**Methyl ferulic acid exerts anti-apoptotic effects on L-02 cells via the ROS-mediated signaling pathway**

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**Abstract.** The present study aimed to investigate the anti-apoptotic effects of methyl ferulic acid (MFA) on L-02 cell apoptosis induced by ethanol, and to elucidate the possible underlying mechanisms. L-02 cells were examined after being soaked in ethanol (400 mM) to allow the ethanol to permeate into the cells for 24 h. Cell survival was measured by MTT assay. Cell apoptosis was assessed by both flow cytometry and single-stranded DNA assays. Intracellular reactive oxygen species (ROS) production was determined using the 2',7'-dichlorofluorescein-diacetate dye. The protein expression levels of p38, p-p38, JNK, p-JNK, NADPH oxidase 4 (NOX4), p22, Bax and Bcl-2 were measured by western blot analysis. The mRNA expression levels of NOX4 and p22 were measured by RT-PCR. It was identified that MFA markedly suppressed the ethanol-induced apoptosis and necrosis of L-02 cells. In addition, MFA decreased the expression levels of superoxide dismutase, catalase and phospholipid hydroperoxide glutathione peroxidase, and downregulated the levels of Bax/Bcl-2 and the cleaved forms of caspase-3 in a dose- and time-dependent manner. This indicated that MFA attenuated the apoptosis of L-02 cells. MFA also decreased the elevated mRNA and protein expression levels of Nox4 and p22phox, and the production of intracellular ROS triggered by ethanol. Further analysis demonstrated that MFA significantly attenuated the phosphorylation of JNK and p38, which are major components of the mitogen-activated protein kinase (MAPK) pathways. On the whole, the findings of this study demonstrated that MFA attenuated the apoptotic cell death of L-02 cells by reducing the generation of ROS and inactivating the MAPK pathways.

**Introduction**

Excessive alcohol consumption triggers a variety of liver disorders, ranging from simple steatosis to severe forms of liver injury, including fatty liver, steatohepatitis, liver fibrosis, cirrhosis and even liver cancer (1). Alcoholic liver disease (ALD) is a major healthcare concern, which inflicts individuals, and society as a whole, with damaging consequences and is a significant cause of economic burden worldwide (2,3). The elucidation of the detailed mechanisms responsible for the development of ALD is important in order to determine an effective treatment. Mounting evidence indicates that oxidative stress plays a key role in ALD. Ethanol-induced oxidative stress directly influences the elevated production of reactive oxygen species (ROS) and increases lipid peroxidation and damage to the antioxidant system, which leads to cell apoptosis and necrosis (4,5). It was recently reported that hepatocyte cell death via apoptosis and necrosis may be a critical process in ALD (6). An increasing number of studies have noted that oxidative stress and superfluous intracellular ROS production induced by ethanol and its metabolites exert a pivotal effect on ethanol-induced cellular apoptosis (7,8), and suggested that apoptosis is mainly induced via the Fas- and mitochondria-mediated pathways (9).

NADPH oxidase 4 (NOX4), is expressed particularly in hepatocytes and hepatic stellate cells (HSCs), and is therefore an important source of ROS in signal transduction, playing a vital role in the physiological and pathological processes of ALD (5,10). It is a reasonable hypothesis that ROS derived from NOX4 on the membrane may be associated with Fas activation (11). Relatively high levels of intracellular ROS induce redox imbalance, causing cell apoptosis via the mitogen-activated protein kinase (MAPK) signaling pathway (12).

It is known that MAPK determines the fate of various cells, and that p38 MAPK and JNK may positively influence the mitochondrial pathways that lead to the apoptosis of ethanol-exposed SK-Hepl cells, suggesting an interaction between apoptosis and MAPK signaling systems (13). L-02 is a new cell line established by the Shanghai Biochemical Institute of Chinese Academy of Sciences and was selected for use in a previous study (14); this cell line was selected for the present study as these cells have alcohol dehydrogenase (ADH) activity (15). Therefore, the results may be more

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**Key words:** ethanol, methyl ferulic acid, L-02 cells, apoptosis, reactive oxygen species, NADPH, mitogen-activated protein kinase

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Apoptosis detection by flow cytometry.

≥3 times. were conducted with strict sterile precautions and repeated Inc.) to measure the absorbance at 570 nm. All experiments
the manufacturer's instructions. Briefly, following treatment of DMSO to dissolve the crystals, the cells were placed in a
5% CO2 and 95% air at 37˚C.

Materials and methods

Drug and reagents. MFA (3,4-dimethoxycinnamic acid) at >98% purity and all other reagents (unless otherwise indi-
cated) were obtained from Sigma-Aldrich; Merck KGaA (Darmstadt, Germany). Ethanol was purchased from JiYu Chemical Factory (Shanghai, China).

Cells and cell culture. L-02, a normal human hepatic cell line (Chinese Academy of Sciences, Shanghai, China), was cultured in DMEM (Invitrogen; Thermo Fisher Scientific, Inc., Waltham, MA, USA) containing 10% (v/v) fetal bovine serum and antibiotics (100 U/ml penicillin and 100 µg/ml strepto-
mycin) (both from HyClone; GE Healthcare Life Sciences, Little Chalfont, UK) and maintained in a humidified atmo-
sphere of 5% CO2 and 95% air at 37˚C.

Cells between generations 3 and 5 were used in the experiments. The L-02 cells were treated under the following conditions unless otherwise stated: Following 12 h of pre-
incubation in serum-free DMEM, MFA (25, 50 and 100 µM) was added to the cultures for pretreatment. Following incuba-
tion with MFA for 1 h, ethanol (400 mM) was added to DMEM and the cells incubated for a further 24 h.

Cell viability assay. Cell viability was ascertained by MTT assay. First, the cells were seeded into 96-well plates (5x10^4 cells/well) 24 h prior to treatment. The cells were then treated with MFA at various concentrations (25, 50, 100, 250, 500, 1,000 and 2,000 µM for 24, 48 or 72 h, respectively) prior to being incubated with a working solution of MTT (5 mg/ml) at 37˚C for 4 h. Following treatment with 150 µl of DMSO to dissolve the crystals, the cells were placed in a microplate reader (Mutiskan FC; Thermo Fisher Scientific, Inc.) to measure the absorbance at 570 nm. All experiments were conducted with strict sterile precautions and repeated ≥3 times.

Apoptosis detection by flow cytometry. To further examined whether MFA could induce apoptosis by ethanol, apoptosis was quantified using an Annexin V-fluorescein isothiocyanate (FITC) Apoptosis Detection kit (Roche Diagnostics GmbH, Mannheim, Germany) in accordance with the manufacturer's instructions. Briefly, following treatment with MFA for 24 h at various concentrations as mentioned above, the cells were harvested and washed twice with cold PBS. Following centrifugation (500 x g, 5 min, 25˚C), the cells were stained with Annexin V-FITC and PI, and then analyzed by flow cytometry (flow cytometer; BD Biosciences, San Jose, CA, USA).

Measurement of aspartate aminotransferase (AST), glutathione peroxidase (GSH-Px), superoxide dismutase (SOD) and catalase (CAT) levels. Firstly 3x10^4 cells were cultured in a dish (diameter, 10 cm) for 24 h, and the DMEM was then discarded. Secondly, the MFA (100, 50 and 25 µM) group was pre-cultured with MFA for 1 h, and the cells in the model group and MAF group were then supplemented with 400 mM ethanol and culture for 24 h. Thirdly, the supernatant from the cultured cells was collected and used immediately for the assay of AST using the aspartate aminotransferase assay kit (C010-2; Nanjing Jiansheng Bio Co., Nanjing, China) according to the manufacturer's instructions. The cells were then split and the active protein was extracted using the One Step Animal Cell Active Protein Extraction kit (C500022; Sangon Biotech Co., Ltd., Shanghai, China). The antioxidant enzyme (SOD and CAT) and GSH-Px activities were then evaluated using different detection kits respectively according to the manufacturer's instructions as follows: The superoxide dismutase assay kit (A001-3), catalase assay kit (A007-2) and glutathione peroxidase (GSH-PX) assay kit (A005) (all from Nanjing Jiansheng Bio Co.).

Measurement of ROS generation. ROS generation was evaluated with an intracellular ROS Assay kit in accordance to the manufacturer's instructions (Cell Bioslabs Inc., San Diego, CA, USA). DCFH-DA itself does not fluoresce, but can freely cross the cell membrane and enter the cell to be hydrolyzed to DCF by esterase in the cell. DCFH cannot permeate the cell membrane, and thus the probe is easily loaded into the cell. ROS in the cell can oxidize the non-fluorescent DCFH to produce the fluorescent DCF, and the amount of DCF fluorescence can be used to identify the level of ROS in the cells. The ROS level was verified with a Tecan Model infinite M200 PRO Microplate reader (Tecan Group, Ltd., Mannedorf, Switzerland) at an emission wavelength and excitation wave-

of 523 and 502 nm, respectively.

RNA extraction and reverse transcription (RT)-PCR. Following the manufacturer's instructions, total RNA was isolated from the cells with the total RNA isolation kit (TriPure Reagent; Mai Bio Co., Shanghai, China), and then reverse transcribed into cDNA with a cDNA synthesis kit (TIANScript cDNA; Tiangen Biotech Co., Ltd., Beijing, China) in accordance with the manufacturer's instructions. Target genes were amplified by the MJ ptc-200 PCR amplification system (MJ Research, Inc., Waltham, MA, USA) with a RT-PCR kit (2X Taq PCR Master Mix; Aidlab Biotechnologies Co., Ltd., Beijing, China) according to the manufacturer's instructions. The specific primers for the target genes, GAPDH and β-actin, were synthe-
sized by Sangon Biotech Co., Ltd. and are listed in Table I. β-actin was used as an internal control. The parameters of the reaction were based on those in our previous study (17). The PCR products were identified using 1.5% agarose gel electrophoresis, and the optical density of the target gene bands in each sample was calculated using the ChemiDoc imaging system with adjustment through β-actin correction to finally obtain the relative expression of the target genes in each sample.
with 82 µl Nucleofector. Approximately 10⁷-10⁸ cells were prepared. A mixed solution of 3 µg of NOX4-cDNA dissolved in 18 µl supplemented P5 primary cell solution were added to the Nucleofector™ kit (Lonza, Basel, Switzerland). For each transfection with NOX4-cDNA with the P5 primary cell Nucleofector™ kit (Lonza, Basel, Switzerland) was used as the internal control.

Table I. Sequences of primers used for the determination of NOX4, p22phox, Bax, Bcl-2 and β-actin gene expression.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Oligonucleotide primer sequence</th>
<th>Annealing temperature</th>
<th>Number of cycles</th>
</tr>
</thead>
<tbody>
<tr>
<td>NOX4 (136 bp)</td>
<td>Forward: 5'-TGTGCGGAACACTCTTGGC-3'</td>
<td>58°C</td>
<td>35</td>
</tr>
<tr>
<td></td>
<td>Reverse: 5'-ATATGCACGCCTGAGAAATA-3'</td>
<td></td>
<td></td>
</tr>
<tr>
<td>p22phox (103 bp)</td>
<td>Forward: 5'-TATTGTGTGCAAGGTGCTCA-3'</td>
<td>58°C</td>
<td>35</td>
</tr>
<tr>
<td></td>
<td>Reverse: 5'-CACAGGCTGTCAGTCACTTC-3'</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bax (155 bp)</td>
<td>Forward: 5'-CCCAGAGGCTCTTTCGCCAG-3'</td>
<td>53°C</td>
<td>35</td>
</tr>
<tr>
<td></td>
<td>Reverse: 5'-CCAGCCCATGATGGTCTGAT-3'</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bcl-2 (89 bp)</td>
<td>Forward: 5'-GGTGGGTCTATGGTGTGG-3'</td>
<td>55°C</td>
<td>35</td>
</tr>
<tr>
<td></td>
<td>Reverse: 5'-CCGTTCCAGTGATCAGTCATCC-3'</td>
<td></td>
<td></td>
</tr>
<tr>
<td>β-actin (199 bp)</td>
<td>Forward: 5'-GGACTCTATGTGGTGACGA-3'</td>
<td>56°C</td>
<td>35</td>
</tr>
<tr>
<td></td>
<td>Reverse: 5'-ACGGTTGGCCTTAGGGTTCA-3'</td>
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**Western blot analysis.** Total proteins were extracted from the cells and the concentration was analyzed with a BCA protein concentration assay kit (Beyotime Institute of Biotechnology, Haimen, China). Sample proteins were separated by electrophoresis by a 12% SDS-PAGE separating gel with a Bio-Rad electrophoresis system (Bio-Rad Laboratories, Inc., Hercules, CA, USA). Equivalent amounts (50 µg) of protein were then transferred onto Pure Nitrocellulose Blotting membranes and blocked with 5% fat-free milk, the proteins were then incubated with the primary antibodies [anti-NOX4 (1:1,000; ab13303), Bel-2 (1:1,000; ab59348) (both from Abcam, Cambridge, UK), caspase-3 (1:1,000; 9662S; Cell Signaling Technology Inc., Danvers, MA, USA), cleaved caspase-3 (1:500; D260009; Sangon Biotech Co., Ltd.), Bax (1:1,000; ab53154), JNK1/2 (1:1,000; ab124956), p-JNK1/2 (1:1,000; ab207477) (all from Abcam), anti-p22phox rabbit antibody (1:500; Bioworld Technology, Inc., Danvers, MA, USA), Bcl-2 (1:1,000; ab207477) (all from Abcam), anti-p38 MAPK antibodies (1:1,000; ab170099) (both from Abcam), and anti-p-p38 MAPK (1:1,000; Bioworld Technology, Inc., Nanjing, China), anti-p-p38 MAPK (1:1,000; ab4822) and anti-p38 MAPK antibodies (1:1,000; ab170099) (both from Abcam)] overnight at 4˚C. The membranes were then incubated with the corresponding horseradish peroxidase-conjugated secondary antibodies (1:50,000; TAI40003; goat anti-rabbit IgG; OriGene Technologies, Inc., Beijing, China) at room temperature for 1 h. The qualitative and quantitative analysis of the blots were estimated with Molecular Imager Chemi Doc XRS (Bio-Rad Laboratories, Inc.) by enhanced chemiluminescence (E002-100; 7Sea Pharmatech Co., Ltd., Shanghai, China) and the JS-780 automatic gel imaging analysis system. β-actin (1:1,000; C640018; Sangon Biotech Co., Ltd.) was used as the internal control.

**Transfection of small interfering (si)RNA.** siRNA directed against NOX4 (NOX4-siRNA) and negative control siRNA (NC-siRNA) were obtained from Gene Pharma (Shanghai, China). The NOX4- or NC-siRNA was transfected into the L02 cells using Lipofectamine® (Invitrogen; Thermo Fisher Scientific, Inc.) in accordance with the manufacturer's instructions. Briefly, the cells were placed in 6-well plates at a density of 2x10⁵ cells/well in 2 ml of complete DMEM. When the cells grew to ~70% confluence transfection with siRNA commenced as follows: A total of 100 pmol of siRNA was mixed with 2 µl of Lipofectamine® in each well, and 500 µl of antibiotic- and serum-free DMEM medium were then added. The cells were incubated with the transfection mixture for 6 h. At the final stage of incubation, 1.5 ml of antibiotic-free complete medium was replenished and the cells were incubated for a further 18 h. Following exposure to ethanol for 24 h, the cells were harvested and the NOX4 protein and mRNA expression levels were evaluated by western blot analysis and RT-PCR, respectively.

**Statistical analysis.** The results are expressed as the means ± standard deviation. All experimental data were analyzed using SPSS software version 17.0 (SPSS, Inc., Chicago, IL, USA). One-way analysis of variance was used to determine significant differences among groups with the Student-Newman-Keuls (SNK-q) post hoc test. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**Effect of MFA on the viability of L-02 cells.** The cytotoxic effects of MFA on the cells were evaluated by MTT assay. The cells were treated with various concentrations of MFA (25, 50, 100, 250, 500, 1,000 and 2,000 µM) for 24, 48 or 72 h, respectively. The percentages of cell growth inhibition for each treatment group were calculated by adjusting the untreated control group to 100%. Only 15% cell growth inhibition was observed at a MFA concentration <250 µM, even after 72 h of treatment (Fig. 1A).

To examine the effects of ethanol on the viability of cells, the L-02 cells were cultured in DMEM with or without (as a control) ethanol. A pre-experimental experiment was conducted on L-02 cells using Lipofectamine® (Invitrogen; Thermo Fisher Scientific, Inc.) in accordance with the manufacturer's instructions. Briefly, the cells were seeded in 6-well plates at a density of 2x10⁵ cells/well in 2 ml of complete DMEM. When the cells grew to ~70% confluence transfection with siRNA commenced as follows: A total of 100 pmol of siRNA was mixed with 2 µl of Lipofectamine® in each well, and 500 µl of antibiotic- and serum-free DMEM medium were then added. The cells were incubated with the transfection mixture for 6 h. At the final stage of incubation, 1.5 ml of antibiotic-free complete medium was replenished and the cells were incubated for a further 18 h. Following exposure to ethanol for 24 h, the cells were harvested and the NOX4 protein and mRNA expression levels were evaluated by western blot analysis and RT-PCR, respectively.

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conducted in which viability for 3, 6, 12, 24 and 48 h was estimated at various concentrations of ethanol (100, 200, 300, 400, 500, 600 and 700 mM). The results demonstrated that the cell inhibition rate of 400 mM ethanol was close to 50% at 24 h (data not shown). It was thus determined that 400 mM ethanol was the appropriate concentration for inducing L-02 cell apoptosis.

In order to obtain the optimal experimental conditions, the concentrations of 1, 2.5, 5, 10, 25, 50, 100 and 200 µM of MFA and 400 mM ethanol were used for the preliminary experiment (Fig. 1B). The concentrations of 100, 50 and 25 µM of MFA and 400 mM of ethanol were selected for use in further experiments as they produced significant changes in the viability of the cells compared with the control.

**MFA attenuates the ethanol-induced apoptosis of ethanol-exposed L-02 cells.** In order to determine whether MFA mediates the ethanol-induced apoptosis or necrosis of L-02 cells, Annexin V-PI flow cytometric analysis, which can differentiate between apoptotic and necrotic cells, was performed. MFA treatment caused a significant increase in the number of Annexin V-positive L-02 cells in a concentration-dependent manner. The results demonstrated that the apoptotic rate in the group exposed to ethanol was significantly increased by 89% (P<0.01) and the apoptotic rate in the MFA group was significantly lower compared with that in the ethanol group (P<0.01). The level of apoptosis of L-02 cells decreased by 29.1, 20.7 and 15.3% following treatment with MFA at 100, 500, 600 and 700 mM. The results demonstrated that MFA attenuates the ethanol-induced apoptosis of L-02 cells (Fig. 2A).

To further evaluate the anti-apoptotic effects of MFA, the effects of MFA on apoptosis-associated proteins that serve as biomarkers of cell apoptosis (Bax, Bcl-2, cleaved caspase-3 and caspase-3) were also examined in vitro. The results from western blot analysis (Fig. 2B) demonstrated that ethanol exposure mainly increased Bax, caspase-3 and cleaved caspase-3 protein expression, but decreased Bcl-2 protein expression compared with the control. Although at the mRNA level, ethanol was also found to increase both the level of Bax and Bcl-2 expression compared with the control (Fig. 2C), the protein level of Bcl-2 decreased with ethanol treatment compared to that of the control. Western blot analysis also revealed that the alcohol-exposed L-02 cells exhibited an elevated Bax/Bcl-2 protein expression (P<0.01). In addition, the protein level of anti-apoptotic Bcl-2 was increased following treatment with MFA in the L-02 cells compared with the control group (P<0.01), while the levels of the apoptotic proteins, caspase-3 and Bax, were decreased (P<0.01). Taken together, these findings suggest that MFA attenuates the ethanol-induced apoptosis of L-02 cells in a dose-dependent manner by regulating the levels of apoptotic proteins. The effect of 100 µM MFA dose is particularly prominent.

**MFA affects the generation of AST, GSH-Px, SOD and CAT in ethanol-exposed L-02 cells.** It has been reported that ethanol-induced cell death is partly mediated by oxidative stress (13). To alleviate the cumulative burden of oxidative stress, cells generally utilize antioxidant defense systems to eliminate ROS. SOD, GSH-Px and CAT are the first line of defense against oxidative stress and can block free radical formation and prevent the cells from oxidative damage by ROS (19). SOD is able to convert the superoxide radical into H₂O₂, which can be broken down to O₂ by CAT and GSH-Px (20). In the present study, to determine whether the protective effects of MFA against ethanol-induced injury are mediated via antioxidant enzymes, the activities of antioxidant enzymes in L-02 cells with or without pretreatment with MFA were investigated. First the L-02 cells were treated with MFA in the presence or absence of ethanol for 24 h, and the activities of AST in the culture medium and the levels of GSH-Px, CAT and SOD in the cell lysates were then determined. As shown in Fig. 3A, the activities of AST in the ethanol-exposed group were significantly increased by 125% compared with those of the control group (P<0.01). However, treatment with MFA (100, 50 and 25 µM) significantly inhibited the ethanol-induced elevation of AST activities by 65.5, 52.8 and 39.7%, respectively. In addition, MFA significantly increased the levels of GSH-Px, CAT and SOD levels compared with those in the ethanol-exposed cells (Fig. 3B-D). MFA (100, 50 and 25 µM) increased the GSH-Px levels by 230, 170 and 110%, respectively compared to that of the control group (P<0.01). However, treatment with MFA (100, 50 and 25 µM) significantly inhibited the ethanol-induced elevation of AST activities by 65.5, 52.8 and 39.7%, respectively. In addition, MFA significantly increased the levels of GSH-Px, CAT and SOD levels compared with those in the ethanol-exposed cells (Fig. 3B-D). MFA (100, 50 and 25 µM) increased the GSH-Px levels by 230, 170 and 110% compared with the ethanol-exposed cells, respectively. MFA (100, 50 and 25 µM) also increased the CAT levels by 56, 412.1 and 223.1%, respectively compared with those in the ethanol-exposed cells, and MFA (100, 50 and 25 µM) also elevated the SOD levels by 631.2, 432.3 and 337.0%, respectively compared with those in the ethanol-exposed cells.
Notably, treatment with MFA significantly reduced ethanol-induced oxidative stress, as demonstrated by the reduction in the ROS levels when compared with the ethanol-exposed cells (Fig. 3E and F).

The anti-apoptotic effects of MFA are mediated via ROS generation. In recent years, a number of studies have demonstrated that oxidative stress can cause cellular apoptosis (6,13). ROS production increases in ethanol-stimulated L-02 cells and leads to apoptosis (14). Therefore, the present study investigated whether the inhibitory effects of MFA on the ethanol-induced apoptosis of L-02 cells were associated with the accumulation of ROS. First, the ROS levels in ethanol-exposed L-02 cells were estimated by determining the oxidative conversion of non-fluorescent DCFH-DA to fluorescent DCF. Consistent with the findings of a previous study (17), the exposure of L-02 cells to ethanol triggered the generation of ROS and apoptosis. The results of the present study demonstrated that ethanol triggered ROS production in the L-02 cells; the level of ROS was significantly increased by about 2 times compared with that of the control group (P<0.01) (Fig. 3E and F). Following treatment with MFA (100, 50 and 25 µM), the levels of ROS generation levels were partly blocked; compared with the ethanol group, the levels of ROS were reduced in all the 3 MFA treatment (all 3 concentrations) (P<0.01).

As mentioned above, the exposure of the L-02 cells to ethanol induced the production of ROS and apoptosis. MFA treatment successfully reduced ethanol-induced ROS...
generation (Fig. 3E and F) and attenuated the ethanol-induced apoptosis of L-02 cells (Fig. 2A). These results suggested that ROS generation acts upstream of apoptosis in ethanol-exposed L-02 cells. A similar association was also confirmed in two other hepatoma carcinoma cell lines (SK-HEP-1 and HepG2; data not shown). Collectively, these findings suggested that the signaling pathway through which ethanol induces the apoptosis of L-02 cells involves ROS as a second messenger. MFA reverses ethanol-induced ROS production and apoptosis.

**MFA inhibits the expression levels of Nox4 and p22phox in L-02 cells.** Recently, studies have emphasized NADPH oxidases (NOX) as a key source of ROS. NOX4 and its subunit p22phox are highly expressed in hepatocytes and HSCs (7,18). In this study, to further understand the mechanisms underlying the MFA-induced reduction in ROS generation, the effects of MFA on NOX4, one of the major sources of ROS generation found in cells, were assessed. NOX4 appears to be the most abundant isoform of NOXs expressed in the liver (18);
therefore, the expression levels of NOX4, as well as its regulatory subunit, p22phox, were measured. It was identified that the protein expression of NOX4 and p22phox was significantly increased (75.3 and 120.5%, respectively) compared with the control group in response to ethanol. By contrast, treatment with MFA significantly decreased the expression of NOX4 by 56.2% and the expression of p22phox by 25.6%, even at the low concentration of 25 µM (P<0.01). A dose-dependent pattern was identified (Fig. 4A). To further determine whether the downregulation of NOX4 and p22phox protein expression induced by MFA was due to alterations in mRNA transcription, RT-PCR was performed to analyze the mRNA levels of NOX4 and p22phox. Following exposure to ethanol, the mRNA levels of NOX4 were increased by 256.2% and those of p22phox were increased by 320.5% compared with the control group. Concomitant with the reduction in the protein expression induced by MFA, treatment with MFA significantly decreased the mRNA expression levels of NOX4 and p22phox. MFA (100, 50 and 25 µM) decreased the mRNA levels of NOX4 by 68.8, 56.2 and 40.6% (P<0.01), respectively, compared with those of the control group. MFA (100, 50 and 25 µM) decreased mRNA levels of p22phox by 75.3, 53.4 and 50.7% (P<0.01), respectively, compared with those of the control group (Fig. 4B).

**MFA treatment attenuates ethanol-induced MAPK phosphorylation in L-02 cells.** ROS has been regarded as a potent regulator of MAPK family members and subsequent cell death. To activate p38 MAPK pathway would accelerate cell apoptosis, JNKs have been considered to be involved in stimulating apoptotic signaling. Oxidative stress can switch on JNK to bring about apoptosis by receptor-initiated extrinsic and mitochondrial intrinsic apoptotic pathways. JNKs also serve an essential role in modulating the functions of pro- and anti-apoptotic proteins located in the mitochondria (12).

MAPK cascades are pivotal signaling pathways involved in the regulation of normal cell proliferation and survival. Hence, the level of phosphorylated MAPKs in the presence and/or absence of MFA was measured in this study. As demonstrated in Fig. 4C, compared with the control, the phosphorylation levels of p38 MAPK and JNK were markedly increased in the L-02 cells following exposure to ethanol, and the levels of p-p38 and p-JNK increased by 50.3 and 76.2%, respectively compared with the control. By contrast, compared with the ethanol-exposed group, treatment with MFA (100, 50 and 25 µM) effectively decreased the level of p-p38 by 70.3, 46.0 and 11.1%, and the level of p-JNK by 59.3, 45.0 and 25.1%, respectively, thereby exhibiting its promising activation of the p38 MAPK and JNK pathways. MFA decreased MAPK activation in a concentration-dependent manner, suggesting that MAPK inhibition by MFA can result in increased survival rates of L-02 cells (Fig. 4C).

**Effects of MFA on cell viability and ROS generation in ethanol-exposed L-02 cells following transfection with NOX4**
LI et al: ANTI-APOPTOTIC EFFECTS OF METHYL FERULIC ACID ON L-02 CELLS

To validate the hypothesis that MFA decreased ethanol-induced ROS generation by blocking the NOX4 signaling pathway, the expression of NOX4 was upregulated by transfection of the L-02 cells with NOX4 overexpression cDNA for 24 h, which increased the protein expression levels of NOX4 and p22phox by 55 and 18%, as verified by western blot analysis (Fig. 5A). Correspondingly, ROS production increased by 25% and the cell activity decreased by 65% compared with the control group (Fig. 5B and C). However, these responses were effectively reversed by treatment with MFA.

In contrast to the results observed for NOX4 overexpression, the expression of NOX4 was downregulated by transfection of the L-02 cells with NOX4-siRNA for 24 h, which reduced the protein expression levels of NOX4 and p22phox by 67 and 34%, as determined by western blot analysis (Fig. 6A). Correspondingly, ROS production was decreased by 49% and cell viability was increased by 65% compared with the ethanol-exposed group (Fig. 6B and C).

Additionally, when treatment with MFA was applied to the NOX4-overexpressing or NOX4-siRNA-transfected L-02 cells, analysis by cell flow cytometry revealed that treatment with MFA further blocked ROS elevation triggered by ethanol (Figs. 5B and 6B). MTT assay demonstrated that when the L-02 cells were exposed to ethanol following treatment with MFA, the decrease in cellular viability induced by ethanol was suppressed, indicating that MFA protected the L-02 cells from ethanol-induced cell toxicity (Figs. 5C and 6C). Taken together, these results suggested that signaling driven by NOX4/p22phox was the major mechanism underlying the anti-oxidative stress activity of MFA against the generation of ROS triggered by ethanol in L-02 cells. These results suggested that MFA protected hepatocytes from ethanol-induced apoptosis through the inactivation of the NOX4/ROS/p38/JNK pathway.

Discussion

As the pharmacological options available for the therapy of liver diseases are limited, proof of effective hepatic protective agents from natural sources is important. Therefore, it is significant to evaluate plant extracts that can help to restore liver function. MFA is a monomer isolated from Securidaca inappendiculata Hassk., which possesses positive antiviral activity via the specific combination with GP120 to prevent the virus reverse transcriptase from interaction with the lymphocytes (21). The present study surveyed the hepatoprotective activity of MFA using an ethanol-exposed L-02 cell model. The results demonstrated that treatment with MFA significantly reduce elevated ROS levels and reversed ethanol-induced L-02 cell apoptosis, indicating that MFA was responsible for a hepatoprotective effect.
The present study explored the possible effects of MFA on the viability and apoptosis induced by ethanol. The results from in vitro experiments revealed that MFA attenuated the ethanol-induced inhibition of the viability and apoptosis of L-02 cells. Previous studies have reported the use of MFA in the therapy of acute or chronic hepatitis and that it demonstrated some inhibitory effect on HBsAg (16,17). Its analog improved antioxidant activity and anti-lipid peroxidation that protects cells against oxidative stress. Ferulic acid (FA) alleviated the oxidative stress and decreased cell apoptosis induced by high glucose in hepatocytes (22). Whether MFA inhibited the apoptosis induced by ethanol in L-02 cells remained to be elucidated. In this study, the results from flow cytometry demonstrated that MFA inhibited the ethanol-induced apoptosis of L-02 cells, suggesting that its antioxidant properties may contribute to its anti-apoptotic activity.

Additionally, the present study identified that culturing the L-02 cells with MFA (25 µM) reduced apoptosis and ROS production (Figs. 2A and 5B). Maruf et al (23) examined FA and demonstrated that it can protect isolated rat hepatocytes against glyoxal- or methylglyoxal-induced cytotoxicity and oxidative stress. FA attenuated hepatocyte apoptosis which induced ischemia/reperfusion (I/R) via the inhibition of JNK activation (24). In addition, the results of the study by Urias-Lugo et al (25) demonstrated that culturing HepG2 cells and primary hepatocytes with phenolic acids exerted an anti-proliferative effect. FA may cause cell cycle arrest in PC-3 cells and leads to the apoptosis of LNCAp cells (26). MFA treatment at a concentration of 25 µM inhibited the proliferation induced by TGF-β in HSC-LX-2 cells (27). These various results indicate that MFA may exert differential effects in different cell types; that is, MFA can be either cytoprotective or cytotoxic depending on the cell type.

In this study, when the L-02 cells were incubated with ethanol at the concentration of 400 mM, ROS generation was increased with the time of incubation. As one of 6 homologues of transmembrane NADPH oxidase, NOX4 has been identified to be involved in ROS generation and highly expressed in the liver (28). In the liver, NOX4 is expressed in hepatocytes and is upregulated by ethanol or TGF-β in vitro as well as in vivo. It is activated in HSCs and, to a small extent, in sinusoidal endothelial cells, although Chuffer cells do not express it (29,30). Furthermore, NOX4 and p22phox are upregulated in patients exposed to ethanol (31). In this study, the pronounced expression of NOX4 was identified by western blot analysis and was confirmed by RT-PCR in L-02 cells. The results of the present study demonstrated that the mRNA levels of NOX4 and p22phox were increased when the L-02 cells were incubated

Figure 6. Effect of methyl ferulic acid (MFA) on ethanol-induced reactive oxygen species (ROS) generation in L-02 cells transfected with NOX4-siRNA. Following transfection with NOX4-siRNA for 24 h, the L-02 cells were treated as designated. (A) Representative images and protein expression levels of p22phox were normalized to β-actin. (B) Measurement of intracellular ROS levels using DCFH-DA in various groups. (C) Cell viability was measured by MTT assay in NOX4-siRNA-transfected L-02 cells. Data are expressed as the means ± SD, n=3. *P<0.05 and **P<0.01 vs. the control group; #P<0.05 vs. the ethanol group; △P<0.05 vs. the control group. Lanes and bars are labeled as follows: 1, control; 2, siRNA-transfected group; 3, siRNA-transfected group exposed to ethanol; 4, siRNA-transfected group treated with MFA; 5, siRNA-transfected group exposed to ethanol and treated with MFA.
with ethanol (Fig. 4B), demonstrating that NOX4 and p22<sub>phox</sub> are required when activating NOX4 on the membrane. The data from western blot analysis of NOX4 further substantiated this at the protein level.

In the NOX4 overexpression and NOX4 siRNA knockdown experiments, it was identified that NOX4 upregulation caused the upregulation of its regulatory subunit, p22<sub>phox</sub> and, simultaneously, NOX4 downregulation caused the downregulation of p22<sub>phox</sub> in L-02 cells exposed to ethanol. The results of the present study support those of previous findings in that SK-Hep1 cells and alveolar macrophage mRNA levels of NOX4 and p22<sub>phox</sub> increased when the cells were incubated with ethanol (13), which also demonstrated that both are required to activate NOX4 on the membrane. However, the mechanisms behind the interaction between NOX4 and p22<sub>phox</sub> remain to be elucidated; further studies are warranted to elucidate these mechanisms.

Based on the results of the present study, the release of ROS was, at least in part, mediated by NOX4 in the ethanol-exposed L-02 cells. The overproduction of ROS is one of the main causes of increasing oxidative stress and triggering apoptosis (32,33). The results of the present study suggested that ROS generation mediated by NOX4 on the hepatocyte membrane was a trigger of apoptosis in ethanol-exposed L-02 cells.

Compared with the ethanol-exposed group, MFA significantly decreased the elevated NOX4 mRNA and protein expression levels induced by ethanol in varying degrees (Figs. 4-6). This result was consistent with the results obtained from the ROS and apoptosis experiments, in that ROS generation and the apoptosis of L-02 cells corresponded with the changes in NOX4 expression. Further investigations into the underlying mechanisms of the anti-apoptotic effects of MFA are required.

Recently, various studies have identified that ROS may play a critical role in the induction of apoptosis (23,34). Oxidative stress can be induced by abnormal ROS release or their constant generation and this is related to apoptosis and other biological events (35). ROS are one of the pivotal regulators of cell signal transduction and are associated with apoptosis, senescence and proliferation (28). A number of drug candidates fulfill a cytotoxic role through the production of ROS as a critical regulator. Studies have demonstrated that the metabolism of ethanol in liver cells can induce ROS production (32,36). It has been demonstrated that ethanol-mediated ROS production can cause alterations in cellular morphology and functions and/or eventually lead to apoptosis (35,36). For example, the dysfunction of the mitochondria induced by excessive ROS production results in apoptosis (37). ROS are also known to be activators of the MAPK signaling pathway (38). The results of the present study demonstrated that the sustained phosphorylation of p38 MAPK and JNK was caused by ROS production following exposure to ethanol (Fig. 4A and B). The suppression of ROS generation by MFA treatment alleviated the effects of ethanol on JNK and p38 MAPK phosphorylation, suggesting that ethanol induced the generation of ROS, which consequently transformed phosphorylated p38 MAPK and JNK, and caused the activation of p38 MAPK and JNK, finally leading to the translocation of Bax to the mitochondria.

Ethanol induced the inhibition of cell growth and triggered the apoptosis of L-02 cells; however, this process was effectively blocked by MFA treatment. Ethanol-induced apoptosis occurs via a decline in Bcl-2 protein synthesis and the transfer of Bax to the mitochondria (39). The present study demonstrated that ROS, JNK, p38 MAPK and Bax participated in the ethanol-triggered apoptotic pathway. In line with this finding, ROS may serve as an upstream signal mediator of the p38 MAPK and JNK signaling pathways in L-02 cells treated with ethanol (Fig. 3E). There is evidence from a previous study to indicate that ROS can aggravate apoptosis induced by a variety of stimuli (40).

One of the important factors for the apoptosis induced by the mitochondrial pathway is the collapse of mitochondrial membrane potential, which leads to the release of cytochrome c and the activation of caspase-9. This event is mediated by anti-apoptotic proteins of the Bcl-2 family. Particularly, the transfer of Bax to the mitochondria leads to the alteration of mitochondrial membrane potential and Bax plays a critical role in triggering apoptosis in response to various stimuli (41). Although Bax is mainly located in the cytoplasm, when stimulated, it can move closer to the mitochondrial membrane (42). When Bax converges onto the mitochondria, combining with other pro-apoptotic Bcl-2 family members, it induces the release of cytochrome c through the pore channels formed in the outer membrane of the mitochondria by oligomerization or by other channels (43-45). Phosphorylation of JNK and p38 MAPK activates Bax, either alone or in combination (46). In the present study, MFA inhibited induction of caspase-3 activation by ethanol in L-02 cells. The increased ratio of Bax and Bcl-2 can alter the mitochondrial membrane potential and lead to the release of cytochrome c, and then further activate caspase-3 to trigger apoptosis (47,48). As shown in Fig. 2B and C, exposure to ethanol upregulated the mRNA expression of Bax and Bcl-2, but also increased the ratio of Bax to Bcl-2 both at the mRNA and protein level; these effects of ethanol were antagonized by MFA. The results indicated that apoptosis plays an important role in ethanol-induced L-02 cell injury and its effect on the mitochondrial pathway. The results of the present study seem to suggest that exposure to ethanol leads to mitochondrial damage and the caspase-dependent apoptosis of L-02 cells.

Although originally identified as an antioxidant, MFA is currently believed to act through different mechanisms in different biological responses. However, the mechanisms through which MFA inhibits ethanol-induced L-02 cell apoptosis are unclear. The present study demonstrated that MFA decreased apoptosis by inhibiting the activation of the p38 MAPK and JNK pathways in L-02 cells. The data identified that MFA treatment caused the persistent inactivation of p38 MAPK and JNK in L-02 cells (Fig. 4C). The signaling proteins in the MAPK family promote a variety of biological reactions in cells and p38 MAPK and JNK play a key role in the signal transduction of apoptosis (49). Since the release of cytochrome c from damaged mitochondria is a key step in the activation of caspases, the finding that JNK and p38 MAPK activities are essential for caspase activation indicates that JNK and p38 MAPK can regulate some of the other mitochondrial-related factors (e.g., Bax). In L-02 cells, MFA treatment caused the downregulation of JNK and p-38 phosphorylation (Fig. 4C), which is in part similar to the results obtained in I/R-induced hepatocytes treated with...
FA (24). Briefly, MFA attenuated the apoptosis of L-02 cells induced by ethanol via the inhibition of the ROS-dependent JNK/p38 MAPK signaling pathway. However, further studies are required to elucidate the mechanisms through which MFA inhibits NOX4 to reduce ROS generation.

In brief, the results of the present study demonstrated that MFA suppressed ethanol-triggered oxidative stress in L-02 cells by inhibiting ROS production and the upregulation of GSH-PX, CAT and SOD. The role of MFA in modulating MAPK phosphorylation, and its established antioxidant effect, suggest that this compound may prove to be a good candidate for use in the treatment of ALD.

In conclusion, the present study demonstrated for the first time, to the best of the authors' knowledge, that MFA exerts beneficial effects against the ethanol-induced lack of viability and apoptosis by inhibiting ROS-dependent JNK, the p38 MAPK signaling pathway and the Bcl-2/Bax signaling pathway in L-02 cells. MFA treatment led to the upregulation of p-JNK, p-p38 MAPK and Bax, and the downregulation of Bcl-2 and cleaved caspase-3. A new understanding of the role of MFA was developed in that it attenuated the apoptosis of L-02 cells induced by ethanol by inhibiting the activation of the MAPK signaling pathway. In the present study, the L-02 cells used in the in vitro experiments were normal human hepatocytes. Whether MFA treatment can inhibit the apoptosis of hepatocytes in vivo requires further investigation. The present study provides a novel theoretical basis for the possible use of MFA in the treatment of liver injury.

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LL and YZ wrote manuscript, performed experiments and analyzed data; CY, CL, LC, HW and MZR performed experiments and analyzed data. ZM, CY, LL, HW and YL analyzed the data. ZM and YL contributed to the discussions and critically edited the manuscript. All authors have reviewed and approved the manuscript. All authors are responsible for the integrity of the data.

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