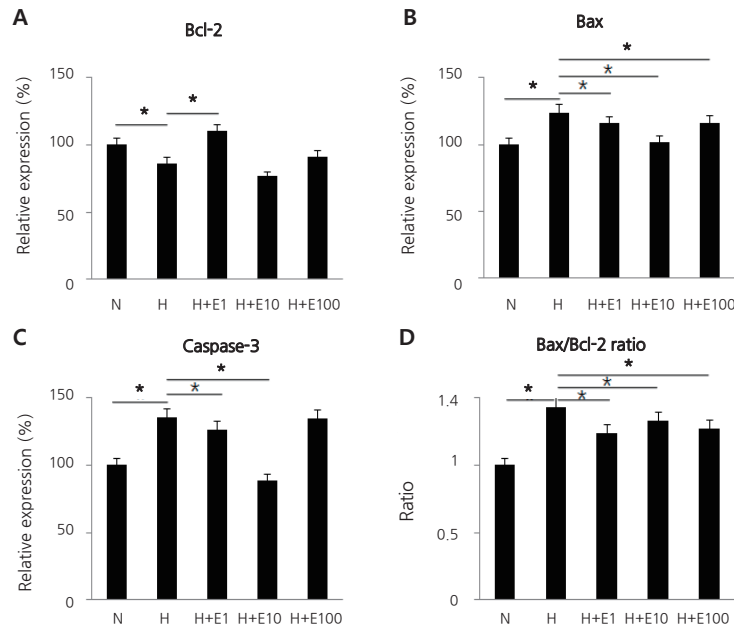


Fig. 7. Real-time PCR of Bcl-2 (A; N, 100 ± 0 ; H, 86.5 ± 0.84 ; H+E1, 110.2 ± 2.1 ; H+E10, 76.8 ± 1.38 ; H+E100, 91.4 ± 0.98), Bax (B; N, 100 ± 0 ; H, 123.9 ± 1.31 ; H+E, 115.7 ± 1.6 ; H+E10, 102.1 ± 1.89 ; H+E100, 116.5 ± 0.82), and caspase-3 (C; N, 100 ± 0 ; H, 135.7 ± 0.79 ; H+E1, 126.6 ± 1.16 ; H+E10, 88.9 ± 0.69 ; H+E100, 134.7 ± 0.74) mRNAs in cultured cortical neuronal cells from 19-day-old rat embryos, and the ratio of Bax/Bcl-2 expression (D). Recombinant human erythropoietin (rHuEPO) was administered at 1, 10, and 100 IU/mL. A, Normoxia group (N, n=4); B, Hypoxia group (H, n=4); C, Hypoxia +1 IU/mL rHuEPO-treated group (H+E1, n=4); D, Hypoxia+10 IU/mL rHuEPO-treated group (H+E10, n=4); E, Hypoxia+100 IU/mL rHuEPO-treated group (H+E100, n=4). * $P < 0.05$ compared with hypoxia.

appeared similar to those in hypoxia.

Both EPO and its specific receptor (erythropoietin receptor, EPO-R) are expressed in different tissues, including the nervous system¹⁹. EPO acts via activation of the specific EPO-R. EPO and EPO-R are present in the developing human brain as early as 5 weeks post-conception, and each protein shows a specific distribution that changes with development²⁹. EPO-R have been identified in the nervous system of rodents, primates and humans^{19, 29} and have been localized on several primary cell lines of neurons, as well as in glial cells. It has been shown that EPO stimulates neurogenesis and prevents apoptosis in the embryonic brain³⁰. EPO-R is widespread throughout fetal brain²⁹. EPO-R binding initiates a signaling cascade that involves multiple antiapoptotic pathways that can account for the rHuEPO-induced reduction²⁶. EPO-R activation protects hippocampal neurons from ischemic neuronal damage through the inhibition of glutamate release via exocytosis in hippocampal slice cultures. In the brain, EPO is released by astrocytes²², triggers dopamine release³¹, promotes neurogenesis³² and vasculogenesis³³, and stimulates proliferation of glia³⁴. EPO is therefore recognized to have both trophic and antiapoptotic mechanisms of action.

EPO might have a direct protective role against a variety of neurotoxic insults such as hypoxic conditions²⁴, glutamate toxicity¹⁹, free-radical injury³⁵ and exposure to neurotoxins³⁶. EPO stimulates dopamine release and tyrosine hydroxylase activity in PC₁₂ cells, through the activation of calcium channels, induces membrane depolarization, stimulates mitogen activated protein kinase activity and increases NO synthesis³⁷. EPO reduces calcium-induced glutamate release from cultured cerebellar granule neurons³⁸.

Apoptosis contributes significantly to cerebral damage during the perinatal period. In neonatal HI model, there is evidence that cerebral ischemia leads to delayed cell death with DNA damage³⁹. EPO has antiapoptotic action on some non-neural cells⁴⁰. There is some evidence suggesting that EPO may also protect neuronal cells that morphologically die via apoptosis both *in vivo* and *in vitro*. And EPO may also exert its neuroprotective effect via the differential regulating of gene expression ruling the balance between the expression of pro- and antiapoptotic molecules. EPO prevents ischemia-induced delayed neuronal death in the hippocampus in animal model of global cerebral ischemia¹¹. EPO decreases apoptotic cell death of neurons under conditions

of hypoxia *in vitro*, suggesting that it might have similar effect *in vivo*²⁹⁾. Antiapoptotic action of EPO has also been confirmed in focal cerebral ischemia and transient spinal ischemia models²⁴⁾. EPO decreases apoptotic cell death of neurons under conditions of hypoxia *in vitro*, suggesting that it might have a similar effect *in vivo*²⁹⁾.

EPO inhibits caspase 8-, caspase 1- and caspase 3-like activities linked to cytochrome c release³⁵⁾. EPO consistently increases the expression of the antiapoptotic gene Bcl-XL and decreases the expression of pro-apoptotic gene bak in PC12 cells. EPO is able to shift the Bcl/Bax ratio towards a net antiapoptotic effect in cultured microglia⁷⁾. It also up-regulates Bcl-XL mRNA and protein expressions in cultured neurons⁸⁾. EPO rescued hippocampal CA1 neurons from lethal ischemic damage and up-regulated the expressions of Bcl-XL mRNA and protein in the hippocampal CA1 field of ischemic gerbils. Thus, EPO prevents delayed neuronal death of hippocampal neurons, possibly through up-regulation of Bcl-XL, which is known to facilitate neuronal survival⁸⁾.

Western blot analysis and real-time PCR both our *in vitro* and *in vivo* models of perinatal HI brain injury revealed that the expressions of Bcl-2 in hypoxia were decreased compared to in normoxia whereas those in rHuEPO-treated groups were increased compared to in hypoxia. The expressions of Bax after hypoxia as well as the ratio of Bax/Bcl-2 in hypoxia were increased compared to normoxia whereas those in rHuEPO-treated groups decreased compared to hypoxia. In addition, the expressions of caspase-3 in hypoxia were increased compared to normoxia whereas those in rHuEPO-treated groups were decreased compared to normoxia.

The above data suggest that rHuEPO's neuroprotective effect against neonatal HI brain injury may have been via an anti-apoptosis mechanism which the ratio of Bax/Bcl-2 and the expressions of caspase-3 in rHuEPO-treated groups were increased compared to hypoxia.

In conclusion, our experiments demonstrate that rHuEPO is able to prevent the degeneration of neonatal cerebral neuronal cells caused by a hypoxic insult. In addition, rHuEPO's neuroprotective effects on HI brain injury in neonatal rats may be via anti-apoptosis. The present study may be useful for the further development of clinical therapies for perinatal HI encephalopathy-induced by cerebral hypoxia.

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