Protective effects of active hexose correlated compound in a rat model of liver injury after hepatectomy

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Running title: AHCC prevents liver injury

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ABSTRACT

Background: Recent evidence has indicated that a functional food, active hexose correlated compound (AHCC), has liver-protective effects via suppression of inflammatory mediators, such as inducible nitric oxide synthase (iNOS) and tumor necrosis factor (TNF)-α.

Objective: This study aimed to investigate whether AHCC has beneficial effects in a rat model of endotoxin-induced liver injury after partial hepatectomy, in addition to clarifying the mechanisms of action of AHCC.

Methods: Rats were treated with 70% of partial hepatectomy and lipopolysaccharide (PH/LPS) to induce acute liver injury. A normal diet with or without 2% AHCC was administered orally 10 days before 70% hepatectomy. Inflammatory mediators were analyzed.
Results: AHCC improved the survival rate by 70% in PH/LPS rats. AHCC prevented an increase in serum transaminase levels, and histopathological changes and apoptosis in the liver. AHCC reduced iNOS mRNA and protein expression in the liver, resulting in inhibition of nitric oxide production. AHCC also reduced TNF-α, cytokine-induced neutrophil chemoattractant-1, and interleukin-6 mRNA expression, but enhanced expression of interleukin-10. An electrophoretic mobility shift assay with hepatic nuclear extracts demonstrated that AHCC reduced the activation of nuclear factor (NF)-κB induced by PH/LPS treatment.

Conclusion: AHCC inhibits induction of inflammatory mediators, including iNOS and TNF-α, in part through inhibition of NF-κB activation in a rat model of liver injury. Our findings suggest that AHCC prevents postoperative liver failure after liver resection.

Keywords: active hexose correlated compound, inducible nitric oxide synthase, liver injury, nuclear factor-κB, tumor necrosis factor-α

INTRODUCTION

Extended hepatectomy is performed for radical resection of hepatobiliary tract malignancies, but the mortality rate is not low. Hepatic failure greatly contributes to death after liver resection, and the pathophysiology involved is similar to that of sepsis [1-5]. Severe septic complications are associated with reduced phagocytic function of the reticuloendothelial system after massive hepatectomy [6]. Although two-thirds resection of the liver is not fatal in rodents, there is increased sensitivity to endotoxin in the early phase after hepatectomy. Furthermore, intravenous injection of a sublethal dose of lipopolysaccharide (LPS) postoperatively induces a high mortality [7].

Active hexose correlated compound (AHCC) is an extract from mycelia of Lentinula edodes. This is a mushroom of the basidiomycete family that is cultured in liquid medium in a large tank. AHCC is one of the functional foods that are most frequently used in cancer patients in clinical practice [8].

Animal liver injury models can be caused by various insults, such as ischemia-reperfusion, partial hepatectomy, and endotoxin shock. Using these models, we have previously reported that drugs showing liver-protective effects inhibit induction of inducible nitric oxide synthase (iNOS) and nitric oxide (NO) production [9-13]. These drugs also decrease production of various inflammatory mediators, such as tumor necrosis factor (TNF)-α, interleukin (IL)-1β, IL-6, and cytokine-induced neutrophil chemoattractant (CINC)-1 (human IL-8 analogue). Furthermore, by using primary cultures of rat hepatocytes, we found that AHCC, its sugar fraction, and one of its components (adenosine) inhibited nuclear factor (NF)-κB activation, resulting in inhibition of iNOS induction and NO production [14-16]. These reports suggest that AHCC has potential effects on prevention of various liver injuries.
However, few studies have reported that AHCC markedly enhances the survival rate in animal models of postoperative liver failure, with greater than 90% mortality. In this study, we investigated whether treatment of AHCC improves the survival rate in a rat model of endotoxin-induced liver injury after partial hepatectomy. Additionally, if the survival rate is improved, we aimed to determine whether AHCC affects expression of various inflammatory mediators, including iNOS and TNF-α.

MATERIALS AND METHODS

Materials: The manufacturing process of AHCC is as follows: When basidiomycetes are cultured in a liquid medium, they proliferate and form globular fungal bodies rather than carpophores [17]. AHCC is produced using these bodies of mycelia of basidiomycetes and contains components of the medium that are modified by the mycelia-produced diverse enzymes. In the actual manufacturing procedure of AHCC, the mycelia of the edible shiitake are subjected to a liquid culture (15,000-L tank). After fermentation, AHCC is produced through manufacturing processes, which include separation, concentration, sterilization, and freeze-drying [18]. AHCC is certified under the “Healthy Do” system (Hokkaido Food Functionality Labelling system) in which the Hokkaido Government in Japan recognizes the conduct of scientific research on the health effects of product components. Lipopolysaccharide (LPS, \textit{Escherichia coli} 0111: B4) was purchased from Sigma-Aldrich (St. Louis, MO, USA).

Experimental design: Male Sprague–Dawley rats (6 weeks old; weighing 200 g) were obtained from Charles River (Tokyo, Japan). The rats were kept at 22°C under a 12-h light-dark cycle and received food and water \textit{ad libitum}. Rats were randomly divided into control (normal diet rat), AHCC, 70% of partial hepatectomy and lipopolysaccharide (PH/LPS), and PH/LPS + AHCC groups. A normal diet with or without 2% (w/w) AHCC (Amino Up Chemical Co. Ltd., Sapporo, Japan) was administered orally with free access 10 days before 70% hepatectomy, which was followed by sham operation (control and AHCC groups). Rats were anesthetized with intraperitoneal injection of pentobarbital, prior to undergoing 70% hepatectomy as reported previously [9, 10]. After 48 h of surgery, LPS (250 µg/kg) was injected into the penile vein (PH/LPS and PH/LPS + AHCC groups). Survival was checked during 72 h after injection of LPS. Blood and liver samples were obtained from individual rats at 1, 3, and 6 h after LPS treatment.

All animal experiments were performed in accordance with the Guidelines for the Care and Use of Laboratory Animals of the National Institutes of Health. Experiments were approved by the Animal Care Committee of Kansai Medical University.

Biochemical analysis in serum: Blood samples were collected at 1, 3, and 6 h after LPS injection. Serum levels of aspartate transaminase (AST) and alanine transaminase (ALT) were determined using commercial kits (Wako Pure Chemical, Osaka, Japan). TNF-α,
CINC-1, IL-6, and IL-10 levels were measured in serum using commercial kits (R&D Systems, Minneapolis, MN, USA). The sum of serum nitrite and nitrate (NO$_2^-$ and NO$_3^-$, respectively; stable metabolites of NO) was measured using the Griess reagent method using a commercial kit (Roche Diagnostics, Mannheim, Germany) [19].

**Histopathological analysis:** Excised liver specimens that were collected 6 h after LPS treatment were fixed in 10% formalin and embedded in paraffin. Sections (3–5 µm) were cut and stained with hematoxylin and eosin. Apoptotic bodies were detected by terminal deoxynucleotidyl transferase-mediated dUTP-digoxigenin nick-end labeling (TUNEL) staining using an apoptosis detection kit (Medical and Biological Laboratories Co., Nagoya, Japan). The number of TUNEL-positive cells per square millimeter was counted. TUNEL-positive cells were quantified in a manner that was blinded to the treatment arm. All histological slides were reviewed by Dr. K. Yoshizawa (Kansai Medical University, Hirakata, Japan), the certificated pathologist by International Academy of Toxicologic Pathologists.

**Western blot analysis:** Frozen liver samples were homogenized in five volumes of cell solubilizing buffer (10 mM Tris-HCl, pH 7.4; containing 1% Triton X-100, 0.5% Nonidet P-40, 1 mM ethylenediaminetetraacetic acid [EDTA], 1 mM ethyleneglycol bis [2-aminoethyl ether] tetraacetic acid, 150 mM NaCl, 1 mM phenylmethylsulfonyl fluoride [PMSF], and protease inhibitor cocktail) (Roche Diagnostics) and centrifuged (16,500 × g for 15 min). The supernatant was mixed with sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) sample buffer (final: 125 mM Tris-HCl, pH 6.8; containing 5% glycerol, 2% sodium dodecyl sulfate, and 1% 2-mercaptoethanol), subjected to a 7.5% gel, and electroblotted onto a polyvinylidene difluoride membrane (Bio-Rad, Hercules, CA, USA). Immunostaining was performed using primary antibodies against mouse iNOS (Thermo Scientific, Rockford, IL, USA) and rat β-tubulin (internal control; Clone TUB2.1; Sigma Chemical Co., St. Louis, MO, USA), followed by visualization with an enhanced chemiluminescence blotting detection reagent (GE Healthcare Biosciences Corp., Piscataway, NJ, USA). The bands corresponding to iNOS protein were quantitated by densitometry (ImageJ software) and normalized to β-tubulin.

**Real-time RT-PCR:** Total RNA was extracted from homogenized liver in TRIzol (Gibco BRL, USA) reagent using the guanidinium-phenol-chloroform method [20]. For strand-specific RT-PCR analysis, complementary DNAs were synthesized from total RNA with strand-specific primers, and step-down PCR was performed as previously described [21]. For iNOS (257 bp), TNF-α (275 bp), IL-6 (286 bp), IL-1β (321 bp), CINC-1 (231 bp), IL-10 (245 bp), and elongation factor-1α (EF; internal control; 332 bp) mRNAs, an oligo (dT) primer was used for the RT reaction. The primer sets that were used for PCR are shown in Table 1. These mRNA levels were measured by real-time PCR using the Rotor-Gene Q 2plex HRM (Qiagen, Tokyo, Japan). The Rotor-Gene SYBR Green PCR Kit (Qiagen) was included
in the reaction mixture, and the following touchdown protocol was applied: 1 cycle at 95°C for 5 min, and 45 cycles at 95°C for 5 s and 60°C for 10 s. Each mRNA was detected by normalizing the copy number by that of EF. The complementary DNAs for the rat iNOS mRNA were deposited in the DNA Data Bank of Japan/European Bioinformatics Institute/GenBank under the accession number AB250951.

**Electrophoretic mobility shift assay:** Nuclear extracts were prepared from frozen liver at −80°C [22] with a minor modification. Liver sections (0.05 g) were homogenized with a Dounce homogenizer in 1 mL of buffer A (10 mM HEPES-KOH, pH 7.9; containing 10 mM KCl, 1.5 mM MgCl₂, 1 mM dithiothreitol, 1 mM PMSF, and 5 µg/mL aprotinin) and allowed to swell for 15 min. The homogenates were then centrifuged (1100 × g for 5 min). The pellet was suspended in 1 mL of lysis buffer (buffer A supplemented with 0.1% Triton X-100), allowed to stand for 10 min, and centrifuged (1100 × g for 10 min). The nuclear pellet was suspended in 200 µL of nuclear extraction buffer (20 mM HEPES-KOH, pH 7.9; containing 0.42 M NaCl, 1.5 mM MgCl₂, 1 mM dithiothreitol, 1 mM PMSF, 5 µg/mL aprotinin, 0.2 mM EDTA, and 25% [vol/vol] glycerol), incubated for 30 min, and centrifuged (16,500 × g for 20 min). Binding reactions were performed by incubating the nuclear extract (5 µg protein) in reaction buffer (20 mM HEPES-KOH, pH 7.9; containing 1 mM EDTA, 60 mM KCl, 10% glycerol, and 1 µg of poly[dI-dC]) with the probe (40,000 dpm) for 20 min at room temperature. Products were electrophoresed on a 4.8% polyacrylamide gel in high ionic strength buffer, and dried gels were analyzed by autoradiography. An NF-κB consensus oligonucleotide (5’-AGTTGAG GGGA-CTTTCCAGGC) from the mouse immunoglobulin light chain was purchased (Promega, Madison, WI, USA) and labeled with [γ-32P]-adenosine triphosphate and T4 polynucleotide kinase. Protein was measured using the method of Bradford [23]. The bands corresponding to NF-κB were quantitated by densitometry (ImageJ software).

**Statistical analysis:** All data are expressed as the mean ± SE. Differences between groups and the rate of survival were identified by a two-way ANOVA and the log-rank test, respectively. P < 0.05 was considered to be statistically significant.

**Table 1. Primers and nucleotide sequences**

<table>
<thead>
<tr>
<th>Primer name</th>
<th>Nucleotide sequence</th>
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<tbody>
<tr>
<td>iNOS F/R</td>
<td>5’CCAACCTGCAGGTCTTCAGTG 3'/5’ GTCGATGCAACAACCTGGGTGAAC 3’</td>
</tr>
<tr>
<td>TNF-α F/R</td>
<td>5’TCCCAACAAGGAGAGAGAAGTTCC 3'/5’ GCCAGCCTTGTCCCTTGAAGAGA 3’</td>
</tr>
<tr>
<td>CINC-1 F/R</td>
<td>5’GCCAAGCCACAGGGGGCCCGGT 3'/5’ ACTTTGGGAGACCCCTTATAGC 3’</td>
</tr>
<tr>
<td>IL-6 F/R</td>
<td>5’GAGAAAAAGAGTTGTCAATGGCA 3'/5’TAGTCTTTTATCTCTTTTGAAGA 3’</td>
</tr>
<tr>
<td>IL-1β F/R</td>
<td>5’TCTTTGAAGAAGAGAGCCCTTCCTC-3'/5’-GGATCCACACTCTCGAGCTGCA 3’</td>
</tr>
<tr>
<td>IL-10</td>
<td>5’GCCAGACTTTAAAGGGTGATCTG 3'/5’CCTTTCTTGGAGACGTTAAA 3’</td>
</tr>
<tr>
<td>EF F/R</td>
<td>5’TCTGGTGGGAATGCGACAACATGC 3'/5’ CCAGGAAGAGCTTCAACTCAAGCTT 3’</td>
</tr>
</tbody>
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(cDNA) to mRNA. iNOS, inducible nitric oxide synthase; TNF-α, tumor necrosis factor-α; CINC-1, cytokine-induced neutrophil chemoattractant-1; IL-6, interleukin-6; IL-1β, interleukin-1β; EF, elongation factor-1α. F/R, forward/reverse.

RESULTS

AHCC enhanced the survival of rats with liver injury

More than 90% of rats died within 72 h after treatment of 70% hepatectomy and LPS (PH/LPS). Pretreatment of normal diet with 2% AHCC in PH/LPS-treated rats enhanced the survival rate by 70% (Fig. 1). Serum levels of transaminases (AST and ALT), which are the markers of hepatocellular damage, were reduced with treatment of AHCC (Fig. 2).

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![Fig. 1](image1.png)

**Fig. 1. Effect of AHCC on survival of rats.** Rats were treated with 70% partial hepatectomy (PH) and lipopolysaccharide (LPS). A normal diet with or without 2% AHCC was administered orally 10 days before PH treatment. Data represent the percentage of survival at the indicated times after LPS. PH/LPS rats: open circles, n = 10; PH/LPS + AHCC rats: filled circles, n = 10. *P < 0.05 versus PH/LPS rats.

![Fig. 2](image2.png)

**Fig. 2. Effect of AHCC on serum AST and ALT levels.** Rats were treated with 70% partial hepatectomy (PH) and lipopolysaccharide (LPS). A normal diet with or without 2% AHCC was administered orally 10 days before PH treatment. Blood samples were collected at the indicated times after LPS. Data represent the mean ± SE (n = 5–6 rats per time point per group). PH/LPS (open), PH/LPS + AHCC (closed). *P < 0.05 and **P < 0.01 versus PH/LPS rats.
AHCC reduced the appearance of pathological changes

Histological alteration of the liver in PH/LPS rats was characterized by inflammatory cell infiltration, hemorrhagic changes, and focal necrosis in the midzone and periportal regions. Focal hepatocyte necrosis with hemorrhagic changes, ballooning degeneration, and inflammatory cell infiltration were observed in the liver 6 h after PH/LPS (Fig. 3A), and AHCC prevented these pathological changes (Fig. 3B). PH/LPS treatment dramatically increased apoptosis of hepatocytes in the liver (Fig. 3C). AHCC markedly reduced apoptosis in hepatocytes and decreased the apoptotic index of the liver (Fig. 3D and 3E). In rats that were treated with AHCC alone (without PH/LPS), the liver showed normal hepatic histology (data not shown).

![Fig. 3. Effect of AHCC on pathological changes in the liver.](image)

Rats were treated with 70% partial hepatectomy (PH) and lipopolysaccharide (LPS). A normal diet with or without 2% AHCC was administered orally before PH treatment. (A and B): histological appearance of the liver after PH/LPS treatment was evaluated. Liver samples of (A) PH/LPS and (B) PH/LPS + AHCC were obtained 6 h after PH/LPS treatment and stained with hematoxylin and eosin (magnification x 200, arrowheads indicate apoptotic cells). Circled areas indicate the salient features (focal necrosis with inflammatory cell infiltration and massive hemorrhagic change). (C, D, and E): the effect of AHCC on apoptosis in the liver was evaluated. Liver sections of (C) PH/LPS and (D) PH/LPS + AHCC that were obtained 6 h after treatment were stained with TUNEL (magnification x 200). Data represent the mean ± SE (n = 5–6 per group). *P < 0.05 versus PH/LPS rats. The number of TUNEL-positive cells per square millimeter was counted (E).
AHCC affected the expression of iNOS and cytokines in the liver

During endotoxemia, excess production of NO caused by iNOS has a crucial role in hepatic dysfunction. PH/LPS treatment increased iNOS mRNA levels and AHCC inhibited its expression in the liver (Fig. 4A). There was a negligible expression of iNOS mRNA in the normal and AHCC alone (without PH/LPS) rat groups and there was no significant difference between the two groups (data not shown). Western blotting showed that PH/LPS treatment increased iNOS protein expression at 3 h and AHCC reduced its expression (Fig. 4B). In support of these observations, serum levels of nitrite and nitrate, which are stable metabolites of NO, increased after PH/LPS treatment at 6 h. AHCC inhibited this increase in nitrite and nitrate levels (Fig. 4C).

**Fig. 4. Effect of AHCC on induction of iNOS and nitric oxide production in the liver.** Rats were treated with 70% partial hepatectomy (PH) and lipopolysaccharide (LPS). A normal diet with or without 2% AHCC was administered orally before PH treatment. (A) Total RNA was obtained at the indicated times after LPS treatment and was analyzed by strand-specific quantitative RT-PCR to detect iNOS. Data were calculated as the fold change versus normal diet rats (control group). Data represent the mean ± SE (n = 5–6 rats per time point per group). (B) For western blotting, liver samples were obtained before or at 3 h after LPS treatment. The cell lysate (100 mg protein) was subjected to SDS-PAGE in a 7.5% gel and immunoblotted with anti-iNOS or anti-β-tubulin antibodies. Representative results of three independent experiments are shown. The bands corresponding to iNOS protein were quantitated by densitometry (n = 3 experiments). (C): blood samples were collected at 6 h after LPS treatment and the levels of NO$_2^-$ and NO$_3^-$ (metabolites of nitric oxide) were measured. Data represent the mean ± SE (n = 5–6 rats per time point per group). PH/LPS (open), PH/LPS + AHCC (closed) at (A) and (C). *P < 0.05 versus PH/LPS rats.
Real-time RT-PCR analysis showed that PH/LPS treatment increased mRNA expression of TNF-α, CINC-1, IL-6, and IL-1β. AHCC decreased mRNA expression of TNF-α, CINC-1, and IL-6 (Fig. 5A, 5B and 5C, respectively), but had no effect on IL-1β mRNA expression (Fig. 5D). In contrast, AHCC increased anti-inflammatory IL-10 mRNA expression (Fig. 5E), whereas PH/LPS treatment had no effect on IL-10 mRNA expression. There was a negligible expression of TNF-α, CINC-1, IL-6, IL-1β and IL-10 mRNA in the normal and AHCC alone (without PH/LPS) rat groups (data not shown). In serum analysis, AHCC increased IL-10 levels (Fig. 6D), but there were no significant differences in other serum cytokines between the two groups (Fig. 6A, 6B, and 6C).

**Fig. 5. Effect of AHCC on mRNA expression of cytokines in the liver.** Rats were treated with 70% partial hepatectomy (PH) and LPS. A normal diet with or without 2% AHCC was administered orally before PH treatment. Total RNA was extracted from the liver at the indicated times after LPS treatment and was analyzed by strand-specific real-time RT-PCR to detect (A) TNF-α, (B) CINC-1, (C) IL-6, (D) IL-1β, and (E) IL-10. Data were calculated as the fold change versus normal diet rats (control group). Data represent the mean ± SE (n = 5–6 rats per time point per group). PH/LPS (open), PH/LPS + AHCC (closed). *P < 0.05 and **P < 0.01 versus PH/LPS rats.
AHCC inhibited the activation of NF-kB in the liver

Recent evidence has indicated that NF-kB is involved in the transcriptional activation of a variety of inflammatory genes, such as iNOS, TNF-α, and CINC-1. We examined the effect of AHCC on NF-kB activation (its nuclear translocation and DNA binding) in the liver. The electrophoretic mobility shift assay with hepatic nuclear extracts showed that PH/LPS treatment stimulated activation of NF-kB at 3 h and AHCC reduced the levels of NF-kB activation (Fig. 7).

Fig. 6. Effect of AHCC on serum cytokine levels. Rats were treated with 70% partial hepatectomy (PH) and LPS. A normal diet with or without 2% AHCC was administered orally before PH treatment. Blood samples were collected at the indicated times after LPS treatment. Serum levels of (A) TNF-α, (B) CINC-1, (C) IL-6, and (D) IL-10 were measured. Data represent the mean ± SE (n = 5–6 rats per time point per group). PH/LPS rats: open circles; PH/LPS + AHCC rats: filled circles. *P < 0.05 versus PH/LPS rats.
Fig. 7. Effect of AHCC on activation of NF-κB in the liver. Rats were treated with 70% hepatectomy (PH) and LPS. A normal diet with or without 2% AHCC was administered orally before PH treatment. Nuclear extracts (4 µg of protein) were prepared from the liver before or at 3 h after LPS treatment, and NF-κB was analyzed by an electrophoretic mobility shift assay (upper panel). Representative results of three independent experiments are shown. The bands corresponding to NF-κB were quantitated by densitometry (lower panel; n = 3 experiments). PH/LPS (open), PH/LPS + AHCC (closed). **P < 0.01 versus PH/LPS rats.

DISCUSSION

In our preliminary study, a normal diet with 1% AHCC demonstrated a tendency to increase the survival rate of rats, but this was not significant (Tanaka and Nakatake, unpublished observation). In the current study, we found that oral intake of a normal diet with 2% AHCC before treatment of PH/LPS attenuated the increase in AST and ALT levels (Fig. 2), and reduced the appearance of pathologic changes, including apoptosis (Fig. 3). This led to marked improvement in the survival rate of rats (Fig. 1).

AHCC is abundant in the α-glucan form of carbohydrates. The presence of α-1,4-glucan, in which the hydroxyl groups of C-2 and/or C-3 position are partially acylated, is particularly reported and is considered to be one of the active ingredients [18]. Based on a survey carried out by the Ministry of Health, Labor and Welfare research group in Japan, AHCC is the second most commonly used health food, followed by Agaricus, which is a fungus, among cancer patients in Japan [8]. AHCC is useful for treating various diseases, including hepatitis and cancer, as a complementary and alternative medicine [24]. We reported that AHCC improved the prognosis of patients with hepatocellular carcinoma postoperatively (a prospective cohort study) [25]. AHCC is effective for treating infectious and inflammatory
diseases, and is a cancer therapy [26-32]. The safety of AHCC has been supported by basic studies, including its interaction with drugs [33-35]. Results of a randomized, controlled trial on healthy volunteers strongly supported the safety of AHCC as a supplement in clinical practice [36].

In this study, AHCC inhibited the expression of iNOS mRNA and its protein, leading to inhibition of NO production (Fig. 4A, 4B, and 4C). AHCC decreased mRNA expression of TNF-α, CINC-1, and IL-6 in the liver (Fig. 5A, 5B, and 5C). However, we did not find a reduction in serum TNF-α, CINC-1, and IL-6 levels. This finding was unexpected and the reasons currently remain unclear. AHCC enhanced IL-10 levels in the liver and serum (Figs. 5E and 6D). Accordingly, part of the protective effects of AHCC in our rat model of endotoxin-induced acute liver injury probably resulted from attenuation of iNOS, TNF-α, CINC-1, and IL-6, and enhancement of IL-10.

One of the earliest events after interaction of LPS with toll-like receptor 4 is activation of transcription factor NF-κB in cells, with its translocation from the cytoplasm to the nucleus and DNA binding [37]. Once activated, NF-κB induces transcriptional upregulation of various inflammatory mediators, such as iNOS, TNF-α, and CINC-1 [38-40]. The electrophoretic mobility shift assay with nuclear extracts from the liver showed that AHCC decreased activation of NF-κB induced by PH/LPS treatment (Fig. 7).

CONCLUSION
AHCC decreased mRNA expression of inflammatory mediators, such as iNOS and TNF-α, in PH/LPS-treated rats. This was achieved in part through inhibition of NF-κB activation, resulting in prevention of liver injury. AHCC may have therapeutic potential for use in prevention of postoperative liver failure after liver resection.

Abbreviations: AHCC, active hexose correlated compound; PH/LPS, 70% of partial hepatectomy and lipopolysaccharide; iNOS, inducible nitric oxide synthase; TNF-α, tumor necrosis factor-alpha; NF-κB, nuclear factor-kappa B. IL, interleukin; CINC-1, cytokine-induced neutrophil chemoattractant-1.

Disclosure of potential conflicts of interest: The authors declare that they have no conflicts of interest.

Author’s contributions: RN participated in the design of the study, data collection, statistical analysis, and drafting of the manuscript. YT and YU participated in supervision and provided advice when the manuscript was being drafted. HM assisted in data collection and analysis. MI, KM, and MK assisted in the design of the study. TO, MN and MKon provided advice regarding protocol development for the study and assisted in the design of the study.

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