In Vivo Neuroprotective Effect of Histidine-Tryptophan-Ketoglutarate Solution in an Ischemia/Reperfusion Spinal Cord Injury Animal Model

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Background: Paraplegia is a devastating complication following operations on the thoracoabdominal aorta. We investigated whether histidine-tryptophan-ketoglutarate (HTK) solution could reduce the extent of ischemia/reperfusion (IR) spinal cord injuries in a rat model using a direct delivery method.

Methods: Twenty-four Sprague-Dawley male rats were randomly divided into four groups. The sham group (n=6) underwent a sham operation, the IR group (n=6) underwent an aortic occlusion, the saline infusion group (saline group, n=6) underwent an aortic occlusion and direct infusion of cold saline into the occluded aortic segment, and the HTK infusion group (HTK group, n=6) underwent an aortic occlusion and direct infusion of cold HTK solution into the occluded aortic segment. An IR spinal cord injury was induced by transabdominal clamping of the aorta distally to the left renal artery and proximally to the aortic bifurcation for 60 minutes. A neurological evaluation of locomotor function was performed using the modified Tarlov score after 48 hours of reperfusion. The spinal cord was harvested for histopathological and immunohistochemical examinations.

Results: The spinal cord IR model using direct drug delivery in rats was highly reproducible. The Tarlov score was 4.0 in the sham group, 1.17±0.75 in the IR group, 1.33±1.03 in the saline group, and 2.67±0.81 in the HTK group (p=0.04). The histopathological analysis of the HTK group showed reduced neuronal cell death.

Conclusion: Direct infusion of cold HTK solution into the occluded aortic segment may reduce the extent of spinal cord injuries in an IR model in rats.

Key words: 1. Spinal cord 2. Reperfusion injury

Introduction

Paraplegia has been reported to account for 5% to 20% of devastating complications following operations on the thoracoabdominal aorta [1-4]. Several methods exist to prevent this postoperative complication, such as (1) distal perfusion with shunts or a partial bypass, (2) hypothermia, (3) drainage of the cerebrospinal fluid, (4) reattachment of the intercostal vessels, (5) evoked-potential monitoring, and (6) the development of pharmacological agents to increase tolerance against ischemic insults. However, it is neces-
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Table 1. Experimental group (n=6)

<table>
<thead>
<tr>
<th>Group</th>
<th>Body weight (g)</th>
<th>Infusion</th>
<th>Ischemia (min)</th>
<th>Reperfusion (hr)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sham</td>
<td>262.5±5.2</td>
<td>No</td>
<td>0</td>
<td>No</td>
</tr>
<tr>
<td>Ischemia/reperfusion</td>
<td>263.3±5.2</td>
<td>No</td>
<td>60</td>
<td>48</td>
</tr>
<tr>
<td>Saline</td>
<td>266.7±7.5</td>
<td>Cold saline 10 mL/kg</td>
<td>60</td>
<td>48</td>
</tr>
<tr>
<td>HTK</td>
<td>264.4±5.8</td>
<td>Cold HTK solution 10 mL/kg</td>
<td>60</td>
<td>48</td>
</tr>
</tbody>
</table>

HTK, histidine-tryptophan-ketoglutarate.

sary to develop a better prevention method, and many studies have been conducted with the goal of developing a mechanism to replicate spinal cord ischemia/reperfusion (IR) injuries in animal models [1,5-9].

Drugs reducing the impact of spinal cord injuries can be categorized into blood-flow maintenance agents and metabolic regulation agents [10]. Examples of the former type of drug include papaverine, nitroglycerine, and nitroprusside. The latter type of drug is subdivided into neuronal metabolic rate reducers (barbiturates and opiates), cell membrane stabilizers (calcium channel blockers and sodium-channel blockers), removers of toxic metabolites (superoxide dismutase and allopurinol), alleviators of metabolite-induced inflammation (steroids, prostaglandin, and adenosine), and excitotoxicity reducers (riluzole, memantine, and magnesium) [10-12].

Bretschneider [13] first developed histidine-tryptophan-ketoglutarate (HTK) solution (Bretschneider’s solution, Custodiol) in the 1970s, and it is still widely used in organ preservation for transplantation (e.g., the kidney, liver, and pancreas) and myocardial protection during open-heart surgery [14]. HTK solution is an effective organ preservation solution due to the powerful buffer action of histidine, the cell membrane stabilization and antioxidant effects of tryptophan, and the role of ketoglutarate as a substrate [13,15].

Despite its use as a preservation solution for various organs, the effect of HTK on the peripheral nervous system has only been studied by Lassner et al. [16] in 1995. Moreover, its protective effect on the central nervous system and its potential role in IR injuries have not been studied. Recent studies have reported neuroprotective effects of histidine analogs (a principal component of HTK), which were enhanced by the excitotoxicity-reducing effect of HTK solution [17,18]. The principal aim of the present study was to confirm the role of HTK solution as a protective solution for spinoplegia. Therefore, an animal model involving direct drug infusion into the occluded aortic segment was developed to determine the effect of HTK in spinal cord nervous tissue.

Methods

1) Experimental animals
Male Sprague-Dawley (SD) rats weighing approximately 250 to 300 g were used in this study. Approval from the Animal Care and Use Committee of Chungnam National University was obtained. All experimental animals were not fed for twelve hours prior to the experiment, and they were confirmed to be neurologically sound.

2) Experimental groups
Twenty-four male SD rats were randomly divided into 4 groups. The sham group (n=6) underwent a sham operation. The IR group (n=6) underwent 60 minutes of aortic occlusion and 48 hours of reperfusion. The saline infusion group (n=6) was infused with 10 mL/kg of 4°C saline into the occluded aortic segment, as well as undergoing 60 minutes of aortic occlusion and 48 hours of reperfusion. The HTK infusion group (n=6) was infused with 10 mL/kg of HTK solution into the occluded aortic segment, as well as undergoing 60 minutes of aortic occlusion and 48 hours of reperfusion (Table 1).

3) Surgical method
1) Development of the aortic occlusion and spinal cord ischemia/reperfusion model: In order to assess the reproducibility of the spinal cord IR model using direct drug delivery in rats, dye was injected and hemodynamic changes were measured in 3 model rats. The model rats were anesthetized through an intraperitoneal injection of 50 mg/kg of pentobarbital. Subsequently, an intraperitoneal injection of 100 IU/kg
Silk string loops were applied at the aorta distally to the left renal artery and proximally to the aortic bifurcation (white arrow). The left common femoral artery was cannulated with a polyethylene-10 tube (black arrow).

The cross-clamp was applied at the aorta distally to the left renal artery and proximally to the aortic bifurcation. When dye was infused, it was visible only in the occluded segment of the aorta (arrow).

The IR spinal cord injury operation was performed under ×16 magnification. The region of operation was aseptically disinfected with betadine. The left common carotid artery was cannulated with a polyethylene (PE)-50 tube (external diameter of 0.97 mm) to continuously monitor real-time hemodynamic changes. The PE-50 tube was connected with a PE-10 tube (external diameter of 0.60 mm) to the dissected left inguinal area. The left femoral artery was cannulated with a PE-10 tube, through which the drug was injected and the blood pressure of the model rat was measured. The same procedure was performed in the dissected right inguinal area, and the right femoral artery was cannulated to measure the distal blood pressure. A median thoracoabdominal incision was made, and the abdominal organs were moved towards the right side of the model rats. The aorta was carefully dissected distally to the left renal artery and tied with 5-0 silk sutures. The proximal portion of the aortic bifurcation was dissected together with the inferior vena cava. The intubated cannula inside the left femoral artery was palpably advanced to the location between the left renal artery and the aortic bifurcation. The loops of both ends were pulled to occlude the aorta for 60 minutes to induce an ischemic spinal cord injury (Fig. 1).

Reperfusion of the aorta was induced after 60 minutes of aortic occlusion, and the femoral artery cannulas were removed. The femoral arteries were ligated for hemostasis. The operative wound was closed with continuous 4-0 nylon sutures, and 50 mg/kg of first-generation cephalosporin and antibiotic cream were applied. The model rats were then moved back to a cage, and were verified to have regained consciousness. The rats were supplied with food and water starting 6 hours after the operation. Urine was drained twice per day by compressing the hypogastrium of the model rats. Tryphan blue dye (2
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mL) was injected in the region around the occluded aorta to confirm that the drug was delivered solely to the site of occlusion (Fig. 2).

(2) Drug delivery in the ischemia/reperfusion spinal cord injury model: The IR spinal cord injury models were prepared as described above, but the left common carotid artery and right femoral artery were not intubated. Instead, the left femoral artery was cannulated with a PE-10 tube and a median thoracoabdominal incision was made to occlude the aorta. The IR group was injected with no drug, the saline group was injected with 10 mL/kg of 4°C cold saline for 5 minutes, and the HTK group was injected with 10 mL/kg of 4°C HTK solution for 5 minutes. The surgical method described above was then followed.

4) Locomotor function test
All experimental animals were tested for locomotor function using the modified Tarlov scoring system [19,20]. In a small portion of the experimental animals, the locomotor function scores of the ipsilateral limb were lower than were observed for the contralateral limb. In such animals, the score of the contralateral limb was used as a reference value. Grade 0 was given to animals with spastic paralysis and no movement of the hind limb. Grade 1 was given to animals with spastic paralysis and slight movement of the hind limb. Grade 2 was given to animals that exhibited good movement of the hind limb but were unable to stand. Grade 3 was given to animals that could stand but were unable to walk normally, and grade 4 was given to animals that made a complete recovery.

5) Postoperative tissue collection
All experimental animals were sacrificed after the locomotor function test, which was performed 48 hours after surgery. Anesthesia was induced through an intramuscular injection of 50 mg/kg of ketamine hydrochloride and 50 mg/kg of xylazine hydrochloride. The animals were put to death by cervical vertebra dislocation. The lower thoracic vertebrae and lumbar vertebrae were extracted. The extracted vertebrae were stored in 10% neutral formalin for histopathological and immunohistochemical tests.

6) Histopathological tests
(1) Hematoxylin-eosin staining: The extracted tissues were fixed in 10% neutral formalin for 3 days. Four slices 2 mm in thickness in the transverse plane of the second and third vertebrae of the lumbar spinal cord were obtained and microscope slides were prepared. The slides were fixed in 10% neutral formalin and paraffin, and sliced to be 5 μm in thickness. The slides were then stained with hematoxylin and eosin dye to study the nuclei and cytoplasm of the cells.

(2) Assessment of pathological grade: A pathologist conducted a blind pathological assessment of each experimental group. The pathological grade was given based on the proportion of red neurons with ischemic injury and the area of infarction [19,20]. The red neurons exhibited cell shrinkage, pyknosis, degradation of the nucleus, Nissl body disappearance, and eosinophilic staining. The images of the slides were acquired using a scanner at ×400 magnification (Aperio Imagescope ver. 11.0; Leica, Rockleigh, NJ, USA). The number of red neurons was measured using an image analysis program (ImageJ; DeNovo software, Los Angeles, CA, USA). The number of red neurons was measured at the anterior horn of the spinal cord at ×200 magnification, and a grade was assigned according to the ratio of red neurons to total neurons. Grade 0 was assigned when red neurons were not detected. Grade 1 was given for a ratio of red neurons to total neurons of <15%. Grade 2 corresponded to a ratio of red neurons to total neurons of >15%. Grade 3 corresponded to focal and mild infarction affecting <25% of the gray matter, and grade 4 was given to samples with diffuse and severe infarction affecting >25% of the gray matter.

7) Statistics
PASW SPSS ver. 18.0 for Windows (SPSS Inc., Chicago, IL, USA) was used to analyze the data and to perform statistical analyses. Continuous variables are expressed as means±standard deviations, and categorical variables are expressed as median values. Analysis of variance was used to compare continuous variables across the experimental groups. The Kruskal-Wallis test was conducted to compare pathological findings and Tarlov scores. Significant results were evaluated using the post hoc test with the Mann-Whitney rank sum law. For hemodynamic variables, the bootstrap-t method or the paired t-test using a bootstrap was
Table 2. Hemodynamic profile before and after aortic occlusion (n=3)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Proximal</th>
<th>Distal</th>
<th>p-value</th>
<th>Proximal</th>
<th>Distal</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Resting</td>
<td>Occlusion</td>
<td></td>
<td>Resting</td>
<td>Occlusion</td>
<td></td>
</tr>
<tr>
<td>Systolic blood pressure</td>
<td>108.3±4.5</td>
<td>120.9±5.6</td>
<td>0.004</td>
<td>121.7±5.9</td>
<td>22.7±0.9</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Diastolic blood pressure</td>
<td>96.2±2.1</td>
<td>105.1±3.2</td>
<td>0.005</td>
<td>105.3±3.2</td>
<td>21.4±0.9</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Mean blood pressure</td>
<td>102.7±3.1</td>
<td>113.3±4.2</td>
<td>0.006</td>
<td>114.1±4.5</td>
<td>22.2±0.8</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Pulse pressure</td>
<td>12.2±3.5</td>
<td>15.7±8.9</td>
<td>0.031</td>
<td>15.9±4.6</td>
<td>1.0±0.4</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Heart rate</td>
<td>343.3±29.6</td>
<td>336.7±22.5</td>
<td>0.252</td>
<td>343.3±29.6</td>
<td>336.7±22.5</td>
<td>0.252</td>
</tr>
</tbody>
</table>

Values are presented as mean±standard deviation.

used to compensate for the low number of samples. All p-values < 0.05 were considered to indicate statistical significance.

Results

1) Overall surgical data

Unexpected cases of hemorrhage that occurred during the intubation of the catheter into femoral artery or aortic dissection were excluded from the total of experimental cases. The average operation time was approximately 90 minutes for non-problematic operations. Every experimental animal regained consciousness from anesthesia without any problems, and no instances of mortality were noted. The average length of PE-10 tubing that was inserted was approximately 3.5–4.0 cm.

2) Changes in hemodynamic indices

When occlusions were made in the aorta distally to the left renal artery and in the aorta at the aortic bifurcation, the proximal pressure increased from 102.7±3.1 mmHg to 113.3±4.2 mmHg (p=0.04) and the distal pressure decreased from 14.1±4.5 mmHg to 22.2±0.8 mmHg (p=0.05). The pulse pressure increased from 12.2±3.5 mmHg to 15.7±8.9 mmHg (p=0.31) and the heart rate decreased from 343.3±29.6 beats/min to 336.7±22.5 beats/min (p=0.25), but no significant change in body temperature was observed (Table 2, Fig. 3). When cold saline was injected after occluding the aorta, a transient decrease of 1.5°C in the body temperature was noted, but the body temperature returned to normal (Fig. 4).

3) Results of motor neuron tests

The sham group exhibited normal lower limb locomotor function in all cases. The IR group showed a significant difference in comparison to the sham group (1.17±0.75, p=0.01). The result of the motor neuron test for the saline group was 1.33±1.03, whereas it was 2.67±0.81 for the HTK group, which was a significant difference (p=0.04) (Fig. 5).
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Fig. 4. Hemodynamic changes before, during, and after occlusion of the aorta with infusion of cold saline in the saline infusion model.

Fig. 5. The Tarlov score of each group. Versus saline group (p=0.04).

4) Histopathological assessment

Red neurons were rarely found in the sham group, numerous red neurons were observed in the IR group and the saline group, while few red neurons were observed in the HTK group (Fig. 6). Using image analysis software, the number of red neurons was determined to be 1.0±0.81 for the sham group, 17.0±2.53 for the IR group, 15.0±5.18 for the saline group, and 5.5±4.73 for HTK group (p=0.02). The histopathological scores were 0.67±0.52 for the sham group, 2.67±0.52 for the IR group, 2.67±0.52 for the saline group, and 1.17±0.41 for the HTK group (p=0.04) (Figs. 7, 8).

Discussion

When blood flow to an organ is blocked and the oxygen supply cannot meet the demand for oxygen, the normal phosphorylation process does not occur in mitochondria, resulting in ATP depletion [10]. The shortage of ATP causes abnormalities in the membrane potential and imbalances in intracellular electrolytes. Influxed calcium ions induce the secretion of various enzymes, denature nucleic acids and proteins, and cause reactive oxygen production and cytotoxicity. An increase in intracellular sodium ions induces cellular swelling [10,18], and the increase of lactic acid in anaerobic conditions causes intracellular acidification [10,12]. An inflammatory reaction is caused by the accumulation of toxic substances, increasing the amount of nitric oxide synthase, which in turn increases free radicals [9,10,20].

Excitotoxicity is a damage process unique to neurons in which N-methyl-D-aspartate (NMDA) or α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid receptors are overexpressed due to glutamate and its analogs [10]. Excitotoxicity interrupts an increase of intracellular sodium ions, exacerbates edema, and inhibits the maintenance of a normal membrane potential. As a consequence, the amount of free reactive oxygen species increases, which in turn increases the recruitment of inflammatory cells to the damaged area [10]. Apoptosis occurs in damaged cells because they continuously secrete inflammatory mediators and
activate neutrophils, macrophages, and T-cells [20].

HTK solution is an organ preservation solution principally composed of histidine, tryptophan, and ketoglutarate, which are substances that can freely pass through the blood-brain barrier [13]. The strong buf-
fering action of histidine prevents a pH drop inside the cell and inhibits anaerobic processes, thereby blocking the accumulation of lactic acid [10]. Tryptophan has a protective effect on the cellular membrane, ketoglutarate acts as a substrate in anaerobic conditions [13], and mannitol blocks cellular swelling [21]. HTK solution has a low viscosity (310 osm) and contains a low amount of K+ (9 mmol) and Na+ (15 mmol). In order to achieve equilibrium of the intracellular electrolytes at the initiation of perfusion, HTK solution is recommended to be administered at a dosage of 1 mL/min/g (per weight of organ) at 4°C to 6°C for more than 5 minutes [13]. However, the allowable dose of HTK solution for SD rats is limited, so the maximum amount (10 mL/kg for 5 minutes) was chosen.

In addition to the widely known organ preservation effect of HTK solution, recent studies have reported that histidine analogs are capable of reducing ischemic injury by decreasing excitotoxicity. Shen et al. [17] verified a significant decrease in NMDA levels mediated by L-carnosine (beta-alanyl-L-histidine) in a spinal cord IR rat model, and Tang et al. [18] reported that macrophages could be modified to display neuroprotective effects. Stvolinsky et al. [22] confirmed that the antioxidant effect and cellular membrane stabilization effect of histidine allow it to have a neuroprotective effect. Research on each component of HTK solution has yet be conducted, but the neuroprotective effect of HTK solution via the buffering action of histidine and various other mechanisms has been elucidated [22].

Although many drug injection pathways exist, oral administration is not an efficient method for simulating actual clinical methods used in emergency settings [10]. The simplest method for drug delivery is via intravenous infusion. An advantage of intravenous infusion is that additional handling of the blood vessels is not needed, but it also has disadvantages. For instance, selective injection to a targeted organ is impossible, and adverse reactions and toxicity due to the systemic circulation of the drug are possible. Another method of drug delivery is retrograde intravenous infusion via the inferior vena cava or the hemiazygos vein. This method requires additional manipulation of the blood vessels, and is also disadvantageous because drug metabolism may be altered when the drug passes through the hepatic circulation and because toxicity due to the systemic circulation of the drug remains possible. A third method is direct injection into the spinal artery, which was performed in the present study. This is considered the ideal method because it causes the least amount of systemic toxicity. Moreover, intrathecal injections are advantageous because they do not require drugs to pass through the blood-brain barrier and the drugs administered in this manner act directly on the spinal cord. The disadvantages of this procedure include the need for the additional manipulation of blood vessels, the possibility of serious adverse effects involving hemorrhage, the risk of nerve injury, and the increase of cerebrospinal pressure induced by the drug injection [10].

Spinal cord IR models have already been developed in various types of experimental animals. Various methods are used to induce spinal cord IR injuries, such as occluding the thoracic aorta using a Fogarty catheter via the femoral artery, occluding the aorta by direct thoracic incision, and occluding the aorta through laparotomy [19,20,23-25]. Another method involves occluding the aorta distally from the left renal artery. This method was used in rabbits during the early stages of development of the spinal cord IR model, but it can now be used in SD rats as well [26].

Different amounts of time for inducing an IR injury have been reported in previous studies [26-28]. In the present study, we determined that inducing an ischemic injury for 30 to 45 minutes caused lower limb paralysis immediately after the experiment, but an almost complete recovery was observed after 48 hours. However, when an ischemic injury was introduced for 60 minutes, lower limb paralysis was maintained in the model rats, and therefore a method involving 60 minutes of ischemic injury was chosen for the spinal cord IR model.

During the early stage of formulating hypotheses and experimental design, systemic adverse effects of HTK solution were a major concern; therefore, a model was developed in which the drug was only delivered to the aortic segment. Using large animals presents problems involving experimental space and financial concerns. With regard to rabbits, the lack of experimental reagents and methods for conducting various molecular biological studies is an issue. In light of factors such as size, safety, reproducibility, and suitability for diverse molecular biological stud-
ies, SD rats were chosen as the experimental animal. The model of creating an occlusion distally to the left renal artery was modified for our purposes. The common femoral artery was cannulated, the aorta proximal to aortic bifurcation was occluded simultaneously, and the drug was selectively delivered to the aortic segment only. Although drug delivery by puncturing the aorta is possible in rabbits, using this method in SD rats leads to problems with thrombosis and hemostasis before and after the operation because the internal membrane of the aorta is thinner than 1 mm. The degree of lower limb movement deterioration due to ligation of the common femoral artery in SD rats has been found to be almost negligible, indicating that a novel animal model could therefore be developed in SD rats [29,30]. In the present study, we measured the locomotor ability of the contralateral limb because our goal was to conduct histological and molecular biological research solely on the spinal cord and to evaluate the contralateral limb.

As preclinical research on the protective effect of HTK solution on the spinal cord in which the drug was delivered to the occluded aorta segment in an SD rat spinal cord IR model, the present research is subject to multiple limitations. First, functional or histopathologically based research to identify the component of HTK that is responsible for its neuroprotective effect has not been conducted. Additionally, the proper dose of HTK solution has not yet been determined. Second, the possibility of reciprocal interactions of HTK with the neuroprotective effects of anesthetic agents such as pentobarbital and ketamine has not yet been studied. Third, the results of this study cannot be generalized due to the small number of experimental animals used in our research. Given these limitations, in order to confirm the neuroprotective effect of HTK solution, it is necessary to conduct large-scale animal experiments incorporating molecular biological research methodologies.

In conclusion, the results of the present study, in which we examined the protective effect of HTK solution on the spinal cord by delivering the drug specifically into the occluded aorta segment in a spinal cord IR SD rat model, allow the following conclusions to be made. First, this new method of producing a spinal cord IR model in SD rats was verified to be reliable. Second, the protective effect of HTK solution on the spinal cord was confirmed, and it is therefore necessary to ascertain the proper injection method and dosage, as well as to conduct further molecular biological research on this topic.

**Conflict of interest**

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

**Acknowledgments**

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**References**

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