Inhibition of heat shock protein 90 alleviates cholestatic liver injury by decreasing IL-1β and IL-18 expression

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Abstract. Severe cholestatic liver injury diseases, such as obstructive jaundice and the subsequent acute obstructive cholangitis, are induced by biliary tract occlusion. Heat shock protein 90 (HSP90) inhibitors have been demonstrated to be protective for various organs. The potential of HSP90 inhibitors in the treatment of cholestatic liver injury, however, remains unclear. In the present study, rat models of bile duct ligation (BDL) were established, the HSP90 inhibitor 17-dimethylamino-ethylamino-17-demethoxygeldanamycin (17-DMAG) was administered, and its ability to ameliorate the cholestasis-induced liver injuries was evaluated. In the BDL rat models and clinical samples, increased HSP90 expression was observed to be associated with cholestatic liver injury. Furthermore, 17-DMAG alleviated cholestasis-induced liver injury in the rat models, as revealed by the assessment of pathological changes and liver function. In addition, 17-DMAG protected hepatocytes against cholestatic injury in vitro. Further assays indicated that 17-DMAG administration prevented cholestasis-induced liver injury in the rats by decreasing the expression of interleukin (IL)-1β and IL-18. Moreover, 17-DMAG also decreased the cholestasis-induced upregulation of IL-1β and IL-18 in liver sinusoidal endothelial cells in vitro. In conclusion, the HSP90 inhibitor 17-DMAG is able to prevent liver injury in rats with biliary obstruction, and this phenomenon is associated with the reduction of IL-1β and IL-18 expression.

Introduction

Obstructive jaundice is a frequently observed condition caused by biliary tract occlusion, which inevitably causes liver injury (1). In biliary obstruction, secondary gram-negative pathogen infection may occur, causing acute obstructive cholangitis, which progresses rapidly and has a poor prognosis (2,3). The liver is the first organ to become involved and is severely injured during biliary obstruction. An important factor affecting prognosis is the extent of liver injury. Therefore, the identification of effective methods to alleviate cholestasis-induced liver injury can improve the prognosis of this disease.

Heat shock proteins (HSPs) are highly conserved proteins that are induced by a wide variety of physiological and environmental insults, such as heat, inflammation, toxic chemicals and oxidative stress. HSP90 is an essential member of the HSP family of chaperone proteins, and is upregulated in various conditions, including inflammation, organ injury and cancer (4-7). According to its subcellular localization, HSP90 comprises cytosolic HSP90 (HSP90AA1), endoplasmic reticulum-localized chaperone HSP90 (HSP90B1, also known as gp96 or grp94) and the mitochondrial member TRAP1 (HSP90L) (8). The potential therapeutic effects of HSP90 inhibitors, such as 17-dimethylamino-ethylamino-17-demethoxygeldanamycin (17-DMAG), have been the focus of a number of studies due to the important roles of HSP90 (9-11). Although studies have shown that 17-DMAG effectively prevents lipopolysaccharide (LPS)- or alcohol-induced liver injury (11,12), the applicability of HSP90 inhibitors to biliary obstruction-induced liver injury remains unclear.

Toll-like receptors (TLRs) are a family of pattern recognition receptors for pathogens with critical roles in innate immunity (13). TLR9 is an essential TLR and regulates pro-inflammatory cytokines, such as interleukin (IL)-1β and IL-18 (14,15). In addition, HSP90B1 is required for the folding of TLR9 and is involved in the biogenesis of TLR9 (16,17).
Liver tissues were snap-frozen in liquid nitrogen or fixed with (1,000 x g for 5 min at 4˚C) and the serum was stored at -80˚C. Prior to analysis, blood samples were centrifuged and liver tissue samples were harvested from the euthanized animals. Prior to analysis, blood samples were centrifuged (1,000 x g for 10 min at 4˚C) and the serum was stored at -80˚C.

In the present study, we hypothesized that HSP90 inhibition may reduce liver injury during biliary obstruction. Therefore, 17-DMAG was used to evaluate the effects of HSP90 inhibition in an animal model of biliary obstruction.

Materials and methods

Animal model. Wistar rats weighing 300-350 g and aged 12 weeks were purchased from Shanghai SLAC Laboratory Animal Co., Ltd. All procedures were approved by the Ethics Committee of Shaoxing People's Hospital and conformed to the Guide for the Care and Use of Laboratory Animals published by the US National Institutes of Health (revised 2011). The rats had ad libitum access to food and water and were maintained at 20˚C, with 50% humidity under 12-h light/dark cycles.

Rats were randomly grouped into sham, bile duct ligation (BDL) and BDL + lipopolysaccharide (LPS) groups. The animals were anesthetized with an intraperitoneal injection of 50 mg/kg pentobarbital. In the BDL and BDL + LPS groups, the distal common bile ducts were dissociated and ligated with 6-0 silk, and PE-10 polyethylene catheters long enough to reach the skin were inserted into the proximal bile ducts as described in our previous study (18,19). Intra-bile duct infusion was performed by injecting 0.2 ml saline or LPS (2 mg/ml, purified from Escherichia coli O111:B4; Sigma-Aldrich; Merck KGaA) into the proximal bile ducts through the catheter. The catheter was sealed with a sealing cap and the abdominal cavity was covered using silk sutures following the introduction of 0.1 ml air. The rats in the sham group underwent a sham procedure, which involved dissecting bile duct carefully without injury and closing the abdominal cavity. The rats in the sham, BDL, and BDL + LPS groups were further divided into three groups according to the intraperitoneal injections they received after surgery. Briefly, in each group, one-third of the rats were intraperitoneally injected with normal saline (NS), another one-third were intraperitoneally injected with 2 mg/kg 17-DMAG (MedChemExpress) and the remaining one-third were intraperitoneally injected with 5 mg/kg 17-DMAG. The intraperitoneal injections were performed daily. A total of 108 rats were randomly classified into the aforementioned three groups according to the surgical procedure (n=36/group). Each of these groups comprised rats treated with NS, 2 mg/kg 17-DMAG and 5 mg/kg 17-DMAG (n=12/treatment), respectively. At 24 and 72 h after the surgery, a fraction (n=6/group/time point) of the rats in each group were euthanized. Caspase-3 activity detection and pathology experiments, including tissue staining and immunohistochemistry, were performed on samples taken at 72 h. Other experiments were performed on samples taken at 24 h. The rats were sacrificed via cervical dislocation under 2% sevoflurane inhalation anesthesia. The death of the animals was confirmed by the cessation of vital signs. Blood and liver tissue samples were harvested from the euthanized animals. Prior to analysis, blood samples were centrifuged (1,000 x g for 5 min at 4˚C) and the serum was stored at -80˚C. Liver tissues were snap-frozen in liquid nitrogen or fixed with 10% formalin.

Clinical tissue samples. Clinical tissue samples were obtained from patients who underwent hepatectomy at Shaoxing People's Hospital (Shaoxing, China) from January 2014 to December 2017. Normal liver tissues (n=5; age range, 42-63 years; inclusion criteria: Definite pathological diagnosis of hemangioma; exclusion criteria: Liver cirrhosis, fatty liver, hepatitis virus positive, liver tumor and hepatolithiasis) were obtained from patients who underwent hepatectomy to treat liver hemangioma; non-hemangioma tissues were selected for pathological examination. Intrahepatic cholangitic liver tissues (n=5; age range: 45-69 years; inclusion criteria: Definite pathological diagnosis of hepatolithiasis; exclusion criteria: Liver cirrhosis, fatty liver, hepatitis virus positive and liver tumor, such as intrahepatic cholangiocarcinoma) were obtained from patients who underwent hepatectomy to treat intrahepatic stones and cholangitis. There were two male patients and three female patients in the two groups, respectively. All participants provided written informed consent, and Shaoxing People's Hospital Institutional Review Board approved the tissue acquisition protocol.

Evaluation of liver function. An Automated Chemical Analyzer (Dimension RxL Max HM; Siemens AG) was used to quantify the levels of aspartate aminotransferase (AST), alanine aminotransferase (ALT) and total bilirubin (TBIL) in rat serum to determine the degree of liver injury.

RNA extraction and reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total RNA was isolated from liver tissue samples or cells using TRIzol (Invitrogen; Thermo Fisher Scientific, Inc.) and reverse transcribed into cDNA using a PrimeScript™ RT reagent Kit (cat. no. RR047A; Takara Bio, Inc.), according to the manufacturer's protocols. Briefly, RT was conducted at 37˚C for 15 min and 85˚C for 5 sec, and the cDNA was stored at 4˚C until further use. qPCR was performed using SYBR® Premix Ex Taq™ II (cat. no. RR820A; Takara Bio, Inc.) and an ABI 7500 Real-time PCR system (Applied Biosystems; Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. The primer sequences used are shown in Table I. β-actin was used as an endogenous control. The qPCR conditions comprised 95˚C for 30 sec, and a total of 40 cycles at 95˚C for 5 sec and extension at 60˚C for 34 sec, followed by a dissociation curve analysis. All assays were performed three times. Relative expression levels were then determined using the 2^ΔΔCq method (20).

Caspase-3 activity detection. A Caspase-3 Activity Assay Kit (cat. no. C1116; Beyotime Institute of Biotechnology) was used according to the manufacturer's protocol to detect the activity of caspase-3 in the rat liver tissues. The absorbance was measured at 405 nm and the relative caspase-3 activity to the activity of the sham group was determined. Six samples from each group were analyzed.

Enzyme-linked immunosorbent assay (ELISA). The liver tissues were weighed and homogenized immediately in 10 volumes of saline after washing in saline. Supernatants were collected to perform ELISAs following centrifugation (12,000 x g for 10 min at 4˚C). ELISA kits (cat. nos. BMS630 and KRC2341, eBioscience; Thermo Fisher Scientific, Inc.)
β-actin were harvested from the rats and prepared using the aforementioned procedure. After 1 week, 'cholestatic culture' assay.

Scope (magnification, x200; Leica Microsystems, Inc.)

were used according to the manufacturer's instructions to measure the content of IL-1β and IL-18 in the liver samples. Each sample was tested in triplicate.

Tissue staining. Liver tissue samples obtained from the rats were fixed with 10% formalin. The formalin-fixed specimens (3 μm) were stained using automatic H&E slide stainer (Leica STS020 Multistainer; Leica Microsystems, Inc.) according to the manufacturer's instructions for histological evaluation. The stained liver tissues were examined for histopathological evidence of pathological damage under light microscope (magnification, x100).

Immunohistochemistry (IHC). HSP90 expression in the rat liver tissues was detected by IHC. Deparaffinization and rehydration of the sections (3 μm thickness) were performed with gradient xylene and a descending ethanol gradient. Antigen retrieval was performed with Citrate Antigen Retrieval solution (1:50; cat. no. P0083; Beyotime Institute of Biotechnology) for 10 min at room temperature. Each stained sample was washed with washing buffer (cat. no. P0106L; Beyotime Institute of Biotechnology) for 1.5 h at room temperature. After thoroughly washing with washing buffer (cat. no. P0106L; Beyotime Institute of Biotechnology), the sections were developed using a 3,3'-diaminobenzidine kit (cat. no. P0202; Beyotime Institute of Biotechnology) and counterstained with hematoxylin staining solution (cat. no. C0107; Beyotime Institute of Biotechnology) for 10 min at room temperature. Each stained sample was observed under high power magnification using a light microscope (magnification, x200; Leica Microsystems, Inc.).

Gene Forward (5'-3') Reverse (5'-3')

| Gene       | Forward (5'-3') | Reverse (5'-3') |
|------------|----------------|----------------|
| β-actin    | ACACCGGCCCAACGATTCG | CCCACGATGGAGGGGAAGA |
| Hsp90a1    | TCAGGCGAAGTTCCGAGT | ATCCAGAGGCTGTCGGAGT |
| Hsp90b1    | ACCGAGAAGACTGACGACT | GCTCTCAAAACCCGAAGGT |
| Trap1      | TGCCACCGCAACATATTT | CGTAGCAGAAGAGCACCCTA |
| Hspa1b     | CTCCTTTGCGTCTGTCGAA | TGCAAAAGACACATCTCAGG |
| Hspa4      | TCAGAGCTTGCTATGTCGTA | GGCATTGGAATACCTGCTGTC |
| Bax        | GGGATCGGTCTCTTCTAC | TTTCCTGGTCTCCTCAT |
| Bcl2       | AGCTTGACCTCTGTTGTA | TCACCTGGGCCCCAGTGAT |
| IL-1β      | CAGCTTTCGACAGTGAGGAGA | TTTCGAGATGTCTGCTGTA |
| IL-18      | ACCGCAGATACCGAGCAT | CTGCGAGATGTCTGCTGTA |

Hsp90a1, mitochondrial HSP90; Hsp90b1, endoplasmic reticulum-localized HSP90; Hspa1b, HSP70 member 1b; Hspa4, HSP70 member 4; IL, interleukin.

Liver tissue samples obtained from the rats were fixed with 10% formalin. The formalin-fixed specimens (3 μm) were stained using automatic H&E slide stainer (Leica STS020 Multistainer; Leica Microsystems, Inc.) according to the manufacturer's instructions for histological evaluation. The stained liver tissues were examined for histopathological evidence of pathological damage under light microscope (magnification, x100).

Immunohistochemistry (IHC). HSP90 expression in the rat liver tissues was detected by IHC. Deparaffinization and rehydration of the sections (3 μm thickness) were performed with gradient xylene and a descending ethanol gradient. Antigen retrieval was performed with Citrate Antigen Retrieval solution (1:50; cat. no. P0083; Beyotime Institute of Biotechnology) at 95°C for 20 min. Endogenous peroxidase activity was quenched using 0.3% hydrogen peroxide (25°C, 10 min). The sections were blocked with 1% BSA (Beyotime Institute of Biotechnology) for 2 h at room temperature. The sections were then incubated with primary anti-HSP90 antibody (1:100; cat. no. 13171-1-AP; ProteinTech Group, Inc.) overnight at 4°C, followed by incubation with horseradish peroxidase-conjugated secondary antibody (1:50; cat. no. A0208; Beyotime Institute of Biotechnology) for 1.5 h at room temperature. After thoroughly washing with washing buffer (cat. no. P0106L; Beyotime Institute of Biotechnology), the sections were developed using a 3,3'-diaminobenzidine kit (cat. no. P0202; Beyotime Institute of Biotechnology) and counterstained with hematoxylin staining solution (cat. no. C0107; Beyotime Institute of Biotechnology) for 10 min at room temperature. Each stained sample was observed under high power magnification using a light microscope (magnification, x200; Leica Microsystems, Inc.).

‘Cholestatic culture’ assay. BDL and sham rat models were prepared using the aforementioned procedure. After 1 week, cholestatic blood samples were harvested from the rats and cholestatic serum was obtained after centrifugation (2,000 x g for 10 min at 4°C). Cholestatic serum samples from all BDL model rats were mixed in one tube and non-cholestatic serum samples from all sham model rats were mixed in another tube. The sera were filtered using a 0.22-µm filter (Millipore; Merck KGaA) for sterilization. The automated chemical analyzer was used to examine the cholestatic and non-cholestatic serum. The cholestatic serum contained 188.02 µmol/l total bilirubin (TBIL) and 146.6 µmol/l direct bilirubin (DBIL), while the non-cholestatic serum contained 2.3 µmol/l TBIL and 1.72 µmol/l DBIL. The two types of serum were used in the subsequent experiments.

The BRL rat hepatic cell line was obtained from the Shanghai Cell Bank of the Chinese Academy of Sciences. Liver sinusoidal endothelial cells (LSECs) were isolated from the rat liver tissues with modifications to the previously described protocol (21). Briefly, the rat liver tissue was perfused with collagenase (Invitrogen; Thermo Fisher Scientific, Inc.). The single-cell suspension was subjected to velocity and density centrifugations (50 x g for 2 min at 25°C) in Percoll gradients to separate hepatocyte and nonparenchymal cell suspensions. The nonparenchymal cell suspension was incubated with CD146 (LSEC)-biotin antibodies (1:50; cat. no. 130-111-844; Miltenyi Biotec, Inc.) for cell surface staining (30 min at room temperature). The LSEC cells were positively selected by magnetic separation using Anti-Biotin MicroBeads (Miltenyi Biotec, Inc.) according to the manufacturer’s instructions.

Prior to stimulation with serum, BRL cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM; Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc.), and LSECs were cultured with a medium comprising 40% MCDB 131 (Gibco; Thermo Fisher Scientific, Inc.), 40% RPMI-1640 (Gibco; Thermo Fisher Scientific, Inc.) and 20% FBS. The ‘cholestatic culture’ was initiated when the FBS was replaced with the cholestatic serum or non-cholestatic serum obtained from the rats.

Cell Counting Kit-8 (CCK-8) assay. BRL cells were seeded into a 96-well plate at a density of 2x10³ cells/well and allowed
to attach overnight at 37°C. The cells were then cultured in a medium containing non-cholestatic serum or cholestatic serum, with or without 17-DMAG (0.5 μM) for 48 h. Cell proliferation was determined using the CCK-8 assay (MedChemExpress). Briefly, a 1/10 volume of CCK-8 solution was added to each well and the cells were cultured for 2 h. Cell viability was then determined by measuring the absorbance at 450 nm using a plate reader (BioTek Instruments, Inc.). Six repeated wells for each cell condition were analyzed.

Flow cytometric analysis. BRL cells were cultured in a cholestatic medium with or without 17-DMAG (0.5 μM) at 37°C. After 24 h, the cells were collected and stained using Annexin V-FITC and propidium iodide (Becton, Dickinson and Company) according to the manufacturer's instructions. Apoptotic cells were quantified by flow cytometry using a Flow Cytometer (Cytomics FC500; Beckman Coulter, Inc.). The data was analyzed with CXP software version 1.0 (Beckman Coulter). The experiments were performed in triplicates.

Statistical analysis. Data are presented as the mean ± SD. The statistical significance of a difference between two groups was determined using the Student's t-test. Two-way ANOVA followed by Bonferroni's adjustment were used to evaluate differences among multiple groups. P<0.05 was considered to indicate a statistically significant result. All statistical analyses were conducted using SPSS 13.0 software (SPSS, Inc.).

Results

HSP90 expression is upregulated during cholestatic liver injury. Serum levels of ALT, AST and TBIL were examined to confirm the successful establishment of the animal model. The significantly increased ALT, AST and TBIL levels in the BDL group compared with sham group were indicative of substantial cholestasis and liver injury in the BDL group (Fig. 1A). In addition to simple biliary obstruction, the gram-negative pathogen LPS is another factor that aggravates biliary obstruction-induced liver injury (22). Therefore, a BDL + LPS rat model was established to simulate frequently occurring clinical cases. As shown in Fig. 1A, LPS administration further aggravated the BDL-induced liver injury.

The mRNA expression levels of three HSP90 family members in the liver tissues of the BDL and BDL + LPS rat models were compared with those in the sham group to clarify the association between HSP90 expression and cholestatic liver injury. In the BDL and BDL + LPS groups, the results showed that Hsp90aa1 and Hsp90b1 mRNA levels were significantly upregulated compared with those in the sham group (Fig. 1B). IHC showed that the HSP90 protein expression was also upregulated in the liver tissues of rats in the BDL and BDL + LPS groups (Fig. 1C). Whether a similar phenomenon exists in human liver tissues was also examined. Liver specimens obtained from patients with hepatolithiasis or intrahepatic cholangitis who had undergone hepatectomy were analyzed using IHC. The results demonstrated that HSP90 expression was upregulated in the patients with intrahepatic cholangitis (Fig. 1D). These results imply that aberrant HSP90 expression is associated with cholestasis and cholangitis.

HSP90 inhibitor 17-DMAG alleviates cholestasis-induced liver injury. Previous studies have supported the potential of 17-DMAG, a water-soluble HSP90 inhibitor, as a therapeutic agent (23,24). The effect of 17-DMAG on cholestasis-induced liver injury was therefore investigated in vivo to identify whether the inhibition of HSP90 enhances the transcription of HSP70, as indicated by previous studies (25-27). The elevated mRNA levels of HSP70 members 1b and 4 (Hspalb and Hspa4, respectively) in the rats following 17-DMAG administration indicated that the activity of HSP90 was effectively inhibited (Fig. 1E).

Following the administration of 17-DMAG or NS to the rats, the serum levels of ALT and AST were assessed. ALT and AST levels in the BDL and BDL + LPS groups exhibited significant reductions upon treatment with 17-DMAG compared with those in the NS-treated rats (Fig. 1A). However, 17-DMAG administration did not significantly mitigate the increase of TBIL levels after the onset of cholestasis (Fig. 1A).

A morphological assessment was performed to evaluate the degree of injury exhibited by the rat liver tissue. Pathological examination revealed dilation of the bile capillaries and increased local necrosis in the livers of rats with BDL-induced cholestasis (Fig. 1F). In addition, following LPS administration, large areas of necrosis and increased inflammatory cell infiltration were observed (Fig. 1F). However, compared with NS treatment, 17-DMAG treatment reduced local necrosis and inflammatory cell infiltration during cholestasis with or without LPS (Fig. 1F).

Caspase-3 activity and the ratio of Bax to Bcl2 are often used to evaluate the apoptotic response (28). Notably, caspase-3 activity increased significantly in the liver tissues of the BDL and BDL + LPS groups compared with the sham group. However, treatment with 17-DMAG effectively attenuated this change (Fig. 1G). Furthermore, the Bax to Bcl2 ratio exhibited similar changes (Fig. 1H). These results indicate that 17-DMAG effectively alleviated cholestasis-induced liver injury.

17-DMAG protects hepatocytes against cholestasis-induced cell damage in vitro. A 'cholestatic culture' experiment was conducted to further clarify the function of 17-DMAG during cholestasis-induced cell damage in vitro. Cholestatic serum from BDL model rats was added to the culture medium of BRL cells to simulate the cholestatic microenvironment in vitro. CCK-8 assays were then performed to evaluate cell viability. Moderate cytotoxicity was observed when BRL cells were treated with cholestatic serum (Fig. 2A). However, 17-DMAG administration significantly alleviated this cytotoxicity, whether or not LPS was present in vitro (Fig. 2A). In addition, the flow cytometric analysis of cell apoptosis indicated that 17-DMAG administration promoted the survival of BRL cells and reduced the number of apoptotic cells in the cholestatic serum-containing medium (Fig. 2B).

17-DMAG administration prevents cholestasis-induced liver injury by decreasing the expression of IL-1β and IL-18 in vivo. Pro-inflammatory cytokines induced by bile acid play an important role in cholestasis-induced liver injury (29,30). IL-1β and IL-18 are important cytokines, as they have a role in the induction of apoptosis during liver injury (15,31). Therefore, they were analyzed in the present study. The results
showed that the mRNA expression levels of IL-1β and IL-18 increased significantly during cholestasis-induced liver injury, and treatment with 17-DMAG substantially attenuated these changes (Fig. 3A). ELISAs of the liver tissues also demonstrated that the cholestasis-induced increases in IL-1β and IL-18 levels were significantly attenuated following treatment with 17-DMAG (Fig. 3B).

17-DMAG administration decreases the expression of IL-1β and IL-18 by LSECs in vitro. LSECs serve an important role in the immune response and liver injury (32,33). LSECs were isolated from rats for in vitro analyses to study the underlying mechanism. The results showed that the addition of cholestatic medium significantly increased the expression of IL-1β and IL-18 by LSECs, compared with non-cholestatic medium (Fig. 3C and D). Notably, the administration of 17-DMAG significantly attenuated the increase in the expression of IL-1β and IL-18 induced by cholestatic medium (Fig. 3C and D).

Figure 1. 17-DMAG alleviates cholestatic liver injury in vivo. (A) ALT, AST and TBIL were examined in rat serum 24 h after BDL or sham surgery (n=6/group). The rapidly increased ALT, AST and TBIL levels demonstrated that cholestasis-induced liver injury occurred in the BDL and BDL + LPS groups. (B) Relative expression of HSP90 mRNA in liver tissues from three rat models (n=6/group). Hsp90aa1 and Hsp90b1 mRNA were upregulated after biliary obstruction. (C) IHC staining shows that HSP90 protein was upregulated 72 h after biliary obstruction (images are representative of 6 rats/group; magnification, x200). (D) IHC staining shows that HSP90 expression was upregulated in patients with intrahepatic cholangitis compared with (images are representative of five rats/group; magnification, x200). (E) Relative mRNA expression levels of HSP70 family members Hspa1b and Hspa4 in rat liver tissues (n=6/group). Elevated mRNA levels of HSP70 indicate that the activity of HSP90 was effectively inhibited. (F) Histological examination (magnification, x100), (G) the activity of caspase-3 and (H) the ratio of Bax to Bcl2 expression levels show that 17-DMAG alleviated cholestasis-induced liver injury (n=6/group). Black arrows indicate inflammatory cell infiltration and yellow arrows indicate necrosis. *P<0.05; **P<0.01. 17-DMAG, 17-dimethylamino-ethylamino-17-demethoxygeldanamycin; ALT, alanine aminotransferase; AST, aspartate aminotransferase; TBIL, total bilirubin; NS, normal saline; HSP90, heat shock protein 90; Hsp90aa1, mitochondrial HSP90; Hsp90b1, cytosolic HSP90; Trap1, endoplasmic reticulum-localized HSP90; IHC, immunohistochemistry; BDL, bile duct ligation; LPS, lipopolysaccharide; HSP70, heat shock protein 70; Hspa1b, HSP70 member 1b; Hspa4, HSP70 member 4; casp3, caspase-3; n.s., not significant.
Discussion

Biliary obstruction is a common clinical symptom in various benign or malignant diseases and often causes severe liver injury. HSP90 inhibitors have been described as protective drugs for various organs due to their anti-inflammatory effects (11,12,35-37). Given that the present study observed that HSP90 expression increased significantly in choledystic human and rat livers, the inhibition of HSP90 may be a potential method for alleviating liver injury during biliary obstruction.

Both simple biliary obstruction and biliary obstruction with bacterial infection cause liver injury in clinical cases. Given that gram-negative bacteria are the most common pathogens associated with biliary infection and LPS is the most important pathogenic factor of gram-negative bacteria, a BDL + LPS group was employed in the present study to simulate biliary obstruction with secondary infection, in addition to a BDL group. The results demonstrated that the HSP90 inhibitor 17-DMAG significantly decreased the levels of ALT and AST and alleviated the pathological changes caused by choledystis with or without LPS. These results strongly imply that HSP90 inhibitors can protect the liver from choledystic injury.

A novel cell experiment named ‘choledystic culture’ was designed and performed in the present study to clarify the protective function of 17-DMAG against choledystis-induced cell damage in vitro. In a preliminary experiment, the dissolution of bilirubin in DMEM was attempted in order to simulate the microenvironment of hepatocytes after biliary obstruction. Although the addition of dimethyl sulfoxide improved the solubility of bilirubin, this method only provided a culture medium containing unconjugated bilirubin. However, following biliary obstruction, the increase in serum bilirubin is often
characterized by a sharp increase in conjugated bilirubin rather than unconjugated bilirubin (1). Therefore, since both BRL cells and LSECs are rat cells, cholestatic serum from BDL rats was used instead. Whether the rat serum influenced the growth of BRL cells in vitro was examined using CCK-8 assays. The BRL cells grew faster in medium supplemented with rat serum than in the medium supplemented with FBS (data not shown). This experiment provides a viable experimental model for the simulation of the cholestatic microenvironment in vitro.

Bile acid and severe liver injury during cholestasis can trigger an innate immune response (29,38). Secondary infection aggravates cholestasis-induced liver injury and further activates the immune response. Several studies have shown that IL-1β and IL-18 can trigger a specific pro-inflammatory type of cell death during an innate response that may be damaging to various organs (39-41). The accumulation of IL-1β and IL-18 has been shown to aggravate alcohol-mediated and acetaminophen-induced liver hepatotoxicity during liver injury (15,34,42,43). However, TLR9 is able to regulate the accumulation of IL-1β and IL-18 (15,42). The present study revealed that IL-1β and IL-18 accumulated in liver tissue while the inhibition of TLR9 attenuated their accumulation during cholestasis. These findings indicate that IL-1β and IL-18 play a key role in cholestasis-induced hepatotoxicity, which is in line with previous findings.

Several studies have demonstrated that the TLR-mediated innate immune response is associated with liver damage and inflammation. In particular, TLR9 has a pathological role in infection-induced liver injury (44-46), as well as in alcohol-or acetaminophen-induced liver injury (15,34). The present study found that HSP90B1 was upregulated in cholestatic liver tissues. A previous study reported the binding affinity of 17-DMAG and HSP90B1 (27). HSP90B1 has been identified as an essential driver for TLR signaling, including TLR9 signaling (17). The present study found that the administration of 17-DMAG decreased IL-1β and IL-18 expression during cholestasis-induced liver injury in vivo and in vitro. Previous studies have also demonstrated that IL-1β and IL-18 facilitate the apoptosis of hepatocytes during liver injury, and that their biogenesis can be regulated by TLR9 signaling (15,31,34). These findings suggest that 17-DMAG alleviates cholestasis-induced liver injury via the inhibition of HSP90s and TLR9 signaling and the subsequent reduction of IL-1β and IL-18 expression, as demonstrated by the schematic in Fig. 3E.

As a critical chaperone, HSP90 helps to regulate the folding, maturation, stabilization and activation of >300 client proteins (47). 17-DMAG is a semi-synthetic HSP90 inhibitor that has several advantages over other geldanamycin derivatives, such as higher water solubility, good bioavailability, reduced metabolism and greater antitumor effects (23). 17-DMAG has been studied in preclinical and clinical trials for certain refractory diseases, including cancer and inflammatory diseases (23). However, some dose-limiting toxicities have been observed during treatment with 17-DMAG, including fatigue, nausea, vomiting, diarrhea and anorexia (48-51). Despite significant improvements, the safety and efficacy of HSP90 inhibitors require further study prior to their clinical translation.

In summary, the results of the present study showed that the HSP90 inhibitor 17-DMAG protected against cholestasis-induced liver injury in rats, via a mechanism involving the reduction of IL-1β and IL-18 expression.

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Availability of data and materials

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

Authors’ contributions

JY and BL designed the present study. CT, JL, WL, WC and JY performed the experiments. WZ and ZZ analyzed the data. JY and BL designed the present study. CT, JL, WL, WC and JY performed the experiments. WZ and ZZ analyzed the data. CT, JL, JY and BL drafted and revised the paper. JY and BL can authenticate the raw data in this study. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The present study was approved by the Ethics Committee of Xiaoxing People's Hospital (Shaoxing, China).

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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