Ursolic acid ameliorates CCl4-induced liver fibrosis through the NOXs/ROS pathway

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Liver fibrosis is a reversible wound-healing response that occurs after liver injury. NADPH oxidases (NOXs) and reactive oxygen species (ROS) which are expressed in hepatocytes (HCs), hepatic stellate cells (HSCs), and Kupffer cells (KCs) play an important role in the development of hepatic fibrosis. In vitro studies, we had shown that ursolic acid (UA) could reverse liver fibrosis by inhibiting the activation of NOX-mediated fibrotic signaling networks in HSCs. In this study, we verified that UA could alleviate CCl4-induced liver fibrosis by reducing the expression of NOXs/ROS in HCs, HSCs, KCs. Meanwhile, the phagocytic index α and clearance index K which represent phagocytosis of KCs were unchanged. Together, all our data demonstrated that UA induced the proliferation of HCs, promoted apoptosis in HSCs, and prevented activation of KCs in vivo by reducing the expression of NOXs/ROS in HCs, HSCs, KCs. Besides, UA had no effect on the host defense function.

KEYWORDS
hepatic stellate cells, hepatocytes, Kupffer cells, ursolic acid

1 INTRODUCTION

Liver fibrosis is a common consequence of chronic liver injury and is caused by a variety of pathogenic processes, including viral infection, alcohol consumption, nonalcoholic fatty liver disease, cholestasis, autoimmune liver disease, and schistosomiasis japonica. Liver fibrosis leads to the accumulation of extracellular matrix or the formation of a fibrous scar (Thompson, Conroy, & Henderson, 2015; Zhang & Friedman, 2012). The transdifferentiation of hepatocyte stellate cells (HSCs) from a quiescent vitamin A storage cell to a proliferative myofibroblast is recognized as the central event in the development of hepatic fibrosis (Thompson et al., 2015). Hepatocytes (HCs) and Kupffer cells (KCs) are involved in the regulation of HSC activation through the release of TGF-β1 (Liang, Kisseleva, & Brenner, 2016; Paik et al., 2014). Therefore, these cells appear to be specific cellular targets for antifibrotic therapy in chronic liver diseases of any etiology. Recent studies have shown that NADPH oxidases (NOXs) and reactive oxygen species (ROS) play an important role in the
development of hepatic fibrosis. Of the seven NOX family isoforms, the expression of NOX1, NOX2, and NOX4 has been widely studied in liver fibrosis. These three NOXs are expressed in HSCs, HCs and KCs, and they play an important role in the pathogenesis of hepatic fibrosis. Studies on the expression of NOXs in those cells and explorations of the drugs that block NOX activation in those cells are expected to achieve the goal of treating hepatic fibrosis (Altenhofer, Radermacher, Kleikers, Wingler, & Schmidt, 2015; Crosas-Molist & Fabregat, 2015).

Ursolic acid (UA), a natural pentacyclic triterpenoid that is derived from berries, leaves, flowers, and fruits of medicinal plants, has been reported to have anti-oxidation, anti-inflammatory, anti-ulcer, antibacterial, anti-virus, anti-tumor, anti-obesity, hypoglycemic, antihypertensive, lipid-lowering, liver protection, and immune regulation activities, among other (Cargnin & Gnoatto, 2017; Kim, Kim, Han, & Kim, 2015; Li, Ren, Luo, & Yang, 2016; Ma, Ding, Zhang, & Liu, 2015; Wang et al., 2011). In vitro studies have shown that UA could reverse liver fibrosis by inhibiting the activation of NOX-mediated fibrotic signaling networks in HSCs, including the NF-κB, PI3 K/Akt, P38MAPK, ERK1/2, JAK2-STAT3, Hedgehog, and other signaling pathways (He et al., 2015). However, no one has reported whether the mechanisms of UA anti-liver fibrosis are related to a reduction in the expression of NOXs in HSCs, HCs, and KCs in vivo. In this study, therefore, we investigated the effects of UA on the activation of NOXs in HSCs, HCs, and KCs in SD rats.

2 | MATERIALS AND METHODS

2.1 | Chemicals and reagents

UA and apocynin were purchased from Sigma Chemical Co. (St. Louis, MO), and a TUNEL assay kit was purchased from Promega Chemical Co. (Madison, WI). CCl4 and olive oil were from Shan-dong Xiya Reagent Bioengineering Institute (Shandong, China). Hydroxyproline, malondialdehyde and glycogen staining assay kit were from Nan-jing Jiancheng Bioengineering Institute (Nanjing, China). Sirius red staining assay kit was from Beijing Solarbio Bioengineering Institute (Beijing, China). Lipopolysaccharides (LPS) ELISA kit was from Wuhan Elabscience Biotechnology Co. (Hubei, China). DCFH-DA cells ROS assay kit was from Beyotime Biotechnology (Shanghai, China). Anti-MMP1 antibody, anti-TIMP1 antibody, anti-TGFβ1 antibody, and anti-F4/80 antibody were from Santa Cruz Biotechnology (Santa Cruz, CA). Anti-caspase3 antibody was from Cell Signaling Technology, Inc. (Boston, MA). All the other antibodies were obtained from Sigma Chemical Co. (St. Louis, MO).

2.2 | Animals and experimental design

All SD rats (160–200 g) were obtained from the department of animal science, Medical College of Nanchang University. All animals received humane care, and the experimental protocol was approved by the Committee of Laboratory Animals and was performed according to institutional guidelines. All animals were kept in an environmentally controlled room (23 ± 2 °C, 55 ± 10% humidity) with a 12 hr light/dark cycle, and they were allowed free access to food and water.

After a 1-week housing period, seventy-two SD rats were randomly divided into six groups: blank control group, liver fibrosis model group, apocynin prevention group, apocynin treatment group, UA prevention group, or UA treatment group. For the liver fibrosis model group, we gave rats an intragastric administration of carbon tetrachloride (CCl4) for 8 weeks at 2 ml/kg (20% olive oil dilution concentration) twice a week. For rats in the prevention groups, 2 weeks after gavaging with CCl4, we gavaged the rats with apocynin or UA (40 mg/kg/day) for the last 6 weeks at the same time. For rats in the treatment groups, we gavaged the rats with apocynin or UA for another four weeks alone, 8 weeks after gavaging with CCl4.

At the end of treatment, eight rats in each group were used for the biochemical analysis and histological evaluations. Rats were sacrificed, and blood samples of approximately 5 ml were drawn through a postcava puncture with vacuum blood collection tubes. The plasma was collected after centrifugation at 5,000 rpm for 10 min and stored in a −70 °C freezer for further analysis. The liver tissues were rinsed in ice-cold 0.9% NaCl solution and stored in a −70 °C freezer for later use.

Four rats in each group were used for extracting primary HSCs, HCs and KCs, in situ perfusion and gradient centrifugation. The rats were anesthetized by an intraperitoneal injection of 10% chloral hydrate. We perfused phosphate buffered saline (PBS) through the inferior vena cava, and cut open the portal vein at the same time. According to various densities in those three kinds of cells, we then separated HCs, HSCs, and KCs using Collagenase IV (Gibco, Grand Island, NY, USA), Pronase E (Sigma Immunochemicals, St. Louis, MO, UA) and Preccoll (Sigma) successively. HCs and HSCs were cultured in Dulbecco's modified essential medium (DMEM, Gibco) supplemented with 10% fetal bovine serum (FBS, Gibco) in humidified air containing 5% CO2 at 37 °C; however, KCs were cultured in 1640 medium (Gibco) supplemented with 10% FBS (Gibco) in humidified air containing 5% CO2 at 37 °C.

2.3 | Blood and liver tests

The alanine aminotransferase (ALT), total bilirubin (TBLI), albumin (ALB), white blood cell (WBC), C-reactive protein (CRP), and procalcitonin (PCT) contents of serum were estimated using an automatic biochemical analyzer (Department of clinical laboratory, the First Affiliated Hospital of Nanchang University, China). The malondialdehyde (MDA) and hydroxyproline (HYP) contents of serum or liver tissue were tested spectrophotometrically using commercial diagnostic kits.

2.4 | Histological evaluations

Liver samples fixed in 10% buffered formalin were embedded in paraffin, cut into 5 μm sections, stained with hematoxylin and eosin (H&E), Sirius red, TdT-mediated dUTP Nick-End Labeling (TUNEL, to detect the apoptosis of HCs in liver tissue), or for immunohistochemistry (IHC, to detect proliferating cell nuclear antigen, PCNA) analysis before being observed and photographed using light microscopy.
For the semi-quantitative of TUNEL and PCNA expression, we used a scoring method as follows. The percentage of positive HCs in liver tissue was determined in at least five areas at ×400 magnification and assigned to one of the following categories: 0 = <5%; 1 = 6–49%; 2 = 50–75%; 3 > 75% (Table A below). The staining intensity of the nucleus were scored as 0 point (uncolored), 1 point (light yellow), 2 point (brown), 3 point (tan) (Table B below). Finally, according to the products of the positive cells percentage and the staining intensity score, the results were decided.

| A The percentage of positive cells |
|-----------------------------------|
| 0–5% | 6–49% | 50–75% | >75.0% |
| 0 | 1 | 2 | 3 |

| B The staining intensity score |
|-------------------------------|
| Uncolored | Light yellow | Brown | Tan |
| 0 | 1 | 2 | 3 |

### 2.5 Western blot analysis

Liver tissue protein was obtained using protein lysate (BestBio, Shanghai, China) for Western blotting. Cells were harvested and re-suspended in PBS. Cell protein extracts were obtained by centrifuging at 12,000 rpm for 10 min. Protein levels were determined using the BCA assay kit (Tiangen, Beijing, China). Protein extracts (10–20 ml) were separated using 6–12% sodium dodecyl sulfate polyacrylamide gel electrophoresis, and electrophoretically transferred onto polyvinylidene fluoride membranes. After blocking with 5% nonfat dry milk in Tris-buffered saline, PVDF membranes were incubated overnight at 4 °C with rabbit anti-NOX1 (1:1000), anti-NOX2 (1:1000), anti-NOX4 (1:1000), anti-TGFβ1 (1:2000), anti-MMP1 (1:2000), anti-TIMP1 (1:2000), anti-p67phox (1:500), anti-p22phox (1:500), anti-p47phox (1:500), anti-caspase-3 (1:1000), anti-caspase-9 (1:1000), anti-GADD45 (1:1000), anti-Bcl2 (1:800), anti-Bax (1:1000), anti-CD95L (1:200), anti-CD95 (1:3000), anti-XIAP (1:2000), or mouse anti-α-SMA (1:200), anti-rac1(1:750), anti-collagenase I (1:1000). Next, a secondary horseradish peroxidase-conjugated anti-rabbit or anti-mouse IgG antibody (Zhongshan Golden Bridge Biotechnology Co., Beijing, China) was applied, and specific bands were visualized using an ECL detection kit (Thermo Fisher Scientific Inc).

### 2.6 RNA preparation

Total liver RNA was prepared from frozen tissue samples or cells using a RNA simple Total RNA Kit and a Fast Quant RT Kit (Tiangen, Beijing, China). The concentration and purity of isolated RNA were determined by measuring optical density at 260 and 280 nm. The integrity of the RNA was verified using agarose gel electrophoresis (AGE).

### 2.7 RT-qPCR

For reverse transcription-polymerase chain reaction (RT-PCR), we used the experimental procedure from the Fast Quant RT Kit and the Super Real Pre-Mix Plus (SYBR Green, tiangen, Beijing, China). The primer sets were as follows (Table 1). All primers were obtained from Beijing Genomics Institute (BGI, Shenzhen, China).

2.8 Extraction of primary HSCs, HCs, and KCs

#### 2.8.1 Liver perfusion

Four rats were taken from each group. Rats were anesthetized by intraperitoneal injection of 10% chloral hydrate. The abdominal cavity was cut open aseptically and the inferior vena cava was punctured. The PBS solution was uniformly perfused by the syringe, at the same time, the hepatic portal vein was cut open till the liver tissue turned into earthy yellow. Injected into the hepatic portal vein with perfusion solution (0.05% Collagenase IV) 50 ml at 37 °C, and digested 10 min.

#### 2.8.2 Extraction of primary HCs

1. The liver tissue was completely cut off into in a plate.
2. Lift the liver with ophthalmal forceps, and remove the capsule, blood vessels, connective tissue.
3. Tear the liver tissue and obtaine suspension of liver cells. The suspension was filtered through a 150 mesh stainless steel mesh, and the filtrate was collected.
4. Centrifuge at 50g for 5 min, and the cell was collected.
5. The cells were re-suspended in PBS, and centrifuged again (50g, 5 min).
6. The cells were suspended in DMEM conditioned medium and cultured overnight in a humidified air containing 5% CO₂ at 37 °C.

#### 2.8.3 Extraction of primary HSCs and KCs

1. The remaining liver tissue in Step 2 was cut into pieces and added with four times volume digestion solution (0.2% Collagenase IV + 0.2% Pronase E) for 30 min with water bath shaking at 37 °C.
2. After digestion, the tissue was mechanical separated softly until there was no obvious tissue mass.
3. Centrifuge at 50g for 3 min, and the supernatant was collected.
4. The supernatant was centrifuged at 500g for 5 min, and the sediment was collected.
5. The sediment was re-suspended in PBS. We slowly added to the upper layer of well-prepared separating medium with 70% and 30% Precoll non-continuous gradient.
6. Centrifuge at 900g for 15 min. All layers of cells were aspirated slowly. The junction of 70% and 30% were KCs and liver sinusoidal endothelial cells, and upper 30% layer were HSCs.
7. For HSCs, re-suspended in DMEM and centrifuge at 500g for 5 min.
8. The sediment was resuspended in DMEM again and cultured overnight in a humidified air containing 5% CO₂ at 37 °C.
9. KCs were re-suspended in 1,640 complete medium and cultured into Petri dishes. Depending on the different speed of the adherent wall, we extracted the primary HSCs, and KCs. After cultured with...
30 min, we discarded unadherent cells, replaced the fresh medium, and the adherent were KCs. KCs were cultured in a humidified air containing 5% CO₂ at 37 °C.

2.9 | Identification of primary HSCs, HCs, and KCs

2.9.1 | Identification of primary HSCs (α-SMA immunofluorescence identification) and KCs (F4/80 immunofluorescence identification)

1. The slides in the culture plate which had climbed with cells were immersed in PBS for three times, 3 min at a time.
2. Fix the slides with 4% paraformaldehyde for 15 min, and then wash the slides with PBS for three times, 3 min at a time.
3. Permeate the slides with 0.5% Triton X–100 (prepared with PBS) at room temperature for 20 min.
4. The slides were immersed in PBS for three times, 3 min at a time. Blot PBS with absorbent paper, and drop normal goat serum on the slides, blocking at room temperature for 30 min.
5. Blot serum with absorbent paper. Each slide was dropped with primary antibody dilution (1:100) and placed in wet box, incubating overnight at 4 °C.
6. Add fluorescent secondary antibody: PBST washed the slides three times, 3 min at a time. Blot PBST with absorbent paper, and add fluorescent secondary antibody dilution to the slides (1:200), incubating in wet box at 37 °C for 1 hr. PBST washed the slides three times, 3 min at a time. (from the step of adding fluorescent secondary antibody, the remained steps were operated as far as possible in the darkroom).
7. Counterstain nuclear: DAPI was added to the slides. After incubation for 5 min, PBST washed the slides four times, 5 min at a time.
8. Blot the liquid on the slides with absorbent paper, and then seal the slide with the antifluorescent quencher. Capture image under a fluorescence microscope.

2.9.2 | Identification of primary HCs (by schiff periodic acid shiff, PAS)

1. The slides in the culture plate which had climbed with cells were immersed in PBS for three times, 3 min at a time.
2. Fix the slides with 4% paraformaldehyde for 15 min, and then wash the slides with PBS for three times, 3 min at a time.
3. Soak the slides with 1% periodic acid aqueous solution for 10 min. Wash the slides with PBS for 2 min.
4. Add Schiff’s reagent on the slides, incubating the slides in the dark at room temperature for 30 min. Dip the slides with PBS for 10 min.
5. Counterstain with Mayer hematoxylin for 2 min, and wash the slides with water until the nucleus turned blue.
6. Dehydration with anhydrous ethanol, and transparention with dimethylbenzene.
7. Seal the slides with neutral balsam after dryness. Capture image under a microscope.

2.10 | Determination of ROS

2.10.1 | For ROS in HCs or HSCs

1. After cell separation, HCs or HSCs were made into a single cell suspension, and the cells were uniformly inoculated into 6-well plates at approximately 4 × 10⁵ cells per well. The cells were cultured overnight in a humidified air containing 5% CO₂ at 37 °C;
2. Digest cells with 0.25% trypsin without EDTA. After termination of digestion, centrifuge at 1,500 rpm for 5 min. Supernatant was discarded and sediments were resuspended in PBS, and the above steps were repeated three times;
3. Detection using DCFH-DA cells ROS assay kit:
   a. Add 1 ml of PBS with diluted DCFH;
   b. Incubate cells at 37 °C for 20 min. Mix every 3 min;
c. Wash the cells three times with serum-free medium; 
d. Add 500 μl of PBS resuspended cells; 
4. Detection with flow cytometer.

2.10.2 For ROS in KCs

1. After cell separation, KCs were made into a single cell suspension, 
and the cells were uniformly inoculated into 6-well plates at 
approximately 4 × 10^5 cells per well. The cells were cultured 
overnight in humidified air containing 5% CO2 at 37 °C; 
2. Prepare 10^{-3} mmol/ml DCFH-DA working solution by adding 
10^{-2} mmol/ml DCFH-DA: PBS = 1: 9; 
3. Digest cells with 0.25% trypsin without EDTA. After termination 
of digestion, use part of the supernatant for protein 
determination. Add the protein concentration that calculated 
from the supernatant into the DCFH-DA working solution, and mix. Set PBS for the control wells. Incubate solution at 
37 °C for 30 min; 
4. Detection of fluorescence intensity: the excitation wavelength 
was 485 nm and the emission wavelength was 525 nm. The

FIGURE 1 Effect of UA on regression of CCl4-induced liver injury. (a) hematoxylin-eosin (HE) staining. (b) Sirius red staining. 
Morphometrical analysis for fibrotic score (c) and area (d) were performed for five random fields in each preparation, and the average percentages of fibrotic area are plotted. Detection of hydroxyproline content in serum (e) and liver tissue (f) by colorimetry. All values are 
expressed as the means ± S.E.M. (n = 8). CCl4 group compared to all other groups, respectively; *p < 0.05, **p < 0.01, ***p < 0.001. Original 
magnification, ×100.
results were expressed as fluorescence intensity/mg protein. ROS (fluorescence intensity/mg protein) = fluorescence intensity/ (protein concentration × 0.19).

2.11 | Test host defense function of KCs

2.11.1 | Carbon particle clearance test detected the phagocytic function of KCs

1. Rats in each group were injected with ink (Carbon suspension, 10 ml/kg) in the tail vein 1 hr before the rats were sacrificed.
2. Then, 20 µl blood was collected from the eyeball of rats at 1, 5, 10 min, respectively. The blood was dissolved in 2 ml 0.1% of Na2CO3 solution.
3. The absorbance (A value) was measured at 650 nm with a spectrophotometer.
4. Weigh the wet weight of liver and spleen, calculate clearance index K and phagocytic index α based the formula:

\[ K = \frac{(\log A_1 - \log A_2)}{(t_2 - t_1)} \]

\[ \alpha = \frac{\text{body weight}}{(\text{liver weight} + \text{spleen weight})} \times \frac{1}{3} K. \]

2.12 | Detection of bacterial infection

Peripheral blood (from eyeball) was taken weekly after drug treatment, and white blood cell (WBC), C-reactive protein (CRP), and procalcitonin (PCT) in serum were estimated. Lipopolysaccharides (LPS) was tested by ELISA kit.

2.13 | Statistical analysis

All statistical analyses were performed using the SPSS software, version 17.0. Data were expressed as the means ± SEM. Statistical differences between means were determined using analysis of variance (ANOVA), Student's t test, Mann-Whitney rank sum test, or Kruskal–Wallis H test. A value of \( p < 0.05 \) was selected prior to the experiments to indicate statistical significance.

3 | RESULTS

3.1 | UA alleviated CCl4-induced liver fibrosis

First, the effect of UA on the regression of CCl4-induced liver injury was observed. As shown in Figures 1a and 1b, CCl4 treatment caused inflammatory cellular infiltration, hepatocellular necrosis, the formation of

**FIGURE 2** Effect of UA intervention on liver function and oxidative stress. (a) Automatic biochemical analyzer estimated ALT in rats’ serum. (b) TBIL in rats serum. (c) ALB in rats’ serum. MDA commercial diagnostic kits were used to estimate MDA in rats serum (d) and liver tissue (e). All values are expressed as the means ± S.E.M. (n = 8). CCl4 group compared to all other groups, respectively; *\( p < 0.05 \), **\( p < 0.01 \), ***\( p < 0.001 \).
fibrous septum, deposition of collagen fibers, and formation of a pseudo-lobe. After UA or AP intervention, the listed visible liver histology injuries were alleviated. Disorder in the hepatic lobular structure, fibrous space and collagen deposition in liver tissue were also significantly lower after prevention or treatment compared to the CCl4 group. Indeed, quantitative analysis demonstrated a significant reduction in the fibrotic score and area (Figures 1c and 1d). The hydroxyproline contents of rat serum and hepatic tissue declined compared to the CCl4 group (Figures 1e and 1f). Nevertheless, there was no significant difference between the UA treatment group and the AP treatment group ($p > 0.05$), nor was there a significant difference between the UA prevention group and AP prevention group ($p > 0.05$). Those results showed that UA could alleviate CCl4-induced liver fibrosis.

### 3.2 The effect of UA intervention on liver function and oxidative stress

We measured the alanine aminotransferase (ALT), total bilirubin (TBIL), albumin (ALB), and malondialdehyde (MDA) contents of rats serum or liver tissue. CCl4 treatment resulted in high expression of ALT (Figure 2a), TBIL (Figure 2b), MDA (Figures 2d and 2f), and low expression of ALB (Figure 2c). After UA or AP intervention, the expression of the listed targets was reduced. Nevertheless, there was no significant difference between the UA treatment group and the AP treatment group ($p > 0.05$), nor was there a significant difference between the UA prevention group and AP prevention group ($p > 0.05$). We concluded that UA could alleviate CCl4-induced hepatic dysfunction and oxidative stress.

### 3.3 The effect of UA intervention on proliferation and apoptosis in HCs

Subsequently, the effect of UA intervention on proliferation and apoptosis of HCs was evaluated. We used immunohistochemistry (IHC) to detect proliferating cell nuclear antigen (PCNA) in liver tissue (Figure 3a). We utilized TdT-mediated dUTP Nick-End Labeling (TUNEL) staining to detect apoptosis in HCs (Figure 3b). CCl4 treatment led to increased apoptosis in HCs, and it led to decreased proliferation of HCs; After UA or AP intervention, the proliferation of HCs was increased, and apoptosis in HCs was decreased. The PCNA

![Image](image-url)
positive score and TUNEL positive score are shown in a table (Figure 3c). Nevertheless, there was no significant difference between the UA treatment group and the AP treatment group \((p > 0.05)\), nor was there a significant difference between the UA prevention group and AP prevention group \((p > 0.05)\). These results showed that UA could reduce apoptosis in HCs, but increased the proliferation of HCs.

### 3.4 The effect of UA intervention on genes and proteins related to fibrosis

The effect of UA on the expression of type I collagen, TGF-\(\beta\)1, MMP-1, and TIMP-1 genes and proteins in liver tissue was tested. As shown in Figure 4a, CCl4 treatment induced high expression of type I collagen, TGF-\(\beta\)1, and TIMP-1 genes, but it induced low expression of the MMP-1 gene. After UA or AP intervention, the expression of type I collagen, TGF-\(\beta\)1, and TIMP-1 genes were reduced, and the expression of the MMP-1 gene was increased. This was the same as the effect of UA on type I collagen, TGF-\(\beta\)1, MMP-1, and TIMP-1 proteins. As shown in Figures 4b and 4c, CCl4 treatment induced high expression of type I collagen, TGF-\(\beta\)1, and TIMP-1 proteins, but a low expression of the MMP-1 protein. After UA or AP intervention, the expression of type I collagen, TGF-\(\beta\)1, and TIMP-1 proteins were reduced, and the expression of the MMP-1 protein was increased. Nevertheless, there was no significant difference between the UA treatment group and the AP treatment group \((p > 0.05)\), nor was there a significant difference between the UA prevention group and AP prevention group \((p > 0.05)\). These results demonstrated that UA could improve the expression of CCl4-induced fibrosis-related genes and proteins.

### 3.5 The effect of UA intervention on NOX proteins in liver tissue

Recent studies have indicated that NADPH oxidases (NOXs) play an important role in the pathogenesis of liver fibrosis (Liang et al., 2016). We speculated that UA relieved liver fibrosis by reducing the
expression of NOXs. The effect of UA on the expression of NOX4, p67phox, NOX2, p47phox, NOX1, p22phox, and Rac1 proteins were detected in liver tissue. As shown in Figure 5a–h, CCI4 treatment induced a high expression of NOX4, p67phox, NOX2, p47phox, NOX1, p22phox, and Rac1 proteins in liver tissue. After UA or AP intervention, the listed proteins was alleviated. Nevertheless, there was no significant difference between the UA treatment group and the AP treatment group (p > 0.05), nor was there a significant difference between the UA prevention group and AP prevention group (p > 0.05). These results demonstrated that UA could ease CCI4-induced liver fibrosis by reducing the expression of NOX proteins.

3.6 The effect of UA intervention on the expression of NOX mRNAs in HSCs, HCs, and KCs

Research shows that NOXs expressed in hepatic stellate cells (HSCs), hepatic cells (HCs), and Kupffer cells (KCs) play a key role in liver fibrosis (Paik et al., 2014). We wanted to explore the effect of UA intervention on the expression of NOX mRNAs in HSCs, HCs, and KCs in vivo. We used in situ perfusion and gradient centrifugation to extract primary HSCs, HCs, and KCs from rats. We then utilized a glycogen staining assay kit to identify primary HCs, and we used Immunofluorescence (IF) to identify primary HSCs and KCs. As shown in Figure 6 (the first row), HCs contain abundant glycogen. After PAS dyeing, the cytoplasm became purple; α-SMA, and DAPI dual-positive cells were detected in primary HSCs (the second row), and F4/80 and DAPI dual-positive cells were detected in primary KCs (the third row). Next, we inspected the expression of NOX mRNAs in HSCs, HCs, and KCs in vivo. As shown in Figure 6, CCI4 treatment resulted in high expression of NOX4, p67phox, NOX2, p47phox, NOX1, p22phox, and Rac1 genes in HCs (B), HSCs (C) and KCs (D). After UA or AP intervention, the expression of the listed genes declined to varying degrees. Nevertheless, there was no significant difference between the UA treatment group and the AP treatment group (p > 0.05), nor was there a significant difference between the UA prevention group and AP prevention group (p > 0.05). These results showed that UA could relieve CCI4-induced liver fibrosis by reducing the expression of NOX mRNAs in HSCs, HCs, and KCs.

**FIGURE 5** Effect of UA intervention on the expression of NOX proteins. (a) WB detected the expression of NOX4, p67phox, NOX2, p47phox, NOX1, p22phox, and Rac1 proteins in liver tissue. Quantitative analysis of NOX4 (b), p67phox (c), NOX2 (d), p47phox (e), NOX1 (f), p22phox (g), and Rac1 (h) proteins in liver tissue. All values are expressed as the means ± S.E.M. (n = 4). CCI4 group compared to all other groups, respectively; *p < 0.05, **p < 0.01, ***p < 0.001.
3.7 The effect of UA intervention on ROS expressed in HCs, HSCs, and KCs

NOXs produce $\text{O}_2^-$ and $\text{H}_2\text{O}_2$. This occurs because NOXs deliver electrons to $\text{O}_2$ molecules, and generate reactive oxygen species (ROS). The agglutination of intracellular ROS can lead to cell structure damage, oxidative stress, and inflammatory injury (Nisimoto, Diebold, Cosentino-Gomes, & Lambeth, 2014; Weiskirchen, 2015). We observed the effect of UA intervention on ROS in HCs, HSCs, and KCs. As shown in Figure 7a, after HCs or HSCs were mixed with DCF-DA working solution, the expression of ROS in primary cells was observed using confocal fluorescence microscopy. We next used flow cytometry to detect the ROS content. Histograms showed that CCl4 treatment led to a reduction in anti-apoptotic proteins like Bcl2 and XIAP, but it led to an increase in pro-apoptotic proteins like Gadd45β, Bax, CD95L, CD95, Caspase3, and Caspase9 in HCs. After UA or AP intervention, the listed effects were reversed, with anti-apoptotic proteins up-regulated and pro-apoptotic proteins down-regulated. As shown in Figure 8b, CCl4 treatment led to increased anti-apoptotic proteins like Bcl2 and XIAP.

3.8 The effect of UA intervention on anti- or pro-apoptosis protein expression on HCs and HSCs

The apoptosis of HCs and the proliferation of HSCs play a key role in liver fibrosis. We tested the effect of UA on the expression of anti-apoptotic proteins, including Bcl2 and XIAP, and pro-apoptotic proteins, like Gadd45β, Bax, CD95L, CD95, Caspase3 and Caspase9 in HCs and HSCs. As shown in Figure 8a, CCl4 treatment led to a reduction in anti-apoptotic proteins like Bcl2 and XIAP, but it led to an increase in pro-apoptotic proteins like Gadd45β, Bax, CD95L, CD95, Caspase3, and Caspase9 in HCs. After UA or AP intervention, the listed effects were reversed, with anti-apoptotic proteins up-regulated and pro-apoptotic proteins down-regulated. As shown in Figure 8b, CCl4 treatment led to increased anti-apoptotic proteins like Bcl2 and XIAP.
but decreased pro-apoptotic proteins like Gadd45β, Bax, CD95L, CD95, Caspase3, and Caspase9 in HSCs. After UA or AP intervention, anti-apoptotic proteins decreased, and pro-apoptotic proteins increased. Nevertheless, there was no significant difference between the UA treatment group and the AP treatment group (p > 0.05), nor was there a significant difference between the UA prevention group and AP prevention group (p > 0.05). Those results showed that UA could alleviate CCl4-induced liver fibrosis by reducing apoptosis in HCs, or increasing apoptosis in HSCs.

3.9 | Safety research about UA to reverse liver fibrosis

We wanted to know the effect of UA on host defense function. We used the carbon particle clearance test to assess the effect of UA on phagocytosis in KCs. In contrast to rats in the CCl4 group, the phagocytic index α (Figure 9b) and clearance index K (Figure 9c), which represent phagocytosis in KCs, were unchanged in the UA or AP prevention group. However, phagocytosis of KCs in the UA or AP treatment group had an advantage over the CCl4 group. There was no significant change in WBC (Figure 9a), LPS (Figure 9d), CRP (Figure 9e) between the CCl4 group and the other groups. CCl4 treatment generated over-expression of PCT (Figure 9f). After UA or AP intervention, PCT was reduced. Nevertheless, there was no significant difference between the UA treatment group and the AP treatment group (p > 0.05), nor was there a significant difference between the UA prevention group and AP prevention group (p > 0.05). These results showed that UA could not weaken phagocytosis of KCs, but could not impair host defense function or increase the chance of bacterial infection.

4 | DISCUSSION

Hepatic fibrosis is caused by various pathogeneses, and it lead to extracellular matrix deposition or scar formation. The sustainable development of liver fibrosis can eventually develop into liver cirrhosis, or even liver cancer, which can threaten human health (Thompson et al., 2015; Zhang & Friedman, 2012). Although, early antiviral therapy can reverse liver fibrosis in patients with HBV and HCV infection (D’Ambrosio et al., 2012; Marcellin et al., 2013), for other causes of liver fibrosis and cirrhosis, there are still no effective anti-fibrosis drugs applied to clinical treatment (Sun & Kisseleva, 2015). Therefore, developing anti-liver fibrosis drugs is highly significant (Lee, Wallace, & Friedman, 2015). We used fibrotic rats as research subjects, and apocynin (AP) (Rahman et al., 2017) as a positive control, and we observe the effect of UA intervention (UA prevention and treatment) on reversing liver fibrosis. Compared to the liver fibrosis group, the fibrous septum and collagen deposition were reduced in the liver tissue.
FIGURE 8 WB tested the effect of UA intervention on anti- or pro-apoptosis protein expression in HCs and HSCs. (a) anti-apoptotic proteins (Bcl2, XIAP), and pro-apoptotic proteins (Gadd45β, Bax, CD95L, CD95, Caspase3, Caspase9) in HCs. (b) anti-apoptotic proteins and pro-apoptotic proteins in HSCs. All values are expressed as the means ± S.E.M. (n = 4). CCl4 group compared to all other groups, respectively; *p < 0.05, **p < 0.01, ***p < 0.001
of the UA prevention and treatment group (Figure 1a–d). There was an obvious reduction in hydroxyproline in rats serum and liver tissue (Figures 1e and 1d). The expression of type I collagen, TGF beta 1, TIMP-1 mRNA, and protein in liver tissue was reduced; However, the expression of the MMP-1 mRNA and protein was increased (Figure 4a–d). These results showed that UA could alleviate CCl4-induced liver fibrosis.

NADPH oxidases (NOXs) are members of a multi-subunit transmembrane enzyme complex that is composed of seven members: NOX1, NOX2, NOX3, NOX4, NOX5, and two double oxidase Duox1, Doux2. The activation of phagocytic NOX2 requires two membrane proteins, NOX2 (gp91phox), and p22phox; three subcellular proteins, p40phox, p47phox, and p67phox; and two guanosine binding. NOX1, NOXA1, p22phox, and Rac GTPase are needed for the activation of NOX1; and NOX4, p22phox, and Poldip2 are co-assembled into activated NOX4. NOXs are the main sources of ROS, and a large number of ROS in the cells may mediate oxidative stress and promote the development of hepatic fibrosis (Liang et al., 2016). Paik et al. (2011) found that NOX1 or NOX2 gene knockout in mice decreased ROS production, and alleviated CCl4 or bile duct ligation-induced liver fibrosis. Lan, Kisseleva, and Brenner (2015) showed that the NOX1/4 double inhibitor GKT137831, or the NOX1/4 gene knockout could reduce the formation of ROS and the expression of inflammation-related genes and block liver fibrosis. Qiu et al. (2016) showed that AP, the non-specific blocking agent of NOXs, could stop atrial remodeling, and prevent myocardial fibrosis,
Jiang et al., 2010) showed that inhibiting the activation of NOX2 could inhibit the activation of HSCs. In this study, the expression of NOX4, p67phox, NOX2, p47phox, NOX1, p22phox, and Rac1 proteins in liver tissue were decreased after UA or AP intervention (Figure 5a–h) compared to the liver fibrosis group. Malondialdehyde (MDA) in rats serum or liver tissue was down-regulated (Figures 6a–h) compared to the liver fibrosis group. This was accompanied by a reduction in activated KCs; The phagocytic index α and clearance index K which represent phagocytosis of KCs were unchanged in the UA or AP prevention group (Figures 9b and 9c); but phagocytosis of KCs in UA or AP treatment group had an advantage over the CCl4 group. These results indicated that UA could lighten liver fibrosis by reducing the activation of KCs, and partially improve phagocytosis of KCs.

In conclusion, UA induces the proliferation of HCs, promotes apoptosis in HCs, and prevents activation of KCs in vivo by reducing the expression of NOXs/ROS in HCs, HSCs, KCs. UA also partially improves phagocytosis of KCs, but it has no effect on the host defense function. The molecular mechanisms that undergird the effect of UA on liver fibrosis require further investigations, and future approaches toward clinical applications of UA need to be done too.

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CONFLICTS OF INTEREST

All the authors who have taken part in this study declared that they have no conflicts of interest to this manuscript.

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