Complement depletion with cobra venom factor alleviates acute hepatic injury induced by ischemia-reperfusion injury

BING WANG, HUA XU, JIAN LI, HONG-MEI GAO, YING-HONG XING, ZHU LIN, HONG-JIE LI, YONG-QIANG WANG and SHU-HUA CAO

Department of Intensive Care Unit and Key Laboratory for Critical Care Medicine of The Ministry of Health, Emergency Medicine Research Institute, Tianjin First Center Hospital, Tianjin 300192, P.R. China

Received March 2, 2018; Accepted August 28, 2018

DOI: 10.3892/mmr.2018.9484

Abstract. Increasing evidence has demonstrated that complement activation is required for ischemia-reperfusion injury (IRI)-induced hepatic damage, and cobra venom factor (CVF) can deplete the complement components. The aim of the current study was to investigate the effect and intrinsic mechanism of CVF pretreatment on IRI-induced acute hepatic injury in rats. Acute hepatic injury in rats was induced by bone fracture to simulate trauma, followed by hemorrhage for 90 min, and then the rats were resuscitated for a period of 20 min of reperfusion. The survival times under different CVF treatment doses and schedules for rats with IRI were evaluated. Hepatic tissues and serum samples were analyzed for acute hepatic injury, complement activation, inflammatory mediator release and apoptosis at predetermined times and compared between the IRI group and the CVF pretreatment + IRI groups. Compared to the rats with IRI alone, the survival times were significantly improved among rats with IRI receiving a high-dose or low-dose CVF pretreatment (all P<0.01). Upon histological examination, severe hepatic damage was observed in the rats with IRI, accompanied by liver function deterioration, complement and membrane attack complex activation, inflammatory mediator release and hepatic cell apoptosis. CVF pretreatment significantly attenuated the hepatic injury through depletion of complement activation and subsequent inhibition of inflammatory mediator release and hepatic cell apoptosis (all P<0.05). The results indicated that CVF pretreatment ameliorates IRI-induced acute hepatic injury. However, further studies are required to determine whether this therapy could be a potential agent for the treatment of IRI injuries in clinical settings.

Introduction

Ischemia-reperfusion injury (IRI) can induce acute hepatic injury, which is associated with a high level of mortality (1,2). The liver is subjected to IRI in different medical conditions, including hepatic resection and liver transplantation (3). Notably, IRI is not only a pathophysiological process involving liver injury; it also affects multiple organs (4). However, the exact intrinsic mechanism of hepatic injury caused by IRI is still unknown.

Recent advances have demonstrated that complement activation is responsible for the pathogenesis of IRI-induced acute hepatic injury (5). The membrane attack complex C5b-9 is formed during complement pathway activation in IRI-induced acute hepatic injury, leading to lysis and destruction of target cells. Additionally, anaphylatoxins C3a and C5a are released and interact with their receptors, causing activation of the inflammatory and apoptotic pathways (6,7). Inhibition of abnormal complement activation potentially alleviates IRI-induced acute hepatic injury (8).

Cobra venom factor (CVF) is a stable anticomplement protein from cobra venom. It is a structural and functional analog of the human complement component C3b (the active fragment of C3). Similar to C3b, CVF binds to factor B, which is then cleaved by factor D to form the bimolecular complex CVF/Bb; and CVF/Bb is a C3/C5 convertase that cleaves both complement components C3 and C5 (9-11). Therefore, CVF/Bb can continuously activate C3 and C5 to deplete complement components and inhibit its activation, achieving the desired anticomplement effect.

Several studies have demonstrated that CVF pretreatment can alleviate acute lung injury, myocardial injury and cerebral injury induced by IRI, and prolong the survival time following organ transplantation (12-15). The aim of the current study was to investigate the effect and mechanism of CVF pretreatment on IRI-induced acute hepatic injury in rats. The results indicated that CVF pretreatment significantly attenuated hepatic injury through depletion of complement activation and subsequent inhibition of inflammatory mediator release and hepatic cell apoptosis. However, further studies are required to determine whether this therapy may be a potential agent for the treatment of IRI injuries in clinical settings.
Materials and methods

Establishment of the rat model of IRI-induced hepatic injury. All of the experiments were performed as approved by the Laboratory Animal Care Committee of Tianjin First Center Hospital (approval no. 20150132). One hundred and seventy-six-week old male and female Sprague Dawley rats (200-250 g; 1:1 ratio) were obtained from the Experimental Animal Center, Academy of Military Medical Sciences (Beijing, China; Animal qualified batch number: SCXK-2007-004). These rats were housed in the rooms of the Laboratory Animal Care of Tianjin First Center Hospital with specified pathogen-free conditions. The room was maintained at room temperature under relative humidity 40-60%. Food and water were provided ad libitum.

To simulate a clinical traumatic hemorrhagic shock situation, the rat model of IRI-induced acute hepatic injury was successfully established, as described previously (16). Briefly, rats were anesthetized by intraperitoneal injection of chloral hydrate (200 mg/kg), as described previously (17,18). An open mid-diaphyseal transverse fracture was created in the left femur to induce trauma, after which a mean arterial blood pressure of 30±5 mmHg was maintained for 90 min via hemorrhage at 2.5 ml/100 g. Subsequently, the rats were resuscitated for a period of 20 min with Ringer's lactate solution at a constant rate to manage the shock. The volume of Ringer's solution was twice that of the blood loss. Finally, the catheters were removed, and the vessels were ligated.

Treatment design. The experimental rats were randomly and equally divided into five groups: i) The control group underwent the same anesthetic process without surgical procedures; and the following four groups all have acute hepatic injury induced by bone fracture to simulate trauma, followed by hemorrhage, and then were resuscitated of reperfusion as described above: ii) the IRI group; iii) the CVF1 group received a high-dose (100 µg/kg) CVF (cat. no. CS8406, Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) pretreatment at 24 h before the surgery + IRI; iv) the CVF2 group received low-dose (50 µg/kg) CVF pretreatment at 24 h before the surgery + IRI; and v) the CVF3 group received low-dose (50 µg/kg) CVF at day 2 after IRI. The CVF frozen powder was dissolved in normal saline.

Survival time analyses. A total 170 rats were used for survival time assessment and the intrinsic mechanism determination. Firstly, the effect of CVF administration on survival time was assessed in the five groups described above (n=10 per group), and survival was defined as the time between the first treatment and the time of rat death or 48 h follow-up for censored rats. Subsequently, 120 rats were used for intrinsic mechanism determined among the control, IRI, and CVF2 + IRI groups. In each group, 10 rats were sacrificed at 1, 3, 6, and 24 h after IRI, and the hepatic tissues and at least 4 ml blood samples were collected and analyzed.

Histological analyses. For hematoxylin-eosin (H&E) staining, the hepatic tissues were fixed with at least 15 ml 10% neutral formaldehyde solution for ≥24 h. Subsequently, the samples were dehydrated, embedded in paraffin, sectioned into 4-µm-thick slices, and mounted onto slides. Following heating at 60°C for 2 h, slicing and dewaxing at room temperature for 10 min, H&E staining was conducted at room temperature for 10 min, and the slides were observed under an Olympus BX40F light microscope (Olympus Corporation, Tokyo, Japan).

For the immunohistochemical assay, the 4-µm-thick hepatic tissue sections were deparaffinized, rehydrated, and treated with 3% H2O2 at 20°C for 10 min. Antigen retrieval was performed by boiling the sections in Tris/EDTA Antigen Retrieval Solution (pH 8.0) for 2 min. Nonspecific antibody binding was blocked using 3% bovine serum albumin at 20°C for 30 min. Subsequently, the tissue sections were incubated with anti-C5b-9 antibody (cat. no. ab55811; 1:500), anti-Bcl-2 associated X, apoptosis regulator (Bax) antibody (cat. no. ab32503; 1:500), anti-Bcl-2 apoptosis regulator (Bcl-2) antibody (cat. no. ab59348; 1:500; all Abcam, Cambridge, UK), or control IgG overnight at 4°C. The bound antibodies were detected with horseradish peroxidase (HRP)-conjugated secondary antibodies (cat. no. ab205718; 1:100; Abcam) at room temperature for 1 h, and the products were visualized using a 3, 3′-diaminobenzidine tetrahydrochloride (DAB) staining kit (Tiangen Biotech Co., Ltd., Beijing, China) at room temperature for 2 min and counterstained with hematoxylin at room temperature for 5 min. The specimens were examined with an inverted fluorescence microscope (IX50; Olympus Corporation, Tokyo, Japan).

In addition, the expression levels of C5b-9 deposition in specimens were assessed by two independent researchers who were double-blinded manner. Their conclusions were in complete agreement for 85% of specimens indicating this scoring method was highly reproducible. If two agreed with the scoring results, the value was selected. If there was a disagreement, then the third researcher worked collaboratively to confirm the score. For evaluation of C5b-9 deposition, a semi-quantitative scoring criterion was used, in which both staining intensity and positive areas were recorded from 0 to 100.

ELISA. For the ELISA, the serum samples were centrifuged at 3,000 rpm (3,000 x g) for 5 min at 4°C and were stored at -20°C until use. Alanine transaminase (ALT; cat. no. EIA-326), total hemolytic complement (CH50; cat. no. EIA-121), C5b-9 (cat. no. EIA-765), tumor necrosis factor (TNF)-α (cat. no. SRTA00), interleukin (IL)-β (cat. no. DY501) and C5a (cat. no. DY2150) were measured using the monoclonal anti-rat capture antibody (ZSGB-BIO, Inc., Beijing, China and R&D Systems, Inc., Minneapolis, MN, USA), according to the manufacturer’s instructions. The colorimetric reaction was measured using a Benchmark microplate reader (Benchmark Electronics, Inc., Scottsdale, AZ, USA).

Semi-quantitative reverse transcriptase-polymerase chain reaction (RT-PCR). Guided by the manufacturer’s instructions, total RNA was extracted using the TRizol reagent (Thermo Fisher Scientific, Inc., Waltham, MA, USA). Following measurement of RNA concentration, reverse transcription was performed using a SuperScript RT kit (Takara Biotechnology Co., Ltd., Dalian, China) under the following conditions: 37°C for 15 min, 55°C for 5 sec, and then maintained at 4°C. RT-PCR was performed in a 25 µl reaction volume: 1 µl diluted cDNA, 1 µl sense primer (10 pmol/µl), 1 µl antisense primer (10 pmol/µl), MgCl2 0.3 µl, 12.5 µl Premix EX Taq II (2X), and 9.2 µl RNase-free d H2O. For the RT-PCR, the following PCR program was applied: 95°C for 30 s, 35 cycles of 95°C for 30 s, 55°C for 30 s, and 72°C for 30 s, followed by 72°C for 5 min.
Following mixing, the cDNAs were synthesized under the following conditions: 94°C for 2 min, 94°C for 30 sec, 60°C for 1 min at 40 cycles, 72°C for 30 sec, 72°C for 10 min, and 10°C for 10 min by PCR. The primers for gene amplification were synthesized by Shenggong Biological Engineering Technology and Services (Shanghai, China): Bax sense, 5'-ACAGATCAT GAAGAGGGG-3' and antisense, 5'-CAAGTGAAGA GCCAACC-3; Bcl-2 sense, 5'-TTCTCTCCAGCCCTGA GGC-3' and antisense, 5'-ATGACCCACCGAATCTCAAG-3'; GAPDH sense, 5'-GGTGAGGGTCTGTGAAGC-3' and antisense, 5'-TCCGCCGTCTCGGAGATGCT-3'. GAPDH was used as an internal control. The fidelity of all RT-PCR assays was assessed by 1.5% agarose gel electrophoresis containing 20 g/l TAE buffer, and the relative gene expression levels were calculated relative to GAPDH.

**Results**

**CVF pretreatment improves the prognosis of rats with IRI.** No rats in the control group died. The survival times were significantly prolonged in the high-dose and low-dose CVF groups (CVF1 and CVF2, respectively), compared with the IRI group (all P<0.05). Additionally, when CVF was administered at day 2 (CVF3 group) after IRI, the survival time was not improved compared with the IRI group (P>0.05) (Fig. 1A).

**CVF pretreatment attenuates the hepatic injury of rats with IRI.** The low-dose CVF pretreatment (CVF2) was used for subsequent experiments because the survival times were not significantly difference between the high-dose and low-dose CVF groups following IRI (P>0.05). In the CVF2 + IRI group, the levels of ALT and AST were significantly attenuated, compared with the IRI group at 3, 6, and 24 h (all P<0.05); however, the levels of ALT and AST in the CVF2 + IRI group remained significantly increased compared with the control group (all P<0.05) (Fig. 1B and C). Furthermore, the H&E staining results of the hepatic tissues of the control, IRI, and CVF2 + IRI groups (24 h) are presented in Fig. 1D. No hepatic cell injury was detected in the control group. However, the IRI group exhibited albuminoid and vacuolar degeneration, eosinophilic variants, spotty liver necrosis, inflammatory cell infiltration and Kupffer cell proliferation; while only focal eosinophilic variants of hepatic cells were observed in the CVF2 + IRI group.

**CVF pretreatment results in complement and membrane attack complex depletion.** To verify that CVF pretreatment induced complement and membrane attack complex depletion, plasma complement CH50 and C5b-9 activities were determined in the control, IRI and CVF2 + IRI groups. The ELISA results demonstrated that IRI significantly increased complement CH50 and C5b-9 expression, and the expression levels reached a peak at 1 h after IRI initiation, compared with the control group (all P<0.05) (Fig. 2A and B). When IRI rats received CVF2 pretreatment, the CH50 and C5b-9 activities were significantly decreased compared with those of the rats in the IRI group within 24 h (all P<0.05) (Fig. 2A and B); however, there was no significant difference in CH50 and C5b-9 levels between the IRI group and the CVF2 + IRI group at >24 h.

Additionally, immunohistochemistry assay demonstrated that the level of C5b-9 deposition was significantly reduced in the CVF2 + IRI group at 3, 6, and 24 h, compared with the IRI group. However, the level of C5b-9 deposition was decreased in the CVF2 + IRI group compared with IRI rats; however, the C5b-9 deposition remained significantly higher than the control group (all P<0.05) (Fig. 2C).

**CVF pretreatment reduces the release of inflammatory mediators.** The ELISA results demonstrated that TNF-α and IL-1β release was significantly increased in the IRI group compared
Figure 1. CVF pretreatment improves the prognosis and attenuates hepatic injury of rats with IRI. (A) Survival times under different CVF treatment doses and schedules for rats with IRI were evaluated (10 rats per group). The liver function markers (B) ALT and (C) AST were assessed using an ELISA at different times following IRI initiation. (D) Pathological changes in the liver at 24 h among the control, IRI and CVF2 + IRI groups. Hepatic tissues stained with hematoxylin and eosin, and observed under a light microscope (magnification, x200). IRI, ischemia-reperfusion injury; CVF, cobra venom factor; ALT, alanine transaminase; AST, aspartate transaminase.

Figure 2. CVF pretreatment resulted in complement and membrane attack complex depletion. (A) Complement CH50 and (B) membrane attack complex C5b-9 were assessed using ELISAs within 24 h after IRI initiation. (C) C5b-9 deposition in liver tissues of the control, IRI and CVF2 + IRI groups were assessed by immunohistochemical staining. Data are representative at 3 h (magnification, x200). IRI, ischemia-reperfusion injury; CVF, cobra venom factor; CH50, total hemolytic complement.
with the control group (all P<0.05). Following CVF2 pretreatment, the levels of inflammatory mediator release were significantly reduced at 3, 6, and 24 h (all P<0.05) (Fig. 3A and B).

Activation of the complement system generates anaphylatoxin C5a. In the IRI group, C5a expression was significantly increased compared with the control group at 6 and 24 h (all P<0.05). By contrast, pretreatment with CVF2 attenuated the elevation of C5a expression in the IRI group at 6 and 24 h (P<0.05) (Fig. 3C). Consistent with the ELISA outcomes, the C5a protein level in the liver tissues within 24 h were also significantly reduced by CVF2 pretreatment, compared with the IRI group, according to western blot analysis (Fig. 3D).

**CVF pretreatment alleviates the apoptosis of liver cells.** The levels of apoptotic hepatic cells were observed using a TUNEL assay. No apoptotic cells were detected in the control group. However, the IRI rats exhibited a marked increase in apoptotic liver cells; whereas CVF2 pretreatment significantly reduced the hepatic cell apoptosis at 3, 6 and 24 h (all P<0.05) (Fig. 4A).

Compared with the IRI group, the Bax expression and Bax/Bcl-2 ratio were significantly lower in the CVF2 group within 24 h (all P<0.05; Fig. 4B and C). Consistent with the reduced Bax expression and Bax/Bcl-2 ratio, pretreatment with CVF2 markedly attenuated the elevation of cleaved caspase-3 expression within 24 h (Fig. 4D).

**Discussion**

IRI can induce acute hepatic injury, which is associated with high mortality rates. The liver is subjected to IRI in several medical conditions, including hepatic resection and liver transplantation. Increasing evidence has demonstrated that complement activation is required for IRI-induced hepatic damage (5-8), and CVF can deplete the components of the complement system (9-11). The current study investigated the effect and intrinsic mechanism of CVF pretreatment on acute hepatic injury induced by IRI in rats, and the results indicated that CVF pretreatment ameliorates IRI-induced acute hepatic injury. However, further studies are required to determine whether this therapy could be a potential agent for the treatment of IRI injuries in clinical settings.

In the present study, the survival time of rats was significantly prolonged in the high-dose and low-dose CVF pretreatment groups (CVF1 and CVF2 groups, respectively), compared with the IRI group. CVF administration at day 2 (CVF3 group) after IRI did not improve the survival time, compared with the IRI group. These results suggested the involvement of complement activation at the early stage, not the late stage, following IRI. In addition, either high-dose or low-dose CVF pretreatment improved the survival time of rats with IRI.

In the current study, when IRI was initiated, the liver function significantly deteriorated compared to the control group; while CVF pretreatment significantly improved the liver function. However, the exact intrinsic mechanism of CVF pretreatment in attenuating hepatic injury following IRI remains unknown. In the present study, the CH50 and C5b-9 levels were significantly increased, and their expression levels reached a peak at 1 h after IRI initiation. These results were in agreement with previous studies (14), suggesting that CVF pretreatment results in complement and
membrane attack complex depletion, and that complement activation is involved in the pathogenesis of hepatic injury following IRI.

Notably, when the IRI rats received CVF pretreatment, the CH50 and C5b-9 activities were significantly decreased within 24 h, but not at >24 h, compared with the rats in the IRI group. These results suggested that systemic complement activity was gradually restored. The return of complement activity may have been due to the development of antibodies against CVF or due to complement regeneration in the liver. Therefore, the window of time available for therapeutic intervention to block complement-mediated events in the post-IRI period may be important.

The mechanism by which complement activity contributes to hepatic injury remains unresolved. Figueroa et al (13) has demonstrated that CVF administration may allow accommodation to take place by impairing the ability of anaphylatoxin C5a to form. Given that the release of inflammatory mediators such as TNF-α and IL-1β was significantly decreased in the IRI rats receiving CVF pretreatment, inflammatory mediator release is potentially actively involved in complement activation (12). In addition, considering that the number of apoptotic liver cells was significantly increased in the rats with IRI, it is likely that apoptosis may participate in complement activation by mediating caspase activity, as reported in Gorsuch et al study (5).

In conclusion, CVF pretreatment ameliorated IRI-induced acute hepatic injury in rats. The results of this study were consistent with those of previous research (12-15) may result in novel therapies based on the strategy of complement inhibition to ameliorate multiple organ injury resulting from IRI. Several clinically applicable complement inhibitors are in development or in preclinical trials for various disorders (19). Continued research may yield data justifying clinical trials of one or more complement inhibitors administered during or shortly after events characterized by IRI.

Figure 4. CVF pretreatment alleviates liver cell apoptosis. (A) The liver cell apoptosis levels were determined by a TUNEL assay (x400) at different times after IRI initiation. The proapoptotic protein Bax, inhibition of apoptotic protein Bcl-2, and the ratio of Bax/Bcl-2 were assessed by (B) RT-PCR and (C) immunohistochemical assays at different times after IRI initiation. (D) The protein expression levels of caspase-3 and cleaved caspase-3 were evaluated by western blotting assays of liver tissues at different times after IRI initiation, and the protein levels were calculated relative to GAPDH. **P<0.01 vs. the IRI group; *P<0.05 vs. the control group; ##P<0.01 vs. the control group. IRI, ischemia-reperfusion injury; CVF, cobra venom factor; HPF, high-powered field; Bax, Bcl-2 associated X, apoptosis regulator; Bcl-2, Bcl-2 apoptosis regulator.
Acknowledgements

Not applicable.

Funding

The present study was supported in part by the Foundation of Tianjin Heath and Family Planning Commission (grant nos. 14KG101 and 2014KR07), and the National Clinical Key Specially Project Foundation of the Ministry of Health in China (grant no. 2011-873) to Professor Yong-Qiang Wang.

Availability of data and materials

The analyzed data sets generated during the study are available from the corresponding author on reasonable request.

Authors' contributions

BW carried out conception and design, performed experimental study, collection and assembly of data, data analysis and interpretation, manuscript writing. HX, JL and HMG performed the experimental study, collection and assembly of data, data analysis and interpretation. YHX, ZL and HJL performed collection and assembly of data, data analysis and interpretation, manuscript writing. All authors read and approved the final manuscript.

Ethics approval and consent to participate

All experimental procedures were performed according to the Animal Care Committee guidelines, and the experimental protocol was approved by the Ethics Committee of Tianjin First Center Hospital (approval no. 2015013Z).

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References


This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International (CC BY-NC-ND 4.0) License.