

effects of the selective mineralocorticoid receptor antagonist eplerenone on hepatic fibrosis induced by bile duct ligation in rat

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Received January 12, 2010; Accepted February 17, 2010

DOI: 10.3892/ijmm_00000417

Abstract. The aim of this study was to examine the effects of eplerenone on hepatic fibrosis induced by bile duct ligation (BDL) in rat. Low- (1.0 mg/kg body weight, BW) and high- (4.0 mg/kg BW) dose eplerenone was administered orally for 21 days immediately following BDL. Fibrosis was assessed by measuring the fibrotic area after Sirius red staining. Immunostaining for α -smooth muscle actin (SMA), 4-hydroxy-2-nonenal (4-HNE) and 8-hydroxy-2-deoxyguanosine (8-OHdG) was also carried out. Gene expression levels of procollagen-I, transforming growth factor- β 1 (TGF- β 1), connective tissue growth factor (CTGF), tissue inhibitor of metalloproteinases-1 (TIMP-1) and matrix metalloproteinase-13 (MMP-13) in the liver were examined by real-time reverse transcriptase polymerase chain reaction. Plasma angiotensin II (ATII) concentration was measured via radioimmunoassay. The area of hepatic fibrosis and α -SMA positivity in the high-dose group was significantly decreased compared with that in the BDL group, but not in the low-dose group. 8-OHdG-positive cells in the low- and high-dose groups were significantly decreased compared with those in the BDL group. Immunostaining of 4-HNE in the high-dose group was significantly lower compared with that in the BDL group. Furthermore, TIMP-1 mRNA levels in the low- and high-dose groups were lower than that in the BDL group. The expression of TGF- β 1, CTGF, procollagen-1 and MMP-13 showed no differences. Plasma ATII concentration in the high-dose group was significantly decreased. Eplerenone attenuated the development of BDL-induced hepatic fibrosis by reducing oxidative stress, suppressing activated hepatic stellate cells and decreasing plasma ATII levels. Eplerenone may prove useful as an alternative treatment for antifibrosis therapy.

Introduction

Angiotensin II (ATII) and aldosterone are the main effector peptides of the renin-angiotensin-aldosterone system (RAAS) and are known to be important mediators of hepatic fibrosis and portal hypertension (1-5). ATII and aldosterone stimulate the proliferation of hepatic stellate cells (HSC) and collagen synthesis, and are involved in the generation of reactive oxygen species (ROS) both *in vitro* and *in vivo* (6-8). ATII type 1 receptor blockers (ARB) and angiotensin-converting enzyme inhibitors (ACEI) have been reported to prevent the development of hepatic fibrosis in numerous animal (6-12) and human (13-15) studies. However, only a few studies examining the antifibrotic effects of aldosterone blockers have been reported to date. The results arising from these studies have proven to be controversial (1,2,5). Fujisawa *et al* (5) reported that spironolactone prevented pig serum-induced hepatic fibrosis in rats, while Oberti *et al* (2) showed that spironolactone did not produce any anti-fibrotic effects during bile duct ligation (BDL)-induced hepatic fibrosis.

The novel aldosterone blocker eplerenone has been shown to exhibit a high specificity for mineralocorticoid receptors (16,17). While the efficacy and safety of eplerenone in the treatment of hypertension have been demonstrated in a number of previous studies (18-20), no studies have reported the effects of eplerenone on hepatic fibrosis. Cholestatic fibrosis induced by BDL has been shown to be a suitable experimental model of human liver diseases including primary and secondary biliary cirrhosis and primary sclerosing cholangitis. Our previous study has demonstrated that components of the RAAS are up-regulated during BDL-induced liver fibrosis, and that RAAS plays an important role in the progression of hepatic fibrosis (6). In addition, oxidative stress and pro-inflammatory cytokinesis play essential roles in the development of cholestatic liver fibrosis (21). In the present study, we examined the effects of eplerenone during cholestatic liver fibrosis induced by BDL in rats.

Materials and methods

Animals. Male Wistar rats were obtained from Japan SLC (Shizuoka, Japan) and housed in a room maintained at a controlled temperature of $24\pm 2^{\circ}\text{C}$ under a 12-h light-dark cycle. Animals were provided standard pellet chow and water

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Key words: bile duct ligation, eplerenone, oxidative stress

ad libitum. BDL operations were performed under anesthesia following a 50 mg/kg intraperitoneal injection of pentobarbital (Dainippon Pharmaceutical, Osaka, Japan). All experiments were carried out in accordance with the Animal Experimentation Guidelines of Tottori University.

Establishment of the BDL model and administration of eplerenone. Wistar male rats (192-231 g body weight) were used in this study and hepatic fibrosis was induced by BDL as previously described (4,6). We used the 3-week-old BDL rat model to study the effects of eplerenone on the development of hepatic fibrosis. Rats in the BDL + low-dose eplerenone group (low-dose group) were administered eplerenone orally at a dose of 1.0 mg/kg, which is an ordinary dose for humans, via a gastric tube every day for 21 days immediately following the operation (survival rate, n=8/10). Rats in the BDL + high-dose eplerenone group (high-dose group) were administered eplerenone orally at a dose of 4.0 mg/kg every day for 21 days immediately following the operation (survival rate, n=9/10). The BDL group received vehicle for the same period (survival rate, n=9/10). Vitamin K (5 mg/kg) was injected intramuscularly once a week to all groups to prevent the occurrence of fatal hemorrhage (22). In addition, six rats were carried out by sham-operation.

On day 21, animals were sacrificed under pentobarbital anesthesia and blood was collected from the inferior vena cava. Serum and plasma samples were frozen and stored at -80°C until required. Total serum bilirubin (T-bil), aspartate aminotransferase (AST) and alanine aminotransferase (ALT) levels were measured using an autoanalyzer (7170, Hitachi, Tokyo, Japan). Plasma renin, ATII and aldosterone levels were measured using a radioimmunoassay. The liver was also collected, weighed, snap-frozen in liquid nitrogen and stored at -80°C until required. Liver specimens were also fixed in 10% buffered formalin and embedded in paraffin for histological analysis.

Assay for hepatic hydroxyproline (Hyp) content. Hepatic tissue (200 mg wet weight) was hydrolyzed in 4 ml of 6N HCl at 105°C overnight. Hydrolysate was evaporated under a vacuum four times. Sediment was redissolved in distilled water, mixed with activated charcoal for decolorization, and then filtered. The solution was kept in the acidic range by adjustment to pH 5.0 and evaporated under a vacuum, and sediment was redissolved in distilled water, supplemented with 2 ml of isopropanol, and incubated with 1 ml of 7% chloramine-T solution for 5 min at room temperature. Ehrlich's solution (2 ml), consisting of 1.76 g of p-dimethylaminobenzaldehyde dissolved in 4.08 ml of 60% perchloric acid and 95.5 ml of isopropanol, was added, and the mixture was incubated at 60°C for 10 min. After cooling, the absorbance measured at 562 nm (23).

Measurement of the hepatic fibrosis area. Formalin-fixed, paraffin-embedded liver (4 µm-thick sections) were stained with picosirius red (Sigma-Aldrich, St. Louis, MO) and counter-stained with fast green (Sigma-Aldrich). The area of hepatic fibrosis was then measured in 10 randomly selected fields/section (magnification x100) per specimen using WinRoof version 5.71 (Mitani Shoji, Fukui, Japan).

Table I. Primers used for real-time PCR.

Gene	Accession no.	Primers
β-actin	BC063166	F: 5'-CCCGCGAGT ACAACCTTCT-3' R: 5'-CGTCATCC ATGGCGAACT-3'
Procollagen-I	Z78279	F: 5'-ATGTTTCAGCT TTGTGGACCTC-3' R: 5'-GCAGCTGAC TTCAGGGATGT-3'
TGF-β1	X52498	F: 5'-CCTGGAAAG GGCTCAACAC-3' R: 5'-CAGTTCCTTCTC TGTGGAGCTGA-3'
TIMP-1	U06179	F: 5'-CAGCAAAAG GCCTTCGTAAA-3' R: 5'-TGGCTGAAC AGGGAAACACT-3'
CTGF	AB023068	F: 5'-GCTGACCTAGAG GAAAACATTAAGA-3' R: 5'-CCCGGTAGG TCTTCACACTG-3'
MMP-13	M60616	F: 5'-GGACAAGC AGCTCCAAAGG-3' R: 5'-GGTCCAGA CCGAGGGAGT-3'

F, forward; R, reverse; TGFβ-1, transforming growth factor β1; TIMP-1, tissue inhibitor of metalloproteinases-1; CTGF, connective tissue growth factor; MMP, matrix metalloproteinase.

α-smooth muscle actin (α-SMA) immunostaining. Immunohistochemical staining of α-SMA was performed using anti-α-SMA antibody (Dako A/S, Glostrup, Denmark). To assess the activation grade of hepatic stellate cells, the area of α-SMA staining was analyzed using WinRoof version 5.71 in 10 randomly selected fields/section (magnification x100) per specimen.

RNA extraction and reverse transcription polymerase chain reaction (PCR) analysis. Tissue samples were homogenized and total RNA was extracted using the RNeasy Mini Kit (Qiagen, Hilden, Germany). RNA concentration was determined by measuring absorbance at 260 nm, and the quality of RNA was verified by electrophoresis on ethidium bromide-stained 1% agarose gels. Total RNA (~2 µg) was then reverse transcribed (RT) in a final volume of 11.5 µl containing 4 µl of 5x standard buffer, 2 µl of 0.1 M dTT, 1 µl of SuperScript II RNase H-Reverse Transcriptase (Invitrogen, Carlsbad, CA), 2 µl of 10 M MdNTPs (Promega, Madison, WI), 1 µl of 50 pmol/µl Random Primer (Promega), 0.5 µl of 100 pmol/µl Oligo (dt)15 Primer (Promega) and 1 µl of 40 U/µl ribonuclease inhibitor (Wako Pure Chemical Industries, Ltd., Osaka, Japan). Samples were then incubated at 37°C for 60 min, 95°C for 5 min and cooled to 4°C for 5 min.

	Sham n=6	BDL group n=9	Low-dose group n=8	High-dose group n=9
Serum T-bil (mg/dl)	0	2.1±1.8	2.1±1.9	3.0±1.9
Serum AST (IU/l)	109±45	331±218	378±262	533±322
Serum ALT (IU/l)	45±18	58±16	59±22	108±53 ^a
Plasma Renin (pg/ml)		33.6±8.5	21.9±9.9	26.7±11.5
Plasma Angiotensin-II (pg/ml)		237±145	170±74	120±45 ^a
Plasma Aldosterone (pg/ml)		750±500	509±274	757±539

Values are presented as the mean ± standard deviation; T-bil, total bilirubin; AST, aspartate aminotransferase; ALT, alanine aminotransferase. ^ap<0.05, compared with the BDL group determined by Mann-Whitney test.

Table III. Hydroxyproline content and expression of procollagen I, TGF-β1, TIMP-1, CTGF and MMP-13 mRNA in the liver at 3 weeks following BDL.

	Sham n=6	BDL group n=9	Low-dose group n=8	High-dose group n=9
Hyp (μg/g wet liver)	129±89	602±152	541±278	446±169 ^b
Procollagen I (x10 ⁻²)		18.1±6.9	5.5±4.3	4.8±4.6
TGF-β1 (x10 ⁻²)		2.7±0.9	2.2±0.7	2.6±1.1
TIMP-1 (x10 ⁻²)		5.1±1.1	2.3±1.7 ^a	3.6±2.1 ^b
CTGF (x10 ⁻²)		5.0±3.0	3.0±3.0	4.0±3.0
MMP-13 (x10 ⁻²)		1.2±1.3	1.1±0.5	1.4±0.7

Values are presented as the mean ± standard deviation. Hyp, hydroxyproline; TGF-β1, transforming growth factor-β1; TIMP-1, tissue inhibitor of metalloproteinases-1; CTGF, connective tissue growth factor; MMP-13, matrix metalloproteinase-13. ^ap<0.01 and ^bp<0.1, compared with the BDL group.

Real-time PCR. Quantitative real-time PCR was performed in a final volume of 10 μl containing 4.1 μl PCR grade water, 1 μl Universal Probe Library probe (Roche, Tokyo, Japan), 0.2 μl forward primer (10 μM), 0.2 μl reverse primer (10 μM), 2 μl Light Cycler TaqMan Master (Roche) and 2.5 μl cDNA. The mRNA levels of transforming growth factor-β1 (TGF-β1) (GenBank, X52498), connective tissue growth factor (CTGF) (GenBank; AB023068), tissue inhibitor of metalloproteinases-1 (TIMP-1, GenBank, U06179), matrix metalloproteinase-13 (MMP-13, GenBank, M60616) and procollagen-type I (GenBank, Z78279) were assessed using the Light Cycler Fast Start DNA Master SYBR Green 1 real-time PCR assay (Roche) with β-actin (BC063166) as the housekeeping gene (Table I). The thermal cycle conditions were as follows, hold at 95°C for 10 min, repeat 45 cycles of 95°C for 30 sec and 60°C for 1 min.

Analysis of oxidative stress and lipid peroxidation. Oxidative stress was evaluated by measuring the levels of 8-hydroxy-2-deoxyguanosine (8-OHdG), a marker of oxidative DNA damage (24), using immunohistochemical staining and enzyme-linked immunosorbent assay (ELISA). Immunohistochemical staining of 8-OHdG was performed using an anti-8-OHdG monoclonal antibody (Nikken Seil, Shizuoka, Japan) in accordance with the manufacturer's instructions. Analysis of immunopositive cells was carried out in 10 intralobular ocular

fields (magnification x400) per specimen, and was expressed as a percentage of fields using the WinRoof version 5.71.

For ELISA, frozen liver tissues were homogenized and DNA was extracted using the Nal method and the DNeasy Mini Kit. After the DNA pellet was dissolved in TE buffer, 50 μg of DNA was digested with nuclease P1 (Sigma-Aldrich) and alkaline phosphatase (Sigma-Aldrich), and centrifuged at 14,000 g for 10 min through a Microcone YM-10 filter (Millipore, Bedford, MA) according to the manufacturer's instructions. The 8-OHdG content in the extracted DNA solution was then determined using the highly sensitive 8-OHdG check ELISA kit (Nikken Seil). The absorbance of each well was measured at 450 nm using a microplate reader.

Thiobarbituric acid reactive substance (TBARS) was quantified as described previously (25). Immunohistochemical staining of 4-hydroxy-2-nonenal (4-HNE) was performed using an anti-4-HNE monoclonal antibody (Nikken Seil) according to the manufacturer's instructions. Histological images of 4-HNE staining were randomly selected in 10 fields/section (magnification x100) per specimen and classified into four grades based on the extent of 4-HNE-positive staining in the cytoplasm (26).

Statistical analysis. The Mann-Whitney test was used to assess statistical significance among all groups. All statistical tests

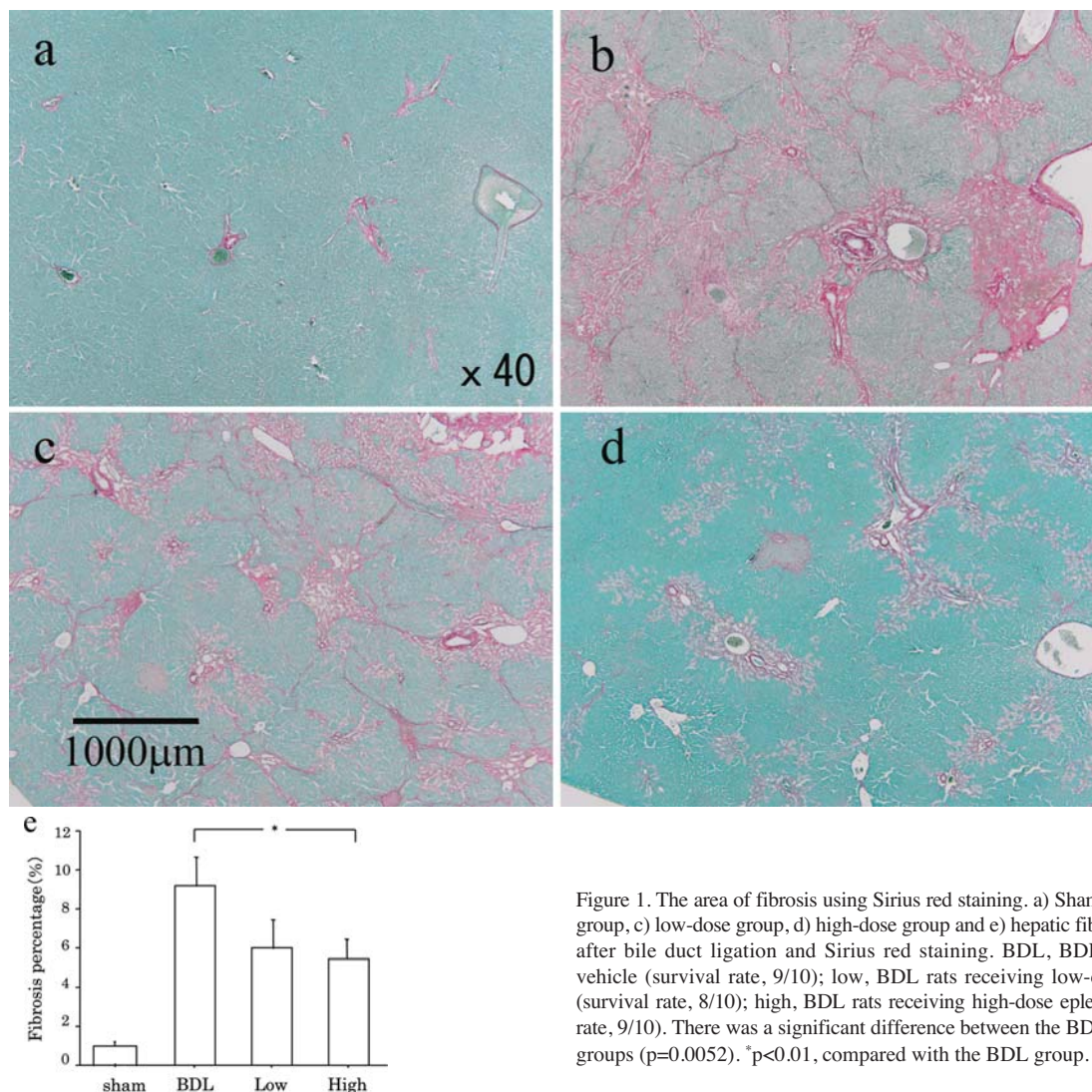


Figure 1. The area of fibrosis using Sirius red staining. a) Sham group, b) BDL group, c) low-dose group, d) high-dose group and e) hepatic fibrosis percentage after bile duct ligation and Sirius red staining. BDL, BDL rats receiving vehicle (survival rate, 9/10); low, BDL rats receiving low-dose eplerenone (survival rate, 8/10); high, BDL rats receiving high-dose eplerenone (survival rate, 9/10). There was a significant difference between the BDL and high-dose groups ($p=0.0052$). * $p<0.01$, compared with the BDL group.

were carried out using StatView for Windows (SAS Institute, Cary, NC). Values of $p<0.05$ were considered significant.

Results

Biochemical data 3 weeks following BDL. Serum ALT in the high-dose group was significantly higher than that in the BDL and low-dose group ($p<0.05$). Serum bilirubin and serum AST levels were not increased in the high-dose group. Plasma ATII concentration in the high-dose group was significantly lower than that in the BDL group ($p<0.05$), while plasma renin and aldosterone levels remained unchanged (Table II).

Effects of eplerenone on BDL-induced hepatic fibrosis. Histological analysis revealed that eplerenone inhibited the progression of hepatic fibrosis when compared with the BDL group. The hepatic hydroxyproline content in the high-dose group tended to be lower than that in the control group (control, 602 ± 152 $\mu\text{g/g}$ wet liver; low-dose group, 541 ± 278 $\mu\text{g/g}$ wet liver; high-dose group, 446 ± 169 $\mu\text{g/g}$ wet liver, $p=0.08$, Table III). Semi-quantitative measurement of the area of fibrosis using Sirius red staining demonstrated that eplerenone administration reduced the progression of liver fibrosis, especially in the high-dose group (sham group, $0.62\pm0.20\%$;

BDL group, $9.8\pm3.6\%$; low-dose group $6.8\pm4.0\%$; high-dose group, $4.8\pm2.5\%$, $p=0.0052$, Fig. 1a-e). Administration of high-dose eplerenone also reduced the area of positive α -SMA immunostaining (sham group, $0.66\pm0.11\%$; BDL group, $4.9\pm2.0\%$, low-dose group $4.6\pm2.1\%$; high-dose group, $2.6\pm2.0\%$, $p=0.032$, Fig. 2a-e).

Eplerenone administration ablated any increases in mRNA levels of procollagen I in a dose-dependent manner. However, these changes were not significant. mRNA expression of TIMP-1 in the low-dose ($p=0.001$) and high-dose ($p=0.08$) groups were significantly lower than that in the BDL group. Gene expression levels of TGF- β 1, CTGF and MMP-13 did not differ among the three groups (Table III).

Eplerenone administration markedly reduced the number of 8-OHdG-positive cells present in the liver samples (Table IV). 8-OHdG-positive cells in the low-dose ($51.6\pm7.9\%$; $p=0.0005$) and high-dose ($53.4\pm7.3\%$; $p=0.0007$) groups were significantly decreased compared to those in the BDL group ($68.7\pm8.7\%$; Fig. 3a-e, Table IV). The total 8-OHdG DNA content also tended to decrease following eplerenone administration (Table IV).

4-HNE protein expression in the high-dose group (2.1 ± 0.69) was lower than that in the BDL group (2.9 ± 0.87) following immunohistochemical staining ($p=0.046$). The low-dose group

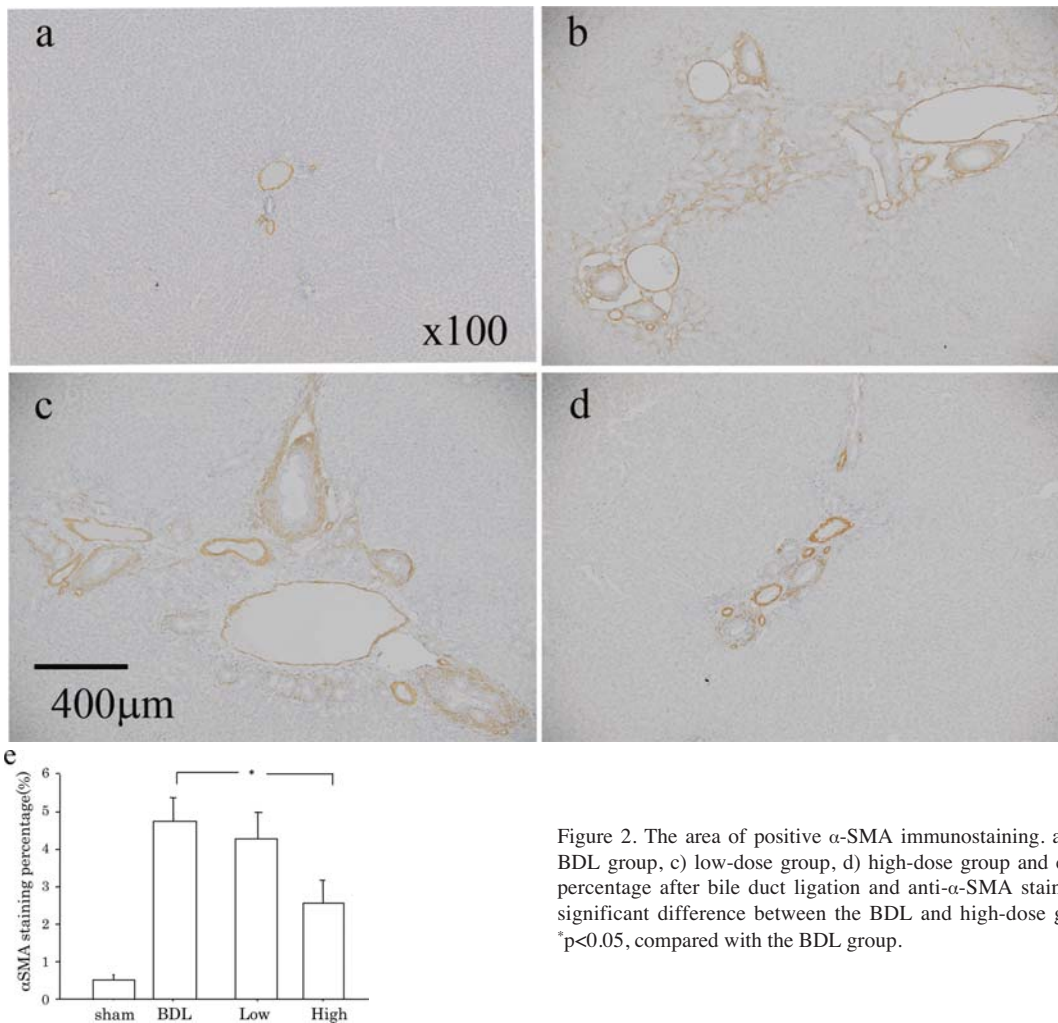


Figure 2. The area of positive α -SMA immunostaining. a) Sham group, b) BDL group, c) low-dose group, d) high-dose group and e) hepatic fibrosis percentage after bile duct ligation and anti- α -SMA staining. There was a significant difference between the BDL and high-dose groups ($p=0.032$). * $p<0.05$, compared with the BDL group.

Table IV. Analysis of lipid peroxidation and oxidative stress.

	Sham n=6	BDL group n=9	Low-dose group n=8	High-dose group n=9
TBARS (nmol/ml)	21.5 \pm 3.3	19.4 \pm 4.1	16.4 \pm 5.3	20.1 \pm 7.1
4-HNE (grade)	0.17 \pm 0.41	2.9 \pm 0.87	2.9 \pm 0.69	2.1 \pm 0.69 ^a
8-OHdG DNA content (pg/ μ g)		0.28 \pm 0.14	0.22 \pm 0.16	0.19 \pm 0.11
8-OHdG IHC (%)	5.2 \pm 2.4	68.7 \pm 8.7	51.6 \pm 7.9 ^b	53.4 \pm 7.3 ^b

Values are presented as the mean \pm standard deviation. TBARS, thiobarbituric acid reactive substance of hepatic tissue; 4-HNE, average 4-hydroxy-2-nonenal staining grade; 8-OHdG DNA, 8-OHdG DNA content of hepatic tissue; 8-OHdG IHC, percentage of 8-OHdG stained cells. ^a $p<0.05$ and ^b $p<0.001$, compared with the BDL group.

(2.9 \pm 0.69) did not demonstrate this reduction (Fig. 4a-e, Table IV). Hepatic TBARS, a marker of lipid peroxidation, did not differ among the three groups (Table IV).

Discussion

The present study demonstrated that high-dose administration of eplerenone, a specific mineralocorticoid receptor blocker, attenuated the progression of BDL-induced hepatic fibrosis, and slightly reduced the expression of collagen-1 and significantly reduced that of TIMP-1. These findings suggested

that eplerenone ameliorated hepatic fibrosis through additive effect of decrease in collagen synthesis and decreased in collagen degradation. Indeed, *in vitro* studies have demonstrated that aldosterone induced the synthesis of procollagen type I in rat stellate cells (27). Canrenone is an anti-aldosterone drug that has been shown to attenuate cell proliferation and migration induced by platelet-derived growth factor (PDGF) and procollagen types I and IV, as well as fibronectin induced by TGF- β in human stellate cells (10). In addition, *in vivo* studies have also demonstrated the antifibrotic effects of spironolactone, a mineralocorticoid receptor antagonist, in pig

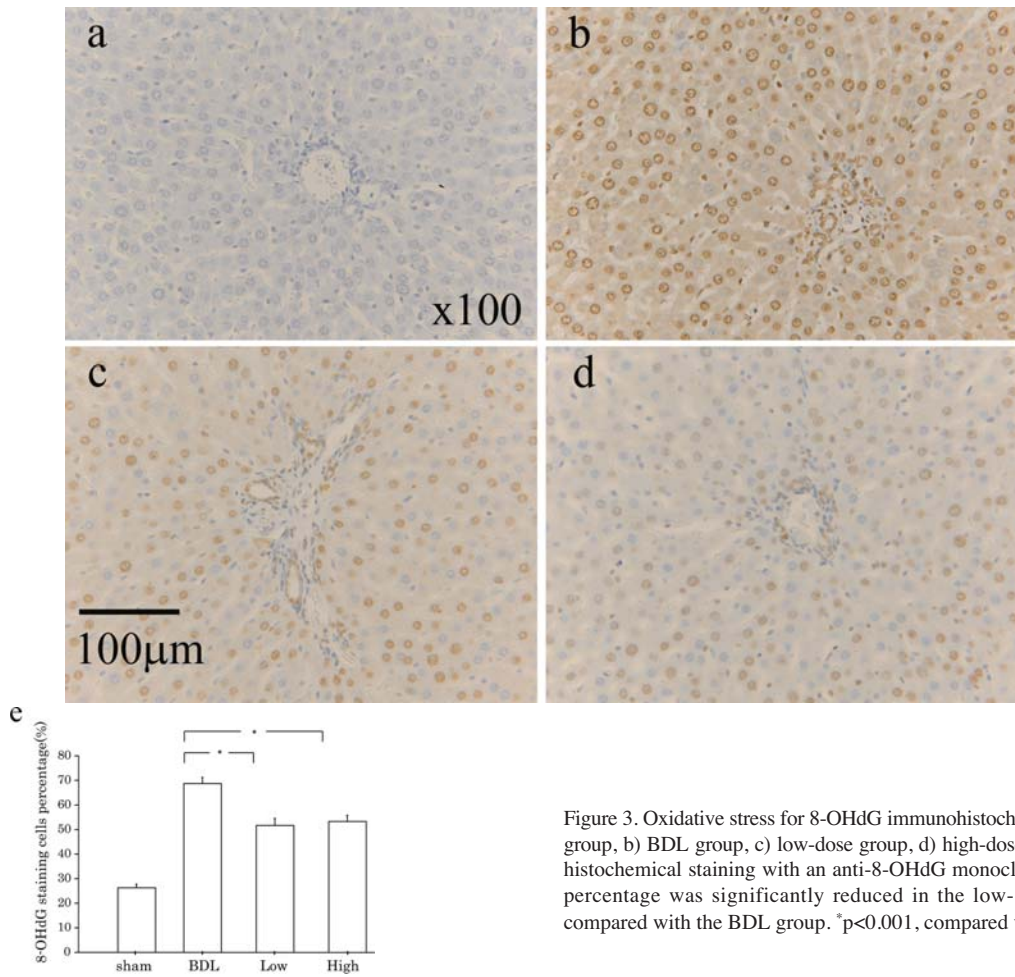


Figure 3. Oxidative stress for 8-OHdG immunohistochemical staining. a) Sham group, b) BDL group, c) low-dose group, d) high-dose group and e) immunohistochemical staining with an anti-8-OHdG monoclonal antibody. 8-OHdG percentage was significantly reduced in the low- and high-dose groups compared with the BDL group. *p<0.001, compared with the BDL group.

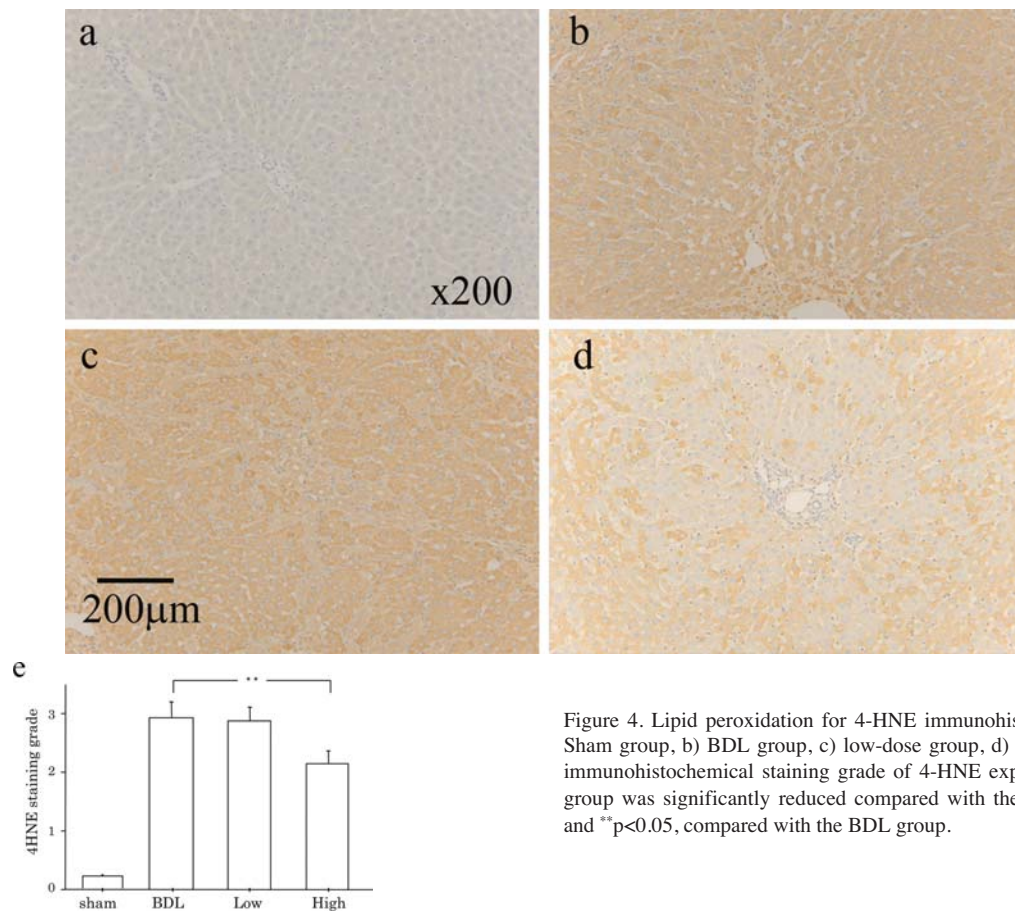


Figure 4. Lipid peroxidation for 4-HNE immunohistochemical staining. a) Sham group, b) BDL group, c) low-dose group, d) high-dose group and e) immunohistochemical staining grade of 4-HNE expression. The high-dose group was significantly reduced compared with the BDL group. *p<0.001 and **p<0.05, compared with the BDL group.



duced hepatic fibrosis. The present study also showed that eplerenone attenuated the activation of HSC. These results suggested that eplerenone affects HSC and exhibits preventive effects on the development of liver fibrosis.

In the present study, we also found that eplerenone administration resulted in a reduction in 8-OHdG expression, a marker of oxidative stress, and a decrease in the staining of 4-HNE, a marker of lipid peroxidation. It is well established that oxidative stress and lipid peroxidation are involved in the pathogenesis of BDL-induced liver fibrosis (28). Indeed, we have showed that 8-OHdG and 4-HNE expression was significantly increased in the liver of BDL rats compared with sham rats as well as our previous reports (6,8). In contrast, antioxidant factors such as glutathione, as well as the activity of GSH peroxidase and catalase, have been reported to be low in the liver of BDL rats (28). Furthermore, it has been reported that aldosterone accelerated oxidative stress in cell culture (29). Therefore, the antifibrotic effects of eplerenone appear to be due to the suppression of oxidative stress and lipid peroxidation.

In addition, we also demonstrated that eplerenone decreased HSC activation, but did not alter mRNA expression levels of TGF- β 1 and CTGF. Recently, inhibitors of ATII have been suggested to function as antifibrotic drugs, while ARB has been shown to inhibit both the activation of HSC and the expression of profibrogenic cytokines such as TGF- β 1 and CTGF (10). These differences in pharmacological action may therefore influence the antifibrotic effects. In fact, it appears that the antifibrotic effects of eplerenone are weaker than those in our previous study demonstrating the antifibrotic effects of ARB (6).

Interestingly, high doses of eplerenone resulted in a reduction in serum ATII levels. ATII exhibits powerful profibrogenic effects, generates ROS and stimulates the proliferation of HSC. Recent studies have reported that aldosterone induces ACE gene expression in rat cardiomyocytes (30) and that aldosterone reduces the gene expression of ACE-2, which compared angiotensin II to angiotensin 1-7 by separating phenylalanine (31). Furthermore, eplerenone administration completely blocked increases in ACE mRNA levels and the reduction in ACE2 mRNA levels due to the presence of aldosterone. Thus, eplerenone may abrogate several harmful effects of ATII via the reduction of serum ATII levels.

In conclusion, we demonstrated that eplerenone treatment attenuated the development of BDL-induced hepatic fibrosis by reducing oxidative stress, suppressing activation of HSC. Thus, eplerenone may prove useful as an alternative drug for antifibrosis therapy.

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