The purpose of the present study was to evaluate the effects of β-carotene on the cell viability and antioxidant status of hepatocytes from chronically ethanol-fed rats. Rats in the ethanol group were given an ethanol-containing liquid diet that provided 36% of total energy as ethanol, while rats in the control group were fed an isonenergetic diet without ethanol. After 4 weeks, hepatocytes were taken out and cultured for 24h. Hepatocytes from the rats in the control and ethanol groups were cultured in medium without (HC, HE) or with β-carotene (HC + B, HE + B). The results showed that lactate dehydrogenase leakage was significantly increased in the HE compared with that in the HC group. However, lactate dehydrogenase leakage of the HE + B group was significantly increased by 61% compared with that in the HE group. When compared with the HC group, activities of glutathione peroxidase and catalase in the HE group were significantly decreased by 54 and 31%, respectively. Catalase activity in the HE + B group was significantly increased by 61% compared with that in the HE group. However, activities of glutathione reductase and superoxide dismutase showed no difference among the groups. The level of glutathione in the HC + B and HE + B groups was significantly increased to 155 and 143% compared with those in the HC and HE groups, respectively. The concentration of lipid peroxides showed no difference among the groups. The present results demonstrate that β-carotene improved the cell viability of hepatocytes, and increased catalase activities and glutathione levels in hepatocytes from chronically ethanol-fed rats.

β-Carotene: Chronic ethanol feeding: Antioxidant status: Hepatocytes

According to the statistics tabulated by the Department of Health, Taiwan (2003), chronic liver disease and cirrhosis together are the sixth leading cause of death in Taiwan. The causation is not only associated with unbalanced dietary patterns but also with excessive consumption of alcohol in recent years. Long-term excessive drinking induces alcoholic liver diseases and psychological damage. Additionally, an unstable mental condition may lead to broken families and several other severe social problems. Therefore, it is important in medical and health research to resolve drinking-induced problems that impact on the health of individuals, families and society.

Ethanol is metabolised to acetaldehyde by some enzymes in the body, including alcohol dehydrogenase, cytochrome P450 2E1, catalase (CAT), xanthine oxidase, etc. Then acetaldehyde is decomposed to acetic acid by acetaldehyde dehydrogenase in the mitochondria. Numerous studies have indicated that an excessive ethanol intake induces the mass production of free radicals in the body, which are considered to be associated with alcoholic liver diseases. Ishii et al. (1997) indicated that the chronic intake of ethanol increases free radical levels, which are produced from the metabolic pathway of ethanol, especially superoxides (O2•−) and H2O2. Bailey & Cunningham (1999) indicated that the exposure of hepatocytes to ethanol resulted in an increased production of reactive oxygen species (ROS), which correlated with decreased cell viability. On the other hand, acetaldehyde stimulates the production of neutrophils, and then neutrophils release a large number of free radicals, which can damage hepatocytes and induce alcoholic liver diseases (Williams & Barry, 1987). Additionally, the chronic intake of ethanol has been shown to induce the production of lipid peroxidation, because of decreasing antioxidant enzyme activities (Rouach et al. 1997). Moreover, many studies have reported that vitamins A and E and β-carotene are reduced in the blood and liver with excessive ethanol intake, and the range of reduction is related to the amount of ethanol administration (Kawase et al. 1989; Leo et al. 1993; Ahmed et al. 1994; Polavarapu et al. 1998). Thus, it was...
thought that antioxidant nutrient supplementation might lower the formation of free radicals produced from excessive ethanol intake by enhancing the body’s antioxidation ability. Lieber (1997) reviewed antioxidant nutrient supplementation and suggested that it serves as a new therapy for alcoholic liver disease.

β-Carotene is viewed as a non-toxic precursor of vitamin A. Also, the strong antioxidant properties of β-carotene have been proven in several studies (Diplock, 1991). It functions as an efficient singlet oxygen quencher and as a radical-trapping antioxidant at low oxygen pressure to reduce the extent of nuclear damage and to inhibit lipid peroxidation (Palozza & Krinsky, 1992). It has been reported that the ethanol-induced hepatic depletion of vitamin A could be corrected by β-carotene supplementation (Ahmed et al. 1994). However, few reports have discussed the antioxidative effects of β-carotene on ethanol-induced oxidative stress in the liver.

The purpose of the present study was to investigate the direct influences of β-carotene on the ethanol-induced hepatic damage of antioxidation status. In order to eliminate the conversion and absorption factors of β-carotene within the intestinal mucosal cells, the primary hepatocytes model was set up to evaluate the effects of β-carotene on cell viability and antioxidative enzyme activities in hepatocytes from rats, which were chronically administered an ethanol-containing liquid diet.

Materials and methods

Animals and diets

The design of the experimental animal model was modified from the studies of Bailey & Cunningham (1999). Twenty male Sprague–Dawley rats weighing 200–210 g (National Laboratory Animal Breeding and Research Center, National Science Council, Taipei, Taiwan) were used in the present study. Rats were individually housed in a room maintained at 23 ± 2°C with 50–70% humidity and a 12 h light–dark cycle. The rats were divided into two groups. One group was not administered ethanol (control group); the other was administered ethanol (ethanol group). The composition of the diets was previously described in detail by Lieber & DeCarli (1994). The rats in the ethanol group were fed liquid diets with free access that provided 36% of total energy as ethanol, and a 12 h light–dark cycle. The rats in the control group received an isoenergetic diet provided 36% of total energy as bovine serum and 1 μM-dexamethasone at a density of 1 × 10⁵ cells/ml. After 20 h incubation at 37°C in 5% CO₂, the medium was replaced by the medium without (HC, HE) or with 1 μM-β-carotene (HC + B, HE + B) and continuously cultured for 24 h. β-Carotene was incorporated with the medium by tetrahydrofuran (0.5%) according to the method of Bertram et al. (1991). Then, cells were collected using a scraper and re-suspended in tri(hydroxymethyl)-aminomethane buffer (50 mm-tri(hydroxymethyl)-aminomethane–HCl, 5 mm-EDTA; pH 7.5) for the following analyses. Thus, hepatocytes from the rats in the control and ethanol groups were cultured in medium without or with β-carotene.

Liver function and pathology

At the end of the experimental period, blood was collected from the tail veins of the rats after 10 h fasting, and glutamate oxaloacetate transaminase (GOT) and glutamate pyruvate transaminase (GPT) activities were measured with commercial kits (RM, 163K; Iatron Laboratories, Tokyo, Japan) to confirm the liver function in each group (Dufour et al. 2000). In addition, one rat in each group was killed and a small piece of liver (5 × 2 mm) was removed at death and fixed in 10% (v/v) formaldehyde. The samples were stained with haematoxylin and eosin for light microscopy. The pathologist who carried out the histological analysis had no prior knowledge of the different experimental groups.

Preparation of isolated rat hepatocytes

Hepatocytes were isolated from the rats according to the two-step collagenase perfusion technique described by Berry & Friend (1969). Isolated cells were cultured as monolayers in William’s medium E with 5% (w/v) fetal bovine serum and 1 μM-dexamethasone at a density of 1 × 10⁵ cells/ml. After 20 h incubation at 37°C in 5% CO₂, the viability of hepatocytes was expressed as the percentage of lactate dehydrogenase (LDH) leakage, which was the LDH content in the culture medium relative to the total LDH content including the culture medium and cytosolic fraction. The LDH level was determined using the method described by Moldeus et al. (1978).

Antioxidative enzyme activities

Cells in tri(hydroxymethyl)-aminomethane buffer were lysed with a sonicator and centrifuged at 4°C, and 10000 rpm for 10 min. Then, supernatant fractions were used to determine the antioxidative enzyme activities. Glutathione peroxidase (GPx) activity was measured spectrophotometrically at 340 nm by the method of Paglia & Valentine (1967) using a commercial kit (RS 504; Randox Laboratories, Crumlin, Antrim, UK). The activity of GPx is expressed as mU/mg protein. GPx activity is expressed as μmol NADPH oxidised/min. Glutathione reductase (GRd) activity was measured spectrophotometrically at 340 nm using a commercial kit (Calbiochem 359962; Calbiochem-Novabiochem, La Jolla, CA, USA). GRd activity is also expressed as μU/mg protein. CAT activity was determined spectrophotometrically at 240 nm by the method of Lück (1963). The activity of CAT is expressed as kU/mg protein. Superoxide dismutase (SOD) activity was determined spectrophotometrically at 525 nm by the method of Nebot et al. (1993) using a commercial kit (Calbiochem 574600; Calbiochem-Novabiochem). The activity of SOD is expressed as U/mg protein. The protein content in the hepatocytes was measured by the modified method of Lowry et al. (1951) using a Bio-Rad DC protein assay kit (Bio-Rad Laboratories, Hercules, CA, USA).
Antioxidants levels

The concentration of GSH in the hepatocytes was measured spectrophotometrically at 400 nm using a commercial kit (Calbiochem 354102; Calbiochem-Novabiochem) according to the method of Anderson (1989).

Concentrations of β-carotene and retinol in the hepatocytes and medium were measured by reverse-phase HPLC (Shimadzu LC-10A HPLC Pump, Shimadzu SPD-10AV UV/Vis Detector; Shimadzu, Tokyo, Japan) using a 4.6 × 250 mm C18 column (5 μm, Vydac 211TP54) according to the method of Wei et al. (1998). The concentration of β-carotene was measured at 452 nm under the mobile phase of methanol, acetonitrile and H2O (88:9:3, by vol.). The concentration of retinol was measured at 325 nm under the mobile phase of methanol. Data were analysed by SISC-LAB chromatographic analysis software (Scientific Information Service Corp., Taipei, Taiwan).

Lipid peroxidation

The concentration of lipid peroxides in the hepatocytes was assessed colorimetrically at 586 nm using a commercial kit (Calbiochem 437634; Calbiochem-Novabiochem) (Esterbauer & Cheeseman, 1990). The cytosolic fraction (200 μl) was mixed with 650 μl reagent 1 (7.7 nmol N-methyl-2-phenylindole/l in 75% acetonitrile and 25% methanol) and 150 μl reagent 2 (15.4 mol methanesulfonic acid/l) at 45°C for 40 min. The levels of malondialdehyde (MDA) and 4-hydroxy-2(E)-nonenal (4-HNE) endproducts derived from the peroxidation of PUFA and related esters were measured at 586 nm.

Statistical analysis

All data are expressed as means and standard deviations. Concentrations of β-carotene and retinol in the hepatocytes were measured colorimetrically at 586 nm using a commercial kit (Calbiochem 437634; Calbiochem-Novabiochem) (Esterbauer & Cheeseman, 1990). The cytosolic fraction (200 μl) was mixed with 650 μl reagent 1 (7.7 nmol N-methyl-2-phenylindole/l in 75% acetonitrile and 25% methanol) and 150 μl reagent 2 (15.4 mol methanesulfonic acid/l) at 45°C for 40 min. The levels of malondialdehyde (MDA) and 4-hydroxy-2(E)-nonenal (4-HNE) endproducts derived from the peroxidation of PUFA and related esters were measured at 586 nm.

Results

Energy intakes in the control and ethanol groups were 257.5 (SD 0.4) and 254.1 (SD 0.4) kJ/d, respectively. And, the average alcohol intake in the ethanol group was 2.62 (SD 0.03) g/d. Initial body weight was 200.4 (SD 8.0) g in the control group and 200.2 (SD 5.1) g in the ethanol group. After 4 weeks, the final body weight of the ethanol group (276.8 (SD 31.0) g) was significantly lower (P<0.05) than that of the control group (313.4 (SD 7.5) g).

GOT activities of the control and ethanol groups were 66 (SD 10) and 82 (SD 1) karmen units. GPT activities of the control and ethanol groups were 18 (SD 4) and 29 (SD 5) karmen units. Compared with the control group, GOT and GPT activities had significantly increased (P<0.05) by 24 and 61% in the ethanol group, respectively. From the light micrograph of liver, fat accumulation was observed in the ethanol group (Fig. 1).

LDH leakage is shown in Table 1. LDH leakage of the HE group had significantly increased (P<0.05) when compared with that of the HC group. In contrast, LDH leakage in the HE + B group was nearly the same as that of the HC group, and significantly lower (P<0.05) than that of the HE group.

Table 2 shows the antioxidative enzyme activities. The HE group had significantly lower (∼54%; P<0.05) GPx activities than the HC group. Furthermore, the HE + B group had significantly decreased (∼49%; P<0.05) GPx activities than the EC + B group, but showed no difference when compared with the HE group. The CAT activity of the HE group was significantly reduced (∼31%; P<0.05) compared with that of the HC group, but it was significantly elevated by 61% in the HE + B group compared with the HE group (P<0.05). However, there was no change in the GRd and SOD activities of any group.

Concentrations of GSH and lipid peroxides are shown in Table 3. The GSH level showed no changes between the HC and HE groups. However, the level of GSH in the HE + B group had significantly increased (P<0.05) by 155 and 143% compared with those of the HC and HE groups, respectively. However, there were no differences in the MDA and 4-HNE levels in any group.

Tables 4 and 5 show the levels of β-carotene and retinol in hepatocytes and medium. β-Carotene levels in the HC + B and HE + B groups had significantly increased (P<0.05) compared with those of the HC and HE groups in both hepatocytes and medium. The retinol concentration of the HE + B and HE + B groups had significantly increased (P<0.05) compared with those of the HC and HE groups in hepatocytes. However, the retinol level of the medium in the HE group was significantly reduced (∼14%; P<0.05) compared with that of the HC group, and those of the HC + B and HE + B groups had significantly increased (P<0.05). Moreover, the retinol level of the medium in the HE + B group was significantly lower (∼12%; P<0.05) compared with that of the HC + B group.

Discussion

Chronic ethanol consumption results in the formation of oxygen free radicals and lipid peroxidation (Reinke et al. 1987). In the present study, β-carotene was used to improve the cell viability and antioxidative status in hepatocytes from chronically ethanol-fed rats.

In the present study, it was found that ethanol consumption in the ethanol group was similar to that reported in other previous reports (Lieber & DeCarli, 1994; Navder et al. 1997). On the other hand, the isoenergetic substitution of carbohydrates by ethanol resulted in a lower weight gain despite similar energy intakes as also previously reported (Pirola & Lieber, 1975). This lower body-weight gain has been attributed to induction of the microsomal ethanol-oxidising system, increased sympathetic tone and associated thermogenesis and/or enhanced...
ATP breakdown secondary to acetate production from ethanol (Cunningham & Spach, 1987). Ethanol consumption for 4 weeks by rats led to high GOT and GPT activities and fatty changes in the liver, which might have been due to an enhanced mobilisation of NEFA from adipose tissue and the increased hepatic biosynthesis of lipids as previously suggested (Fig. 1) (Lieber, 1991).

LDH is an enzyme that exists in many tissues and organs, such as the heart, muscle, kidney, liver, etc. When those tissues or organs are damaged, LDH is released into the blood from cells. Therefore, LDH leakage can be used to indicate cell viability. In the present study, the viability of hepatocytes was monitored by the release of LDH into the medium; that is to say, higher LDH leakage was interpreted as lower viability of hepatocytes. Hepatocytes from the ethanol-fed rats showed higher LDH leakage than those from the control rats (Table 1; HC v. HE). This suggests that chronic ethanol administration might reduce the cell viability of hepatocytes, which might be due to the oxidative stress from ethanol (Bailey & Cunningham, 1999). However, β-carotene significantly decreased LDH leakage when it was added to the medium (Table 1; HE v. HE + B). Similar to the present results, previous in vitro studies have shown that β-carotene can decrease LDH leakage of hepatocytes by means of reducing ROS production (Ek et al. 1994; Martin et al. 1996; Lawlor & O’Brien, 1997).

The metabolism of ethanol is believed to result in an increased production of ROS, especially superoxide and H₂O₂, and the removal of these toxic species is thought to be a vital initial step in ensuring cell survival during ethanol intoxication (Nordmann et al. 1992). Four major antioxidant enzymes available to the cell during ethanol-induced oxidant stress include GPx, GRd, CAT and SOD. Schisler & Singh (1989) reported that GPx, CAT and SOD activities in ethanol-treated animals, including five inbred strains of mice and Sprague–Dawley rats, were generally reduced in comparison with those of their
matched controls. Polavarapu et al. (1998) also suggested that GPx, CAT and SOD activities showed an inverse correlation with severity of pathological injury in rats fed ethanol. However, Oh et al. (1998) reported that chronic ethanol feeding resulted in a lower activity of GPx with significantly higher activities of GRd and CAT.

The present results indicated that GPx and CAT activities showed significant reductions, but GRd and SOD showed no changes in the hepatocytes from the chronically ethanol-fed rats (Table 2; HC v. HE). Both GPx and CAT can react with H$_2$O$_2$. GPx is thought to be more active in the removal of H$_2$O$_2$ because of its dual location (mitochondria and cytosol) (Diplock, 1991). Therefore, this theory suggests that GPx should be more sensitive and more easily alterable than CAT in a situation of chronic ethanol intake.

Also, it has been reported that chronic ethanol feeding results in a lower activity of GPx with significantly higher activities of GRd and CAT.

<table>
<thead>
<tr>
<th>Group†</th>
<th>Mean</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>HC</td>
<td>41$^a$</td>
<td>2</td>
</tr>
<tr>
<td>HC + B</td>
<td>69$^a$</td>
<td>9</td>
</tr>
<tr>
<td>HE</td>
<td>40$^a$</td>
<td>4</td>
</tr>
</tbody>
</table>

Table 1. Effect of β-carotene on lactate dehydrogenase (LDH) leakage in primary rat hepatocytes*

<table>
<thead>
<tr>
<th>Group†</th>
<th>Mean</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>HC</td>
<td>0·21$^a$</td>
<td>0·10</td>
</tr>
<tr>
<td>HC + B</td>
<td>0·51$^b$</td>
<td>0·18</td>
</tr>
</tbody>
</table>

*Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).

Table 2. Effects of β-carotene on antioxidant enzymes glutathione peroxidase (GPx), glutathione reductase (GRd), catalase (CAT) and superoxide dismutase (SOD) activities in primary rat hepatocytes*

<table>
<thead>
<tr>
<th>Group†</th>
<th>Mean</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>HC</td>
<td>42·97$^{abc}$</td>
<td>8·63</td>
</tr>
<tr>
<td>HC + B</td>
<td>39·69$^{abc}$</td>
<td>15·63</td>
</tr>
</tbody>
</table>

Livers of rats chronically fed ethanol might have originated from differences in the strains of rats used and the dose, duration and route of ethanol administration among different studies. The most acceptable concept is that chronic ethanol intake significantly increases the rate of GSH turnover (Morton & Mitchell, 1985). Therefore, the GSH:GSSG ratio must be determined in future studies.

Lipid peroxidation induced by long-term ethanol intake results not only from an increased ROS production but also from the enhanced generation of acetaldehyde (Müller & Sies, 1982). However, another mechanism that promotes lipid peroxidation is GSH depletion. Many reports have demonstrated that chronic ethanol intake promotes the accumulation of lipid peroxidation (Shaw et al. 1981; Müller & Sies, 1982; Polavarapu et al. 1998), but the present results did not show an alteration of the MDA + 4-HNE contents of hepatocytes from the etha-

Table 3. Effect of β-carotene on reduced glutathione and lipid peroxide concentrations in primary rat hepatocytes*

<table>
<thead>
<tr>
<th>Group†</th>
<th>GSH (μM)</th>
<th>MDA + 4-HNE (μM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HC</td>
<td>0·27*</td>
<td>6·21*</td>
</tr>
<tr>
<td>HC + B</td>
<td>0·27*</td>
<td>6·21*</td>
</tr>
</tbody>
</table>

*Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).
Table 4. Effects of β-carotene on β-carotene concentrations of hepatocytes and medium in each group (Mean values and standard deviations of three replicate assays)

<table>
<thead>
<tr>
<th>β-Carotene concentration</th>
<th>Hepatocytes (nmol/10⁷ cells)</th>
<th>Medium (10⁻¹⁰mol/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Group</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HC</td>
<td>0⁰</td>
<td>0</td>
</tr>
<tr>
<td>HC + B</td>
<td>5·96⁺</td>
<td>0·82</td>
</tr>
<tr>
<td>HE</td>
<td>0⁰</td>
<td>0</td>
</tr>
<tr>
<td>HE + B</td>
<td>5·49⁺</td>
<td>0·58</td>
</tr>
</tbody>
</table>

a,b Mean values within a column with unlike superscript letters were significantly different (P<0·05).

† Hepatocytes from the control group were cultured in medium without (HC) or with β-carotene (HC + B), and hepatocytes from the ethanol group were also cultured in medium without (HE) or with β-carotene (HE + B).

Table 5. Effects of β-carotene on retinol concentrations of hepatocytes and medium in each group (Mean values and standard deviations of three replicate assays)

<table>
<thead>
<tr>
<th>Retinol concentration</th>
<th>Hepatocytes (10⁻¹⁰mol/10⁷ cells)</th>
<th>Medium (nmol/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Group</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HC</td>
<td>5·32⁺</td>
<td>0·34</td>
</tr>
<tr>
<td>HC + B</td>
<td>7·50⁺</td>
<td>0·31</td>
</tr>
<tr>
<td>HE</td>
<td>5·32⁺</td>
<td>0·15</td>
</tr>
<tr>
<td>HE + B</td>
<td>6·20⁺</td>
<td>0·71</td>
</tr>
</tbody>
</table>

a,b Mean values within a column with unlike superscript letters were significantly different (P<0·05).

† Hepatocytes from the control group were cultured in medium without (HC) or with β-carotene (HC + B), and hepatocytes from the ethanol group were also cultured in medium without (HE) or with β-carotene (HE + B).

levels were significantly reduced in the ethanol-fed rats’ hepatocytes whether treated with β-carotene or not (Table 5; HC v. HE, HC + B v. HE + B). The present results suggest that β-carotene might enter hepatocytes and be converted to retinol, and the hepatocytes from the ethanol-fed rats consumed more retinol than those from the control rats. This phenomenon agrees with previous studies, that is, ethanol intake induces the depletion of hepatic vitamin A (Kawase et al. 1989; Leo et al. 1993; Ahmed et al. 1994; Polavarapu et al. 1998).

In previous studies (Leo et al. 1993; Ahmed et al. 1994), it has been demonstrated that β-carotene could alleviate ethanol-induced liver damage by correcting the retinol depletion. However, the present study indicates that cell viability, CAT activities, GSH levels and retinol contents were improved when β-carotene was directly supplied to the hepatocytes of the chronically ethanol-fed rats. This means that β-carotene could attenuate the oxidative stress induced by chronic ethanol intake. On the basis of the present study, the dosage of β-carotene must be reconsidered, when β-carotene is provided in food or supplements during chronic ethanol ingestion, and an in vivo study is essential in the future.

References


β-Carotene and alcoholic liver disease


