The angiotensin-converting enzyme inhibitor, captopril, suppressed hepatic stellate cell activation via NF-kappaB or wnt3α/β-catenin pathway

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ABSTRACT

Activation of hepatic stellate cells (HSC) is associated with hepatic fibrogenesis, which is one of complications of diabetes mellitus. Captopril possesses potent anti-inflammation, oxidative stress and fibrosis effects. However, the specific molecular mechanism of captopril in high glucose (HG)-induced hepatic stellate cells has not been elucidated. Following the treatment of HG or captopril treatment for rat hepatic stellate cells (HSC-T6), cell activities were detected by Cell Counting Kit-8 (CCK8) assay. Reactive oxygen species (ROS) levels were determined by ROS staining. The expression of inflammation-related proteins (Interleukin (IL)-β, IL-6 and IL-8) and fibrosis-related proteins (fibronectin (FN), collagen I, collagen III, collagen IV, matrix metalloproteinase (MMP-2 and MMP-9) were determined by Western blot. Captopril significantly decreased HSC-T6 cell viability induced by HG in a dose-dependent manner, as well as decreased levels of malondialdehyde (MDA), ROS, pro-inflammatory markers and fibrosis-related proteins, while upregulated superoxide dismutase (SOD) activities. We further found that captopril decreased the ratio of p-IkBα/IκBα and the ratio of p-p65/p65. Intriguing, phorbol myristate acetate (PMA) or LiCl was able to significantly reverse the captopril-induced alteration of oxidative stress-, inflammation- and fibrosis-marker levels. In conclusion, in HG-stimulated HSC-T6 cells, captopril displayed a potent ability to inhibit oxidative stress, inflammation and hepatic fibrogenesis via NF-kappaB or wnt3α/β-catenin. These results demonstrated the mechanism of captoprill as well as the role of the NF-kappaB or wnt3α/β-catenin on HSC-T6 activation induced by HG.

INTRODUCTION

Hepatic fibrosis is one of the many complications of diabetes mellitus. Hepatic stellate cells (HSCs) are the main cell source for the generation of extracellular matrix when liver injury occurs and play a central role in the formation of liver fibrosis [1,2]. The activation process of hepatic stellate cells can be divided into initiation stage and continuous activation stage. If the activation process can be blocked in time, the further development of the disease can be blocked, and the result of hepatic fibrosis can be terminated or even reversed [3–5].

Captopril is an inhibitor of angiotensin converting enzyme (ACE), which is commonly used to low blood pressure in patients with high blood pressure [6]. Captopril combined with gliclazide has been found to reduce vascular and renal complications and control blood glucose in streptozotocin-induced diabetic rats [7]. In addition, captopril has a protective effect on renal injury caused by ischemia reperfusion in diabetic rats [8]. These studies suggest that captopril may work against diabetes. In addition, captopril alleviates renal injury and inflammation in spontaneously hypertensive rats through NF-κB signaling pathway [9], which also is reported to improve fibrosis and oxidative stress of lipopolysaccharide-induced pneumonia [10]. In addition, some studies have reported that captopril plays a protective role in diabetic liver injury [11], and can reduce TAC-induced heart failure by inhibiting Wnt3A/β-catenin and JAK2/STAT3 pathways [12]. It is reported that the activation of Wnt/β-catenin signaling pathway can promote liver fibrosis [13]. However, the specific molecular mechanism of captopril in HG induced hepatic stellate cells has not been studied. Taken together, we assume that NF-κB and Wnt/β-catenin signaling pathways could be involved in the effects of captopril on HG-induced hepatic fibrosis.
stellate cells. Therefore, we further explore the mechanism of captopril in HG-induced HSCs, and speculate that captopril can reduce the abnormal proliferation, inflammatory response and fibrogenesis of HG-induced HSCs by inhibiting NF-κB and Wnt/β-catenin signaling pathways.

**Method**

**Cell culture**

HSC-T6 cells were purchased from ATCC (USA) and cultured in DMEM medium (Hyclone, USA) containing 10% FBS (Gibco, USA) in 37 °C with 5% CO2. HSC-T6 was subjected to glucose treatment at a concentration of (0, 10, 30, 50, 70, 100 mM) to observe cell activity, inflammatory cytokines and ROS production. In the next experiment, HSC-T6 cells were pretreated with different concentrations of captopril (0, 0.1, 0.3, 0.5, 1, 3 mM) for 12 h, and incubated with HG (50 mM) for 48 h.

**CCK8 assay**

HSC-T6 was seeded into 96-well plates with 2 × 10^4 cells per well. After high glucose or captopril treatment, or pretreatment with different concentrations of captopril for 12 h and then incubated with HG (50 mM) for 48 h, the cell viability was detected using CCK8 kit according to manufacturer’s guidance (ThermoFisher, Inc). 10 μL CCK8 solution was added to each well for incubation for 2 h. The absorbance at 450 nm was detected using a microplate reader (ThermoFisher, Inc).

The detection of SOD activity and MDA levels

HSC-T6 cells were collected and washed with PBS of 4°C. The SOD activities were detected using Total SOD activity detection kit (Beyotime, Shanghai, China). For the detection of MDA, cells were collected and lysed using RIPA lysis buffer (ThermoFisher, Inc). Then, the supernatant was collected after centrifugation at 12000 g for 10 min. The MDA activities were detected using MDA detection kit (Beyotime, Shanghai, China).

**ROS staining**

The ROS was detected using Reactive oxygen species assay kit (Beyotime, Shanghai, China) according to manufacturer protocol. The cells were collected and suspended in DCFH-DA. After incubation at 37°C for 20 min, the cells were observed under a fluorescence microscope (OLYMPUS).

**Quantitative reverse transcription PCR (RT-qPCR) assay**

The total RNA of cells was extracted with FAST200 kit. The concentration and purity of RNA were determined by spectrophotometer and cDNA was synthesized according to the Fermentas Reverse Transcription Kit instructions. The cDNA was amplified by SYBR* Premix Ex Taq (TAKARA, Japan). The primers were designed using the gene sequences provided in NCBI Genebank and synthesized by Beijing Aoko Biological Engineering Co., Ltd. The used primers used in this study are as following: IL-6 forward 5’-CCACCAGGAACGAAGTCAAC-3’, reverse 5’-GGCAGTGGCTGTCAACAAC-3’; IL-1β forward 5’-GACCTGTCTTTGAGGCTGAC-3’, reverse 5’-TCCATCTTCCTCTTGGGTATTGTT-3’; IL-8 forward 5’-TGTTCTCAGCCACCCGC-3’, reverse 5’-TCCACGACATCTCAGCA-3’;

β-actin forward 5’-GCTGTCGTCGACAA CGGCTTG- 3’, reverse 5’-CAAACATGATCTGC GGTCATCTTTTC-3’. β-actin was used to be as an internal reference.

**Enzyme Linked Immunosorbent Assay (ELISA)**

The cells were seeded into 6-well plate. After the intervention of high glucose, the supernatant of cell culture medium was collected and centrifuged at 3000 r/min for 15 min to collect supernatant. The experiment was performed to detect the levels of IL-1β, IL-6 and IL-8 in strict accordance with the kit instructions (Beyotime Biotechnology, Shanghai, China).

**Western blot**

Total protein was extracted by protein extraction kit, and then electrophoresis was performed in polyacrylamide gel. The protein was electrically transferred to PVDF membrane, blocked with 5% skimmed milk powder and incubated with primary antibodies (FN, ab268020; collagen I, ab270993; collagen III, ab275881; collagen IV, ab270993; MMP-2, ab92536; MMP-9, ab76003; GAPDH, ab8245. Abcam, England) at 4 °C overnight, followed by the incubation with HRP-conjugated secondary antibodies (ab7090, Abcam, England) for 2 h at 37°C. GAPDH was used as an internal reference. The optical density ratio of target protein to corresponding internal reference was calculated by Image J soft ware 1.46 r.
Statistical analysis
Prism 7.0 software was used for statistical analysis of experimental data. Data were represented by mean ± standard deviation (SD). One-way analysis of variance was used to determine the difference among different groups, followed by turkey’s test. p < 0.05 indicates that the difference is statistically significant.

Result
Captopril suppressed HSC-T6 cell activities induced by HG
To investigate the effects of captopril on HSC-T6 cell subject to HG, we assessed HSC viability after captopril treatment using CCK8 assay. Different concentrations of HG were used to stimulate HSC. The results showed that with the increase in HG concentration, the cell viability of HSC was significantly increased (Figure 1a). In order to evaluate the cytotoxic effect of captopril on HSC-T6, HSC-T6 cells were treated with different concentrations of captopril. There were no significant effects for captopril on cell viability (Figure 1b). Then, we further found that captopril reduced viability of HSC challenged with HG in concentration-dependent manner (Figure 1c). In subsequent experiments, 1 mM of captopril was used for deeper mechanism exploration for its effects on HSC in response to HG.

Captopril suppressed oxidative stress, inflammation and fibrosis in HSC-T6 cells exposed to HG
To analyze the mechanism of captopril in HSC-T6 cells challenged with HG, the effects of captopril on oxidative stress, inflammation and fibrosis in HSC-T6 cells cotreated with HG and captopril were determined. The results showed that captopril exhibited markedly higher SOD activities and SOD levels than HG treatment alone, while reverse effects were observed in MDA levels (Figure 2 A). We observed an obvious decrease in fluorescence intensity for ROS staining after captopril treatment (Figure 2b). By testing the expression of inflammation-related factors through qPCR and ELISA assay, the results showed that captopril treatment markedly lessened the levels of IL-1β, IL-6 and IL-8 (Figure 2c-d) as relative to HG group. The levels of FN, collagen-I, collagen-III, collagen-IV, MMP-2 and MMP-9, which have been recognized to be as markers for liver fibrosis, were also analyzed by Western blot assay in HG-challenged HSC-T6 cells following captopril treatment. As shown in Figure 2e, these protein levels were markedly decreased when captopril was used to co-treat cells with HG.

Captopril inhibits HG-induced activation of the NF-kappaB and wnt3a/β-catenin pathway in HSC-T6 cells
To analyze the role of the NF-kappaB and wnt3a/β-catenin after captopril treatment, we

![Figure 1](image-url)  
*Figure 1.* Captopril reduced HSC-T6 cells viability. (a) After HSC-T6 cell was exposed to glucose of different concentration for 48 h, cell viability was detected through CCK8 assay. (b) CCK8 assay. (c) CCK8 assay. *p < 0.05, **p < 0.01, *** p < 0.001.
looked at expression levels of NF-kappaB pathway-related proteins and Wnt3α/β-catenin. The phosphorylation of IκBα makes it possible that NF-κB is released from the cytoplasmic NF-κB/IκBα complex and activated to expose the nuclear localization domain and p65 is confirmed to be the main transcriptional factor. Increased ratio of p-IκBα/IκBα and the ratio of p-p65/p65 are considered to be related to be involved in NF-κB activation [14–16]. As shown in results, captopril markedly decreased the ratio of p-IκBα/IκBα and the ratio of p-p65/p65 when compared with HG alone (Figure 3). In addition, the expression of Wnt3A and β-catenin were detected. The results showed that the expression of Wnt3α/β-catenin protein was increased after HG treatment, while that of Wnt3α/β-catenin was decreased after captopril treatment (Figure 3). The above results indicated that captopril could inhibit HG-induced activation of NF-kappaB pathway and Wnt3α/β-catenin pathway in HSC-T6 cells.

Captopril inhibited HG-induced oxidative stress, inflammatory response, and fibrosis in HSC-T6 cells by blocking the NF-κB pathway and Wnt3α/β-catenin pathway

Next, whether the effects of captopril on oxidative stress, inflammatory response, and fibrosis in HSC-T6 cells were related to NF-κB pathway and Wnt3α/β-catenin pathway was further investigated. We used NF-κB activator or Wnt signaling pathway activator to determine whether captopril affected NF-κB signaling pathway and Wnt signaling pathway to suppress oxidative stress, inflammation and fibrosis. Accumulating evidences have confirmed that PMA is able to activate NF-κB pathway and LiCl is an activator of Wnt signaling pathway [17–22]. Treatment with PMA (NF-κB activator) or treatment with LiCl Wnt signaling pathway activator) resulted in a significant decrease in SOD activities and an increase in MDA levels compared with the cotreatment group of HG and captopril (Figure 4a). DCFH-DA staining for ROS demonstrated marked...
attenuation in green fluorescence intensity after PMA or LiCl treatment (Figure 4b). Next, we examined the effects of PMA or LiCl on inflammatory markers. PMA or LiCl treatment significantly partly recovered captopril-treated effects on levels of IL-1β, IL-6 and IL-8 in HG-induced HSC-T6 cells (Figure 4c-d). We also studied FN, collagen-I, collagen-III, collagen-IV, MMP-2 and MMP-9 levels, which are markers for liver fibrosis. The results demonstrated a marked increase after treatment with PMA or LiCl (Figure 4e), which implied that captopril inhibited fibrosis through NF-κB and Wnt signaling pathway.

**Discussion**

Our results demonstrate that captopril is a potentially effective anti-fibrosis reagent in HG-induced HSC-T6 cells and its effects are mediated by suppressing NF-kappaB and wnt3a/β-catenin. Additionally, captopril significantly increases cell viability and suppresses inflammation and oxidative stress. Use of NF-κB signaling pathway activator or Wnt signaling pathway activator confirms that NF-κB pathway and wnt3a/β-catenin pathway mediate the effects of captopril on oxidative stress, inflammation and fibrosis [23].

Liver fibrosis is a chronic process that usually progresses over decades and can eventually lead to cirrhosis [24]. Recently, more and more studies pay attention to the identification of drugs, which can produce inhibitory effects on the progression of hepatic fibrosis [25]. Captopril’s potent ant-fibrosis properties suggest that it is expected to be a potential anti-fibrosis agent. Captopril also suppresses oxidative stress and inflammation in HSC-T6 cells. Treatment of captopril significantly reduced MDA and ROS levels and increased SOD activities, as well as a decrease in inflammatory marker levels in HG-induced HSC-T6 cells. The mechanism of antioxidant activity and anti-inflammatory effects of captopril could be related to the inhibition of NF-kappaB and wnt3a/β-catenin. In previous research, captopril significantly increased Nrf2 and HO-1 protein levels, which could be involved in the modulation for oxidative stress in vitro [23]. We predicted that Nrf2 signaling pathway could be also engaged in mediating the effects of captopril on oxidative stress.

The roles of captopril in anti-oxidative stress and inflammation, as well as anti-fibrosis have been revealed in some studies [10,26,27]. Besides the regulatory role of NF-kappaB in inflammation and oxidative stress in diabetes, wnt3a/β-catenin was also reported to be involved in these processes [28–30]. Wnt3a activation in response to glucose was related to increased β-catenin level [31]. A study found that wnt3a/β-catenin pathway was engaged in hepatoprotective effects of hesperidin, which could be related to oxidative imbalance and inflammation [32]. Additionally, Wnt3a/β-catenin pathway was confirmed to modulate oxidative stress and inflammation [33–35]. Treatment with
Captopril contributes to the decrease in protein levels of FN, collagen I, collagen III, collagen IV, MMP-2 and MMP-9 in HG-induced HSC-T6 cells, which plays key roles in the formation of ECM through inducing the activation of HSCs [36–39]. In summary, captopril plays a protective role in protecting hepatic stellate cell from oxidative stress, inflammation and fibrogenesis injury caused by HG via NF-kappaB pathway and Wnt3a/β-catenin pathway.

Conclusion
Liver injury is a typical inducement for the development of liver fibrosis and cirrhosis. Our study demonstrated that inhibition of the NF-kappaB pathway and Wnt3a/β-catenin by captopril under these HG conditions could exert inhibitory effects on fibrogenesis. The study provided experimental basis to confirm that captopril could be a potential therapeutic drug for the treatment of hepatic fibrosis. How captopril plays a role in vivo still requires amounts of experiments to explore, which will be considered in the next step of the study.

Highlights
1. Captopril reduces increased viability of rat hepatic stellate cells induced by HG stimulation.
2. Captopril alleviates oxidative stress, inflammation and fibrogenesis in HG-induced HSC-T6 cells.
3. These effects of captopril could be mediated by NF-κB pathway and Wnt3a/β-catenin pathway.
Data availability statement

The datasets used and/or analyzed during the current study are available from the corresponding author or first author on reasonable request.

Disclosure statement

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