Anti-fibrotic effects of Acremoniumterricola milleretal mycelium on immunological hepatic fibrosis in rats

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Abstract. Acremoniumterricola milleretal mycelium (AMM) exerts numerous protective effects on organs, and has been used in Chinese herb prescriptions to treat refractory diseases. The aim of this study was to investigate the effects of AMM on immunological hepatic fibrosis induced by porcine serum (PS) in rats. Male Sprague Dawley rats were administered 0.5 ml sterile PS by intraperitoneal injections twice a week for 18 weeks. AMM (175, 350 or 700 mg/kg) and colchicine (0.1 mg/kg) were administered intragastrically each day until the rats were sacrificed. PS administration resulted in marked hepatic fibrosis, as assessed by increased oxidative stress and hepatic collagen content, as well as α-smooth muscle actin (α-SMA) expression. AMM significantly reduced liver damage and fibrosis. In addition, AMM decreased the elevation in hydroxyproline, hyaluronic acid, laminin and procollagen type III; increased the activity of superoxide dismutase and glutathione peroxidase; decreased α-SMA expression; and eliminated hepatic collagen deposits. Furthermore, AMM inhibited Smad2/3 phosphorylation and Smad7 expression. These results indicate that AMM is able to reduce oxidative stress, inhibit collagen synthesis and block the transforming growth factor-β/Smad signaling pathway in a dose-dependent manner.

Introduction

Hepatic fibrosis is a common pathology in China that severely impacts the health of affected individuals and poses a significant risk of morbidity. Fibrosis of the liver occurs predominantly through the accumulation of collagen, proteoglycans and other macromolecules within the extracellular matrix (ECM). Both the quantity and composition of the ECM are often markedly altered in patients with liver fibrosis (1,2). Chronic liver disease is known to progress from mild to severe inflammation, and then to fibrosis and cirrhosis. Hepatic fibrosis therefore occurs between chronic liver disease and cirrhosis, and constitutes a dynamic and bidirectional stage (3). In the present study, porcine serum (PS)-induced hepatic fibrosis was used as a unique model of liver fibrosis in the absence of obvious hepatocyte injury or inflammatory cell infiltration (4,5). This liver fibrosis model, induced by an immunological reaction, was similar to models generated by repeat injections of equine serum, egg yolk and human albumin, but not rat serum (6). Previous research has indicated that hepatic fibrosis is a complex pathological process involving numerous cytokines and cell signaling pathways (7). The activation of hepatic stellate cells (HSCs), which produce collagen, is considered to be induced in a paracrine manner through the mediation of various factors released from necrotic hepatocytes, Kupffer cells or endothelial cells (8). Domitrovic and Jakovac (9) suggested that liver fibrosis decreased through the inactivation of HSCs under the control of fibrogenic cytokines. Transforming growth factor (TGF)-β1 has a pivotal role in liver fibrosis, and can activate HSCs, the principal cellular source of excess ECM during hepatic fibrosis, though the TGF-β/Smad signaling pathway (10).

Colchicine has been widely used in clinical practice for the treatment of acute gout and other immunological diseases, and therefore was used as a positive control in the present study. In China, it is believed that patients with liver fibrosis have been treated with herbal medicines for thousands of years. Traditional Chinese medicine is still extensively used for the treatment of liver disease. In recent years, considerable attention has been paid to the use of traditional Chinese medicine in the treatment of liver fibrosis and cirrhosis (11). Acremoniumterricola milleretal mycelium (AMM) is isolated from Acremonium terricola, in submerged fermentation. AMM consists of a number of beneficial components, including Cordyceps polysaccharide, palmitic acid and unsaturated fatty acids, and has been used for its anti-inflammatory and antioxidant effects, as well as for regulation of the immune...
system. This Chinese medicine has been reported to exert various pharmacological effects on animals both in vitro and in vivo (12). Our previous study demonstrated that AMM has protective effects on carbon tetrachloride-induced liver fibrosis in rats (13). The present study was performed to determine whether AMM has any beneficial effects on PS-induced immunological hepatic fibrosis.

Materials and methods

Materials. AMM was supplied by Anhui Haikui Biotechnology Company (Hefei, China). Colchicine was obtained from Sigma-Aldrich (St. Louis, MO, USA). PS was purchased from Zhengzhou YiKang Biotechnology Company (Zhengzhou, China). Commercial kits used to assay alanine transaminase (ALT), aspartate transaminase (AST), superoxide dismutase (SOD), glutathione peroxidase (GSH-Px) and hydroxyproline (Hyp) were obtained from Nanjing Jiancheng Institute of Biotechnology (Nanjing, China). The hyaluronic acid (HA), laminin (LN) and procollagen type III (PCH) radioimmunoassay kits were purchased from Beijing North Institute of Biotechnology (Beijing, China). The α-SMA and anti-β-actin antibodies, as well as rabbit anti-phosphorylated (p) Smad2, anti-Smad2, anti-p-Smad3, anti-Smad3 and anti-Smad7 antibodies were purchased from Bioworld Technology, Inc., (St. Louis, Park, MN, USA).

Animals. Male Sprague Dawley rats, weighing 130-150 g, were provided by the Experimental Animal Center of Anhui Province (Hefei, China). Animals were housed five per cage, with access to water and food ad libitum, and maintained at a constant temperature (22±1°C) and humidity (50±20%) under a 12-h light/dark cycle. Animal treatment and maintenance were carried out in accordance with the guidelines for the humane treatment of animals set by the Association of Laboratory Animal Sciences and the Center for Laboratory Animal Sciences, Anhui Medical University (Hefei, China).

Experimental design. After 1 week of acclimation, the animals were subjected to experimentation. Sixty adult male Sprague Dawley rats were randomly divided into six groups, as follows: Control, model (PS-treated), PS plus colchicine (0.1 mg/kg) and PS plus AMM (175, 350 and 700 mg/kg, respectively). All groups, with the exception of the control, received PS intraperitoneally twice per week for 18 weeks. The colchicine and AMM groups were treated with colchicine (0.1 mg/kg, intragastrically) and AMM (175, 350 or 700 mg/kg, intragastrically) each day, respectively, at the beginning of the injection of PS. The control and model groups were administered the same volume of vehicle.

The procedure of dividing the animals into groups and for the generation of the PS-induced model of liver fibrosis was based on the method described previously, with certain modifications (14,15). Animals were weighed once per week, and 24 h after the final injection of PS all animals were sacrificed under anesthesia with ether. Blood samples were collected from the abdominal aorta and centrifuged (3,000 x g for 10 min), and the serum was stored at -80°C until further analysis. The liver was subsequently washed in situ with ice-cold isotonic saline, then removed and divided into two portions; one portion was fixed for histopathology and the other was immediately frozen at -80°C until required.

Hepatotoxicity studies. To assess hepatotoxicity, the serum levels of ALT and AST were measured using commercial kits according to the manufacturer’s instructions (Nanjing Jiancheng Institute of Biotechnology).

Hepatic Hyp. The Hyp content in fresh liver samples was measured in accordance with the methods described by Jamall et al (16). The content of hepatic Hyp was determined using a Hyp kit following the manufacturer’s instructions (Nanjing Jiacheng Bioengineering Institute).

Oxidative stress. The livers were thawed and washed with normal saline to remove blood and clots. Homogenates were centrifuged (1,000 x g for 10 min, 4°C) and aliquots of the supernatants were then used to assay the expression levels of SOD and GSH-Px.

Serum fibrotic markers. The serum levels of HA, LN and PCH were assayed using a radioimmunoassay kit (North Institute of Biotechnology, Beijing, China) according to the manufacturer’s instructions.

Histopathology. A portion of the liver specimens of each rat was fixed in 10% neutral buffered formalin and embedded in paraffin. Sections measuring 4 µm were then cut and stained with hematoxylin and eosin (H&E) and Sirius Red. The stained slides were examined independently by two pathologists with no prior knowledge of their source. The histological grade of hepatic fibrosis was assessed based on the New Inuyama staging system (17): 0, no fibrosis (normal liver and absence of fibrosis); I, fibrosis present (collagen fibers present that extend from the portal triad or the central vein to the peripheral region); II, mild fibrosis (mild collagen fiber presence with extension without compartment formation); III, moderate fibrosis (mild collagen fibers present with moderate pseudo-lob formation); IV, severe fibrosis (severe collagen fiber presence with thickening of the partial compartments and frequent pseudo-lob formation) (18). The percentage of area occupied by collagen fibrosis was calculated by dividing the number of red collagen fibroses by the total number of collagen fibroses. At least five fields were selected under light microscopy at x200 magnification to determine positive collagen fibrosis, and the average was calculated.

Immunohistochemistry. Immunohistochemical analyses were performed on paraffin sections incubated with primary mouse monoclonal α-SMA antibody, diluted 1:100, at 4°C overnight. The sections were then incubated with biotinylated secondary goat anti-mouse immunoglobulin G antibody (Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA), prior to the sections being counterstained with hematoxylin. Normal mouse antiserum was used as a negative control in place of the primary antibody, and consistently demonstrated no antigenic reaction. The numbers of α-SMA-positive and -negative HSCs were visualized using diaminobenzidine (Dako, Tokyo, Japan). Five fields were randomly selected from each section and the number of positive cells for each antibody was counted under...
light microscopy at x400 magnification. The percentage of α-SMA-positive HSCs was calculated by dividing the number of HSCs counted in each slide by the total number of cells, and an average number for each group was calculated.

**Western blotting.** Frozen liver tissue (100 mg) was washed several times in distilled water and homogenized in extraction buffer (25 mM HEPES, 400 mM KCl, 1 mM EDTA and 1.5 mM MgCl₂). The homogenate was then centrifuged at 12,000 x g for 10 min at 4˚C, and the supernatant was transferred to a fresh tube and stored at -80˚C. The concentration of protein homogenate was determined using a bicinchoninic acid protein assay kit according to the manufacturer's instructions (ZSGB-BIO, Beijing, China). Proteins were assayed using a Tanon-4200 automatic digital gel imaging system (Tianneng, Shanghai, China) (19). Protein samples (50-150 µg) were separated by SDS-PAGE and transferred to polyvinylidene difluoride membranes for 2 h at 100 V (Millipore, Bedford, MA, USA). The membranes were then incubated overnight at 4˚C under agitation with the following primary antibodies (diluted 1:800 in Tris-buffered saline with Tween): Anti-p-Smad2, -p-Smad3, -Smad2, -Smad3, and -Smad7 (Bio-world Technology, Inc.). The membranes were subsequently incubated with secondary antibody (Santa Cruz Biotechnology, Inc.) for 2 h at room temperature, and the immunoreactive proteins were visualized using enhanced chemiluminescence (Amersham Pharmacia Biotech, Piscataway, NJ, USA) and autoradiography. Gel reverse zymography was performed on the liver of five randomly selected rats per group. Quantity One® software (Bio-Rad, Hercules, CA, USA) was used to quantify the band densities.

**Statistical analysis.** Data were expressed as the mean ± standard deviation. Data were analyzed by a one-way analysis of variance, and the Student’s t-test was used for two-group comparisons. A Ridit test was used for the statistical analysis of qualitative data. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**Serum markers of liver damage and antioxidant status.** The effects of treatment with AMM on the PS-induced elevation of serum markers due to hepatic damage are shown in Table I. The serum activities of AST and ALT were both increased in the PS-treated group as compared with those in the control group, but the difference did not reach statistical significance (P>0.05). The activities of SOD and GSH-Px in the PS-treated group were significantly lower as compared with those in the control group (P<0.001). AMM (350 and 700 mg/kg) only moderately reduced the AST and ALT activities relative to the control group (P<0.05) but significantly enhanced SOD and GSH-Px activities as compared with the PS-treated group (P<0.05 and P<0.01, respectively) (Table I).

**Hepatic Hyp content and serum fibrotic marker levels.** The Hyp content and HA, LN and PCⅢ levels were markedly increased in the PS-treated group as compared with those in the control group (P<0.01). AMM (350 and 700 mg/kg) significantly attenuated the increase in the content of Hyp, and reduced the levels of serum HA, LN and PCⅢ as compared with the PS-treated group (P<0.01 and P<0.001) (Table II).

**Histopathological changes in the liver.** The histopathological changes in the liver are shown in Fig. 1. The structure of the liver tissue remained relatively normal (Fig 1A). The livers from the control group stained with Sirius Red stain showed traces of collagen only in the walls of major blood vessels (Fig. 1G). PS administration resulted in an extensive accumulation of collagen in the liver tissue. Fibrosis predominantly developed around the central veins near the hepatic capsule, and complete septal fibrosis was observed; furthermore, pseudo-lobate formation was apparent in severe cases. Only a small number of inflammatory cells were found around the portal area and central vein, without notable hepatocyte necrosis (Fig. 1B and H). AMM (350 and 700 mg/kg) treatment resulted in a dose-dependent decrease in fibrotic deposits with short fibrous septa, and markedly reduced the pathological changes as compared with the PS-treated group. The structure of the liver tissue remained relatively normal (Fig. 1E, F, K and L).

The PS-treated group showed a high degree of fibrosis. AMM (350 and 700 mg/kg) and colchicine (0.1 mg/kg) treatment resulted in a marked improvement in the histological scores in comparison with the PS-treated group (Table III).

### Table I. Effects of AMM on serum markers of liver damage and antioxidant status in PS-induced hepatic fibrosis.

<table>
<thead>
<tr>
<th>Group</th>
<th>Dose (mg/kg) of AMM/colchicine</th>
<th>ALT (U/l)</th>
<th>AST (U/l)</th>
<th>SOD (U/mg)</th>
<th>GSH-Px (U/mg)</th>
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<tbody>
<tr>
<td>Control</td>
<td>-</td>
<td>56.08±12.26</td>
<td>64.50±15.76</td>
<td>134.02±17.23</td>
<td>123.03±16.18</td>
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<tr>
<td>PS-treated</td>
<td>-</td>
<td>67.35±13.68</td>
<td>81.13±18.84</td>
<td>97.61±15.25</td>
<td>64.95±10.89</td>
</tr>
<tr>
<td>PS+AMM</td>
<td>175</td>
<td>63.31±14.43</td>
<td>74.59±13.52</td>
<td>108.35±11.05</td>
<td>71.04±15.78</td>
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<tr>
<td></td>
<td>350</td>
<td>61.31±12.51</td>
<td>71.23±12.94</td>
<td>117.43±19.85</td>
<td>85.32±18.06</td>
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<tr>
<td></td>
<td>700</td>
<td>59.74±14.62</td>
<td>69.35±14.12</td>
<td>125.45±26.21</td>
<td>100.73±17.56</td>
</tr>
<tr>
<td>PS+colchicine</td>
<td>0.1</td>
<td>62.16±15.26</td>
<td>73.36±17.72</td>
<td>127.67±22.58</td>
<td>83.79±13.34</td>
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</tbody>
</table>

*a*P<0.001 vs. the control group; *b*P<0.05 and *c*P<0.01 vs. the PS- treated group. n=8 per group. Data are presented as the mean ± standard deviation. AMM, Acremonium terricola milleretal mycelium; PS, porcine serum; ALT, alanine transaminase; AST, aspartate transaminase; SOD, superoxide dismutase; GSH-Px, glutathione peroxidase.
Immunohistochemical expression of α-SMA. The immunohistochemical expression of α-SMA is shown in Fig. 2. Few cells were detected to be positively stained for α-SMA in the control group (Fig. 2A). The percentage of α-SMA-positive cells was significantly increased in the PS-treated group as compared with that in the control group (P<0.001), with most...
of the α-SMA-positive cells detected around the periportal fibrotic band areas, central vein and fibrous septa (Fig. 2B). AMM (350 and 700 mg/kg) significantly decreased the percentage of α-SMA-positive HSCs among the groups, (A) Control group; (B) PS-treated group; (C) PS and colchicine-treated group (0.1 mg/kg colchicine); (D-F) PS and AMM-treated groups (175, 350 and 700 mg/kg AMM, respectively). Arrows show α-SMA immunopositive cells. *P<0.01 vs. the control group; **P<0.001 vs. the model group. Original magnification, x400. AMM, Acremoniumterricola milleretal mycelium; PS, porcine serum.

Table III. Effects of AMM on pathological grading in PS-induced hepatic fibrosis.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Dose (mg/kg) of AMM/colchicine</th>
<th>Pathological grading of hepatic fibrosis (n)</th>
<th>Collagen area (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>0</td>
<td>I</td>
</tr>
<tr>
<td>Control</td>
<td>-</td>
<td>8</td>
<td>0</td>
</tr>
<tr>
<td>PS-treated</td>
<td>-</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>PS+AMM 175</td>
<td>0.1</td>
<td>0</td>
<td>2</td>
</tr>
<tr>
<td>PS+AMM 350</td>
<td>0.1</td>
<td>0</td>
<td>2</td>
</tr>
<tr>
<td>PS+AMM 700</td>
<td>0.1</td>
<td>0</td>
<td>2</td>
</tr>
<tr>
<td>PS+colchicine</td>
<td>0.1</td>
<td>0</td>
<td>3</td>
</tr>
</tbody>
</table>

Results are the average from 10 fields of vision. *P<0.001 vs. the control group; ªP<0.05 and ñP<0.001 vs. the PS-treated group. Data are presented as the mean ± standard deviation. AMM, Acremoniumterricola milleretal mycelium; PS, porcine serum.

Figure 2. Expression and tissue distribution of α-SMA. Representative photomicrographs of five liver sections and a comparison of the number of α-SMA-positive HSCs among the groups. (A) Control group; (B) PS-treated group; (C) PS and colchicine-treated group (0.1 mg/kg colchicine); (D-F) PS and AMM-treated groups (175, 350 and 700 mg/kg AMM, respectively). Arrows show α-SMA immunopositive cells. *P<0.01 vs. the control group; **P<0.001 vs. the model group. Original magnification, x400. AMM, Acremoniumterricola milleretal mycelium; PS, porcine serum; α-SMA, α-smooth muscle actin; HSCs, hepatic stellate cells.

Expression of Smad2/3 phosphorylation and Smad7. A higher expression of p-Smad2/3 was observed in the PS-treated group as compared with the control group (P<0.001) (Fig. 3A and B). Conversely, the expression of Smad7 was significantly decreased in the PS-treated group (P<0.001) (Fig. 3C). AMM (350 and 700 mg/kg) and colchicine (0.1 mg/kg) significantly decreased the expression of p-Smad2 and p-Smad3.
and increased the expression of Smad7 (P<0.05 and P<0.01) (Fig. 3A-C). These results demonstrated that AMM (350 and 700 mg/kg) may act to prevent hepatic fibrosis by blocking Smad2/3 phosphorylation and enhancing expression of the inhibitor Smad7 in the TGF-β/Smad signaling pathway.

Discussion

Hepatic fibrosis is a common consequence of almost all causes of chronic liver disease. HSCs, the predominant ECM-producing cells, are activated by fibrogenic cytokines, such as TGF-β1, angiotensin II and leptin. It was originally believed that hepatic fibrosis was an irreversible process due to disruption of the normal liver architecture (20); however, hepatic fibrosis is more recently considered to be a reversible wound-healing response to chronic liver injury (21). The development of antifibrotic drugs to restrain fibrogenesis is of particular importance. The present study showed by H&E staining that low numbers of inflammatory cells were present around the portal area and central vein without notable hepatocyte necrosis, in the PS-treated (model) group. This was consistent with the data showing no significant elevation in ALT and AST levels. Sirius Red staining can be used to detect varying degrees of hepatic fibrosis and to score this fibrosis (22). The results showed that AMM (350 and 700 mg/kg) markedly attenuated the pathology grading, which was consistent with the findings for Hyp content. SOD and GSH-Px are two primary enzymes of the antioxidant defense system (23). The results presented in this study indicated that AMM could significantly enhance the levels of SOD and GSH-Px. The analysis of Hyp content was used to directly determine the quantity of collagen in the tissue and can be used as a measure to reflect the degree of hepatic fibrosis (24). Additionally, HA, LN and PCIII in serum are considered to be important biomarkers of hepatic fibrogenesis (25). In this study, AMM (350 and 700 mg/kg) significantly decreased the Hyp content and the levels of HA, LN and PCIII in PS-induced hepatic fibrotic rats, suggesting that this may be the mechanism underlying its hepatoprotective effects.

HSCs have a key function in the pathogenesis of fibrosis (26,27). Upon the activation of HSCs, levels of fibrillar collagen, particularly types I and II, become markedly increased. α-SMA is an indicator of activated HSCs (28). Therefore, the expression level of α-SMA was assessed among the groups. Few α-SMA-positive HSCs were detected in the AMM (350 and 700 mg/kg) treatment groups. Thus, AMM could inhibit the expression of α-SMA-positive HSCs and subsequently prevent hepatic fibrosis.

TGF-β1 activates HSCs and promotes ECM production, which regulates the fibrogenic process. TGF-β1 protein is stored in an inactive form. Once activated, TGF-β1 signals via its cognate receptors to Smad proteins (29). The TGFβ1/Smad signaling pathway has a central role in hepatic fibrosis. Through this pathway, TGFβ1 predominantly activates HSCs, which can result in hepatic fibrosis (30,31). Activated TGF-β1 binds to the constitutively active TGF-β Type II (TβRII) receptor in the cell membrane, which leads to the recruitment of the TβRI receptor. A heterotetrameric complex is formed and results in the phosphorylation and activation of the TβRII receptor. A heterotetrameric complex is formed and results in the phosphorylation and activation of the TβRI receptor. Activated TβRI in turn phosphorylates Smad2 and Smad3, known as receptor-activated Smads (R-Smads), which form a hetero-oligomeric complex with a common mediator, Smad4 (Co-Smad). This complex translocates to the nucleus, where it can regulate the transcription of target genes (32). Smad7 is an inhibitory Smad (I-Smad) that acts differently to the signal-transducing R-Smads and Co-Smads. Smad7 competitively associates with TβRII to prevent the
phosphorylation of Smad2 and Smad3 and therefore inhibit the signal transduction of TGFβ1. A previous study showed that Smad7 overexpression inhibited TGFβ1/Smad signaling in rat liver fibroblasts (33). The present study results showed that AMM (350 and 700 mg/kg) treatment markedly suppressed the expression of p-Smad2, and p-Smad3 and increased that of Smad7, indicating that AMM had a significant effect on TGFβ1/Smad signaling.

In conclusion, the present findings indicate that AMM plays an important role in the inhibition of PS-induced immunological hepatic fibrosis. The molecular mechanism of this therapeutic effect could be due to a decrease in oxidative stress, a reduction in the accumulation of collagens and the inhibition of TGFβ1/Smad signaling.

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