Original Article

Hyperbaric oxygen treatment promotes antioxidative effects on rats with acute carbon monoxide poisoning via NF-E2-related factor 2 activation

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Abstract: Oxidative stress plays an important role in brain injuries after carbon monoxide (CO) poisoning. The aim of this study was to investigate the antioxidative effects of hyperbaric oxygen (HBO) treatment on rats with acute CO poisoning and elucidate underlying mechanisms associated with this event. Sprague-Dawley male rats were subjected to CO poisoning and HBO treatment. Their cognitive function was tested to show the effects of CO poisoning and HBO treatment. Antioxidative effects were investigated by determination of antioxidant enzymes. Expression of NF-E2-related factor 2 (Nrf2) and its cytoplasm binding protein kelch-like ECH-associated protein 1 (Keap1) was evaluated by immunohistochemical staining and Western blot. Additionally, Nrf2 transfer into nuclear and combination with ARE were assessed by Western blot and Electrophoretic Mobility Shift Assay (EMSA). Results showed that HBO treatment increased levels of T-AOC, T-SOD, CuZnSOD, GPx, GR, and CAT, decreased by CO poisoning, and decreased MDA that had been increased by CO poisoning. Nrf2 disassociated from Keap1 in cytoplasm and transferred into nuclear to combine with ARE in the brains of HBO-treated CO-poisoned rats. These results suggest that HBO treatment increased antioxidant enzymes via the Nrf2-Keap1-ARE pathway. Therefore, Nrf2-mediated antioxidative effects may be one of the cellular mechanisms responsible for HBO treatment of brain injuries after acute CO poisoning.

Keywords: Carbon monoxide poisoning, hyperbaric oxygen, Nrf2-Keap1-ARE pathway

Introduction

Acute carbon monoxide (CO) poisoning is a common disease with high incidence and mortality. It is estimated that 3%-30% of patients will develop delayed neuropathological sequelae (DNS), despite adequate treatment [1]. It has been reported that the major causes of long-term mortality, after CO poisoning, are mental and psychiatric disorders [2]. The mechanism of brain injuries and psychiatric disorders, after acute CO poisoning, has been associated with free radicals [3], oxidative stress [4], apoptosis [5], and immune reaction [6]. Hyperbaric oxygen (HBO) treatment is an effective traditional therapy for acute CO poisoning and its delayed neuropathology [7]. However, it has been reported that HBO has the potential for significant free radical generation, inducing oxidative stress [7-9].

Nucleic factor erythroid 2-related factor 2 (Nrf2), belonging to the “cap’-n’-collar” (CNC) family, has recently been demonstrated to be an important transcription factor regulating oxidative stress [10-12]. The Nrf2 system regulates expression of various cytoprotective enzymes including NADPH quinone oxidoreductase 1 (NQO1), heme oxygenase 1 (HO-1), GPX1, GPX2, GR, and CAT, thus, exerting antioxidant effects [13-15]. Kelch-like ECH-associated protein 1 (Keap1) is a substrate adaptor protein of a Cullin 3 (Cul3)-Rbx1-E3 ubiquitin ligase complex. Under physiological conditions, the complex binds Nrf2 and regulates ubiquitination and degradation of Nrf2, resulting in low
levels of Nrf2. In contrast, Nrf2 and Keap1 separate and combine with ARE in the nucleus, thus, regulating cytoprotective genes [16].

The aim of this study was to further explain the mechanisms of HBO treatment association with antioxidative effects by establishing a rat model of acute CO poisoning and observing learning and memory abilities, antioxidant capacity, and expression of associated protein in Keap1-Nrf2-ARE pathways.

Materials and methods

Experimental groups

After a week of adaptive feeding, 99 SD male rats (2-month-old) were randomly divided into three groups: normal control (NC) group (9 rats), acute carbon monoxide poisoning (CO) group (45 rats), and hyperbaric oxygen treatment (HBO) group (45 rats). The CO group and HBO group had 5 time points (1 day, 3 days, 7 days, 14 days, and 21 days after CO poisoning). NC group had no poison and no HBO treatment. CO group had carbon monoxide poisoning. HBO group had CO poisoning followed by daily bouts of HBO treatment. Each group had 9 rats. Six rats in each group were used for cognitive function testing, antioxidant enzyme determination, and Western blot, while the other 3 rats were used for histopathological experiments after perfusion-fixation. All rats were provided by the Experimental Animal Center of the Fourth Military Medical University. This study was approved by the Animal Ethics Committee of the Fourth Military Medical University. All experimental procedures were conducted in conformance with guidelines for the care and use of laboratory animals formulated by the Ministry of Science and Technology of the People’s Republic of China.

Animal manipulations

CO exposure was performed, according to published protocol, in a small temperature-controlled (22–24°C) animal hyperbaric chamber [15]. CO detectors in the chamber were used to determine the concentration of CO gas. Rats breathed 1,000 ppm CO gas for 40 minutes, then 3,000 ppm for 20 minutes. Rats were then removed to room air. Rats of the HBO group were given HBO treatment immediately after CO exposure. For HBO treatment, rats were put into the same hyperbaric chamber used for CO exposure. HBO sessions began with a progressive increase in pressure for 30 minutes, followed by 60 minutes of continuous exposure to 100% oxygen at 2.5 ATA. After 60 minutes of exposure, pressure in the chamber was slowly reduced over a 30 minute period. After the procedure, animals were placed in single cages in the animal care room. Control rats and CO-poisoned rats were placed outside the hyperbaric chamber in the same room. After cognitive function tests, rats were anesthetized with 2% pentobarbital sodium and sacrificed by left ventricle perfusion of 0.9% physiological saline. Brains were removed and washed with physiological saline and deionized water, repeatedly. All surgery was performed under pentobarbital sodium anesthesia and all efforts were made to minimize suffering.

Tissue preparation

Tissue preparation for Western blotting and antioxidant capacities: The rats were anesthetized and sacrificed by left ventricle perfusion of 0.9% physiological saline. To determine antioxidant capacities in brain tissues, brains were removed from the skulls and washed with physiological saline and deionized water, repeatedly. Isolated brains were homogenized in 1 mL ice-cold saline. After centrifugation (2000 rev/minute for 5 minutes), the pellet was discarded and clear supernatant liquid was obtained. Total antioxidant capacities (T-AOC) activities, glutathione peroxidase (GSH-PX) activities, glutathione reductase (GR) activities, catalase (CAT) activities, superoxide dismutase (SOD) activities, and malondialdehyde (MDA) concentration in the brains were evaluated by spectrophotometric method. For Western blot, total protein of the cerebral cortex and hippocampus were obtained using Tissue Protein Extraction Reagent (Thermo Scientific), following manufacturer instructions. Nuclear protein of the cerebral cortex and hippocampus were extracted using Minute Cytoplasmic and Nuclear Extraction Kit (Invent Biotechnologies), following manufacturer instructions. Protein concentrations were determined using a BCA kit (Pierce) with albumin standard ampules as a standard. Total protein and nuclear protein were denatured with NuPAGE LDS sample buffer (Invitrogen), containing NuPAGE sample.
reducing agent (Invitrogen), and boiled at 70°C for 10 minutes, prior to use. These proteins were separated by gel electrophoresis using NuPAGE Novex 4-12% Bis-Tris precast gels and transferred to polyvinylidene fluoride membranes blotted with primary antibodies (dilution: anti-Nrf2 polyclonal antibody 1:500, anti-Keap1 polyclonal antibody 1:500 and anti-beta-actin monoclonal antibody 1:1000). Horseradish peroxidase-conjugated goat anti-mouse/rabbit IgG (1:5000) was used as secondary antibody, followed by detection with enhanced chemiluminescence substrate for 5 minutes. After exposure for 1 minute, the film was developed and total density of sample proteins was quantified with Image-Pro Plus software. Results were calculated as: density of sample band/density of control band.

Tissue preparation for histopathological experiment: Rats were anesthetized and sacrificed by left ventricle perfusion of 0.9% saline followed by 4% paraformaldehyde in 100 mM phosphate buffer. Brains were then removed from the skulls and postfixed in 4% paraformaldehyde at 4°C for 48 hours. Afterward, the brains were cut into 2 mm blocks. The 2 mm sections were then paraffin-embedded and cut into 5 mm sections, using a microtome, and mounted on poly-L-lysine-coated slides. Prior to staining, sections were de-waxed in xylene and dehydrated through graded alcohol. For immunohistochemical staining, sections were incubated in 0.5% hydrogen peroxide (H₂O₂) in methanol to block endogenous peroxidase activity and then incubated in 0.2% Triton X-100 for 30 minutes. They were rinsed and incubated in phosphate-buffered saline, containing 1% bovine serum albumin and 10% normal calf serum, for 1 hour at room temperature. Incubation with primary antibodies (anti-Nrf2 polyclonal antibody 1:500 and anti-Keap1 polyclonal antibody 1:500) was performed for 24 hours at 4°C. The sections were then incubated with biotinylated goat anti-mouse/rabbit IgG (1:100) followed by ABC complex with 3, 3’-diaminobenzidine. Negative controls were included in each experiment where the primary antibody was omitted. No immunoreactivity was detected in brain sections from the CO group in the absence of primary antibodies. The sections were then dehydrated, cleared, and mounted for microscopic analysis.

Cognitive function tests

Morris water maze: The Morris water maze experiment is based on the fact that rodents placed in water have a strong motivation to escape the water environment using the fastest and most direct route. Rats escaping from the water environment reflect an ability to learn. Rats swimming to a safe place (platform) in the surrounding environment reflect an ability to use spatial memory. Morris water maze experiment was performed in a tank, 122 cm in diameter, with water temperature maintained at 21°C. The water was tinted with black ink to obscure the platform. A 10 cm x 10 cm platform was hidden 1 cm below the surface of the water in quadrant 1. Entry points were randomly assigned to quadrant 2, 3, or 4. Briefly, the familiarization procedure was performed 1 day and 2 days before CO poisoning. Acute carbon monoxide poisoning occurred after the familiarization session and testing procedure was performed 1 day, 3 days, 7 days, 14 days, and 21 days after CO poisoning. Each trial lasted until the rats found the fixed platform or for a maximum of 60 seconds. Rats were allowed to rest on the platform for 10 seconds and each rat was given four trials per day. Place navigation ability was determined by recording the latency period before searching for the platform. After place navigation testing (21 days after CO poisoning), the platform was removed to create a spatial probe test. Swimming times and swimming distances in searching for the platform in quadrant 1 were recorded, representing spatial probe ability.

Open field test

Open Field test is based on the exploratory behavior and spontaneous activity of rodents entering an open environment. Movement distances and speed of the rats reflect their fear. Rats activities in the central zone reflect their anxiety. Open Field testing was performed on the 10th day after CO poisoning. Briefly, the apparatus consisted of a gray square of 70 cm x 70 cm x 40 cm. It was divided into 35 x 35 equal squares, which had been drawn in the floor of the arena. The testing room was dimly illuminated. A single rat was placed in the center of the floor and, after 30 seconds of adaptation, total distance, average speed, distance in...
the central zone, and stay period in the central zone were counted, automatically, for 15 minutes by an image collecting and processing system. After each test, the arena was cleaned with 90% alcohol solution.

High plus maze

Elevated plus maze (EPM) consists of a device with two opposite open arms (50 × 10 cm) and two closed arms (50 × 10 cm), elevated 45 cm from floor level and a central platform (10 × 10 cm). The high plus maze was performed on the 6th day after CO poisoning. Rats were placed, individually, in the center of maze with the head turned to one of the closed arms. Their behavior was evaluated for 5 minutes. Each trial was video recorded by a digital camera linked to a computer in an adjacent room. The recordings were analyzed, offline, by a highly-trained researcher using X-PLO-RAT (version 3.3). Behavioral parameters evaluated in this test were the number of entries and length of stay of the animals in the open arms, closed arms, and center area of the equipment. Results are expressed as mean ± SD for each area and analyses were performed by one-way ANOVA. Between groups, variance was determined using LSD-T test. \( P \) value < 0.05 was considered statistically significant.

Electrophoretic mobility shift assays (EMSA)

ARE binding activity of Nrf2 was measured with electrophoretic mobility shift assays. Nuclear fraction was isolated, as above. A double-stranded 27-mer oligonucleotide with the Nrf2 consensus sequence 5'-TGGGGAACCTGTGCTGAGTCACTGGAG-3' (Santa Cruz) was used for Nrf2-ARE binding activity. These probes were end-labeled using \([\gamma\text{-}32P]\) ATP (3000 Ci/mmol, Amersham) and T4 polynucleotide kinase, and purified with a G-25 Sephadex column.

Statistical analysis

The number of staining cells in each field was counted at high magnification \((\times 400)\). Five high-power fields of the cerebral cortex and hippocampus were photographed in each section. These high-power fields were separated by at least 60 mm to avoid double counting of cells. Positive cells, in each sample, were taken as the average count from five brain sections and number of values are presented as mean ± SD. Statistical analysis was performed with SPSS Statistics 17.0, using one-way analysis of variance to establish whether differences were statistically significant. Between groups, variance was determined by least significant difference t-test. \( P \) value < 0.05 was considered statistically significant.

Results

HBO treatment increases cognitive function of rats with acute CO poisoning

HBO treatment increases learning and memory ability of rats with acute CO poisoning: In the place navigation experiment with the Morris water maze, escape latency in the CO group was longer than the NC group on each recording day. Compared with the CO group, HBO treatment shortened escape latency. Times of the HBO group were not different, however, from that of the NC group (Figure 1A). Escape latency and swimming locus of the three groups (Figure 1B) showed that HBO treatment improved place navigation ability that had been lessened by acute CO poisoning.

In spatial probe testing with the Morris water maze, swimming times and distances of the CO group in quadrant IV were shorter than that of the NC group. In the HBO group, swimming times and distances were longer than that in the CO group but differences between the HBO group and NC group were not significant (Figure 1C). Swimming times and swimming locus of the three groups (Figure 1D) showed that HBO treatment enhanced their spatial probe ability which had been lowered by acute CO poisoning.

HBO treatment increases adaptive capacity of rats with acute CO poisoning in a novel environment: In open field testing, the total distance and distance in the central zone in the CO group were both longer than those in the NC group. Average speeds were faster in the CO group than the NC group. Stay periods in the central zone in the CO group were longer than the NC group. In the HBO group, total distance was shorter, average speed was slower, and stay period in the central zone was shorter than those in the CO group. Differences in distance in the central zone between the HBO group and CO group were not significant (Table 1 and
HBO promotes antioxidative effects via Nrf2
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Figure 1. Results of cognitive function in each group. In the place navigation test, escape latency in CO group was longer than that in NC group and HBO treatment made it shorter than CO group (A, B). In the spatial probe test, the swimming time in quadrant 1 in the CO group was shorter than that in the NC group and the HBO group was longer than that in the CO group (C, D). In open field test, total distance and distance in the central zone in the CO group were both longer than that in the NC group. In the HBO group, total distance was shorter, average speed was slower, and stay period in the central zone was shorter than in the CO group (E). In the elevated plus maze test, CO poisoned rats significantly approached less in open arms compared with control rats. After HBO treatment, open arm retention time and distance were significantly greater than that of CO group, the number of entry open arm was significantly more than that of CO group (F, G). Compared with NC group, *p < 0.05; compared with the CO group, #p < 0.05.

Table 1. Results in the open field test (n=6)

<table>
<thead>
<tr>
<th>Group</th>
<th>Total distance (cm)</th>
<th>Average speed (cm/s)</th>
<th>Distance in central zone (cm)</th>
<th>Stay period in central zone (s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>NC</td>
<td>2271.35±551.25</td>
<td>2.52±0.61</td>
<td>133±67</td>
<td>26.02±12.80</td>
</tr>
<tr>
<td>CO</td>
<td>3540.42±592.44</td>
<td>4.26±0.52</td>
<td>133.75±86.71</td>
<td>26.02±12.80</td>
</tr>
<tr>
<td>H₂</td>
<td>2865.98±794.86</td>
<td>3.19±0.88</td>
<td>21.95±20.82</td>
<td>3.69±2.71</td>
</tr>
</tbody>
</table>

* P < 0.05, compared with NC group; # P < 0.05, compared with CO group.

Table 2. Activities of antioxidant enzymes and content of MDA in brain (n=6, x±s)

<table>
<thead>
<tr>
<th>Group</th>
<th>T-AOC (U/mg protein)</th>
<th>T-SOD (U/mg protein)</th>
<th>Cu-Zn SOD (U/mg protein)</th>
<th>GSH-PX (U/mg protein)</th>
<th>CAT (U/mg protein)</th>
<th>GR (U/mg protein)</th>
<th>MDA (nmol/mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>NC</td>
<td>1.25±0.40</td>
<td>344.57±24.10</td>
<td>280.53±25.60</td>
<td>182±53</td>
<td>5.2±1.3</td>
<td>14.5±3.0</td>
<td>5.43±0.81</td>
</tr>
<tr>
<td>CO1d</td>
<td>0.68±0.09</td>
<td>218.46±12.65</td>
<td>131.85±10.15</td>
<td>42±13</td>
<td>1.6±0.8</td>
<td>4.3±0.7</td>
<td>7.64±0.94</td>
</tr>
<tr>
<td>HBOT1d</td>
<td>0.97±0.31</td>
<td>251.11±22.20</td>
<td>173.37±42.97</td>
<td>203±63</td>
<td>5.2±2.2</td>
<td>13.9±3.3</td>
<td>6.35±1.08</td>
</tr>
<tr>
<td>C03d</td>
<td>0.45±0.17</td>
<td>170.93±34.22</td>
<td>114.52±32.46</td>
<td>106±46</td>
<td>4.3±1.6</td>
<td>2.6±0.5</td>
<td>11.48±1.67</td>
</tr>
<tr>
<td>HBOT3d</td>
<td>0.97±0.39</td>
<td>225.01±47.77</td>
<td>144.66±56.04</td>
<td>325±86</td>
<td>8.8±2.8</td>
<td>4.3±1.0</td>
<td>6.57±1.57</td>
</tr>
<tr>
<td>C07d</td>
<td>0.71±0.18</td>
<td>66.97±15.84</td>
<td>47.62±15.68</td>
<td>197±49</td>
<td>3.9±1.0</td>
<td>3.0±1.2</td>
<td>19.85±2.32</td>
</tr>
<tr>
<td>HBOT7d</td>
<td>1.45±0.15</td>
<td>287.96±24.56</td>
<td>164.22±72.59</td>
<td>389±29</td>
<td>5.3±1.0</td>
<td>3.9±0.7</td>
<td>8.67±1.94</td>
</tr>
<tr>
<td>C014d</td>
<td>0.69±0.29</td>
<td>102.08±22.81</td>
<td>79.13±16.39</td>
<td>173±42</td>
<td>8.5±2.6</td>
<td>1.8±0.8</td>
<td>12.45±3.78</td>
</tr>
<tr>
<td>HBOT14d</td>
<td>1.40±0.25</td>
<td>256.11±41.05</td>
<td>193.09±41.8</td>
<td>385±10</td>
<td>9.2±2.1</td>
<td>4.8±0.9</td>
<td>6.84±0.68</td>
</tr>
<tr>
<td>CO21d</td>
<td>0.48±0.29</td>
<td>115.94±14.81</td>
<td>84.40±20.55</td>
<td>429±58</td>
<td>5.4±1.7</td>
<td>3.2±1.9</td>
<td>6.64±1.15</td>
</tr>
<tr>
<td>HBOT21d</td>
<td>1.20±0.20</td>
<td>300.37±27.22</td>
<td>173.34±46.01</td>
<td>453±32</td>
<td>14.1±3.8</td>
<td>4.6±0.9</td>
<td>6.92±0.48</td>
</tr>
</tbody>
</table>

* P < 0.05, compared with NC group; # P < 0.05, compared with CO group.

Figure 1e). HBO treatment enhanced their adaptive capacity in a novel environment.

HBO treatment decreases anxiety status of rats with acute CO poisoning: In the elevated plus maze test, CO-poisoned rats spent less time and contributed less distance in the open arms, compared with controls. As for the number of entries, CO-poisoned rats significantly approached less in open arms, compared with control rats. After HBO treatment, open arm retention times and distances were significantly greater than that of CO poisoning group (P < 0.05) and number of entry of open arms was significantly more than the CO poisoning group (P < 0.05) (Figure 1F and 1G). HBO treatment decreased rat anxiety status. Antioxidant enzyme activities and MDA concentration in brain tissues might determine antioxidant capacities after acute CO poisoning. This idea raised the question of how HBO treatment affects antioxidant capacities. Compared with the control group, the CO group had lower T-AOC activities and higher MDA content of brain tissue (P < 0.05). GSH-PX, GR, CAT, T-SOD, and CuZnSOD activities in the CO group decreased to different degrees (P < 0.05). Compared with the CO group, the HBO group increased T-AOC activity and decreased MDA content of brain tissue. GSH-PX, GR, CAT, T-SOD, and CuZnSOD activities in the HBO group increased to differ-
HBO promotes antioxidative effects via Nrf2

**Figure 2.** Immunohistochemical staining and Western blot of Keap1. Keap1-positive cells in the cerebral cortex tissue and hippocampus from rats of the CO group were not significant different with that of the NC group at 1st day (B) and 7th day (A, D) and less at 3rd day (C) compared with the NC group. After HBO treatment, Keap1-positive cells were significantly more than that in CO group at 1st day (B) and 3rd day (C). Samples from the HBO group showed less than that from the CO group in 7th day (D). In Western blot samples in the CO group, Keap1 protein was not significantly different relative to samples from either the NC group or the HBO group at any time point (E, F). Compared with N group, *p < 0.05; compared with the CO group, #p < 0.05.
HBO promotes antioxidative effects via Nrf2

A

Nrf2

NC  CO3d  HBOT3d

cortex

E  G

hippocampus

F

Nrf2  Actin

NC1d  CO1d  HBOT1d

cerebral cortex tissue

G

Nrf2  Actin

NC3d  CO3d  HBOT3d

brain of hippocampus

D

NC  CO7d  HBOT7d

cortex

H  N

hippocampus

C

20  10  5  0

#  #

NC  CO  HBOT

cell number

cortex  hippocampus

E

20  10  5  0

#  #

NC  CO  HBOT

cell number

cortex  hippocampus

Result of Nrf2 in cortex and hippocampus at 1 day

Result of Nrf2 in cortex and hippocampus at 3 day

Result of Nrf2 in cortex and hippocampus at 7 day

HBO promotes antioxidative effects via Nrf2

Figure 3. Immunohistochemical staining and Western blot of Nrf2. Nrf2-positive cells in the CO group were significantly less than that of the NC group, at all time points, and HBO treatment increased expression of Nrf2 protein (A-E). However, the results of Western blot for Nrf2 showed that Nrf2 protein in the CO group was not significantly different in cerebral cortex and hippocampus compared to the NC group, at all time points, expect for the 7th day. At 7th day, the Nrf2 from brain of hippocampus of CO poisoned rats was significantly decreased compared to that of the NC group. HBO treatment increased Nrf2 in cortex in 7th day and hippocampus in 1st and 3rd days (F, G). Compared with the NC group, *p < 0.05; compared with the CO group, #p < 0.05.

HBO treatment changed Keap1 and Nrf2 protein expression in brains of CO poisoned rats

Protein expression levels of Keap1 and Nrf2 were examined by immunohistochemical and Western blot analyses. In immunohistochemical analysis, cells that stained positive for Keap1 and Nrf2 were found in the brain isolated from NC, CO-poisoned, and HBO rats.

Keap1-positive cells, in the cerebral cortex tissue and hippocampus from rats of CO group, were not significantly different than that of NC group on the 1st day (Figure 2B) and 7th day (Figure 2D). However, samples from the CO group showed less staining of Keap-1 on the 3rd day compared with NC group (P < 0.05) (Figure 2C). After HBO treatment, Keap1-positive cells in the cerebral cortex and hippocampus from brain sections of rats were significantly more than that in the CO group on the 1st day (Figure 2B) and 3rd day (Figure 2C) (P < 0.05). In contrast, samples from HBO group showed less than that from the CO group (P < 0.05) on the 7th day (Figure 2A and 2D). Total protein of the cerebral cortex and hippocampus was used for Western blot analysis. In samples from the cerebral cortex and hippocampus of rats after CO exposure, Keap1 protein was not significantly different relative to samples from either NC group or HBO, at any time point (Figure 2E and 2F). The results indicate that there was no significant tendency of Keap-1 among each group.

Nrf2-positive cells in the cerebral cortex and hippocampus of CO group were significantly less than the NC group, at all time points, and HBO treatment increased expression of Nrf2 (P < 0.05, Figure 3A-E). However, results of Western blotting for Nrf2 showed that Nrf2 proteins in CO group were not significantly differ-
HBO promotes antioxidative effects via Nrf2

Figure 4. Nuclear translocation of Nrf2 and ARE binding. Nrf2 protein in nucleus of brain tissues in the CO group was not significantly different in cerebral cortex and hippocampus compared to the NC group, at all time points, except for the 1st day (C-F). At 1st day, the Nrf2 in nuclear protein in hippocampus of rats after CO exposure was significantly increased than that of the NC group (A, B). HBO treatment increased Nrf2 in nucleus in cortex and hippocampus, at all time points, except for hippocampus at the 1st day. In EMSA, the CO rats displayed decrease in activity of Nrf2 binding to ARE over the level observed in control rats. HBO treatment significantly increased Nrf2 binding activity in CO poisoned rats (G, H). Compared with the NC group, *p < 0.05; compared with the CO group, #p < 0.05.
HBO promotes antioxidative effects via Nrf2

HBO-treated rats. As shown in Figure 4G and 4H, CO rats displayed a decrease in activity of Nrf2 binding to ARE over levels observed in control rats. However, HBO treatment significantly increased Nrf2 binding activity in CO-poisoned rats. Specific binding of Nrf2 in CO rats was confirmed by the addition of a 100-fold molar excess of unlabeled Nrf2 competitor DNA into the EMSA reaction. Unlabeled Nrf2 DNA competed for binding in brain tissue extracts prepared from CO rats. Furthermore, an antibody super shift assay was used to demonstrate the presence of Nrf2 in protein complexes, as Nrf2 antibody binding significantly shifted major Nrf2 binding complexes.

Discussion

The purpose of this study was to assess the effects of HBO treatment on cognitive function, antioxidant ability, and Nrf2 pathways in rat models with acute carbon monoxide poisoning.

Data from human studies has revealed that up to 44% of patients with carbon monoxide poisoning have cognitive problems with memory, attention, and concentration [6]. These problems could be improved by HBO treatment [16]. This present study used the Morris Water Maze, Open Field Test, and High Plus Maze to reflect neurophysiological and cognitive deficits. Results of this study revealed that the CO group demonstrated significant impairment in learning and memory ability, adaptive capacity, and anxiety status. The results also suggest that HBO treatment can improve these cognitive deficits.

Under normal conditions, CO is an endogenous generated gas, in humans, and a cytoprotective molecule [17]. It has been reported that low concentrations of CO play an important role in treatment of ischemic strokes and its beneficial effects could be partially mediated by activation of Nrf2 pathways [18]. However, since CO can promote the formation of ROS, overdose of CO may result in overproduction of ROS, possibly playing a critical role in the reduction of antioxidant capacity of brain tissues from rats with CO poisoning. In this study, comparing antioxidant abilities between the NC group and CO group, it was revealed that rats with CO poisoning had lower T-AOC, T-SOD, GSH-PX, GR, and CAT activities and higher MDA concentration in brain tissues. This present study also demonstrated that overdose of CO decreased Nrf2 protein accumulation in the nucleus after CO poisoning and that low Nrf2 protein levels in the nucleus is associated with lower antioxidant capacity.

Additionally, high concentrations of oxygen may exacerbate ROS damage after HBO treatment [8]. In this study, CO promoted the generation of ROS. Rats were then given HBO treatment. It was found that T-AOC and activities of antioxidant enzymes were significantly improved and MDA concentrations reduced after HBO treatment. These results indicate that HBO treatment increases the antioxidant capacity and reduces oxidative stress of brain tissues in rats. Furthermore, nuclear accumulation of Nrf2 was observed in the HBO group. In this study, it was also found that changes of Nrf2 proteins only occur at the 85~115 kDa range with no alteration in levels of other nonspecific bands or any migratory species of 55~65 kDa. This phenomenon had been mentioned, previously, in other studies [19, 20].

As mentioned above, Keap1-Nrf2-ARE is an important pathway which can regulate oxidative stress. This present study demonstrated that HBO-induced ROS activated Nrf2 and made Nrf2, released from Keap1 and transferred to nucleus to combine with ARE, to activate expression of antioxidant enzymes. In conclusion, HBO treatment improves the antioxidant ability of rats with CO poisoning by activating Keap1-Nrf2-ARE pathways.

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Disclosure of conflict of interest

None.

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References


