



Effects of carnosine and hypothermia combination therapy on hypoxic-ischemic brain injury in neonatal rats

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Background: Carnosine has antioxidative and neuroprotective properties against hypoxic-ischemic (HI) brain injury. Hypothermia is used as a therapeutic tool for HI encephalopathy in newborn infants with perinatal asphyxia. However, the combined effects of these therapies are unknown.

Purpose: Here we investigated the effects of combined carnosine and hypothermia therapy on HI brain injury in neonatal rats.

Methods: Postnatal day 7 (P7) rats were subjected to HI brain injury and randomly assigned to 4 groups: vehicle; carnosine alone; vehicle and hypothermia; and carnosine and hypothermia. Carnosine (250 mg/kg) was intraperitoneally administered at 3 points: immediately following HI injury, 24 hours later, and 48 hours later. Hypothermia was performed by placing the rats in a chamber maintained at 27°C for 3 hours to induce whole-body cooling. Sham-treated rats were also included as a normal control. The rats were euthanized for experiments at P10, P14, and P35. Histological and morphological analyses, *in situ* zymography, terminal deoxynucleotidyl transferase-mediated dUTP nick end-labeling (TUNEL) assays, and immunofluorescence studies were conducted to investigate the neuroprotective effects of the various interventional treatments.

Results: Vehicle-treated P10 rats with HI injury showed an increased infarct volume compared to sham-treated rats during the triphenyltetrazolium chloride staining study. Hematoxylin and eosin staining revealed that vehicle-treated P35 rats with HI injury had decreased brain volume in the affected hemisphere. Compared to the vehicle group, carnosine and hypothermia alone did not result in any protective effects against HI brain injury. However, a combination of carnosine and hypothermia effectively reduced the extent of brain damage. The results of *in situ* zymography, TUNEL assays, and immunofluorescence studies showed that neuroprotective effects were achieved with combination therapy only.

Conclusion: Carnosine and hypothermia may have synergistic neuroprotective effects against brain damage following HI injury.

Keywords: Carnosine, Hypothermia, Hypoxia-ischemic brain injury

Key message

Question: How can we further improve the neuroprotective effects of hypothermia in newborns with hypoxic-ischemic (HI) brain injury?

Finding: Combination carnosine and hypothermia therapy effectively reduced brain damage in neonatal rats. The *in situ* zymography, terminal deoxynucleotidyl transferase-mediated dUTP nick end-labeling assay, and immunofluorescence study results showed that neuroprotective effects were achieved with combination therapy only.

Meaning: Carnosine and hypothermia have synergistic neuroprotective effects against brain damage following HI injury.

Introduction

Hypothermia is known to have protective effects against hypoxic-ischemic (HI) brain injury in neonatal animal studies.¹⁻³ Additionally, as a result of large-scale clinical studies of newborns with HI encephalopathy in the early 2000s, therapeutic hypothermia was found to have effects against brain damage caused by perinatal asphyxia.^{4,5} Subsequently, in 2010, there was an international consensus regarding the application of hypothermia for newborns with HI encephalopathy.⁶ Currently, therapeutic hypothermia has been established as a standard treatment strategy for HI encephalopathy, and it is used for full-term and some near-term infants with moderate to severe HI encephalopathy.^{7,8} The neuroprotective actions of hypothermia comprise multifactorial mechanisms, including the reduction of the cell metabolic rate, prevention of high-energy phosphorus compound deficiency, inhibition of excitatory amino acids and inflammatory cytokines, and suppression of programmed cell death and microglial activation at different stages of neuronal cell damage.^{7,9} Hypothermia is the only treatment method of newborns with HI brain injury, but the therapeutic benefits are

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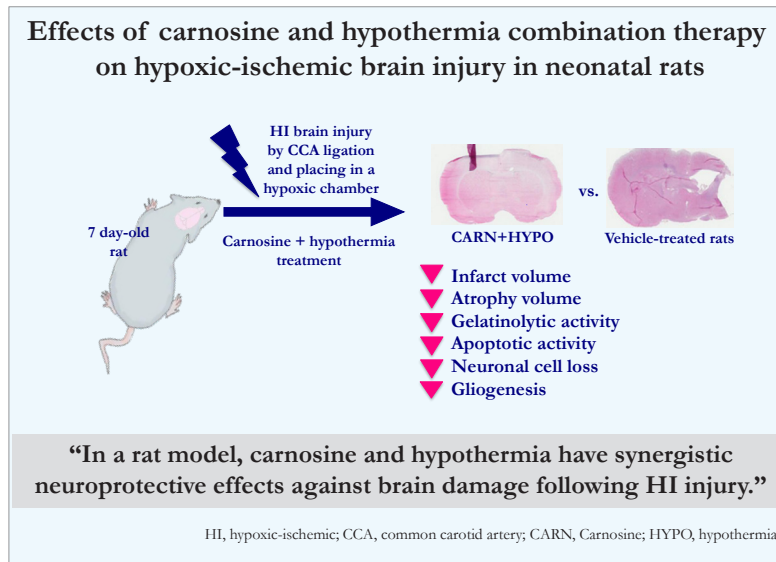
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Received: 4 November, 2020, Revised: 22 January, 2021, Accepted: 5 February, 2021

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Graphical abstract

still limited.¹⁰⁻¹²⁾ Recently, combination therapies involving hypothermia and adjunctive agents, including phenobarbital,¹³⁾ topiramate,¹⁴⁾ erythropoietin,¹⁵⁾ and xenon,¹⁶⁾ have been reported to have better therapeutic effects than hypothermia alone.

Carnosine is an endogenous dipeptide (β -alanyl-L-histidine) that is distributed in several parts of the body, including the skeletal and cardiac muscles and nervous tissues.^{17,18)} Recently, carnosine has been shown to have neuroprotective effects against HI brain injury in laboratory and animal studies. In cultured cortical astrocytes under ischemic conditions, carnosine showed the ability to preserving mitochondrial functions by decreasing the intracellular metabolism during the HI phase and improving cell viability by increasing energy production during the reperfusion phase.¹⁹⁾ In cultured neurons/astrocytes exposed to oxygen glucose deprivation/recovery, carnosine exhibited neuroprotective effects by regulating extracellular glutamate and gamma-aminobutyric acid levels and mitochondrial energy metabolism.²⁰⁾ During the study of the mouse model, carnosine alleviated brain damage by mechanisms such as inhibiting reactive oxygen species (ROS) and matrix metalloproteinase (MMP) expression of HI lesions and maintaining glutathione activity.²¹⁾

Thereafter, this study investigated whether carnosine has additional neuroprotective effects on induced HI brain injury in neonatal rats when used in combination with hypothermic therapy.

Methods

1. Experimental protocol

The study protocol and all experimental procedures were approved by the institutional animal research and ethics committee of Keimyung University School of Medicine (approval number: KM-2014-15). Neonatal Sprague-Dawley rats (Koatech-Harlan, Pyeongtaek, Korea) with a body weight of 14 to 18 g at

postnatal day 7 (P7) were used during this study. Anesthesia with isoflurane was administered to induce ischemic brain damage in the rats. After exposing the right common carotid artery of the rat, it was isolated and cut between the double ligations. After surgery, the rats were allowed to recover with the dam for 60 minutes in a cage. Hypoxic injury was caused by placing rats in a chamber at 37°C with a flow of mixed gas comprising 8% humidified oxygen and 92% nitrogen for 90 minutes under positive pressure.

The study included 2 protocols: protocol I was involved the administration of carnosine or vehicle after HI injury and hypothermic therapy and protocol II involved the administration of carnosine or vehicle after HI injury without hypothermic therapy (Fig. 1). After inducing brain damage, the rats were transferred to a cage; after 15 minutes, carnosine or vehicle was administered; and rats were allowed to recover with the dam for 60 minutes. Carnosine (Sigma-Aldrich, Seoul, Korea) was dissolved in distilled water and intraperitoneally administered immediately after HI injury, at a dose of 250 mg/mL/kg. Additionally, after 24 hours and 48 hours, carnosine or vehicle was administered a total of 3 times. Hypothermia was performed by placing the rats in a chamber maintained at 27°C for 3 hours to induce whole-body cooling. The study was divided into the following 4 groups (n=8-9 per group): vehicle-treated HI group; carnosine-treated HI group; HI group with vehicle and hypothermia; HI group with carnosine and hypothermia. Sham-treated rats, they were not subjected to any HI injury and included in the study as normal control.

2. Histological and morphological studies

Experimental rats were euthanized by decapitation at P10 (3 days after HI injury). Brains were dissected out immediately, and coronal-sectioned specimens were prepared with a thickness of 1 mm using razor blades. They were then stained with 2% 2,3,5 triphenyltetrazolium chloride (TTC) solution for 30 minutes at 37°C, fixed with formalin solution, and photographed with a

digital camera; the cerebral infarction size was measured using an Image J program (version 1.45; <http://rsbweb.nih.gov/ij/download.html>; National Institute Health, Bethesda, MD, USA). The extent of brain infarction was expressed as a percentage of the corresponding area on the undamaged cerebral hemisphere to correct brain edema by the injury.

Experimental rats were also euthanized by decapitation at P35 (4 weeks after HI injury). Brains were dissected out and immediately cooled. Subsequently, 14- μ m-thick coronal slices were prepared using a cryostat (MEV, Mainz, Germany). Sample slides were fixed in formalin for 30 minutes; then, hematoxylin and eosin (H&E) staining was performed. The brain damage

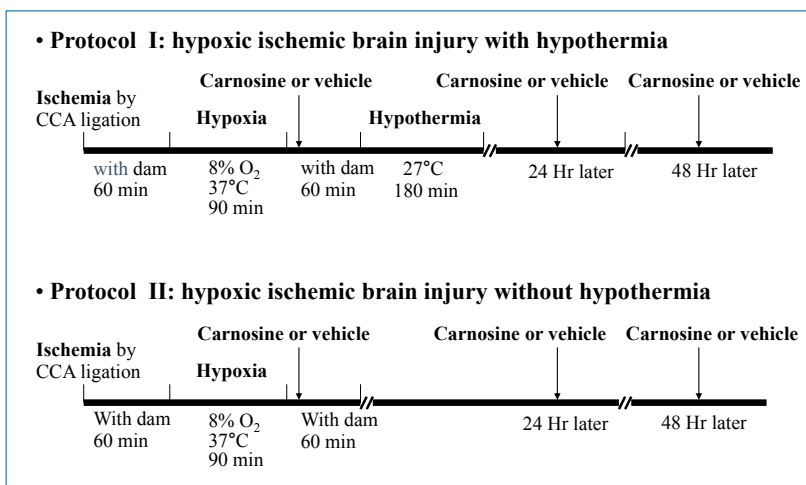


Fig. 1. The diagram shows the experimental processes for protocols I and II. CCA, common carotid artery.

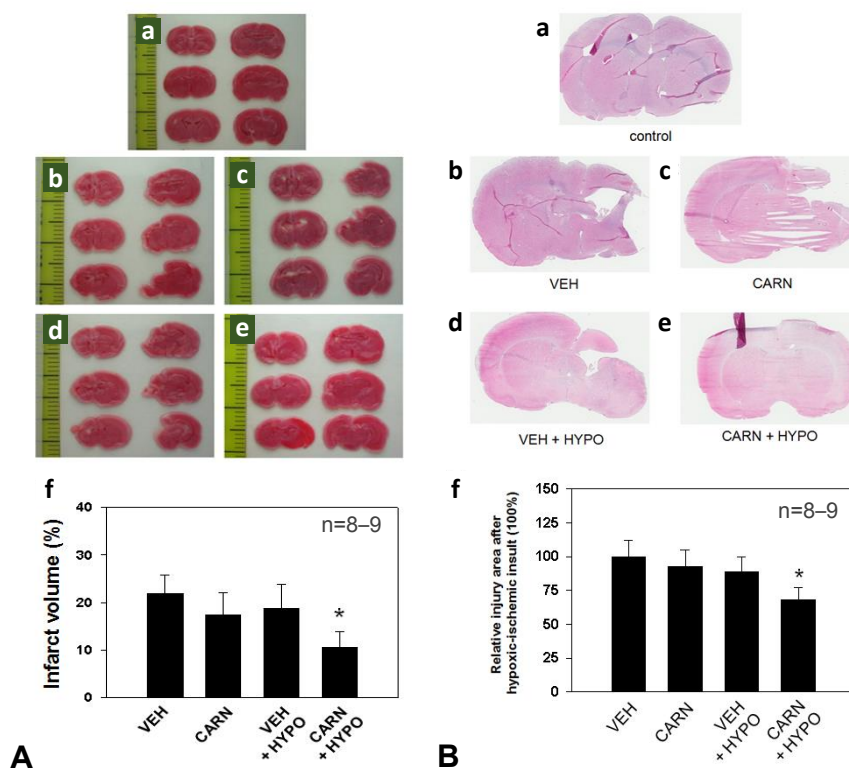


Fig. 2. Histological analysis of hypoxic-ischemic (HI) brain injury in experimental rats. (A) Representative sections of the brain stained with triphenyltetrazolium chloride (TTC) showing infarction areas in P10 rats. Compared to the vehicle (VEH) group, cerebral infarction was significantly reduced with combined carnosine (CARN) and hypothermia (HYPO) therapy. (B) Brain damage findings observed using hematoxylin and eosin staining in P35 rats. Compared to VEH group, brain atrophy was significantly reduced according to the image analysis when a combination of CARN and HYPO was used. P10, postnatal day 10; P35, postnatal day 35; a, sham-treated; b, VEH-treated HI group; c, CARN-treated HI group; d, HI group treated with VEH and HYPO; e, HI group treated with CARN and HYPO. * $P < 0.05$ vs. VEH-treated HI group.

image was obtained with an optical microscope and digital camera. The brain damage contour was examined using an image analyzer, and the extent of cerebral atrophy was evaluated and compared with the undamaged hemisphere.

3. *In situ* zymography

After euthanizing the rats, they were perfused with phosphate-buffered saline (PBS) and brains were rapidly dissected out and frozen in 2-methylbutane and liquid nitrogen. Brain tissue was prepared as section specimens of 14 μm using a cryostat, placed in an incubator at 37°C for 18 hours according to the instructions of the gelatinase activity assay kit (Enz-Check kit; Invitrogen, Waltham, MA, US), and then examined by fluorescence microscopy.

4. Terminal deoxynucleotidyl transferase-mediated dUTP nick end-labeling assay

After euthanizing the rats, they were perfused with PBS and brains were dissected out and frozen in 2-methylbutane and liquid nitrogen. Brain tissue was prepared as section specimens with a thickness of 14 μm using a cryostat. Tissue samples were fixed with 10% formalin solution and washed 3 times with PBS. After following the instructions of the terminal deoxynucleotidyl transferase-mediated dUTP nick end-labeling (TUNEL) staining

kit (Roche Diagnostic, Indianapolis, IN, USA) and, adding the reaction solution to the sample tissue, diaminobenzidine substrate solution was added to detect chromogenic apoptosis. Then, the samples were examined using microscopy.

5. Immunofluorescence

After euthanizing the rats, they were perfused with PBS, and 4% paraformaldehyde (PFA) solution were used to fix the brain tissue. Brains were rapidly dissected out and further fixed with 4% PFA solution; they were soaked with 15% and 30% sucrose solution for 24 hours before freezing. Brain tissue was prepared as section specimens with a thickness of 20 μm using a cryostat. Slides were prepared and stored in a deep freezer. Samples were blocked with PBS containing 0.3% Triton X-100 and 3% normal goat serum. Subsequently, primary antibodies, such as anti-neuronal nuclear antigen (anti-NeuN) (1:100; Millipore, Bedford, MA, USA) or anti-gial fibrillary acidic protein (anti-GFAP) (1:200; Millipore), were added to the sample and incubated at 4°C for 18 hours. During the next step, a secondary antibody, anti-mouse tetramethylrhodamine-5-isothiocyanate (1:100; Jackson ImmunoResearch, West Grove, PA, USA) was added to the sample and incubated for 30 minutes. Then, the sample was washed with PBS and examined using fluorescence microscopy.

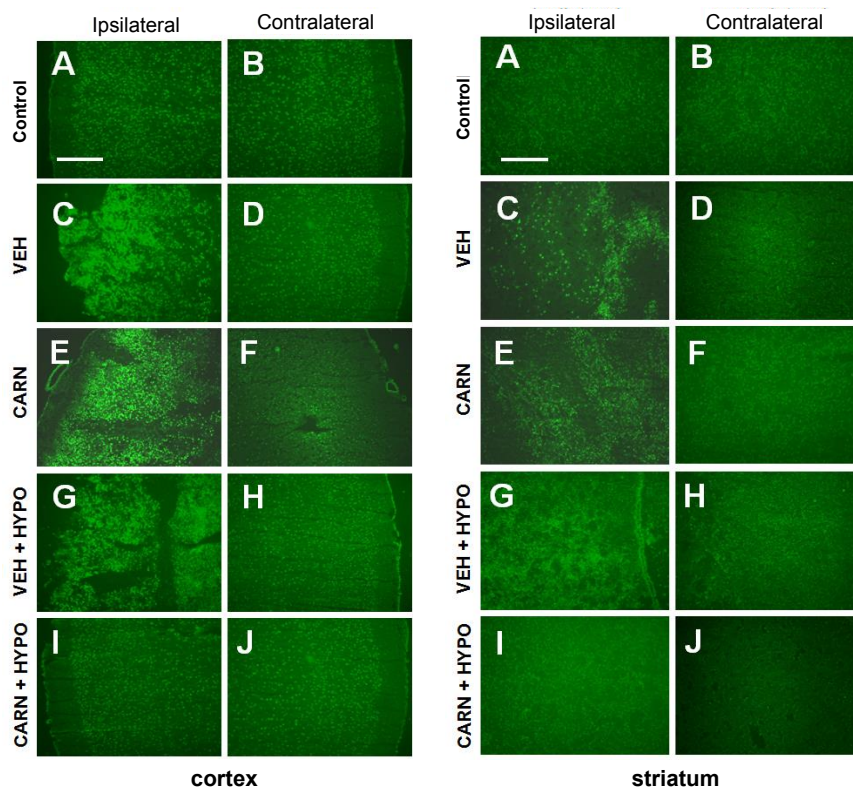


Fig. 3. Gelatinolytic activity of hypoxic-ischemic (HI) brain lesions detected with *in situ* zymography in experimental rats. Gelatinase activity of the affected cerebral cortex and corpus striatum increased compared to that of the contralateral sides of the vehicle (VEH) group of P14 rats. Compared to the VEH group, gelatinolysis did not decrease with carnosine (CARN) or hypothermia (HYPO) alone. However, combined CARN and HYPO therapy reduced enzyme activity. P14, postnatal day 14; A and B, sham-treated; C and D, VEH-treated HI group; E and F, CARN-treated HI group; G and H, HI group treated with VEH and HYPO; I and J, HI group treated with CARN and HYPO. Scale bar = 500 μm .

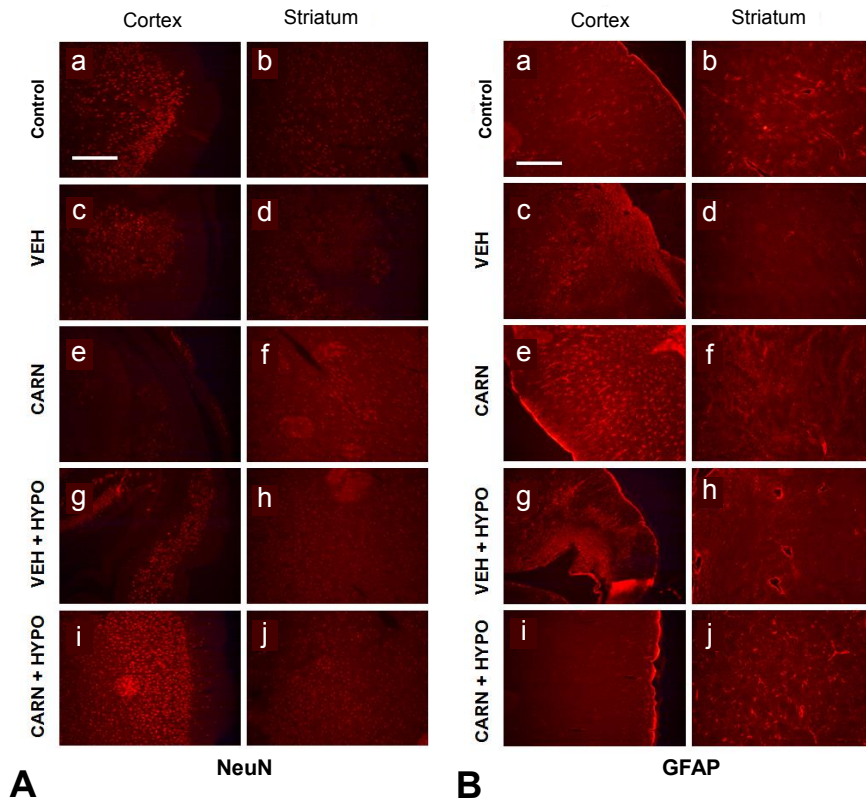


Fig. 5. Immunofluorescence findings of hypoxic-ischemic (HI) brain lesions in experimental rats. (A) Expression of neuronal nuclear antigen (NeuN). Neuronal cells of the cerebral cortex and corpus striatum in the P35 vehicle group were decreased compared to sham-treated rats. Compared to vehicle (VEH) group, carnosine (CARN) or hypothermia (HYPO) alone did not prevent NeuN-positive cell loss. However, combined CARN and HYPO therapy reduced neuronal cell loss. (B) Expression of glial fibrillary acidic protein (GFAP). Glial cells of the cerebral cortex and corpus striatum in the P35 VEH group were increased compared to sham-treated rats. Compared to the VEH group, CARN or HYPO alone did not prevent an increase in GFAP-positive cells. However, combined CARN and HYPO therapy suppressed gliogenesis. P35, postnatal day 35; a and b, sham-treated; c and d, VEH-treated HI group; e and f, CARN-treated HI group; g and h, HI group treated with VEH and HYPO; i and j, HI group treated with CARN and HYPO. Scale bar = 500 μ m.

to sham-treated rats. Carnosine alone or hypothermia alone did not prevent NeuN-positive cell loss. However, combined carnosine and hypothermia therapy reduced neuronal cell loss compared with vehicle (Fig. 5A).

Expression of GFAP-positive cells of the cerebral cortex and corpus striatum in the P35 vehicle group was increased compared to sham-treated rats. Although carnosine alone or hypothermia alone did not prevent an increase in GFAP-positive cells, combined carnosine and hypothermia therapy suppressed gliogenesis compared with vehicle (Fig. 5B).

Discussion

When comparing the brain development status between rats and humans according to postnatal age, 7-day-old rats are equivalent to term infants, and 10-day-old rats are closely related to infants with 40–42 weeks gestational age.^{22,23} Moreover, rats at 15-day-old are equivalent to a few months to 1 year old infants, and rats at 28–33 days-old are correspond to 2-year-old infants.^{22,24} In this study, combination therapy for P7 rats

with HI brain injury started with the first dose of carnosine administered immediately after the injury. Subsequently, hypothermia was applied for 3 hours; then, secondary and tertiary doses of carnosine were administered. After these interventional treatments, the therapeutic effect against brain damage was evaluated in rats at P10, P14, and P35 based on the morphological and histological findings of the affected parts. In morphological evaluation, carnosine alone and hypothermia alone did not show any protective effects against HI brain injury. However, a combination of carnosine and hypothermia morphologically reduced the extent of brain lesions compared with vehicle. Additionally, the results of *in situ* zymography, TUNEL assays, and immunofluorescence studies showed that neuroprotective effects were achieved with only combined therapy.

According to studies of neonatal rats, neuroprotective effects of hypothermia against HI brain damage vary depending on the severity of HI injury, and is also affected by the onset of cooling and the duration of hypothermia.^{25,26} Sabir et al.¹² reported that 5-hour hypothermic therapy for 7-day-old rats with a moderate HI injury is most effective when applied immediately after injury, when applied up to 6 hours after injury, the infarct area increases

by 1.8% per hour. However, when hypothermic therapy is applied at 12 hours or more after injury, has no neuroprotective effects. Additionally, for rats with a severe HI injury, hypothermia had no protective effects immediately or within 6 hours after injury. Neuroprotective effects of 3-hour hypothermia after inducing HI insults in 7-day-old rats differ according to the severity of the injury. Tresche et al.¹⁾ reported that there were neuroprotective effects in rats with a mild HI injury, whereas Liu et al.¹⁴⁾ pronounced that there were no therapeutic effects was expressed in rat with moderate HI injury. In the experimental protocol of this study, neonatal rats were exposed to hypoxia for 90 minutes after ligation of a carotid artery, resulting in moderate HI brain damage,¹²⁾ and then the rats were exposed to the environment of 27°C to lower the target rectal temperature to 28°C,¹⁴⁾ and the cooling is applied for 3 hours. Therefore, it is considered that neuroprotective effects were not observed with hypothermia alone because cooling was performed for only 3 hours for moderately severe brain injury. In future studies, if a cooling time of 5 hours or more is applied, hypothermia alone is expected to have therapeutic effects.¹²⁾

During a study of a neonatal rat model by Zhang et al.,²⁴⁾ carnosine was administered as a dose of 250 mg/kg intraperitoneally 3 times immediately after HI injury, 24 hours later, and 48 hours later. During the evaluation of brain tissues approximately 72 hours after the injury, administration of a series of carnosine doses showed neuroprotective effects by lowering the concentration of prostaglandin F2 α in brain tissue and reducing expression levels of neuronal apoptosis and mitochondrial caspase-3 protein. In this study, carnosine was administered according to an existing reported protocol,²⁴⁾ but the use of carnosine alone had no neuroprotective effects. Other studies of rodent models have reported that the neuroprotection of carnosine after HI injury is concentration-dependent and the protective effect is evident when at high doses of 500 to 1,000 mg/kg were used or when doses were administered prior to brain damage.^{18,21,27)} Therefore, it may be necessary to increase the dosage of carnosine and evaluate the changes in neuroprotective effects bases on the blood concentrations of the therapeutic agent.

To enhance the brain-protective effects of hypothermia after HI injury, several studies have been conducted to determine whether the administration of combined hypothermia and adjuvant agents has synergistic effects on neuroprotection. Liu et al.¹⁴⁾ reported that topiramate administered immediately after HI injury extends the therapeutic window for hypothermia-mediated neuroprotection in a neonatal rat model. In their study that evaluate the brain volume showing cerebral infarction and atrophy after HI injury, topiramate alone and hypothermia alone had no neuroprotective effects; however, a combination of topiramate and hypothermia significantly reduced the extent of cerebral lesions. Martin et al.¹⁶⁾ studied a neonatal rat model with HI injury and reported that asynchronous administration of xenon after hypothermia had a synergistic effect on neuroprotection. In their study, the extent of cerebral infarction after

HI injury did not decrease with hypothermia alone or xenon alone, but the lesion size was significantly reduced when xenon was administered asynchronously at 1- or 5-hour intervals after hypothermia.

In this study, interventional measures performed after the injury had no therapeutic effects with hypothermia alone or adjuvant alone, and protective effects were observed with combined administration, similar to other studies of neonatal rat model of HI brain injury.^{14,16)} After combined administration of carnosine and hypothermia, gelatinolysis in the brain tissue of the affected side decreased. Neuronal apoptosis was reduced approximately 1 week after the injury, and this reduction was observed even after 4 weeks. Additionally, loss of neuronal cells and gliogenesis of the lesion were suppressed by a combined therapy. This study had some limitations. First, according to the protocol, hypothermia was applied to decrease the target temperature to 28°C, but the rectal temperature of all rats used in this study was not measured. Furthermore, quantitative analyses of histological changes in brain lesions following therapeutic measures were not performed.

In this study, carnosine was repeatedly administered before and after hypothermia; however, the cellular or molecular mechanisms of carnosine that would increase the neuroprotection of hypothermia is uncertain. Presumably, it is likely related to various biological roles of carnosine against HI injury, including inhibiting ROS and MMP expression and suppressing microglial activation and neuronal apoptosis.^{27,28)}

In conclusion, the combined carnosine and hypothermia therapy in a neonatal rat model of HI injury synergistically reduced the extent of brain injury. Therefore, it could be a therapeutic strategy for HI encephalopathy of newborns. Further research is needed to elucidate the molecular mechanisms associated with their neuroprotective functions.

Footnotes

Conflict of interest : No potential conflict of interest relevant to this article was reported.

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How to cite this article: Byun JC, Lee SR, Kim CS. Effects of carnosine and hypothermia combination therapy on hypoxic-ischemic brain injury in neonatal rats. *Clin Exp Pediatr* 2021;64:422-9. <https://doi.org/10.3345/cep.2020.01837>