Ethanol in low chronic dose level attenuates major organic effects in malnourished rats

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ABSTRACT

The aim of the present study was to evaluate the chronic toxicity of ethanol low blood levels in malnourished rats. Female Wistar rats (220 g) were subjected to either an ad libitum diet (W, well-nourished, n=10) or food restriction (M, malnourished, n=10). Water (WW and MW) or ethanol solution (W5% and M5%) was offered to half of each nutritional group (n=5) as the only fluid source. The treatment was continued for two months. After sacrifice, blood biochemical parameters and macroscopic, histologic and morphometric evaluation of the liver were performed. Results indicated that: Ethanol consumption was higher in malnourished rats and minimized body weight loss in malnourished rats, while it decreased the body weight gain in well-nourished ones. Behavioral ethanol intoxication was more severe in malnourished rats. Malnutrition decreased hematocrit and hemoglobin but, on the other hand, ethanol was a protective factor of that effect (hemoglobin: MW 10.6 mg/dl / ME 13.02 mg/dl, p< 0.05). Ethanol increased the relative liver weight of both well-nourished and malnourished rats. Ethanol intake minimized iron pigment, collagen area and binuclear hepatocyte/field increased by malnutrition. These data are in accordance with previous reports which showed ethanol as an important source of calories and, even chronically, ethanol still attenuates the effects of malnutrition.

Key terms: ethanol, low dose, malnutrition, toxicity

1. INTRODUCTION

Alcoholism may be considered as a chronic illness with a multifactorial etiology combining several components, genetic as well as organic, psychological and environmental (Addolorato et al., 2005).

As ethanol may be a source of calories to malnourished organisms and because of its anxiolytic action (Almeida et al., 1996), which may alleviate the stress and discomfort brought from hunger, it is possible to consider malnutrition as a risk factor for alcoholism. In fact, malnourished rats do drink more ethanol than well-nourished rats (Stiglick & Woodworth, 1984).

Nonetheless, previous studies carried out by our group have shown that low doses of ethanol (5% v/v, ad libitum as the only available fluid source) had challenging effects on malnourished rats. Instead of increasing toxicity, rats kept under restricted food presented beneficial effects such as less mortality and better offspring psychomotor performance. (Da-Silva et al., 2000).
Increased body weight in malnourished rats was detected after 90 days of exposure to ethanol at 20% v/v (Da-Silva et al., 1982).

Aguiar et al (2004) evaluated the threshold dose in which ethanol would be useful as an energy source to malnourished rats. Ethanol solutions at 5, 10 and 20% v/v decreased body weight loss in malnourished rats and did not change biochemical parameters (hematocrit, hemoglobin, total protein and albumin). On the other hand, ad libitum rats presented body weight loss when solution was 10% v/v, with worse effects at 40% v/v. However, the malnourished rats that benefited from ethanol calories showed more behavioral intoxication than the well-nourished ones (Aguiar et al., 2006). One limitation was that the ethanol treatment lasted only 7 days. Thus, the effects of chronic exposure were not investigated. Besides, increased body weight is not a guarantee of absence or decrement of drug toxicity.

The pathologic consequences of acute and chronic alcohol abuse are multifactorial and multisystemic (Molina et al., 2003). Ethanol toxicity has been extensively studied but most research has been performed with high doses of the drug. For instance, chronic consumption of high ethanol concentrations (11 - 12g/ Kg) has negative effects in the liver and impairs the function of a wide range of cells and tissues (Iumuro et al., 1997). The few studies with lower dose levels indicate that ethanol may induce morphological and functional changes in hepatocytes (Puzziferri et al., 2000). As far as we know, the effects of malnutrition when ethanol exposure occurs in low dose levels and chronically have not been investigated. Studies designed to evaluate toxicity in target organs under the effect of low doses of ethanol, especially in long term exposure, are necessary. This represents a research area of great medical interest, mainly considering that most cultures approve moderate alcohol consumption and scientific evidence seems to support it (Dufour, 1999).

The food restriction model is closer to what occurs to millions of people exposed to poverty. About 8% of the adult Brazilian population is malnourished, which is thought to happen due to poverty resulting from under-employment and low wages (FAO, 2006). This situation could be favorable to alcohol intake and consequently to alcohol addiction.

Our hypothesis is that ethanol, in spite of being a caloric source to malnourished rats, will still be toxic to target organs and that this toxicity could be modified by malnutrition. The aim of the present study was to evaluate the toxicity of chronic exposure (two months) to low doses of ethanol (5% v/v) in two different nutritional states (food ad libitum and restricted to 50%).

2. MATERIAL AND METHODS

2.1. Animals and treatment

All animal work was carried out in accordance with the Brazilian Animal Protection Law and was approved by the Local Ethics Committee for Research at Hospital Universitário Antônio Pedro, Universidade Federal Fluminense, Brazil (n 172/05).

Twenty female Wistar rats (Rattus norvegicus) weighing between 200-220g were placed in individual polyethylene cages (41x 34x 16 cm) and randomly assigned to groups of either water or ethanol, which were divided in two nutritional groups, thus generating four experimental groups. All the procedures were performed between 10:00 and 11:00 a.m. Groups were treated as described below:

- Well-nourished Water (WW) (n=5): receiving food (a commercial diet Nuvital Purina® Brazil) and water ad libitum;
- Malnourished Water (MW) (n= 5) food restricted to 50% of consumption of WW and water ad libitum;
- Well-nourished ethanol 5% v/v (W5%) (n= 5): receiving food ad libitum and ethanol solution at 5%v/v as the only drinking source;
- Malnourished ethanol 5% v/v (M5%) (n= 5): food restricted to 50% and ethanol solution at 5%v/v as the only drinking source.
The treatment lasted two months, which is equivalent to 10 years of a human being’s lifetime.

During the first three weeks of the experiment, all groups received water as liquid source, and the MW and M5% ones were submitted to food restriction of 50% of the ad libitum consumption until they reached a weight loss of 20% (Stiglick & Woodworth, 1984). Only after this criterion had been achieved did ethanol treatment begin.

2.2. Intoxication test

Rats were evaluated weekly through a modified behavioural intoxication scale proposed by Majchrowicz (1975). Before being weighed weekly, rats were observed in individual cages for 30 seconds and received the corresponding grade: 0: neutrality; 1: hypoactive, mildly ataxic; 2: ataxic, elevated abdomen; 3: ataxic, non elevated abdomen; 4: loss of righting reflex, retain eye blink reflex; 5: loss of righting reflex, loss of eye blink reflex.

2.4. Biochemical analyses

Blood was collected by intracardiac puncture, under ether anaesthesia (Brzóska et al., 2003; Aguiar et al., 2006). Laboratory analysis of total proteins, albumin, hematocrit and hemoglobin were conducted using standard protocols, according to the manufacturers’ instructions (Kits BIOCLINâ, QUIMASA QUÍMICA BÁSICA, MG, Brazil).

2.5. Procedures for histopathological and histomorphometrical analyses

Rats were killed by cervical dislocation. The liver was quickly removed, cleaned in cold saline, dried with filter paper and weighed. Then, a portion was dipped in Gendre’s fixative for 6 hours and then maintained in 10% phosphate buffered formalin, dehydrated and embedded in paraffin. Sections of 5 μm were prepared and stained with hematoxylin and eosin (H&E) for histopathological routine, Periodic Acid- Schiff (PAS) for hepatocyte area (Kiernan, 1990), Picrosirius Red (PS) for collagen (Junqueira et al., 1979) and Perls and Stein techniques were performed in order to detect iron pigments and bilirubin pigments, respectively (Kiernan, 1990). All sections were analysed by light microscopy (Leitz DFV) under the supervision of a pathologist (PJSF).

Histomorphometry was performed using an image analysis system composed of a digital camera (Evolution MP Color Media Cybernetics) coupled to a light microscope (Nikon Eclipse E400). Five fields of liver parenchyma (10x objective lens), considering avoiding liver capsule, portal spaces or central veins/ animal, were captured for hepatocyte binuclearity assessment. Ten fields of liver parenchyma from the perivenular (Zone III) and periportal (Zone I) region/ animal were also captured by a 40x objective lens. PAS stained slides were used to capture images for hepatocyte parameters quantification and PS for collagen. Quantification was estimated on captured high quality images (2048 x 1536 pixels buffer) by considering the total field area using the software Image-Pro® Plus 4.5.1 (Media Cybernetics, USA).

2.3. Ethanol blood determination

An independent group of rats kept in the same experimental conditions was used for ethanol blood level measurement. These rats were exposed to one week of continuous 5% v/v ethanol. Blood was collected from the tip of the tail. Ethanol blood was measured using a head space technique (Da-Silva & McLean, 1988). Briefly, 10 μL of blood was placed in a 1.9 mL vial together with 10 μL of the internal standard (100 mg% propanol) and the vial was sealed. After 60 min of equilibration in a water bath at 38 ± 1°C, 500 μL of the air present in each vial was injected into a gas-chromatograph (Hewlett-Packard 573 A) connected to a C-R 1B Shimadzu integrator. The ratios of the peak areas derived from ethanol and the internal standard were compared with a standard curve of ethanol solutions at 40, 20, 10, 5 and 2.5 mg% (prepared with ethanol absolute PA Merck® + saline).
2.6. Statistical analysis

Experimental groups were compared by non-parametric tests (U- Mann Whitney) and whenever data fulfilled criteria for parametric statistics, the test was performed by using SPSS for Windows 12.0. Analyses of variance (ANOVA) of one factor (one-way), followed by the Student’s t test, were used to compare body weight data. The two-way ANOVA was used to compare the effects of two independent factors: the nutritional state (well-nourished or malnourished) and the treatment (water and ethanol). Intoxication and biochemical parameters were analyzed with the U-Mann Whitney Test (WW x MW and W5% x M5%). Data were presented as mean ± standard deviation (SD) or median (maximus and minimal values), and p values below 0.05 (two tailed) were considered significant.

3. RESULTS

3.1. Ethanol intake, water intake and ethanol blood levels

Ethanol consumption in g/kg body weight was higher (approximately twice) in the malnourished group throughout the experiment. (Means in g/kg for each week: W5% 6.3; 8.6; 7.9; 7.0; 5.8; 6.5; 6.3; 6.2/ M5%:11.3; 16.4; 22.1; 17.5; 16.3; 14.5; 15.1; 14.1 g, p<0.05 U- Mann Whitney Test for all weeks W5% x M5%). In spite of this increased consumption, there were no statistically significant differences between groups concerning ethanol blood levels (mean: 1.43 mg% ± 1.86).

There were no statistically significant differences on daily water intake between WW and MW groups throughout the experiment. Means for the last week of experiment were as following: WW: 32 ± 1.92 mL and MW: 23.3 ± 1.15 mL.

3.2. Body weight

Ethanol had opposite effects according to the nutritional state. Malnourished rats treated with ethanol had the severe effects of malnutrition minimized and increased survival. Two malnourished rats which were not exposed to ethanol died before the end of the experiment. At the time, their body weight was around 120 g. On the other hand, rats kept with food ad libitum decrease the body weight gain significantly when exposed to ethanol at 5% (p<0.05) (Figure 1).

3.3. Intoxication

Significant differences related to intoxication grades between groups were detected only on the second week. (Intoxication grade: W5%: 0 x M5% 2.0 p< 0.05 Mann Whitney U test).

3.4. Biochemical analyses

The chronic ethanol intake (5% v/v) did not change the biochemical parameters of the well-nourished rats. Malnutrition decreased hematocrit and hemoglobin levels (Hematocrit: WW: 36% x MW: 31% / Hemoglobin WW: 13.5 g/dl x MW: 10.6 g/dl p< 0.05 Mann Whitney U test). However, ethanol consumption at 5% v/v attenuated the effects of malnutrition (Hematocrit: M5% 37.2% and hemoglobin M5%: 13.2 g/dl). No significant differences between groups were found in total protein and albumin (Table I).

3.5. Liver and histopathology

Malnutrition decreased liver weight (g/100g body weight). Ethanol increased this parameter in both groups. The two-way ANOVA revealed significant effects of nutrition and ethanol, without significant interaction between factors. WW: 2.99 ± 0.25 W5%: 3.38 ± 0.30 MW 2.61 ± 0.10 M5% 3.02 ± 0.17 (p< 0.05 for nutritional state F: 10.4 and ethanol intake F: 12.06 - ANOVA two-way).

Histological examination of the liver was normal in all well-nourished rats (WW and W5%). There were no significant effects of ethanol in liver histology. Briefly, the expected inflammatory portal infiltrate was observed in all animals (WW, W5%, MW and M5%), the trabecular structure of the lobules was preserved and bilirubin
**Figure 1:** Body weight (g) of well-nourished and malnourished rats drinking either ethanol at 5% v/v or water for eight weeks

* First to third week: water for all groups until malnourished rats reached a weight loss of 20%.

† Two rats died in group MW.

# * p< 0.05 (ANOVA one-way followed by Student’s t test)

### TABLE I

Hematocrit, hemoglobin, total protein and albumin values of the well-nourished and malnourished rats exposed to water or ethanol at 5% v/v for eight weeks

<table>
<thead>
<tr>
<th></th>
<th>Hematocrit (%)</th>
<th>Hemoglobin (g/dL)</th>
<th>Total Protein(g/dL)</th>
<th>Albumin(g/dL)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Well-nourished</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water (WW)</td>
<td>36.6 a (34-39)</td>
<td>13.5 b (12.5-14.5)</td>
<td>6.9 (6.7-7.4)</td>
<td>4.2 (3.9-4.8)</td>
</tr>
<tr>
<td>Ethanol (W5%)</td>
<td>38.2 (37-40)</td>
<td>13.4 (12.5-14.5)</td>
<td>7.5 (7.0-7.9)</td>
<td>4.6 (4.2-5.0)</td>
</tr>
<tr>
<td><strong>Malnourished</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water (MW)</td>
<td>31 a (28-34)</td>
<td>10.6 b c (10-11.5)</td>
<td>6.6 (5.9-7.3)</td>
<td>4.0 (4.0-4.1)</td>
</tr>
<tr>
<td>Ethanol (M5%)</td>
<td>37.2 (33-40)</td>
<td>13.0 c (12.3-13.5)</td>
<td>6.7 (6.2-7.2)</td>
<td>4.1 (3.8-4.6)</td>
</tr>
</tbody>
</table>

* Values are significantly different (p< 0.05 U-Mann Whitney) according to paired superscripts

Median (minimal - maximus values)

WW (n=5) / WE5% (n=5) / MW (n=3) / M5% (n=5)
pigment was not observed. In malnourished rats (MW), important cell atrophy and iron pigmentation were observed, but the iron pigmentation was minimized by ethanol intake (M5%). The hepatocyte area quantification was sensible to detect the cell atrophy observed in malnourished rats (MW), thus certifying that the method was sensible to detect that chronic ethanol intake did not change the cell area (Table II). The cell atrophy observed in malnourished rats (MW) was confirmed by the increased number of the hepatocyte cell/field (F= 12.28 p=0.004 two-way ANOVA) (Table III). Ethanol intake interacted with malnutrition only when the number of binuclear hepatocytes was analysed. (F=11.02 p= 0.006 two-way ANOVA).

Malnutrition (MW) increased the collagen content in both regions. Ethanol intake minimized this effect only in the perivenous region (Table IV).

TABLE II

Hepatocytes area (μm²) in perivenous and periportal regions of well-nourished and malnourished rats exposed to water or ethanol at 5% v/v for eight weeks

<table>
<thead>
<tr>
<th></th>
<th>Perivenous</th>
<th>Periportal</th>
</tr>
</thead>
<tbody>
<tr>
<td>Well-nourished</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water (WW) (n=5)</td>
<td>203.80 ± 64.87</td>
<td>232.50 ± 77.47</td>
</tr>
<tr>
<td>Ethanol (W5%) (n=5)</td>
<td>227.80 ± 69.96</td>
<td>236.63 ± 82.42</td>
</tr>
<tr>
<td>Malnourished</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water (MW) (n=3)</td>
<td>123.15 ± 38.86</td>
<td>160.28 ± 53.01</td>
</tr>
<tr>
<td>Ethanol (M5%) (n=5)</td>
<td>127.21 ± 47.13</td>
<td>118.58 ± 47.12</td>
</tr>
</tbody>
</table>

* Field area: 0.175 mm² (H&E 100x)
Mean ± SD
Image pro-Plus 4.0 (PAS 400x) Program was used.

TABLE III

Number of hepatocyte cells and binuclear hepatocytes in each histological field evaluated by optical microscopy of the liver of well-nourished and malnourished rats exposed to water or ethanol at 5% v/v for eight weeks

<table>
<thead>
<tr>
<th></th>
<th>Binuclear</th>
<th>Hepatocyte cell/ field</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Hepatocyte</td>
<td></td>
</tr>
<tr>
<td>Well-nourished</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water (WW)</td>
<td>16.92 ± 4.49</td>
<td>295.40 ± 21.80</td>
</tr>
<tr>
<td>Ethanol (W5%)</td>
<td>16.96 ± 3.63</td>
<td>299.60 ± 38.78</td>
</tr>
<tr>
<td>Malnourished</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water (MW)</td>
<td>37.26 ± 3.30</td>
<td>378.0 ± 107.76</td>
</tr>
<tr>
<td>Ethanol (M5%)</td>
<td>20.80 ± 6.18</td>
<td>358.4 ± 55.86</td>
</tr>
</tbody>
</table>

* Field area: 0.175 mm² (H&E 100x)
Mean ± SD
ab p< 0.05 for nutritional state (*F= 22.08/ b F= 155.83 - Two-way ANOVA)
TABLE IV

Collagen area (μm²) of perivenular and periportal regions of well-nourished and malnourished rats exposed to water or ethanol at 5% v/v for eight weeks

<table>
<thead>
<tr>
<th></th>
<th>Perivenousa,c</th>
<th>Periportalb</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Well-nourished</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water (WW)</td>
<td>494.12 ± 289.63</td>
<td>703.99 ± 528.19</td>
</tr>
<tr>
<td>Ethanol (W5%)</td>
<td>492.11 ± 303.15</td>
<td>560.16 ± 442.96</td>
</tr>
<tr>
<td><strong>Malnourished</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water (MW)</td>
<td>1531.72 ± 904.74</td>
<td>1435.76 ± 1370.93</td>
</tr>
<tr>
<td>Ethanol (M5%)</td>
<td>1006.52 ± 529.13</td>
<td>1010.23 ± 690.56</td>
</tr>
</tbody>
</table>

Mean ± SD

\[ a, b \) \ p< 0.05 for nutritional state (a F= 29.16/ b F= 10.27 - Two-way ANOVA)

\[ c \] \ p<0.05 for ethanol intake (F= 5.03 - Two-way ANOVA)

WW (n=5) / W5% (n=5) / MW (n=3)/ M5% (n=5)

Image pro-Plus 4.0 Program was used.

DISCUSSION

Data of the present study are indicative that ethanol in low doses can be toxic but still attenuates the effects of malnutrition. This assumption was derived from the exposure to low levels of ethanol chronically. Thus, well-nourished rats exposed to ethanol presented a decrease in body weight gain and increase in relative liver weight without important toxic effects on liver morphology or blood biochemistry. This could be suggestive of the known malnourishing effect of ethanol, which, in this case, was mild. In spite of being a malnourishing drug, ethanol decreased the severe effects of food restriction. Ethanol at 5% v/v in drinking solution was beneficial to malnourished rats in all parameters evaluated in this study (body weight, biochemical parameters and liver histopathology). This is in accordance with Aguiar et al (2004), who reported that malnourished rats utilized ethanol calories for body weight maintenance up to ethanol concentrations in drinking water at 20% v/v.

Malnutrition significantly increased ethanol consumption in g/kg body weight. This finding is in agreement with the study reported by Stiglick & Woodworth (1984), in which food restricted rats displayed higher ethanol consumption. The higher intake of ethanol resulted in eventual higher degrees of intoxication (only on the second week of the experiment). Water intake was not different between malnourished and control rats. Therefore, increased ethanol intake cannot be considered as non-specific increased fluid intake. Different mechanisms might be envolved in this increased ethanol intake, which has also been observed by other authors (McGregor et al, 1999). Previous studies have demonstrated a close relationship between changes in body weight and voluntary ethanol intake (Aguiar et al, in press).

Differences in ethanol blood levels were not significant. As ethanol distributes in water, volume of distribution is decreased proportionally to fat content. This could explain why even drinking more, blood levels were the same in both nutritional conditions.

Food restriction resulting in malnutrition decreased relative liver weight. This could be because of the atrophy and hepatocyte hipoplasia (Parra et al, 1995). On the other hand, the association of ethanol in malnourished rats resulted in increase of liver weight of 15% (compared to group
MW), while in well-nourished rats, the increase was of 13%. Chronic consumption of ethanol alters Na\(^+\) and K\(^+\) channels, thus increasing intracellular water (Wondelgem & Davis, 1994, Häussinger & Schliess, 1995) and cell size as well as promoting fat and proteins increase (Lieber, 2004). In the present study, the increase in relative liver weight was an important marker of ethanol toxicity even when light microscopy was unable to detect significant changes.

There has been little research directed to the study of toxicity of chronic exposure to ethanol in low doses. Puzziferri et al (2000) reported that chronic exposure at 3% v/v ethanol with blood levels of 4 mg% in Wistar rats resulted in a slight, although significant increase in lipidic mytochondrial peroxidation. This could be indicative of more extensive damage previously observed in rats exposed to high doses of that substance. A higher dose of ethanol, (40 % v/v) results in steatosis, hepatitis, necrosis of hepatocytes and fibrosis, thus being suggested as an experimental model of the hepatic alcoholic disorder in Wistar rats (Keegan et al., 1995). Other authors who evaluated the toxicity of low doses of ethanol. Brzóska et al (2003) reported that Wistar rats exposed to 10% v/v ethanol for 12 weeks did not show any body weight changes. However, the authors reported important changes in hepatic trabecular structure and increased hepatocytes with cytoplasmatic vacuoles in zones I and III. Ito et al (2007) reported no changes in liver injury markers in Wistar rats drinking ethanol at 10% v/v. Histological examination of the liver showed microvesicular steatosis.

The characteristic Liver Alcoholic Disorder (steatosis, hepatitis and fibrosis-Harrison & Burt, 1993), is not observed when ethanol concentrations are low. However, all the authors reported some changes in liver parameters that could be predictive of toxicity. In our study, the most sensitive parameter was the liver weight.

Our study also demonstrated that malnutrition induces several effects in liver histopathology especially in zone III. Ethanol did not reverse all these changes but minimized them. Apparently, the known effects of malnutrition (Parra et al., 1985) are attenuated by the caloric effect of the drug, which at this dose level would be more beneficial than its slight and, perhaps, initial toxicity. Of great importance was the observation that malnourished animals presented increased deposition of collagen in the perivenular region, which was attenuated by ethanol. The most commonly associated characteristic of ethanol is an increased deposition of collagen (Gutiérrez-ruiz & Gomez-Quiroz, 2007). One can dare to speculate that in malnourished animals ethanol would diminish the potential danger of induced fibrosis. In low dose levels, the toxic effects of the drug would not be as important as its caloric protection against the severity of malnutrition. Perhaps the concentration of 5% or blood levels of 1.4mg% would be the threshold for this paradoxical effect. In humans, these blood levels are observed after the ingestion of 30 mL of beer, which is unlikely to occur in usual conditions. Therefore, we are speculating about protective effects in much lower levels than the low risk levels established by World Health Organization (around 2-3 bottles a day). However, if a protective effect is unlikely to occur, in human conditions, it is extremely important that even very low levels of ethanol might already be toxic. Mild and perhaps initial signs of toxicity were detected in liver morphology and brain function (intoxication).

Many people in developing countries drink ethanol and are already malnourished. Ethanol is not an ideal source of calories, but, unfortunately, can be the cheapest. If it is possible that its calories may diminish the deleterious effects of malnutrition, but certainly in very restricted dose levels and conditions. Nevertheless, it must be considered that chronic exposure to ethanol, even at low doses (3% v/v), may have important brain effects. Dopamine receptors, which are related to the complex events that result in drug dependence, have been described to change their density or affinity (Lograno et al., 1993). Whether ethanol can be attractive to the malnourished organism for its caloric value and it may indeed be protective at some
blood levels, it is still relevant to consider that ethanol dependence will be more likely to occur. Malnutrition would be a risky factor for alcoholism because it favours the consumption of higher volumes of ethanol, which may lead to the development of ethanol dependence. Future research should address this hypothesis.

In summary, ethanol is a complex drug and its effects may vary depending on dose levels and nutritional state. It is not just a quantitative difference, but a qualitative one as well (toxic to the well-nourished, protective to the malnourished). However, because of the complexity of the dependence phenomenon and of the effects of ethanol in target organs, brain is still vulnerable to ethanol effects even at low doses. In our study, this could be seen in higher and intermittent grades of intoxication. Although and because it is attenuated by ethanol, malnutrition may not change this hypothesis.

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