Comparison of murine cirrhosis models induced by hepatotoxin administration and common bile duct ligation

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Abstract

AIM: To build up the research models of hepatic fibrosis in mice.

METHODS: Inbred wild-type FVB/N mice were either treated with alpha-naphthyl-isothiocyanate (ANIT), allyl alcohol (AA), carbon tetrachloride (CCl₄), 3,5-diethoxycarbonyl-1,4-dihydrocollidine (DDC), and silica, or subjected to common bile duct ligation (CBDL) to induce hepatic injury. Liver biopsies were performed every 4 wk to evaluate hepatic fibrosis over a period of 6 mo. Cumulative cirrhosis and survival curves were constructed by life table method and compared with Wilcoxon test.

RESULTS: Under the dosages used, there was neither mortality nor cirrhosis in AA and silica-treated groups. DDC and ANIT caused cirrhosis within 4-12 and 12-24 wk, respectively. Both showed significantly faster cirrhosis induction at high dosages without significant alteration of survival. The duration for cirrhosis induction by CCl₄ ranged from 4 to 20 wk, mainly dependent upon the dosage. However, the increase in CCl₄ dosage significantly worsened survival. Intraperitoneal CCl₄ administration resulted in better survival in comparison with gavage administration at high dosage, but not at medium and low dosages. After CBDL, all the mice developed liver cirrhosis within 4-8 wk and then died by the end of 16 wk.

CONCLUSION: CBDL and administrations of ANIT, CCl₄, and DDC ensured liver cirrhosis. CBDL required the least amount of time in cirrhosis induction, but caused shortened lives of mice. It was followed by DDC and ANIT administration with favorable survival. As for CCl₄, the speed of cirrhosis induction and the mouse survival depended upon the dosages and the administration route.

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INTRODUCTION

Being the final outcome of chronic liver damage regardless of the underlying etiologies, hepatic fibrosis demands thorough study. The studies of hepatic fibrosis rely upon the availability of animal models. Although no experimental model exactly reproduces human liver fibrosis by etiologies, animal models serve to improve our understanding of pathogenetic mechanisms of liver fibrosis. In view of the cost-effectiveness in inducing hepatic injuries, hepatotoxin administration and common bile duct ligation (CBDL) were considered as useful methods[3].

In recent years, advances in molecular biology have helped to generate a transgenic mouse model of liver fibrosis[3]. This represents a new form of perturbation analysis whereby selective expression of novel or altered genes are used to perturb a complex system so that the information about the changes during development, functioning and malfunctioning can be deduced[9]. This approach potentially allows development of animal models with liver fibrosis that are inducible and genetically similar to that of man. A reliable and reproducible model of hepatic fibrosis in wild-type control mice will aid to unveil the pathologic mechanisms deduced from genetic and molecular approaches. However, informative data of mouse hepatic fibrosis generated in the laboratory remained lacking in the literature because rats had long been the most common animal model for studying hepatic fibrosis induced by various chemical or surgical hepatic injuries[11]. Since different animal models display distinct characteristics in the nature of pathogenesis of fibrosis or cirrhosis, the data acquired from rats could not apply to mice without bias[12]. This urged us to set up a mouse model of hepatic fibrosis using CBDL or the administration of various hepatotoxins including alpha-naphthyl-isothiocyanate (ANIT)[16,19], silica[19], carbon tetrachloride (CCl₄)[12-14], 3,5-diethoxycarbonyl-1,4-dihydrocollidine (DDC)[18,21], and allyl alcohol (AA)[17,18]. This study mainly focused on the strain of FVB/N inbred mice, which have been widely used to generate a variety of transgenic lines owing to their high reproduction rate and easy access in microinjection[20,21].

Key words: Common bile duct ligation; Fibrosis; Hepatotoxin; Liver; Mice

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MATERIALS AND METHODS

Mice

Inbred FVB/N mice (8-10 wk old) were obtained from the National Laboratory Animal Center (Taipei, Taiwan). All animals received care as outlined in the “Guides for the Care and Use of Laboratory Animals” prepared by the Committee of Animal Research, Chang Gung Memorial Hospital. Mice were supplied with food and water and exposed to a 12-h light-dark cycle. Ten mice (five males and five females) composed a group to which surgical CBDL or a hepatotoxin via intraperitoneal (IP) or oral gavage (OG) administration at the same dosage was given during a period of 6 mo.

Methods to induce hepatic injuries

ANIT (Sigma Chemical Co., St. Louis, MO, USA) ANIT was dissolved in corn oil and administered via IP or OG route, respectively. For each route of administration, there were three subgroups according to the maintenance dose as follows: 25 mg/kg (low dosage), 50 mg/kg (medium dosage) and 125 mg/kg (high dosage). After a single loading dose of 100 mg/kg, the maintenance doses were given twice a week. Additionally, there was a modified method of OG administration with a single loading dose of 100 mg/kg and subsequent maintenance doses of 50 mg/kg for the 1st and 2nd mo, 75 mg/kg for the 3rd and 4th mo, and 100 mg/kg for the 5th and 6th mo.

AA (Sigma Chemical Co.) AA was prepared in double distilled water and administered via OG route for five subgroups according to the dosages as follows: 0.2, 0.4, 0.6, 0.8, and 1.0 mmol/kg twice a week.

CCl₄ (Sigma Chemical Co.) The mice were fed via OG or IP route with CCl₄ (density = 1.594 kg/L) diluted in double distilled water. As for each route of administration, they were further subgrouped into three according to the maintenance doses as follows: 0.8 g/kg (low dosage), 1.6 g/kg (medium dosage) and 2.0 g/kg (high dosage). After a single loading dose of 2.4 g/kg, the maintenance doses were then administered twice a week.

DDC (Sigma Chemical Co.) The mice were fed via OG with DDC in corn oil twice a week. They were subgrouped by four different dosages as follows: 0.25, 0.5, 2.5, and 5 mg. In addition, 0.1% DDC contained diet (Purina Mills TestDiet, St. Louis, MO, USA) was used as the 5th subgroup.

Silica (US Silica Company, Berkeley Springs, WV, USA) Silica was dissolved in Dulbecco’s modified eagle media and given subcutaneously at the dose of 3.5 g/kg, or intraperitoneally at the dose of 1.6 g/kg twice a week.

CBDL The surgical procedure was performed under sterile conditions. The mice were anesthetized with isoflurane. Midline laparotomy was performed for exploring the hepatic hilum and identifying CBD. Under the dissecting microscope, CBD was then isolated, doubly ligated and transected between two ligatures.

Liver biopsy

Liver biopsy demanded survival procedures and sterile techniques. After anesthesia with isoflurane, midline laparotomy was performed to expose the mouse liver. A liver block sized 0.5 cm in diameter was taken and then the cut surface was sutured for hemostasis. Serial biopsy was preformed monthly (per 4 wk) for each mouse till cirrhosis was well documented by histopathology. At the first liver biopsy, ascites, and intra-abdominal adhesion were examined and recorded.

Histopathology

Mouse livers from biopsies were fixed in 10% neutral buffered formaldehyde solution and embedded in paraffin. Sections, 4 μm in thickness, were stained with hematoxylin and eosin. Fatty change of hepatocytes was recorded when hepatocytes showed fat accumulation, balloon-like swelling or formation of fat cysts. For liver necrosis, there were the disorganization of liver cell cords, homogeneous cytoplasm, pyknotic nuclei and periportal infiltration of inflammatory cells. Fibrosis was defined as the presence of fibrous scarring band in the liver parenchyma by Mason’s trichrome collagen stain. Once the irregular zones of fibrosis developed and surrounded the remaining parenchymal tissues to display a pattern of nodularity under microscope, cirrhosis was diagnosed.

Statistical analysis

For survival and cirrhosis rate analyses, life table method was used. The length of time intervals used in this study was 4 wk. Any mouse which died, or whose liver was pathologically documented to be cirrhotic was considered to have experienced an event. When a mouse died before cirrhosis was confirmed at cirrhosis rate analysis, it was counted as a withdrawal and treated the same way as cases lost to follow-up. Plots were constructed with weeks after inducing liver injury to survival or cirrhosis rate. The Wilcoxon test was employed to compare the curves. Differences were regarded as significant if a P value was less than 0.05.

RESULTS

Mouse survival

During the period of 6 mo, none of the 260 mice enrolled in this study died of the procedures for CBDL, liver biopsy or administration of hepatotoxins. There was no mortality case in the AA and silica groups. As for the other groups, 71 mice died in the course of this study. Among the 71 deaths, 20 occurred before cirrhosis could be histopathologically confirmed (pre-cirrhetic) and the remaining 51 after cirrhosis was found (post-cirrhetic). Nineteen of the 20 pre-cirrhetic deaths occurred in the CCl₄ group. As for the post-cirrhetic deaths, there were 4 in the ANIT group, 26 in the CCl₄ group, 11 in the DDC group and 10 in the CBDL group. The case numbers of mortality in each subgroup were listed in Table 1. Mortality had been found in all the CCl₄ subgroups, ranging from 40% to 100%. In the DDC group, mortality was mainly post-cirrhetic, and seen in 2.5 and 5 mg and 0.1% diet subgroup. All the mice in CBDL group had post-cirrhetic death by the end of 16 wk. The overall comparison for the survival curves in the ANIT and DDC groups could not reach any statistical significance (P = 0.0823 and 0.1227, respectively), whereas there was a significant difference in the CCl₄ subgroup (P<0.0001). Pairwise comparisons for survival curves of the six CCl₄ subgroups showed that the
low dosage (IP0.8 and OG0.8) was significantly better than the medium dosage (IP1.6 and OG1.6, Figure 1, \( P = 0.0010-0.0317 \)), and the medium dosage was also superior to the high dosage (IP2.0 and OG2.0, Figure 1, \( P = 0.0002-0.0044 \)). Considering the administration routes, IP route compared favorably in survival with OG route at the high dosage (\( P = 0.0221 \)), but not at the medium and low dosages (\( P = 0.7899 \) and 0.5902, respectively).

|                | Pre-cirrhotic death | Post-cirrhotic death |
|----------------|---------------------|----------------------|
| ANIT OG 25 mg/kg | 0                   | 0                    |
| ANIT OG 50 mg/kg | 0                   | 0                    |
| ANIT OG 125 mg/kg| 0                   | 3                    |
| ANIT IP 25 mg/kg | 0                   | 0                    |
| ANIT IP 50 mg/kg | 0                   | 0                    |
| ANIT IP 125 mg/kg| 0                   | 0                    |
| ANIT modified    | 0                   | 1                    |
| AA OG from 0.2 to 1.0 mmol/kg | 0 | 0 |
| CCl4 OG 0.8 g/kg | 0                   | 0                    |
| CCl4 OG 1.6 g/kg | 2                   | 7                    |
| CCl4 OG 2.0 g/kg | 5                   | 5                    |
| CCl4 IP 0.8 g/kg | 4                   | 0                    |
| CCl4 IP 1.6 g/kg | 6                   | 1                    |
| CCl4 IP 2.0 g/kg | 2                   | 8                    |
| DDC OG 0.25 mg  | 0                   | 0                    |
| DDC OG 0.5 mg   | 0                   | 1                    |
| DDC OG 2.5 mg   | 0                   | 3                    |
| DDC IP 0.8 g/kg | 0                   | 4                    |
| Silica (SC 3.5 g/kg or IP 1.6 g/kg) | 0/0 | 0/0 |
| CBDL            | 0                   | 10                   |

OG: Oral gavage; IP: intraperitoneal; SC: subcutaneous.

**Figure 1** Survival curves of six CCl4 subgroups are shown. The number immediately behind the IP or OG indicates the dosage of CCl4 used in g/kg.

**Ascites and intraabdominal adhesion**

Ascites was universally found at laparotomy when the hepatotoxins were administered via IP route. In the CBDL group, there were only three mice presenting with ascites at the subsequent laparotomy. Intraabdominal adhesion existed in over 50% of cases (ranging from 50% to 100%) when IP route or CBDL was used. Additionally, granuloma-like masses were intraperitoneally identified in all the cases of silica IP subgroup and in nine cases of ANIT IP subgroups (two for 25 mg/kg, one for 50 mg/kg and six for 125 mg/kg).

**Histopathology**

The case numbers of histopathological findings in each subgroup were listed in Table 2. Fatty change was seen mainly in the ANIT and CCl4 groups, and sporadically in the AA group (1 for 0.8 mmol/kg and 2 for 1.0 mmol/kg). Bile lakes were the common findings in the DDC and CBDL groups. Liver necrosis and fibrosis were demonstrated almost in all the cases of the ANIT, CCl4, DDC, and CBDL groups, two cases of the AA group (one for 0.8 mmol/kg and the other for 1.0 mmol/kg), but none of the silica group. The necrosis and fibrosis were detected at the first or second liver biopsy (4-8 wk after liver injury). Liver fibrosis in the two mice from the AA group did not progress to cirrhosis by the end of this study. Cirrhotic nodules could be identified in 30 mice (42.9%) of ANIT group (\( n = 70 \)), 32 mice (53.3%) of CCl4 group (\( n = 60 \)), 39 mice (78.0%) of DDC group (\( n = 50 \)) and all the mice (100%) of CBDL group (\( n = 10 \)). DDC subgroup with higher dosages (2.5 and 5 mg) and 0.1% diet could substantially reach 100% successful rate of cirrhosis induction as seen in the CBDL group.

**Durations required for cirrhosis induction**

Results from life table analysis for the durations were presented in the curves with weeks after inducing liver injury vs cirrhosis rate. CBDL represented a rapid method to induce liver cirrhosis with 70-100% of successful rate within 4-8 wk (Figure 2A). DDC caused liver cirrhosis within 4-12 wk (Figure 2B). Statistical analysis revealed that the lower the DDC dosage, the longer the time spent by DDC in inducing cirrhosis among 0.5, 2.5 and 5 mg subgroups (DDC0.5 vs DDC2.5, \( P = 0.0387 \) and DDC2.5 vs DDC5, \( P = 0.0004 \)). There was no significant difference between 0.25 and 0.5 mg DDC subgroups (\( P = 0.1311 \)). DDC 0.1% diet required significantly less time to induce cirrhosis than DDC 0.25 mg (\( P = 0.0001 \)), 0.5 mg (\( P = 0.0008 \)) and 2.5 mg (\( P = 0.0032 \)), but was not significantly different from 5 mg subgroups (\( P = 0.3415 \)). As for CCl4, the duration for inducing cirrhosis was quite variable, ranging from 4 to 20 wk (Figure 3A). The IP and OG high dosages (2.0 g/kg) of CCl4 spent significantly less time in cirrhosis induction than IP and OG low dosages (0.8 g/kg) of CCl4 (\( P = 0.0005-0.0209 \)). There was no difference of the cirrhosis curves between IP and OG administrations at the high (\( P = 0.1760 \)) and low (\( P = 0.1661 \)) dosages, whereas OG administration compared favorably in cirrhosis induction than IP administration at the medium dosage (1.6 g/kg) of CCl4 (\( P = 0.0060 \)). The OG medium dosage (OG1.6) had no significant difference from the high dosage (\( P = 0.1852 \) for OG2.0 and \( P = 0.7241 \) for IP2.0), and the IP medium dosage (IP1.6) had no significant difference from the low dosage (\( P = 0.8822 \) for OG0.8 and \( P = 0.1514 \) for IP0.8). Cirrhosis caused by ANIT did not show up until 12-24 wk (Figure 3B). The high dosage of ANIT significantly spent less time in cirrhosis induction than the medium (\( P = 0.0002-0.0263 \)) and low dosages (\( P = 0.0002-0.0027 \)), whereas the medium dosage could not be significantly superior to the low dosage (\( P = 0.0904-1.0000 \)). At the same ANIT dosage used, there was no significant difference of cirrhosis curves between OG and
Table 2: Histopathological changes after inducing liver injuries

|                | Fatty change | Necrosis/fibrosis | Cirrhosis | Bile lake |
|----------------|--------------|-------------------|-----------|-----------|
| ANIT OG 25 mg/kg | 0            | 10                | 2         | 0         |
| ANIT OG 50 mg/kg | 4            | 10                | 4         | 0         |
| ANIT OG 125 mg/kg | 5           | 10                | 8         | 0         |
| ANIT IP 25 mg/kg | 0            | 9                 | 1         | 0         |
| ANIT IP 50 mg/kg | 1            | 10                | 1         | 0         |
| ANIT IP 125 mg/kg | 4           | 10                | 9         | 0         |
| ANIT modified   | 4            | 10                | 5         | 0         |
| AA OG from 0.2 to 1.0 mmol/kg | 3\(^1\) | 2\(^2\)            | 0         | 0         |
| CCl\(_4\) OG 0.8 g/kg | 4            | 10                | 5         | 0         |
| CCl\(_4\) OG 1.6 g/kg | 5           | 10                | 8         | 0         |
| CCl\(_4\) OG 2.0 g/kg | 8           | 10                | 5         | 0         |
| CCl\(_4\) IP 0.8 g/kg | 3           | 10                | 2         | 0         |
| CCl\(_4\) IP 1.6 g/kg | 5           | 10                | 4         | 0         |
| CCl\(_4\) IP 2.0 g/kg | 6           | 10                | 8         | 0         |
| DDC OG 0.25 mg | 0            | 10                | 3         | 6         |
| DDC OG 0.5 mg  | 0            | 10                | 6         | 9         |
| DDC OG 2.5 mg  | 0            | 10                | 10        | 10        |
| DDC OG 5 mg    | 0            | 10                | 10        | 10        |
| DDC 0.1% diet  | 0            | 10                | 10        | 9         |
| Silica (SC 3.5 g/kg or IP 1.6 g/kg) | 0/0        | 0/0              | 0/0       | 0/0       |
| CBDL           | 0            | 10                | 10        | 10        |

OG: oral gavage; IP: intraperitoneal; SC: subcutaneous. \(^1\) for 0.8 mmol/kg and 2 for 1.0 mmol/kg; \(^2\) for 0.8 mmol/kg and 1 for 1.0 mmol/kg.

Figure 2: Cirrhosis rates of CBDL and DDC groups. A: The cirrhosis curve for CBDL group is shown; B: The cirrhosis curves for 5 DDC subgroups are shown. The number immediately behind DDC indicates its dosage used in mg per mouse. DDC diet represents 0.1% DDC contained diet.

Figure 3: Cirrhosis rates of CCl\(_4\) and ANIT groups. A: The cirrhosis curves for six CCl\(_4\) subgroups are shown. The number immediately behind the IP or OG indicates the dosage of CCl\(_4\) used in g/kg; B: The cirrhosis curves for seven ANIT subgroups are shown. The number immediately behind the IP or OG indicates the dosage of ANIT used in mg/kg.

IP administrations \((P = 0.2444\) for high dosage, \(P = 0.0904\) for medium dosage, and \(P = 0.5032\) for low dosage). The modified method for ANIT administration compared favorably in cirrhosis induction only with IP low dosage and IP medium dosage \((P = 0.0349,\) for both\), but poorly with OG high dosage \((P = 0.0303)\).
DISCUSSION

In this 6-mo study of experimental cirrhosis induction, AA and silica were ineffective in inducing liver fibrosis let alone cirrhosis. AA administration in the drinking water (50 ppm equivalent to 0.08-0.11 mmol/kg per d) for 15 wk has no effect on the histopathological examination of rat livers[19]. Moreover, IP injection of AA in rats only produced variable periportal liver necrosis predominantly at 6-12 h, followed by restitutive proliferation of periportal necrosis to repopulate the necrotic zone. Eventually, the liver architecture was essentially restored by 1 wk[19]. It indicated that AA merely induced short-term toxicity in rats. Although we had observed liver necrosis and fibrosis in two mice with OG administration of 0.8 and 1.0 mmol/kg AA, this liver damage was transient and insufficient to result in cirrhosis. It suggested that rats and FVB/N mice had similar hepatotoxic resistance to AA. As for silica, subcutaneous or IP administration was reported to cause liver fibrosis and cirrhosis in nude mice at 12 mo and in (C57BL/6×BALB/c) F submice at 18 mg[23]. Thus, a 6-mo duration might not be long enough for silica to cause hepatic injury with subsequent fibrosis and cirrhosis in FVB/N mice. As a result, a longer-term administration of silica might be required for inducing liver cirrhosis in FVB/N mice. A major drawback with silica administration was the granuloma formation at the injection site. This might be of concern to subsequent laparotomy for liver biopsy when the IP route is used.

The IP or OG administration of ANIT, CCl, or DDC could effectively lead to liver fibrosis within 4-8 wk. All the liver fibrosis in mice did not always progress to microscopic cirrhotic nodules by the end of 24 wk. It suggested that restitutive response of mice to the injuries by those hepatotoxins sometimes restore liver architecture. ANIT has been used for years to study cholangiolitic hepatotoxicity in laboratory animals[9,10]. This study showed that ANIT usually led to mouse cirrhosis within 12-24 wk after administration. Mortality was only seen in the subgroups with high dosage and modified administration. However, there was no significant difference among the survival curves of all the ANIT subgroups. The high dosage of ANIT could more rapidly induce liver cirrhosis than the medium or low dosage. At the same dosage of ANIT, it made no significant difference between IP and OG administrations in cirrhosis induction. The modified method by gradual increase of ANIT dosage could not substantially provide any benefit in cirrhosis induction in this study. Overall, ANIT model required a longer period for the formation of cirrhosis. This might provide more time for researchers to investigate the progression of liver fibrosis.

CCl, has been used extensively for decades to induce liver injury in various experimental models to elucidate the mechanisms behind hepatotoxicity[23]. Also, it was commonly used as a hepatotoxic agent in transgenic mice to evaluate liver fibrosis and cirrhosis on the basis of the selective expression of some novel or altered genes[11,12]. We experienced that the major problem in association with the use of CCl, to induce liver fibrosis or cirrhosis had been the high mortality rates of mice, accounting for 40-100 % in those subgroups. Animals might die before or after cirrhosis was found. The higher CCl, dosage was associated with the worse survival curve. The duration required for cirrhosis induction by CCl, was quite variable, ranging from 4 to 20 wk, mainly depending upon the dosage. The high dosage (2.0 g/kg) of CCl, could rapidly cause liver cirrhosis within 4-12 wk, but led to poor survival. Under the circumstances, IP administration will be suggested since IP route posed better survival than OG route only at the high dosage of CCl,. The low dosage (0.8 g/kg) of CCl, required 12-24 wk to successfully induce cirrhosis. At the low dosage, it was shown that the administration routes had no influence on the survival and the cirrhosis induction. At the medium dosage (1.6 g/kg) of CCl, OG administration was more rapid to induce cirrhosis than IP administration, but did not significantly differ in survival curves from IP administration. Nevertheless, the high mortality rate along with the use of CCl, must be taken into consideration since it will prematurely terminate the sequential studies or observations of liver fibrosis or cirrhosis. Furthermore, CCl, might not be an ideal solvent for investigating hepatocarcinoma since liver carcinogenesis might be due to the genotoxic chemical carcinogen effect of CCl, itself as well as CCl-induced liver cirrhosis[25,26].

DDC did not result in pre-cirrhotic death, but tended to lead to post-cirrhotic death at higher dosage (2.5 and 5 mg) and 0.1% DDC diet. The overall comparison for all survival curves of those DDC subgroups led to the insignificant difference. It took about 4-12 wk for DDC to successfully induce cirrhosis. Except 0.25 mg DDC subgroup, the successful rate of cirrhosis induction was quite satisfactory. The higher the dosage of DDC we used, the lesser the time we needed to induce cirrhosis. DDC 0.1% diet could not be superior to 5 mg DDC subgroup in cirrhosis induction. A major concern of DDC use was the high proportion of bile lake formation. This might cause some biases in observing the relationship between cirrhosis and its sequelae.

CBDL represented a rapid and consistent method to induce liver fibrosis and cirrhosis. Hepatocellular injuries were caused by cholestasis, leading to liver dysfunction, promoting fibrogenesis and ultimately resulting in liver failure and death. All the mice with CBDL rapidly progressed to cirrhosis within 4-8 wk after operation, and died by the end of 16 wk. The rapid progression to cirrhosis and death by CBDL might limit its application in chronic fibrosis studies and make it difficult to carry out a long-term investigation of liver cirrhosis. A major technical problem was the reversibility of histological changes due to recanalization of bile ducts[1]. In our tested animals, the bile ducts remained totally obstructed till their death in view of bile lakes at final pathology and the coloration of peritoneum and auricle. The successful obstruction by CBDL might be due to the fact that CBD was doubly ligated and transected between two ligatures[23]. Besides, intraabdominal adhesion could be an obstacle for subsequent survival laparotomy to study CBDL livers.

In summary, CBDL and hepatotoxins of ANIT, CCl, and DDC could be effective in causing liver fibrosis of mice. CBDL rapidly and irreversibly led to liver fibrosis and subsequently to cirrhosis within 4-8 wk, but inevitably caused mortality by the end of 16 wk. As a result, CBDL was suitable for creating a mouse model for the short-term study of liver fibrogenesis. Although CCl, had been commonly used for inducing liver fibrosis and cirrhosis, the duration
required for cirrhosis induction varied (4-20 wk) and mainly depended upon the dosage. It could be used in mice for studying both acute and chronic liver fibrogenesis. A major drawback in association with CCl4 use was a relatively high mortality. DDC could quite similar to that of CBDL, but the survival experience was better in the DDC group. ANIT slowly induced liver cirrhosis in mice. The duration from initial liver inflammatory damage to cirrhosis formation by ANIT was more than 12 wk after administration. Thus, ANIT was quite a good toxin for a long-term study of fibrosis progression into cirrhosis.

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REFERENCES

1. Wu J, Norton PA. Animal models of liver fibrosis. Scand J Gastroenterol 1996; 31: 1137-1143
2. Sanderson N, Factor V, Nagy P, Kopp J, Kondaiah P, Wakefield L, Roberts AB, Sporn MB, Thorgeirsson SS. Hepatic expression of mature transforming growth factor beta 1 in transgenic mice results in multiple tissue lesions. Proc Natl Acad Sci USA 1995; 92: 2572-2576
3. Hanahan D. Transgenic mice as probes into complex systems. Science 1989; 246: 1265-1275
4. Quick DJ, Shuler ML. Use of in vitro data for construction of a physiologically based pharmacokinetic model for naphthalene in rats and mice to probe species differences. Biotechnol Prog 1999; 15: 540-555
5. Lindstrom AB, Yeowell-O’Connell K, Waidyanatha S, McDonald TA, Golden RT, Rappaport SM. Formation of hemoglobin and albumin adducts of benzene oxide in mouse, rat, and human blood. Chem Res Toxicol 1998; 11: 302-310
6. Cunningham ML, Bucher JR. Pharmacodynamic responses of F344 rats to the mouse hepatocarcinogen oxazepam in a 90-d feed study. Toxicol Appl Pharm 1998; 149: 41-48
7. Watt KC, Buckpitt AR. Species differences in the regio- and stereoselectivity of 1-nitronaphthalene metabolism. Drug Metab Dispos 2000; 28: 376-378
8. Dill JA, Lee KM, Bates DJ, Anderson DJ, Johnson RE, Chou BJ, Burka LT, Roycroft JH. Toxicokinetics of inhaled 2-butoxyethanol and its major metabolite, 2-butoxyacetic acid, in F344 rats and BeC3F1 mice. Toxicol Appl Pharm 1998; 153: 227-242
9. Palmeira CM, Ferreira FM, Rolo AP, Oliveira PJ, Santos MS, Moreno AJ, Cipriano MA, Martins MI, Seica R. Histological changes and impairment of liver mitochondrial bioenergetics after long-term treatment with alpha-naphthyl-isothiocyanate (ANIT). Toxicology 2003; 190: 185-196
10. Ohta Y, Kongo M, Sasaki E, Harada N. Change in hepatic antioxidant defense system with liver injury development in rats with a single alpha-naphthylisothiocyanate intoxication. Toxicology 1999; 139: 265-275
11. Williams AO, Knappion AD. Hepatic silicosis, cirrhosis, and liver tumors in mice and hamsters: studies of transforming growth factor beta expression. Hepatology 1996; 23: 1268-1275
12. Natsume M, Tsuji H, Harada A, Akizawa Y, Yano T, Ishikura H, Nakanishi I, Matsushima K, Kaneko S, Mukaida N. Attenuated liver fibrosis and depressed serum albumin levels in carbon tetrachloride-treated IL-6-deficient mice. J Leukoc Biol 1999; 66: 601-608
13. Brenner DA, Veloz L, Jaenisch R, Alcorn JM. Stimulation of the collagen alpha 1 (I) endogenous gene and transgene in carbon tetrachloride-induced hepatic fibrosis. Hepatology 1993; 17: 287-292
14. Inagaki Y, Truter S, Bou-Ghariou G, Garret LA, de Crombrugghe B, Nemoto T, Greenwel P. Activation of Proalpha2(I) collagen promoter during hepatic fibrogenesis in transgenic mice. Biochem Biophys Res Commun 1998; 250: 606-611
15. Preisegger KH, Factor VM, Fuchsbichler A, Stumptner C, Denk H, Thorgeirsson SS. Atypical ductular proliferation and its inhibition by transforming growth factor beta1 in the 3,5-diethoxyxycarbonyl-1,4-dihydrocollidine mouse model for chronic alcoholic liver disease. Lab Invest 1999; 79: 103-109
16. Fickert P, Trauner M, Fuchsbichler A, Stumptner C, Zatloukal K, Denk H. Bile acid-induced Mallory body formation in drug-primed mouse liver. Am J Pathol 2002; 161: 2019-2026
17. Alam K, Nagi MN, Al-Shabanah OA, Al-Bekairi AM. Beneficial effect of nitric oxide synthase inhibitor on hepatotoxicity induced by allyl alcohol. J Biochem Mol Toxicol 2001; 15: 317-321
18. Yavorkovsky L, Lai E, Ilic Z, Sell S. Participation of small intraportal stem cells in the restitutive response of the liver to periporal necrosis induced by allyl alcohol. Hepatology 1995; 21: 1702-1712
19. Carpanini FM, Gaunt IF, Hardy J, Gangolli SD, Butterworth KR, Lloyd AG. Short-term toxicity of allyl alcohol in rats. Toxicology 1978; 9: 24-45
20. Taketo M, Schroeder AC, Mobbraaten LE, Cunning KB, Hanten G, Fox RR, Roderick TH, Stewart CL, Lilly F, Hansen CT. FVB/N: an inbred mouse strain preferable for transgenic analyses. Proc Natl Acad Sci USA 1991; 88: 2063-2069
21. Perret-Gentil ML, Murray L, Bird DJ, Ladiges WC. Evaluation of FVB/N mice as recipients for transgenic embryos. Lab Anim Sci 1999; 49: 427-428
22. Ebbesen P. Chirality of quartz. Fibrosis and tumour development in dust inoculated mice. Eur J Cancer Prev 1991; 1: 39-41
23. Plaa GL. Chlorinated methanes and liver injury: highlights of the past 50 years. Anna Rev Pharmacol Toxicol 2000; 40: 42-65
24. Schuur J, Olah J, Szepesi A, Nagy P, Thorgeirsson SS. Thioacetamide-induced hepatic fibrosis in transforming growth factor beta-1 transgenic mice. Eur J Gastroenterol Hepatol 2004; 16: 127-133
25. Zalatanai A, Lapis K. Simultaneous induction of liver cirrhosis and hepatocellular carcinomas in F-344 rats: establishment of a short hepatocarcinogenesis model. Exp Toxicol Pathol 1994; 46: 215-222
26. Reddowes EJ, Faux SP, Chipman JK. Chloroform, carbon tetrachloride and glutathione depletion induce secondary genotoxicity in liver cells via oxidative stress. Toxicol 2003; 187: 101-115
27. Prado IB, dos Santos MH, Lopasso FP, Iriya K, Laudanna AA. Cholestasis in a murine experimental model: lesions include hepatocyte ischemic necrosis. Rev Hosp Clin Fac Med Sao Paulo 2003; 58: 27-32

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