Hepatocellular apoptosis after hepatectomy in obstructive jaundice in rats

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Abstract

AIM: To investigate the hepatocellular apoptosis after hepatectomy in obstructive jaundice and biliary decompression rats.

METHODS: After bile duct ligation for 7 days, rats were randomly divided into OB group in which the rats underwent 70 % hepatectomy, OB-CD group in which the rats underwent hepatectomy accompanied by choledochoduodenostomy, CD-Hx group in which the rats underwent choledochoduodenostomy and then received 70 % hepatectomy on the fifth day after biliary decompression. The control group (Hx group) only underwent hepatectomy.

RESULTS: The level of total serum bilirubin and serum enzymes was significantly lower in CD-Hx group than in OB-CD and OB groups on day 1, 3 and 5 after hepatectomy. The apoptotic index was significantly lower in CD-Hx group than in OB-CD and OB groups on day 3 and 5. The oligonucleosomal DNA fragments and Caspase-3 activity were also lower in CD-Hx group than in OB-CD and OB groups 3 days after hepatectomy, without differences between CD-Hx and Hx groups.

CONCLUSION: Hepatocellular apoptosis plays vital roles in jaundice rats, and biliary decompression is more effective in treatment of patients with severe jaundice before operation.

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INTRODUCTION

Obstructive jaundice is often a clinical manifestation of the disease of extrahepatic biliary system or pancreas. In jaundiced patients with hilar bile duct carcinoma or gallbladder carcinoma, hepatectomy is one of the surgical regimens, but surgical procedures are associated with increases of morbidity and mortality rates, mainly due to postoperative complications such as hepatic failure, sepsis, bleeding and renal failure[11-12]. The significance of biliary drainage before surgery is controversial. Preoperative biliary drainage decreased the mortality and morbidity rates in patients with obstructive jaundice in some studies but not in others[3-5]. Experimental and clinical studies have identified several etiological factors including hypotension, impaired nutritional status, depressed immune function, hepatic dysfunction and the presence of toxic bile salts in circulation[6-8]. A high serum bilirubin concentration in jaundiced patients undergoing operation has been recognized as a predictor of mortality[9].

Hepatocyte injury and progression of liver disease are due to direct chemical damage to hepatocytes by toxic hydrophobic bile salts[9]. Although toxic bile salts are known to cause hepatocyte toxicity by inducing apoptosis, the precise mechanisms responsible for bile salt-mediated apoptosis remain to be established. This information is important because it could help provide rational strategies for the treatment of cholestatic liver disorders and decreasing postoperative complications. The aim of this study was to investigate the hepatocellular apoptosis after hepatectomy in obstructive jaundice and biliary decompression rats.

MATERIALS AND METHODS

Animals and experimental design

Male Wistar rats (Shionogi Aburabi Laboratory, Shiga, Japan) weighing 190-240 g were used for this study. Animals were housed in a controlled environment with a 12 h light/dark cycle. Rats had free access to normal rat chow and water before surgery. The care and handling of the animals were made in accordance with the National Institutes of Health Guidelines for the Care and Use of Laboratory Animals. The rats were divided into four groups each containing 6 animals. Hx group in which normal rats underwent 70 % hepatectomy, OB group in which the rats underwent 70 % hepatectomy after bile duct ligation for 7 days, OB-CD group in which the rats underwent 70 % hepatectomy accompanied by choledochoduodenostomy after bile duct ligation for 7 days, CD-Hx group in which the rats underwent choledochoduodenostomy after bile duct ligation for 7 days and then received 70 % hepatectomy on the fifth day after biliary decompression, as described below.

Surgical procedures

All surgical procedures were carried out under ether anesthesia. All rats were weighed before surgery and at the end of the study. Laparotomy was performed through an upper midline incision. After ligation of the proximal and distal bile ducts, the common bile duct was divided to prevent recanalization[10]. The ligatures were placed in the same position in all rats. The 70 % hepatectomy was done according to the method of Higgins and Anderson[11] on the seventh day after biliary obstruction in OB and OB-CD groups and on the fifth days after the biliary decompression in CD-Hx group.

Choledochoduodenostomy was performed according to the method of Ryan et al with minor modifications[12]. After biliary obstruction for 7 days, the abdomen was reopened through the previous incision. The dilated common hepatic bile duct was freed from its surrounding tissues and the contents were aspirated. A small incision was made in the bile duct and a silicone tube with an outer diameter of 0.9 mm was inserted and secured in position with a 7-0 silk suture. The free end of
the tube was inserted into duodenum approximately 1.5 cm from the pylorus, and then tied in position with a 6-0 PDS purse-string suture. Then the suture was carried out between the bile duct and the duodenal wall to protect the separation of the anastomosis.

**Measurement of bilirubin and enzymes in blood**

Blood samples were collected from the tail vein with a 27-gauge needle. Serum total bilirubin, activities of alkaline phosphatase (ALP) and aspartate aminotransferase (AST) levels were measured enzymatically using a commercial kit (Spotchem Co., Kyoto, Japan).

**Quantification of oligonucleosomal (histone-associated) DNA fragments**

An aliquot from the liver homogenate was centrifuged at 13 000×g. The supernatant was diluted 1 000-fold and subjected to a sandwich ELISA designed to detect DNA/histone fragments using a test kit from Boehringer Mannheim, according to the manufacturer’s instructions. Briefly, a 96-well microtiter plate was coated with antibody against histones. Diluted samples were then applied to the wells for 90 min, followed by incubation with peroxidase-conjugated anti-DNA antibody for 90 min. Color substrate was added and absorbance was measured at 405 nm at 5, 10, and 15 min using a 96-well microtiter plate reader.

**In situ TdT-UTP nick-end labeling (TUNEL staining)**

On the third and fifth day after hepatectomy, a section of liver tissue measuring approximately 5 mm in thickness was cut from the right lateral lobe and fixed in 10 % buffered formalin, processed by standard techniques and embedded in paraffin. Liver tissues were immediately frozen by liquid nitrogen and stored at -80°C. The total serum bilirubin, which was measured fluorometrically. Liver samples were homogenized in 500 µl of lysis buffer. Samples were centrifuged at 16 000g in a microfuge for 10 min at 4°C and normalized to protein using the Bradford assay (Bio-Rad). Fluorescent substrate, DEVD- AFC was incubated with 50 µl supernatants for liver extracts at 37°C for 60 min. The fluorescence of cleaved substrates was measured using a fluorometer (Molecular Dynamics Inc.) at an excitation wavelength of 400 nm and an emission wavelength of 505 nm. An AFC calibration curve was established by serial dilution of a pure AFC solution provided in the kit.

**Statistical analysis**

All data were expressed as mean ±SEM. Statistical analysis was done by statistical package of SPSS for Window release 9.0. Differences between mean were tested with the Mann-Whitney test. A probability value less than 0.05 was considered statistically significant.

**RESULTS**

**Bilirubin and enzymes in blood**

The level of total serum bilirubin reached more than 7 mg/dl after 7 days’ biliary obstruction, and the high bilirubin level rapidly decreased to 0.8 mg/dl only on day 1 after biliary decompression. We therefore thought that our models functioned well for biliary decompression. The total serum bilirubin concentration on the first day after hepatectomy in OB-CD group was 1.6 mg/dl, which was higher than that in CD-Hx group. The level of bilirubin in OB group was increased, while there was no statistically significant differences among other three groups on day 5 after hepatectomy (Figure 1).

The ALP concentration of OB group was significantly higher than that of other groups, but no differences of serum ALP level were found among the other groups after hepatectomy (Figure 2).

**Caspase-3-like protease activity assay**

Liver tissues were immediately frozen by liquid nitrogen and stored at -80°C on the third and fifth days after hepatectomy. The Caspase 3-like activity was measured using a commercially available kit (Clontech Laboratories, Palo Alto, CA) according to the manufacturer’s instructions. Caspase enzymatically cleaved the substrate and released free 7-amino-4-trifluoromethyl coumarin (AFC) that produced a blue-green fluorescence, which was measured fluorometrically. Liver samples were diluted 1 000-fold and subjected to a sandwich ELISA designed to detect DNA/histone fragments using a test kit from Boehringer Mannheim, according to the manufacturer’s instructions. Briefly, a 96-well microtiter plate was coated with antibody against histones. Diluted samples were then applied to the wells for 90 min, followed by incubation with peroxidase-conjugated anti-DNA antibody for 90 min. Color substrate was added and absorbance was measured at 405 nm at 5, 10, and 15 min using a 96-well microtiter plate reader.

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The ALP concentration of OB group was significantly higher than that of other groups, but no differences of serum ALP level were found among the other groups after hepatectomy (Figure 2).

**Figure 1** Changes of total serum bilirubin after hepatectomy. Blood was withdrawn from the tail vein while each rat was under ether anesthesia, and serum was obtained by means of centrifugation. Each point represents the mean ± SEM. ▲ <0.01 CD-Hx group vs OB group, ▼ <0.05 CD-Hx group vs OB-CD group, ▼ <0.01 OB-CD group vs OB group.

**Figure 2** Changes of serum ALP after hepatectomy. Six rats were used for each group. Results were the mean ±SEM. ▲ <0.05 CD-Hx group vs OB group.
The level of serum AST peaked on the first day after hepatectomy and decreased gradually (Figure 3). There was no difference of AST concentration between CD-Hx and Hx groups on the third and fifth days after hepatectomy. The serum AST concentration in OB group on the fifth day was nearly 6 times that of the concentration in CD-Hx group.

Caspase-3 activity in CD-Hx group was significantly lower than that in OB and OB-CD groups on the third day after hepatectomy. But on day 5, the level of Caspase-3 activity in OB-CD group decreased rapidly, without differences between OB-CD and CD-Hx groups (Figure 6).

**Figure 3** Changes of serum AST after hepatectomy. Six rats were used for each group. Results were the mean ±SEM. *P* < 0.01 CD-Hx group vs OB group, *P* < 0.01 OB-CD group vs OB group.

**Figure 4** Quantification of oligonucleosomal DNA fragments. The cytoplasmic levels of DNA (histone-associated) oligonucleosomal fragments in hepatocytes after hepatectomy were determined. The presence of these fragments in the cytoplasm reflected the extent of DNA fragmentation and nuclear disruption that are characteristic of apoptosis. The level of DNA fragments in CD-Hx group was significantly lower than that in OB and OB-CD groups on the third day after hepatectomy (Figure 4). There was no significant difference between CD-Hx and OB groups.

**Figure 5** Changes of hepatocellular apoptotic index after hepatectomy. Morphometric analysis of the positive cells in tissue stained by TUNEL method was performed under high power magnification (×400) in a blinded fashion. Each group consisted of 6 rats. The number of positive cells was counted and expressed as the total number of apoptotic cells seen in every ten high-power fields sections. Values were expressed as mean ±SEM. *P* < 0.01 CD-Hx group vs OB group, *P* < 0.01 CD-Hx group vs OB-CD group, *P* < 0.05 OB-CD group vs OB group, *P* < 0.01 OB-CD group vs OB group.

**Figure 6** Changes of Caspase-3-like protease activity after hepatectomy. 50 µM DEVD-AFC was added to whole cell lysates (50 µg of total protein) for 1 h. The activity of Caspase-3-like protease was measured by determining the cleaved fluorogenic substrates using a spectrofluorometer at 400/505 nm. Each group consisted of 6 rats. Values shown represented mean ±SEM. *P* < 0.01 CD-Hx group vs OB group, *P* < 0.05 CD-Hx group vs OB-CD group, *P* < 0.05 OB-CD group vs OB group.

**DISCUSSION**

The perioperative mortality rate in patients with obstructive jaundice was reported to be 8 % to 28 % [13]. Perioperative complications in patients with obstructive jaundice included infections, endotoxemia, renal failure, stress ulceration, disseminated intravascular coagulation, and wound dehiscence [14,15]. Numerous reports suggested that in obstructive jaundice there was not only impairment of hepato cellular function but also depression of Kupffer cell activity and ultimately hepatic structural damage, leading to the development of portal hypertension [16,17]. In addition, it could result in depressed wound healing, significant portal and systemic endotoxaemia, decreased reticuloendothelial cell function, reduced T cell responses, depressed non-specific cell-mediated immunity, decreased bacterial clearance and increased bacterial translocation [18,19].
This animal model was designed to simulate the clinical treatment procedures used for patients with obstructive jaundice undergoing hepatectomy. Choledochoduodenal drainage was performed before or at the same time as 70 % hepatectomy. Obstruction at the choledochoduodenostomy occurred in about 2 % rats. These rats were excluded from further studies. The mortality rate was about 20 % following choledochoduodenostomy or hepatectomy in jaundice rats. No rats died during the procedure and all deaths occurred in the first 3 days postoperatively, usually secondary to bleeding.

During obstructive jaundice, the retention of biliary constituents and high biliary pressure caused hepatocellular injury[20]. This is supported by the fact that serum AST and ALP levels increased during obstructive jaundice and decreased after choledochoduodenostomy (not data shown). Kanno[21] reported that biliary obstruction for even 5 days suppressed rat liver function, as assessed with aminopyrine breath test and galactose tolerance test. These results suggested that obstructive jaundice could cause serious impairment of pivotal liver functions, consequently resulting in poor prognosis after partial hepatectomy. The present study clearly showed that biliary drainage before hepatectomy significantly improved liver functions and decreased hepatocellular apoptosis. Serum liver function tests showed that choledochoduodenal drainage was effective in both OB-CD and CD-Hx groups. Furthermore, serum enzyme levels, apoptotic index, DNA fragments and Caspase-3 activity were significantly lower in CD-Hx group than in other groups. These data suggested that biliary drainage before hepatectomy had beneficial effects in severe jaundice patients.

Accumulation of these toxic bile salts within hepatocytes has been thought to play a key role in liver injury during obstructive jaundice. Indeed, hepatic levels of toxic bile salts chenodeoxycholate and deoxycholate correlated with the degree of liver damage[22]. Because widespread necrosis was not prominent in most cholestatic liver diseases, it has become apparent that the occurrence of hepatocyte death during cholestasis was rather due to apoptosis than due to necrosis[23]. The mechanisms of apoptosis by toxic bile salts could be partially elucidated in recent years. Toxic bile salts could directly cause apoptosis in cultured rodent hepatocytes[24]. Bile acid-induced toxicity is typically characterized by hepatocyte swelling, disruption of plasma membrane integrity, and release of intracellular constituents. The apoptotic pathway in hepatocytes could be initiated by a ligand-independent activation of Fas by bile salt[25]. Apoptosis was triggered by the Fas pathway, at least in part, by the activation of intracellular enzymes called Caspases. Caspases have been found to be cysteine-containing, aspartic acid-specific proteases which exist as zymogens in the soluble cytoplasm, mitochondrial intermembrane space, and nuclear matrix of virtually all cells[26]. Upon activation of the apoptotic pathway, initiator Caspases (i.e., Caspases 8 and 9) were converted to their active forms, which in turn activated downstream Caspase 3, which could activate other Caspases and ultimately cleave various substrates[27]. One of these substrates has been found to be a Caspase-dependent endonuclease that is freed from its inhibitor by Caspase-3 in the cytoplasm, and subsequently enters the nucleus, where it cuts DNA into oligonucleosomal (180 bp) fragments[28]. Additionally, Bid, a protein responsible for the release of mitochondrial cytochrome C and amplification of the apoptotic cascade, could also be activated by Caspases[29]. Ultimately, this pathway would ensure the self-destruction of cells with the characteristic morphological features of apoptosis, including nuclear fragmentation, cellular shrinkage, and acidophilic staining of the cytoplasm[30]. Bile acid induced apoptosis also appeared to require an activation and translocation of protein kinase C, which then could lead to an increase of intracellular magnesium[31,32]. This would cause an activation of Mg²⁺-dependent endonucleases, leading to DNA cleavage.

In conclusion, hepatocellular apoptosis plays a vital role in jaundice rats. Biliary drainage is preferable, as evidenced in rats by significantly better liver function and decrease of hepatocellular apoptosis. Biliary decompression is an effective preoperative treatment for patients with severe jaundice.

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