LPS-TLR4/MD-2–TNF-α signaling mediates alcohol-induced liver fibrosis in rats

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Abstract: Liver fibrosis results from liver inflammation and progresses to liver cirrhosis or liver cancer. It is known that nonalcoholic liver disease is mediated by the Toll-like receptor 4 (TLR4)/myeloid differentiation factor-2 (MD-2)–tumor necrosis factor-alpha (TNF-α) signaling pathway. This study aimed to investigate whether alcoholic liver disease is also mediated by this pathway. To this end, we first established rat models of liver fibrosis by administering alcohol. Next, the rats were injected with anti-TLR4 and anti-MD-2 antibodies. Real Time Quantitative PCR (RT-qPCR) and Western blotting were used to detect the activation of the TLR4/MD-2–TNF-α signaling pathway and hepatic stellate cells (HSCs). Moreover, the expression of molecules related to liver fibrosis was estimated. The morphology of rat liver tissue was observed through hematoxylin–eosin staining and Masson staining. For in vitro studies, Kupffer cells (KCs) isolated from the liver were transfected with si-TLR4 and si-MD-2 and co-cultured with HSCs to determine the activity of HSCs. It was found that alcohol treatment activated the TLR4/MD-2–TNF-α signaling pathway and upregulated the molecules associated with liver fibrosis. However, inhibition of TLR4 and MD-2 partially reversed this trend. Notably, in vitro studies indicated that knockdown of TLR4 and MD-2 in KCs partially inhibited LPS-induced activation of KCs and HSCs. Overall, this study showed that alcohol induces liver fibrosis via the LPS-TLR4/MD-2–TNF-α signaling pathway. (DOI: 10.1293/tox.2021-0018; J Toxicol Pathol 2022; 35: 193–203)

Key words: hepatic fibrosis, TLR4/MD-2, Kupffer cells, hepatic stellate cells

Introduction

Liver fibrosis results from the healing reaction of inflammatory injury of the liver caused by alcohol, drugs, and autoimmune1, 2. In the absence of timely treatment, liver fibrosis progresses to liver cirrhosis, hepatic failure, or even liver cancer3, 4. Fibrosis is characterized by abnormal deposition of the extracellular matrix (ECM) in the liver, which is usually reversible at a specific stage. However, once cirrhosis is formed, the change is almost irreversible5, 6. Worldwide, the number of deaths from liver cirrhosis and cancer is rising, with a very high mortality rate due to late diagnosis7. Therefore, to avoid the transformation of liver fibrosis into advanced cirrhosis or cancer, it is vital to develop effective anti-fibrosis methods for the clinical management of liver fibrosis.

Inflammation is a pivotal process in the occurrence and development of liver fibrosis, where hepatic macrophages play a major role8. Kupffer cells (KCs), a type of hepatic macrophages, produce pro-inflammatory factors, such as tumor necrosis factor-alpha (TNF-α), interleukin (IL)-1β, and MIP-1α, that promote the activation of hepatic stellate cells (HSCs)9, 10. Previous studies have shown that activated HSCs are the key effector cells that cause an abnormal increase in ECM deposition in liver fibrosis11, 12. The pathogenesis of liver fibrosis involves cell damage and the release of inflammatory factors that prompt the activation and transformation of resting HSCs into astrocyte-like cells13–15. Moreover, HSCs have higher proliferation rates and release inflammatory and pro-proliferative factors. Notably, during the recovery of liver fibrosis, the apoptosis of HSCs increases significantly16, 17. Tissue inhibitor of metalloproteinase-1 (TIMP-1) plays an important role in the pathogenesis of liver fibrosis, and its expression in HSCs is enhanced by TNF-α18, 19. Some studies have shown that after CCL4 treatment, liver fibrosis in transgenic mice overexpressing TIMP-1 in the liver increased20. Until now, liver biopsy is the most common and effective method to identify liver fibrosis and cirrhosis21. However, liver biopsy is characterized by high consumption, low patient acceptance, increased risk of complications, and low
replicability21. Therefore, the molecular mechanism of liver fibrosis has been studied extensively to find a safer and more feasible detection method. Fortunately, significant progress has been made in this regard. For example, IL-33 was identified as an activator of HSCs that induce hepatic fibrosis, inhibition of the NF-κB and MAPK signaling pathways improves liver fibrosis, and antiplatelet drugs prevent liver fibrosis.22–24 LPS is a known endotoxin found in the outer membrane of many intestinal gram-negative bacteria25. Previous studies have shown that LPS induces the expression of Toll-like receptor 4 (TLR4) and myeloid differentiation factor-2 (MD-2, co-receptor of TLR4) in KCs and produces TNF-α in nonalcoholic liver disease26, 27. Moreover, the TLR4/MD-2–TNF-α signaling pathway that mediates the activation of KCs is induced by LPS28, 29. However, it is not known whether alcohol-induced alcoholic liver disease is linked to the LPS-TLR4/MD-2–TNF-α signaling pathway. Therefore, this study aimed to investigate the effect of ethanol on liver disease via the modulation of the LPS-TLR4/MD-2–TNF-α signaling pathway.

Materials and Methods

Animals and treatment

Twenty-four male Wistar rats, weighing 200 ± 20 g, were purchased from Beijing Vital River Laboratory Animal Technology Co., Ltd. (Beijing, China) This study was performed according to the experimental protocol approved by the Ethics Committee of Heilongjiang Province Hospital. After 1 week of adaptation to general meals, the rats were randomly divided into four groups (n = 6 rats in each group): the control group (Ctrl) and the ethanol groups (EtOH). The rats in the EtOH groups were subjected to intragastric administration of ethanol solution (40% v/v, dissolved in 40 g anhydrous ethanol in distilled water to prepare 100 mL; 10 mL/kg/d, Aladdin), once a day for 12 weeks. In this study, the method of establishing a rat alcoholic liver injury model was improved on the basis of existing methods.30. The two groups of ethanol-fed rats were injected with anti-TLR4 (ab217274, dilution 1/300, 0.2 mg/kg, once per week, Abcam, Cambridge, UK) and anti-MD-2 antibodies (ab24182, dilution 0.5 µg/mL, 0.2 mg/kg, once per week, Abcam Cambridge, UK) via the caudal vein to verify whether TLR4/MD-2 plays a role in alcoholic liver fibrosis from the fifth week until the end of the experiment, which lasted for 8 weeks. The antibody dilution was recommended by the manufacturer, and the dosage was the optimal dosage selected after the pre-experiment. Meanwhile, the rats in the control group were administered with saline (NaCl, Merck, Shanghai, China) by gavage once a day for 12 weeks. The rats were intraperitoneally injected with 4% pentobarbital (Merck, 30 mg/kg) and killed by cervical dislocation. The plasma samples were collected and stored at −80°C for further analysis. The liver tissue was quickly removed from the abdominal cavity and divided into three equal parts. One part was fixed in 10% formalin at room temperature (RT) for 24 h for histopathological examination, the other part was used for the separation of KCs and HSCs and the detection of cytokines, and the last part was rapidly frozen in liquid nitrogen and stored at −80°C.

Cell isolation

KCs were isolated from the rat liver through collagenase digestion in vitro and differential centrifugation.31, 32 First, 12 male SPF-Wistar rats were anesthetized through the intraperitoneal injection of pentobarbital sodium (50 mg/kg). After abdominal incision, 500 mL PBS buffer solution (37°C, 30 mL/min) was infused through the hepatic portal vein to remove the red blood cells, and then the liver was removed. Next, the minced liver was digested in MDEM with 0.1% type IV collagenase in a 37°C water bath for 30 min, and the liver homogenate was filtered through a 200-gauge mesh. The filtrate was centrifuged at 50 × g for 2 min to collect the suspension of KCs and HSCs. The suspension was recentrifuged at 500 × g for 7 min at 4°C to collect the cell pellet. The cell precipitate was added to a mixture of Nycodenz and PBS and centrifuged at 1,500 × g for 17 min at 4°C. HSCs were collected from the interface and washed three times with PBS. These were cultured in MDEM with 10% FBS and 100 U/mL penicillin/streptomycin at 37°C and 5% CO₂. The culture medium was changed every 24 h.

After the removal of HSCs, the remaining part of the sample was washed three times with PBS and centrifuged to purify the KCs. The cells were seeded in a 16-well plate and cultured in serum-free DMEM. After 2 h, the nonadherent cells were removed by washing with PBS. The purified KCs were cultured in MDEM with 10% FBS and 100 U/mL penicillin/streptomycin at 37°C and 5% CO₂. The culture medium was changed every 24 h.

Finally, KCs and HSCs were identified using cell markers and flow cytometry, and the trypan blue exclusion assay and the ink particle phagocytosis test were used to estimate the viability and phagocytic activities of the KCs and HSCs, respectively. Antibodies for flow cytometry included the anti-CD68 antibody (EPR23917-164] (ab283654, dilution 1/500, Abcam, Cambridge, UK) and anti-α-SMA antibody (14A) (ab7817, dilution 1.137 µg/mL, Abcam, Cambridge, UK).

Co-culturing of the cells

After 1 day of culture, KCs were divided into four groups. In the control group, KCs were cultured in DMEM supplemented with 10% FBS and normal saline. In the LPS group, the KCs were stimulated with LPS (100 ng/mL) for 6 h. In the LPS + si-TLR4 and LPS + si-MD-2 groups, the KCs were stimulated with LPS and transfected with the siRNA plasmids siTLR4/si-MD-2 using the Lipofectamine 2000 reagent (Invitrogen, Barcelona, Spain). The plasmids were synthesized commercially by Sangon. After 2 days, each group was co-cultured with HSCs in the Boyden chamber and the cell culture inserts, polyethylene terephthalate membrane (pore size 8 µm). The HSCs (1.5 × 10⁵ cells) were plated in the bottom plates and cultured in DMEM with 10% FBS. After 1 day, the medium was removed. KCs (4.5 × 10⁵ cells)
cells) were plated on cell culture inserts and transferred onto the HSCs. Empty inserts were used as co-culture controls. Next, 4 mL EMDM without serum was added to the chamber. After 2 days, the HSCs in the bottom plates were used for the MTT assay.

Liver histopathology

The rat liver tissue was fixed with 10% formalin and cut into 4-μm-thick slices using the paraffin slice technique. To observe the histological changes in liver injury, one part of the section was stained with hematoxylin and eosin (H&E), whereas the other part was used to investigate hepatic collagen deposition through Masson’s trichrome staining. The degree of liver injury was estimated based on the infiltration of inflammatory cells under a light microscope (Olympus, Tokyo, Japan).

Western blotting

Western blotting was used to detect the protein levels linked to hepatic fibrosis, including collagen I, α-SMA, TIMP-1, TLR4, MD-2, TNF-α, IL-1β, and MIP-1α. The total protein fraction was extracted from the liver homogenate or KC cell lysate using the Total Protein Extraction Kit (BestBio, Shanghai, China). The protein concentrations of the samples were estimated using the BCA Protein Assay Kit (Haigen Gene, Harbin, China). Protein samples (10 µg each) were separated through electrophoresis using 10% SDS-PAGE and transferred onto polyvinylidene difluoride membranes. After blocking with 5% nonfat milk at 25°C for 1 h, the membranes were incubated overnight with primary antibodies against collagen I (ab34710, dilution 1/10,000, Abcam), α-SMA (#149685, dilution 1/1,000, Cell Signaling Technology, Shanghai, China), TIMP-1 (ab211926, dilution 1/1,000), TLR4 (ab13556, dilution 1/500), MD-2 (ab241182, dilution 1 µg/mL), TNF-α (ab215188, dilution 1/1,000), IL-1β (ab2105, dilution 1/2,000), and MIP-1α (sc-365691, dilution 1:800, Santa Cruz Biotechnology, Shanghai, China) at 4°C. Subsequently, the membranes were incubated with secondary antibodies anti-goat anti-rabbit IgG H&L (HRP) (ab205718, dilution 1/50,000) or goat anti-mouse IgG H&L (HRP) (ab6789, dilution 1/10,000) for 1 h at 37°C. The protein bands were visualized with the ECL Western blotting kit (Thermo Fisher Scientific, Waltham, MA, USA) and immediately exposed to autoradiography film. β-A CTin was used as an internal control.

ELISA

ELISA was used to estimate the levels of hydroxyproline and alanine aminotransferase (ALT), which are markers of liver function. The assay was performed according to the instructions of the Hydroxyproline ELISA Kit (ASB-OKEH025, ENZO, Shanghai, China) and ALT ELISA Kit (ab234579, Abcam). Standards/samples (50 µL/well) were added to the appropriate wells of 96-well plates. Subsequently, a 50 µL antibody cocktail was added to all wells and co-cultured for 1 h at 25°C. Next, all wells were washed three times with 350 µL of 1X wash buffer. Next, 100 µL TMB development solution was added to each well and incubated for 10 min. The reaction was stopped with 100 µL stop solution. Finally, the optical density was recorded at 450 nm using a microplate reader.

Reverse transcription quantitative polymerase chain reaction

Reverse transcription quantitative polymerase chain reaction (RT-qPCR) was used to estimate the expression of TLR4, MD-2, and TNF-α in liver tissue and CD68, TNF-α, IL-1β, and MIP-1α in KCs. Liver tissue (100 mg) was ground with liquid nitrogen, and total RNA was extracted from the homogenate using TRIzol (Invitrogen). Next, the total RNA was reverse-transcribed to cDNA using SuperScript™ II Reverse Transcriptase (Invitrogen) according to the manufacturer’s instructions. Then, a 20 µL PCR reaction system including cDNA, primers, and PCR reaction solution was set up for RT-qPCR with QuantStudio™ 5 Real-Time PCR System (Thermo Fisher Scientific, Shanghai, China) and PowerTrack™ SYBR Green Master Mix (Thermo Fisher Scientific, Shanghai, China). RT-qPCR reaction conditions were as follows: 95°C for 8 min, 95°C for 4 s (45 cycles), 65°C for 6 s, and 72°C for 1 min. GAPDH was used as an internal control, and the 2−ΔΔCt method was used to estimate the relative expression levels. Each experiment was repeated thrice. The primer sequences used in RT-qPCR are listed in Table 1.

MTT 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide assay

The viability of HSCs was determined using the MTT assay. Briefly, 2 × 10⁵ cells/mL was inoculated into a 96-well plate. Then, 100 µL MTT or control (dimethyl sulfoxide, DMSO) and 900 µL DMEM with 10% FBS and 100 U/mL penicillin/streptomycin were added to each well plate. Then, 100 µL MTT or control (dimethyl sulfoxide, DMSO) and 900 µL DMEM with 10% FBS and 100 U/mL penicillin/streptomycin were added to each well plate. After 2 days, the HSCs in the bottom plates were used for the MTT assay.

| Gene    | Forward                     | Reverse                     |
|---------|------------------------------|-----------------------------|
| TLR4    | 5'-AGGATGGAGCAGCTGGGTAAGGA-3'| 5'-CTGGAGTGAAGTGCGTGGGAC-3'|
| MD-2    | 5'-GAACGAGTTTTGGTCTGCAA-3'  | 5'-TTGAGGACATTCTGCTGGCA-3'  |
| TNF-α   | 5'-CCAGAGGCCAGGGACCTCC-3'   | 5'-GCTAACACGCTGGGTTACC-3'  |
| CD68    | 5'-GCTTGTACATCTCCCTGTG-3'   | 5'-TGGATCCGGGTCTCTAACC-3'  |
| IL-1β   | 5'-GACGATTCTACGAGCACGAC-3'  | 5'-GACCGGAAACCTCATGGAG-3'  |
| MIP-1α  | 5'-GCTGACCCCAAGGACCTG-3'    | 5'-CCCTGGATGCTGGATTTGG-3'  |
| GADPH   | 5'-GACCGTTCAAGGCTGAAGC-3'   | 5'-ATGGGTGGTGGACGCAGG-3'   |
were allowed to grow for 3 h at 37°C. MTT was reduced to blue crystals by metabolically active cells, which were dissolved in 100 µL DMSO. Finally, absorbance was measured at 570 nm using a UV–Vis spectrophotometer. The GraphPad Prism 7.0, GraphPad Software, San Diego, CA, USA program was used to calculate the viability of KCs.

Immunofluorescence studies

An immunofluorescence assay was used to observe the expression of CD68, a marker of KCs in the rat liver. First, the liver sections were infiltrated with PBS containing 0.5% Triton X-100. Next, the sections were incubated with 10% normal goat serum at RT. After 1 h, the sections were incubated overnight with mouse anti-CD68 antibody (Abcam, ab201340, dilution 1–2 µg/mL) at 4°C. Later, the cells were washed thrice with PBS and incubated with the goat anti-mouse IgG Alexa 488 conjugated fluorescence secondary antibody (Invitrogen, R37120, dilution 1/1,000) for 1 h at RT. Finally, the sections were stained with 4',6-diamidino-2-phenylindole (DAPI) and observed under a confocal microscope. The DAPI1 group was used as a control group, and the PBS replaced the primary antibody as a negative control.

Statistical analysis

All data are presented as the mean ± standard deviation. Statistical analysis was performed using GraphPad Prism software (version 7.0). One-way ANOVA followed by Sidak’s multiple comparisons test or Student’s t-test was used to perform statistical evaluation. Statistical significance was set at p<0.05. Each experiment was performed in triplicates.

Results

Alcohol administration induced liver fibrosis and activated TLR4/MD-2–TNF-α pathway in the model group of rats

To establish a model of hepatic fibrosis, rats were subjected to the intragastric administration of alcohol. Liver tissue was sectioned and used for histological analysis (H&E and Masson). Meanwhile, using the liver homogenate, Western blot, ELISA, and RT-qPCR assays were performed to determine the levels of molecules associated with liver fibrosis. In the control group, the structure of hepatic lobules was clear, and the hepatocytes were orderly arranged (Fig. 1A). In the model group, a large number of collagen fibers and pseudolobules in the portal area were observed (Fig. 1B). Meanwhile, Western blotting analysis showed that the levels of proteins related to liver fibrosis, such as collagen I, α-SMA, and TIMP-1<sup>34</sup>, were higher in the model group than that in the control group (Fig. 1C). Similarly, ELISA revealed that hydroxyproline (Fig. 1D) and ALT (Fig. 1E) were upregulated in the model group. Importantly, RT-qPCR and Western blotting analyses revealed that TLR4, MD-2, and TNF-α were significantly increased in the model group compared to that in the control group (Fig. 1F–G). Overall, these results indicated liver fibrosis and activation of the TLR4/MD-2–TNF-α pathway in the model group of rats.

Inhibition of TLR4/MD-2 alleviated partially alcohol-induced liver fibrosis

To substantiate the regulatory effect of TLR4/MD-2 in alcoholic liver fibrosis, the rats were injected with antibodies against TLR4/MD-2 to inhibit their expression. The results of RT-qPCR and Western blot assays showed that inhibition of TLR4 or MD-2 could partially reduce the levels of TLR4, MD-2, and TNF-α induced by alcohol (Fig. 2A–B). In addition, the antibodies reduced the expression of TLR4 and MD-2 by 53% and 33%, respectively (Fig. 2B). Moreover, compared to that in the control group, HE staining of liver tissue indicated that the liver was severely damaged with a large number of pseudolobules in the EtOH group (Fig. 2C). Interestingly, inhibition of TLR4 or MD-2 partially alleviated alcohol-induced liver damage in rats (Fig. 2C). In addition, Masson staining showed that a large number of collagen fibers were deposited around the central vein, hepatocytes, and portal area, the portal area was expanded, and some pseudolobules were surrounded by fibrosis to form a fibrous septum in the EtOH group (Fig. 2D). In the anti-TLR4 and anti-MD-2 groups, collagen fiber deposition slightly decreased, the area of portal area had a slight reduction, and the fibrous septum was narrowed compared with that in the EtOH group (Fig. 2D). Next, the levels of proteins linked to liver fibrosis were detected, and it was found that TLR4 or MD-2 inhibition could also partially reverse the upregulation of collagen I, α-SMA, and TIMP-1 induced by alcohol (Fig. 2E). ELISA results also indicated that the upregulation of hydroxyproline and ALT induced by alcohol was partially reduced by inhibiting TLR4 or MD-2 (Fig. 2F–G). The above results suggest that inhibition of TLR4 (53%) and MD-2 (33%) could partially alleviate alcohol-induced liver fibrosis in the model group of rats.

Inhibition of TLR4/MD-2 could partially reduce the activation of KCs in liver fibrosis

Next, using immunofluorescence, the levels of CD68, a biomarker of KCs in the rat liver, were determined. The staining results showed that after alcohol treatment, CD68 was highly expressed in the portal area. However, inhibition of TLR4 and MD-2 partially reduced the levels of CD68 (Fig. 3A). Western blotting analysis also revealed that alcohol-induced overexpression of CD68 and the inflammatory...
Fig. 1. Alcohol treatment induced liver fibrosis and activated the TLR4/MD-2–TNF-α pathway in model rats.
In the alcohol-damaged liver of model rats, the molecules associated with liver fibrosis were upregulated, and the signaling pathway of
liver fibrosis was activated. (A) Hematoxylin and eosin staining and (B) Masson staining were used to detect liver injury. The arrows
indicate the portal area. Compared with that in the control group, the livers of rats in the EtOH group showed obvious pathological damage.
(C) The levels of proteins related to liver fibrosis were detected by Western blotting, and data quantification was performed using ImageJ
software; compared with that in the control group, the levels of proteins related to liver fibrosis in the EtOH group showed an obvious
increase. (D–E) ELISA was used to determine the collagen-related markers (D) hydroxyproline and (E) ALT. The results showed that
hydroxyproline and ALT levels significantly increased in the EtOH group. (F–G) The levels of TLR4, MD-2, and TNF-α were detected
by (F) Western blotting and (G) RT-qPCR, and quantitation was performed using ImageJ software. The expression levels of TLR4, MD-
2, and TNF-α were significantly increased in the EtOH group. The scale bar is 20 μm at high power and 80 μm at low power. **p<0.01,
***p<0.001, vs. the control group.
factors IL-1β and MIP-1α was partially reversed upon treatment with antibodies against TLR4 and MD-2 (Fig. 3B). Furthermore, the results of the RT-qPCR test were also consistent with the Western blotting findings, suggesting regulation at the mRNA level (Fig. 3C). Overall, based on CD68 levels, the above results suggest that inhibition of TLR4 and MD2 could partially alleviate the activation of KCs in the liver of fibrotic rats.

Knockdown of TLR4/MD-2 in KCs could partially reduce the activity of LPS-induced HSCs

KCs were isolated and cultured as described in the Methods section. Next, RT-qPCR and Western blotting were used to estimate the levels of TLR4, MD-2, and TNF-α. The results indicated that LPS upregulated the levels of TLR4, MD-2, and TNF-α. However, the increase was partially reversed upon treatment with si-TLR4 and si-MD-2 (Fig. 4A–B). In addition, siRNA reduced the expression of
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TLR4 and MD-2 by 34% and 37%, respectively (Fig. 4B). When the levels of CD68 and pro-inflammatory factors IL-1β and MIP-1α were estimated using RT-qPCR, it was found that LPS-induced increase in CD68, IL-1β, and MIP-1α was also partially reduced by si-TLR4/si-MD-2 treatment (Fig. 4D). Similar results were obtained through Western blot analysis (Fig. 4D). HSCs are the key effector cells of liver fibrosis. Therefore, we next studied the effect of TLR4 and MD-2 knockdown in KCs on the activity of HSCs. For this purpose, KCs and HSCs were co-cultured. The MTT assay revealed that LPS overactivated HSCs, whereas the inhibition of TLR4 and MD-2 partially restrained this overactivation (Fig. 4E). Next, we estimated the levels of α-SMA and desmin, a biomarker of HSCs, through Western blotting. The results showed that LPS-induced increase in α-SMA and desmin levels was partially inhibited by the knockdown of TLR4 and MD-2 (Fig. 4F). In brief, in vitro knockdown of TLR4 and MD-2 in KCs partially inhibited the activation of KCs and reduced the expression of pro-inflammatory factors, suggesting reduced activity of HSCs.

Discussion

This study established a rat model of hepatic fibrosis by treating rats with alcohol, which activated the TLR4/MD-2–TNF-α pathway. In a further study, it was found that inhibition of TLR4 and MD-2 partially decreased the activity of KCs and HSCs and partially relieved hepatic fibrosis in the model rats. The in vitro experiments indicated that LPS-induced activation of the TLR4/MD–TNF-α signaling pathway was partially inhibited by the knockdown of TLR4 and MD-2 in KCs. In addition, the release of pro-inflammatory factors and the activity of HSCs partially decreased. In other words, inhibition of TLR4 and MD-2 could partially alleviate alcohol-induced liver fibrosis in rats.

TLR4, a pattern recognition receptor, is mainly expressed in macrophages. It is linked to macrophage activation and the release of inflammatory factors. MD-2 is a helper protein of TLR4, which assists TLR4 in recognizing LPS and interferes with LPS-induced inflammation. A previous study in mice showed that monoclonal antibodies against TLR4/MD-2 could reduce LPS-induced acute hepatitis by inhibiting TNF-α. Similarly, deficiency of TLR4
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and MD-2 was reported to alleviate nonalcoholic liver fibrosis in mice. Yin et al. reported that blocking the expression of TLR4 and MD-2 improved LPS-induced liver failure. This evidence suggests that nonalcoholic liver disease is caused by LPS through the TLR4/MD-2–TNF-α signaling pathway. To explore whether alcoholic liver disease is also mediated by this pathway, a rat liver fibrosis model was established through the intragastric administration of alcohol, and in vitro studies in KCs were performed.

Earlier studies suggested that long-term drinking upregulates TLR4 in the liver and makes the organ more sensitive to LPS through the overexpression of TNF-α and IL-6. Our study found that TLR4/MD-2 levels in the rat liver were increased after the intragastric administration of alcohol, along with the expression of TNF-α. Notably, the inhibition of TLR4 reduced the secretion of TNF-α and IL-6, thus reducing alcohol-induced liver injury. Our study also showed that upon TLR4/MD-2 inhibition, the degree of alcohol-induced liver injury was partially reduced, and the expression of proteins related to liver fibrosis was partially decreased. Akashi-Takamura et al. showed that mAb Sa15-21 protects rats from LPS/D-GalN-induced liver failure by activating NF-κB protective signaling through the TLR4/MD-2 pathway and influencing TNF-α-induced hepatocyte apoptosis. Our study showed that the inhibition of TLR4/MD-2 could inhibit the expression of TNF-α to reduce cell apoptosis, thus partially alleviating alcoholic liver injury in rats. Meanwhile, the mRNA levels of TLR4 and MD-2 were also partially reduced by the TLR4 and MD-2 antibodies. We speculate that this phenomenon may be related to a negative feedback regulation mechanism, that is, TLR4/MD-2 reduction inhibits the NF-κB signaling pathway and activates the β-catenin pathway, whereas the activation of the β-catenin pathway inhibits the TLR4/MD-2 pathway.

In the future, studies on the specific mechanisms may be needed. Besides, numerous evidences suggest that TLR4 induces activation of HSCs, which is an important transformation step for liver inflammation into liver fibrosis. Alcoholic-induced TLR4 expression activates HSCs via the TLR4/MD-2–TNF-α signaling pathway and promotes chemokine secretion. This increases the recruitment of KCs to the liver injury site, which releases pro-inflammatory factors that cause liver inflammation. Our research indicates that KCs are activated by the alcohol-induced TLR4/MD-2–TNF-α signaling pathway, which promotes pro-inflammatory factors. Notably, this phenomenon was partially suppressed by inhibiting TLR4 and MD-2. Moreover, in vitro experiments showed that inhibition of TLR4 and MD-2 in KCs partially
reduced the activation of KCs and HSCs and partially suppressed the LPS-induced release of pro-inflammatory factors. It has been found that ethanol can affect the integrity and permeability of the intestinal mucosa and destroy the barrier structure. Therefore, we infer that ethanol increases the possibility of LPS translocation from rat intestine, and LPS transported from the intestine of rats enters the systemic circulation through the gastrointestinal lymphatic vessels and portal vein and then participates in the production and development of liver inflammation. In addition, Akashi-Takamura et al. also proved that the monoclonal antibody sa15-12 of TLR4/MD-2 can alleviate the activation of KCs and inhibit hepatocyte apoptosis. The antibodies used in our study have been shown to alleviate KC activation. However, whether these factors affect hepatocyte apoptosis needs to be confirmed in subsequent studies.

It is well known that reactive oxygen species (ROS) play an important role in ethanol metabolism in the liver. The Nrf2/ARE signaling pathway is a hot topic in the field of liver fibrosis. Certain cell stimulations from ROS combine the nuclear transcription related factor-2 (Nrf2) with the antioxidant response element (ARE) promoter sequence and activates γ-glutamylcysteine ligase (γ-GCS). The synergistic effect of Nrf2 and γ-GCS is conducive to the body against free radicals, avoids hepatocyte damage, and then slows down the process of liver fibrosis. Studies have shown that the Wnt/β-catenin signaling pathway is involved in the formation of alcoholic liver fibrosis, which activates...
and proliferates HSCs, resulting in liver fibrosis. It is possible that the activation of HSCs can be inhibited and the expression level of various liver fibrosis indices can be reduced by inhibiting the Wnt/β-catenin signaling pathway to promote the recovery of liver fibrosis. A multipronged approach may completely alleviate liver injury caused by alcohol, which may be our future research direction.

In conclusion, alcohol activates KCs through the LPS-TLR4/MD-2–TNF-α signaling pathway, and KCs release inflammatory factors to activate HSCs, which led to liver fibrosis in model rats. However, inhibition of the LPS-TLR4/MD-2–TNF-α signaling pathway partially relieved liver fibrosis in model rats. Overall, this study provides a scientific basis for the prevention and treatment of alcoholic liver disease by suggesting new therapeutic targets.

Disclosure of Potential Conflicts of Interest: None.

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