The decrease in uroporphyrinogen decarboxylase activity induced by ethanol predisposes rats to the development of porphyria and accelerates xenobiotic-triggered porphyria, regardless of hepatic damage

Abstract

We evaluated the porphyrinogenic ability of ethanol (20% in drinking water) per se, its effect on the development of sporadic porphyria cutanea tarda induced by hexachlorobenzene in female Wistar rats (170-190 g, N = 8/group), and the relationship with hepatic damage. Twenty-five percent of the animals receiving ethanol increased up to 14-, 25-, and 4.5-fold the urinary excretion of δ-aminolevulinate, porphobilinogen, and porphyrins, respectively. Ethanol exacerbated the precursor excretions elicited by hexachlorobenzene. Hepatic porphyrin levels increased by hexachlorobenzene treatment, while this parameter only increased (up to 90-fold) in some of the animals that received ethanol alone. Ethanol reduced the activities of uroporphyrinogen decarboxylase, δ-aminolevulinate dehydrase and ferrochelatase. In the ethanol group, many of the animals showed a 30% decrease in uroporphyrinogen activity; in the ethanol + hexachlorobenzene group, this decrease occurred before the one caused by hexachlorobenzene alone. Ethanol exacerbated the effects of hexachlorobenzene, among others, on the rate-limiting enzyme δ-aminolevulinate synthetase. The plasma activities of enzymes that are markers of hepatic damage were similar in all drug-treated groups. These results indicate that 1) ethanol exacerbates the biochemical manifestation of sporadic hexachlorobenzene-induced porphyria cutanea tarda; 2) ethanol per se affects several enzymatic and excretion parameters of the heme metabolic pathway; 3) since not all the animals were affected to the same extent, ethanol seems to be a porphyrinogenic agent only when there is a predisposition, and 4) hepatic damage showed no correlation with the development of porphyria cutanea tarda.
Introduction

Porphyrias are diseases associated with alterations of heme metabolic pathway enzymes (1). Chronic alcoholism has been recognized as the main factor triggering porphyria cutanea tarda (PCT), the most common form of porphyria in humans, which is clinically characterized by severe cutaneous photosensitivity. PCT patients can accumulate porphyrins in the skin which undergo photoexcitation, leading to cell injury (1,2).

Biochemically, PCT is associated with a deficiency in uroporphyrinogen decarboxylase (UroD) activity and with excessive urinary excretion and hepatic accumulation of higher carboxylated porphyrins (3-6). There are two main types of PCT: sporadic PCT (type I), and familial PCT (type II). The familial disease is inherited as an autosomal dominant defect in the UroD gene, and is manifested in all tissues (4,5). Patients with the sporadic type present the defective UroD only in the liver, and no related hereditary component is known (2). In both types of PCT, exogenous factors seem to be responsible for the clinical manifestation of the disease. Among these, alcohol and estrogens are considered to be the most important (2,7).

Hexachlorobenzene (HCB), a xenobiotic with recognized porphyrinogenic activity (8), mimics human PCT when administered to rats.

Studies have been carried out to investigate the response of induced PCT to chronic alcohol treatment. Several investigators have reported alterations in different parameters of the heme pathway, but the situation remains unclear (2). Thus, the effects of combined exposure to other xenobiotics cannot be predicted since they generally vary according to the system studied. These variations may be attributed to differences between animal groups (9), the methodology applied in the determinations, or animal handling.

The present study was undertaken in order to obtain more information about this subject by evaluating: 1) the porphyrinogenic ability of ethanol, 2) the effect of ethanol intake on the development of experimental PCT induced by HCB, and 3) the possible relationship between porphyria and hepatic damage. To minimize the variations cited above, an attempt was made to identify as many key parameters as possible in the same animal. These parameters included: a) urinary excretion of porphyrins and precursors, b) hepatic levels of cytochrome P-450 and porphyrins, c) hepatic enzyme activities of the heme metabolic pathway, and d) the plasma activities of enzymes that are markers of hepatic damage.

We used a model involving chronic alcohol intake in order to avoid malnutrition, treatment rejection and/or severe hepatic damage.

Material and Methods

HCB (commercial grade), composed of 95% HCB and 5% tetrachlorobenzene and pentachlorobenzene, was a gift from Compañía Química S.A., Buenos Aires, Argentina. Uroporphyrin III was from Porphyrin Products, Inc. (Logan, UT, USA); uroporphyrinogen was prepared by reduction with sodium amalgam (10). Aminolevulinate (ALA), porphobilinogen (PBG), Dowex 1-X8-400, and Dowex 50-X8-400 were from Sigma (St. Louis, MO, USA). Sephadex G-25 was purchased from Pharmacia (Uppsala, Sweden). Protoporphyrin IX was a generous gift from Dr. Moises Grinstein (Faculty of Sciences, University of Buenos Aires, Buenos Aires, Argentina). All other chemicals were of analytical grade.

Animals and treatments

Female Wistar rats weighing 170-190 g at the beginning of the experiments were used. Procedures involving animals were
conducted in accordance with international guidelines (Guide for Care and Use of Laboratory Animals, National Research Council, USA, 1996, and the Council of the European Communities Directive, 86/609/ECC). Animals were maintained on a 12-h light/dark cycle, at 20-24°C and 45-75% humidity, fed Purina 3 diet and water ad libitum, and randomly divided into four groups of 8 animals each. The reported data are the result of three independent experiments, since all controls were homogeneous.

The following groups were used: control, untreated group; ethanol, treated with 20% (v/v) ethanol in drinking water; HCB, treated with 1 g/kg body weight HCB, suspended in water (40 mg/ml) with the aid of Tween 20 (0.5%, v/v) and administered by gavage 5 days a week, and (ethanol + HCB), treated with ethanol and HCB as described for the ethanol and HCB groups. Feeding bottles were placed in the inverted position to reduce ethanol evaporation. Moreover, the solution was replaced daily in order to reduce possible variation in ethanol concentration. Generally no deaths or external alterations resulted from treatment at this dose of HCB alone. The low level of toxicity was indicative of the minimal absorption of HCB across the intestinal tract when administered in water (11). In previous studies, no changes were observed in any of the parameters studied between animals treated with vehicle and untreated animals (data not shown). For this reason, vehicle-treated controls were omitted from this study.

**Assays**

Treatment lasted 1, 3, 6, or 8 weeks. At the end of treatment, animals from each group were anesthetized with ether, and arterial blood was collected into heparin and used for hepatic function tests. Animals were then sacrificed by decapitation and livers removed. After separating 1 g from each liver for δ-aminolevulinate synthase (ALA-S) determination, a whole homogenate in 0.154 M KCl (1:5, w/v) was prepared for further determinations.

**Hepatic porphyrin content**

Total porphyrins were estimated spectrophotometrically in 5% HCl(w/v) as described by San Martín de Viale et al. (13).

**Hepatic cytochrome P-450**

The content of hepatic cytochrome P-450 was determined by the method of Omura and Sato (14).

**Preparation of enzyme extracts**

The whole homogenate (see Assays) was submitted to differential centrifugation. Mitochondria obtained by centrifugation at 11,000 g were washed with saline and used to measure ferrochelatase activity. Aliquots of the 11,000 g supernatant were used to measure ALA dehydrase (ALA-D) and UroD activities, while another aliquot was centrifuged for 1 h at 105,000 g to separate microsomes for cytochrome P-450 determination. All procedures were carried out at 4°C.

**Enzyme activities**

ALA-S activity was measured in whole homogenates as described by Maines et al. (15) and reported as nmol ALA/h per g liver.

ALA-D activity was measured in the 11,000 g supernatant and is reported as nmol ALA/h per mg protein (15).
UroD activity was measured in the 11,000 g supernatant using uroporphyrinogen III as substrate, and the reaction products were quantified by HPLC as methyl ester porphyrins (12). Enzyme activity is reported as nmol decarboxylated uroporphyrinogen (sum of 7-, 6-, 5-, and 4-COOH porphyrinogen formed)/30 min per mg protein.

Ferrochelatase activity was measured using mitochondria disrupted by thermal shock as the source of enzyme and iron and protoporphyrin IX as substrates according to Ríos de Molina el al. (16). Activity is reported as nmol heme/30 min per mg protein.

Protein concentration

Protein concentration was determined by the method of Lowry et al. (17) using bovine serum albumin as standard.

Liver function tests

Alanine aminotransferase, aspartate aminotransferase, alkaline phosphatase, γ-glutamyl transeptidase, lactate dehydrogenase and pseudocholinesterase were measured using commercial kits (Wiener Lab., Rosario, Santa Fe, Argentina).

Statistical analysis

Results are reported as median and range and were analyzed statistically by nonparametric methods. Kruskal-Wallis ANOVA and the Mann-Whitney test for unpaired samples were employed since there was neither homoscedasticity nor normal distributions.

Results

During the experimental period, treated animals did not suffer from dehydration or malnutrition; weight was similar in all animals which, in general, appeared to be healthy. However, some rats receiving both drugs (2.5% of this group) showed a significant loss of weight and weakness, and they died even though treatment was immediately interrupted. All the other animals were in a healthy condition, and gained weight at the same rate during treatment.

The volume of aqueous alcohol ingested was 16.8 ± 2.8 ml/day and that of the animals receiving only water was 17.3 ± 1.0 ml/day. Therefore, the presence of ethanol in water did not affect the daily volume intake. The daily amount of ethanol ingested by each animal that received alcohol, expressed as g ethanol/kg body weight, was 14.4 ± 2.1 and 14.8 ± 1.3 g/kg in the ethanol and ethanol + HCB groups, respectively.

The evolution of the urinary excretion of ALA, PBG, and porphyrins is shown in Figure 1A-C which shows data representative of three independent experiments. The majority of the rats exposed to ethanol were not different from controls in terms of ALA, PBG and porphyrins. However, 25% of the animals (filled circles in Figure 1A-C) showed higher values than control for the heme intermediates measured. These values were 50-300 vs 21.3 ± 13.3 µg ALA/day (2.5- to 14-fold increase), 18-75 vs 3.0 ± 2.5 µg PBG/day (6- to 25-fold increase) and 6-13 vs 2.9 ± 2.0 µg porphyrins/day (2- to 4.5-fold increase). Excretion of metabolites indicative of porphyria by animals treated with both drugs (ethanol + HCB), or with HCB alone, was considerably elevated beginning from the third week of exposure, and increased in a time-dependent way. In these groups, maximum values were 696 µg ALA/24 h (33-fold increase), 2,547 µg PBG/24 h (849-fold) and 164 µg porphyrins/24 h (16-fold) in the ethanol + HCB group, and 339 µg ALA/24 h (16-fold), 1,420 µg PBG/24 h (473-fold) and 189 µg porphyrins/24 h (67-fold) in the HCB group.

Although the excretion of the precursors ALA and PBG was higher in the ethanol + HCB group than in the HCB group, this was not observed for porphyrin excretion. Some animals (12.5%) of the HCB-treated groups
showed normal excretion values of all heme intermediates, even at the longest exposure time (data not shown).

The liver weight/body weight ratio of animals receiving HCB or ethanol + HCB treatments was 25-60% higher than controls (4.4 ± 0.4%) from the first week of exposure to the end of the experiment (Table 1). In addition, there were no significant differences between the two experimental groups receiving HCB. Animals receiving only ethanol were within the normal range.

HCB, alone or with ethanol, caused hepatocyte enlargement, whereas ethanol alone did not have this effect (data not shown).

The values of hepatic porphyrin of both HCB-treated groups were significantly higher than those of control animals, even during the first week of treatment (Table 1) and these increases were time-dependent for both groups. However, the extent of increase was related to treatment since it was lower in animals receiving both drugs than in those treated only with HCB. The median values obtained for the different times in the ethanol group did not differ statistically from the age-matched control. It is important to note that when the behavior of each individual animal was analyzed some alcohol-treated animals showed increases that reached 16- and 90-fold the control values by the sixth and eighth weeks, respectively.

Hepatic cytochrome P-450 is involved in the metabolism of a wide range of xenobiotics, HCB among them. The results obtained (Table 1) show that the group receiving only alcohol exhibited a slight, transient increase at the beginning of exposure, which was significantly different during the first and third weeks. Both HCB-treated groups displayed similar profiles even when the extent of the alterations differed according to the treatment and intoxication times. In these groups, cytochrome P-450 was rapidly induced since significant increases (two-fold) were found during the first week, with a subsequent reduction over time, but without attaining normal values.

The first, and rate-limiting, enzyme of the heme metabolic pathway, ALA-S, was

![Figure 1. Daily urinary excretion of precursors and porphyrins as a function of treatment time. A, Excretion of δ-aminolevulinate (ALA); B, excretion of porphobilinogen (PBG); C, excretion of porphyrins. Female Wistar rats received 20% ethanol in drinking water and hexachlorobenzene (HCB) by gavage as described in Material and Methods. The figure shows a representative excretion pattern corresponding to one of the three independent experiments. Each point indicates the mean urinary excretion of 4 animals per group. Control (lozenges); ethanol (circles); HCB (triangles); ethanol + HCB (squares); individual rats of the ethanol group with excretion values above the controls (filled circles). The range of control values is represented between dotted lines.](https://example.com/f1.png)
Table 1. Effect of chronic ethanol and/or hexachlorobenzene (HCB) treatment on different hepatic parameters.

<table>
<thead>
<tr>
<th>Group</th>
<th>Week</th>
<th>Lw/Bw (%)</th>
<th>Porphyrins (µg/g liver)</th>
<th>Cytochrome P-450 (nmol/mg protein)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>1</td>
<td>4.2 (3.9-4.8)</td>
<td>0.9 (0.5-1.9)</td>
<td>0.91 (0.64-1.10)</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>4.5 (3.6-5.3)</td>
<td>1.1 (0.7-1.6)</td>
<td>0.98 (0.83-1.25)</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>4.2 (3.2-5.3)</td>
<td>1.0 (0.6-2.4)</td>
<td>0.99 (0.68-1.26)</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>4.7 (3.2-4.8)</td>
<td>1.2 (0.7-2.4)</td>
<td>0.94 (0.94-1.27)</td>
</tr>
<tr>
<td>Ethanol</td>
<td>1</td>
<td>4.4 (3.4-5.1)</td>
<td>0.8 (0.4-1.4)</td>
<td>1.29* (0.65-1.76)</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>4.2 (3.4-5.1)</td>
<td>1.2 (0.7-1.6)</td>
<td>1.30* (0.76-1.59)</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>4.1 (3.4-4.8)</td>
<td>1.0* (0.5-1.6)</td>
<td>1.10 (0.72-1.43)</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>4.0 (3.2-4.6)</td>
<td>1.4* (0.7-9.0)</td>
<td>0.95 (0.89-1.40)</td>
</tr>
<tr>
<td>HCB</td>
<td>1</td>
<td>6.7** (4.6-10.5)</td>
<td>3.5** (1.4-25)</td>
<td>2.13* (0.81-3.56)</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>6.1** (4.0-8.4)</td>
<td>71.0** (13.0-250)</td>
<td>2.51** (1.22-2.62)</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>5.4** (4.3-6.3)</td>
<td>191.0** (38.5-370)</td>
<td>1.68** (0.93-1.73)</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>6.2** (4.4-6.4)</td>
<td>249.0** (119-632)</td>
<td>1.31** (1.23-1.47)</td>
</tr>
<tr>
<td>Ethanol + HCB</td>
<td>1</td>
<td>7.0** (5.3-7.7)</td>
<td>2.5** (1.2-6.5)</td>
<td>1.70** (1.20-2.67)</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>5.6** (5.1-7.6)</td>
<td>89.0** (24-159)</td>
<td>1.98** (1.20-3.70)</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>4.8** (4.8-7.8)</td>
<td>89.0** (23-338)</td>
<td>1.24** (0.98-1.80)</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>5.4** (4.8-7.2)</td>
<td>148.0** (73-569)</td>
<td>1.45** (1.13-1.80)</td>
</tr>
</tbody>
</table>

Data are reported as median and range (in parentheses). N = 6-8 animals per group. Lw/Bw: liver weight/body weight ratio. *0.05<P<0.1, **P<0.05 compared to the respective control (Mann-Whitney test). Treatment conditions were as described in the legend to Figure 1. Porphyrins and cytochrome P-450 were measured as described in Material and Methods.

Table 2. Effect of treatments on hepatic heme metabolic enzymes.

<table>
<thead>
<tr>
<th>Group</th>
<th>Week</th>
<th>ALA-synthase (U/mg protein)</th>
<th>ALA-dehydrase (U/mg protein)</th>
<th>Ferrochelatase (U/mg protein)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>1</td>
<td>7.3 (4.7-8.7)</td>
<td>11.4 (9.1-13.5)</td>
<td>1.8 (1.5-2.2)</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>7.0 (4.8-12.1)</td>
<td>13.8 (11.6-16.2)</td>
<td>1.5 (1.4-2.2)</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>8.6 (7.5-11.4)</td>
<td>12.1 (10.0-14.4)</td>
<td>1.2 (1.1-2.6)</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>9.6 (5.8-11.0)</td>
<td>10.8 (7.6-16.6)</td>
<td>1.7 (1.1-2.0)</td>
</tr>
<tr>
<td>Ethanol</td>
<td>1</td>
<td>7.3 (5.8-15.6)</td>
<td>13.8 (12.1-16.6)</td>
<td>1.7 (1.5-1.8)</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>8.3 (6.2-12.5)</td>
<td>13.0 (9.9-17.7)</td>
<td>1.4* (1.3-1.5)</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>9.8 (6.7-14.7)</td>
<td>12.3 (11.5-15.7)</td>
<td>1.2* (1.0-1.8)</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>8.1 (5.0-12.5)</td>
<td>8.7* (6.0-9.0)</td>
<td>1.2** (0.7-1.5)</td>
</tr>
<tr>
<td>HCB</td>
<td>1</td>
<td>8.3 (4.2-10.4)</td>
<td>9.9 (8.8-12.0)</td>
<td>1.9 (1.5-2.7)</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>8.8 (5.2-22.3)</td>
<td>12.5 (8.1-14.6)</td>
<td>1.8 (1.6-2.4)</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>13.5** (8.3-20.8)</td>
<td>14.7 (4.8-30.8)</td>
<td>2.3* (1.3-4.9)</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>6.2** (6.4-41.6)</td>
<td>20.2 (7.1-27.8)</td>
<td>1.9* (1.3-2.4)</td>
</tr>
<tr>
<td>Ethanol + HCB</td>
<td>1</td>
<td>7.8 (4.2-14.8)</td>
<td>12.1* (9.5-13.2)</td>
<td>1.9 (1.4-2.4)</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>8.3 (4.2-18.7)</td>
<td>12.4* (8.6-13.5)</td>
<td>1.6 (1.4-1.8)</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>22.0*** (15.6-46.8)</td>
<td>15.4* (8.8-30.3)</td>
<td>2.6* (1.4-4.4)</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>26.3** (7.5-54.9)</td>
<td>17.7* (12.3-22.0)</td>
<td>2.3* (1.4-2.9)</td>
</tr>
</tbody>
</table>

Data are reported as median and range (in parentheses). N = 6-8 animals per group. *0.05<P<0.1, **P<0.05 compared to the respective control (Mann-Whitney test). *0.05<P<0.1, **P<0.05 compared to the hexachlorobenzene (HCB) group (Mann-Whitney test). Treatment conditions were as described in the legend to Figure 1. ALA = δ-aminolevulinate.
Ethanol and porphyria cutanea tarda measured in hepatic tissue (Table 2). Until the third week, there were no significant differences between the groups studied. After this time, ALA-S was induced in the HCB- and ethanol + HCB-treated groups and began to show differences between them, with significant and greater increases for the ethanol + HCB-treated group. Exposure to alcohol alone did not elicit any ALA-S induction.

The second enzyme of the pathway, ALA-D, was measured in the hepatic cytosol (Table 2). The ethanol treatment only produced the expected reduction of this enzyme activity by the eighth week, and when administered together with HCB, the maximal increase was lower than that observed with HCB alone.

The fifth enzyme of heme metabolism and the target for HCB in sporadic induced porphyria, UroD, was measured using uroporphyrinogen III as substrate (Figure 2). A 20 to 30% loss in UroD activity was observed in both groups treated with HCB from the first week, and the decrease reached 70% by the end of treatment. During the intermediate weeks, treatment with both drugs, ethanol + HCB, produced a more marked decrease in UroD activity than that observed in rats receiving only HCB. In the group treated only with ethanol, UroD activity was significantly lower than control only during the first week. At the other times assayed, while there were no significant differences between the medians for the two groups (control and ethanol), the lower limits for the ethanol group were always below control, with UroD reductions of 18 and 30% compared to control at the third and sixth weeks, respectively.

Ferrochelatase, the last enzyme of the heme pathway, was reduced in a time-dependent manner after treatment with ethanol. The effect was significant from the third week. Both groups treated with HCB showed statistically significant increases in this enzyme activity after the sixth week of treatment.

The enzymes aspartate aminotransferase and alanine aminotransferase (biomarkers of hepatic cell damage), lactate dehydrogenase...
ase (sensitive to cellular injury, and a marker of hemolysis), γ-glutamyl transpeptidase (a marker of biliary system obstruction), pseudocholinesterase (a marker of exposure to xenobiotics), and alkaline phosphatase were measured as indicators of hepatic injury (Figure 3). The profiles of all enzyme activities for the three treated groups, ethanol, HCB and ethanol + HCB, were similar, and the values showed time-dependent differences with respect to the control, particularly noticeable at the end of treatment. All the enzymes assayed showed increases as a consequence of single (ethanol or HCB) or combined (ethanol + HCB) treatment, with the exception of pseudocholinesterase whose activity tended to decrease in response to the three treatments. In general, the greatest differences from control values were found in the group receiving both drugs (ethanol + HCB).

**Discussion**

In the present study, ingestion of 20% (v/v) alcohol did not cause any alterations in body weight or in the general appearance of the rats. For this reason, we believe that the metabolic alterations observed can be ascribed to specific effects on hepatocytes generated by the drugs, and are not of nutritional origin. Possibly, rats that received both drugs and died represent a group of more susceptible animals. In previous work we have identified three different groups of rats; one that is highly susceptible and responds to HCB treatment within a very short time, another that responds moderately and a third one that is less susceptible and could remain biochemically and biophysically stable even after long treatment (18). It has been suggested that this difference may be of multigenic origin (9), as proposed by Elder (2) for

![Figure 3. Hepatic enzymes as markers of liver injury. Data points indicate the median of 6-8 animals per group. Ethanol (circles); HCB (triangles); ethanol + HCB (squares). The range of control values is represented between the dotted lines. *P<0.05 compared to controls (Mann-Whitney test). HCB = hexachlorobenzene; AST = aspartate aminotransferase; AP = alkaline phosphatase; PCh = pseudocholinesterase; ALT = alanine aminotransferase; GGT = γ-glutamyl transpeptidase; LDH = lactate dehydrogenase.](image-url)
human acquired PCT. In this respect, the author defines as susceptible the subjects that can develop ethanol-induced PCT, with the mechanism involved being unknown.

Ibrahim et al. (19) also found different degrees of susceptibility to the inhibitory effect of ethanol on the heme metabolic pathway of rabbits.

The lack of alcohol-mediated induction of ALA-S activity observed (Table 2), in accordance with the results of Kondo and Shimizu (20), may be the consequence of the insufficient ethanol-induced decrease in UroD, which did not reach the 50% level necessary to deregulate the pathway (12). The decreased ferrochelatase activity observed for this group supports the interpretation of Doss et al. (7,21), who attributed this reduction to the lack of the enzyme’s cofactor, pyridoxal phosphate. The decrease in ferrochelatase and UroD activities elicited in the ethanol-treated group would not be sufficient to reduce the ALA-S regulatory heme pool. The decreases observed here in these heme pathway enzymes agree with the reported in vivo effect of ethanol on them (22).

Nevertheless, the small but sustained block produced by ethanol at the level of UroD might be the cause of hepatic porphyrin accumulation exhibited by all animals during the experiment, and of the increase in the excretion of porphyrins and their precursors observed in some of the exposed animals. In the most susceptible animals, these increases exceeded 14, 25 and 4.5 times control values for ALA, PBG and porphyrins, respectively. Although they did not reach the maximum values obtained with HCB, these significant increases may indicate an alteration in the heme metabolic pathway, specifically at the UroD level. This hypothesis is also supported by the increase in the hepatic porphyrin content shown by some animals in the ethanol group. However, the maximum values of ALA and PBG excretion in the ethanol + HCB group were approximately twice as high as those of the HCB group. Again, this would suggest that ethanol affects the development of HCB-induced porphyria.

These results agree with those of Doss (7), who found that patients who ingest more than 60 g/day of alcohol show a 3- to 6-fold increase over normal values in urinary excretion of ALA, PBG, and porphyrins. Since an increase in urinary excretion of porphyrins and precursors was not observed in most of the animals that received only ethanol, we suggest that the animals that do show increases might be predisposed to developing sporadic porphyria. Similarly, in humans, ethanol and various porphyrinogenic agents were reported to trigger porphyria only in susceptible subjects (2).

Since ethanol can generate free radicals (23), the early decrease in UroD activity produced by this drug may be due to UroD inhibitor formation through oxygen free radical species (24).

The initial increase in P-450 content observed in the three treated groups corresponds to a primary response to administration of the drug in the form of an induction of this hemoprotein due to its detoxifying function. This is followed by a mechanism of adaptation to the treatment that leads to the normalization of cytochrome content (25). The most significant increases detected were observed in the group treated with both drugs at one week, and can be attributed to the sum of the effects of ethanol and HCB on the different P-450 isoforms involved in their metabolism (26,27).

With regard to the effects of alcohol on the development of HCB-induced porphyria, ethanol administered together with HCB was found to increase urinary excretion of the precursor metabolites ALA and PBG (Figure 1A,B) as a consequence of the significantly greater induction of ALA-S observed during the final weeks of treatment when compared to HCB alone (Table 2). Despite the fact that the most marked de-
crease in UroD occurred earlier, and in some weeks was greatest in the ethanol + HCB group, enzyme activity values were not sufficiently different from those of the HCB-treated animals to produce differences at the level of urinary porphyrin excretion (Figures 2 and 1C).

The lower increase in ALA-D activity observed in the group with double treatment compared to HCB alone shows that ethanol is able to modify the response to HCB.

Alcohol exacerbated the increase produced by HCB in ferrochelatase levels at longer treatment times. Since ferrochelatase is the last enzyme of the pathway, this increase could be a compensatory response to the decrease in UroD.

ALA-S, ferrochelatase, and UroD were the heme metabolic pathway enzymes most affected in the group receiving both drugs, suggesting a synergistic effect of ethanol on the induction of experimental PCT by the fungicide. The differences in the levels of these enzyme activities did not result in increased porphyrin excretion or hepatic accumulation. This result can be attributed to a competition by the two drugs for some type of cytochrome which is responsible for the tetrapyrrole oxidation and excretion.

The liver weight/body weight ratio, a marker of macroscopic hepatic alteration, was modified by the fungicide HCB, but not by ethanol. Consequently, the enlarged hepatocytes in HCB-treated rats were probably due to changes in the reticuloendoplasmic system, as well as to the proliferation of small lipid droplets, as previously reported (28). The similar alterations in liver weight/body weight ratio observed in both HCB groups, but not in ethanol-exposed rats, did not parallel the changes in plasma enzyme activities, since plasma enzyme levels were affected similarly in all treated groups. Therefore, hepatic damage was not correlated with development of porphyria, and we consider the alteration of the liver weight/body weight parameter to be related to the onset of sporadic porphyria in experimental animals. Moreover, since alcohol treatment did not affect this ratio under our experimental conditions, ethanol would not generate porphyria per se, but instead would predispose to, and/or exacerbate, the disease induced by porphyrinogenic drugs such as HCB.

The mild effects caused by the alcoholic treatment applied here might be due to the dose of ethanol used. This treatment, however, allowed us to conclude that this dose of alcohol may induce a pre-porphyric state without causing hepatic damage.

Although alcohol may induce sporadic porphyria only in genetically predisposed humans (2), the present results indicate that ethanol does exacerbate the biochemical manifestations in overt porphyrías. Consequently, since slightly porphyrinogenic compounds such as ethanol might produce a pre-porphyric state, it is advisable to avoid the intake of alcohol in all cases of suspected hepatoporphyría.

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References


