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Preprint del documento publicado en Alcohol, 2011, 45, 183-192

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Cómo citar el documento:


(Se recomienda indicar fecha de consulta al final de la cita. Ej: [Fecha de consulta: 19 de agosto de 2010]).
Continuous vs. discontinuous drinking of an ethanol liquid diet in peripubertal rats:
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lymphocyte subset populations.

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Abstract

Excessive alcohol consumption continues to be a major public health problem, particularly in the adolescent and young adult populations. Generally, such a behavior tends to be confined to the weekends, to attain frequently binge drinking. This study in peripubertal male rats compares the effect of the discontinuous feeding of a liquid diet containing a moderate amount of ethanol (6.2 % wt/vol) to that of continuous ethanol administration or a control diet, taking as end points the 24-h variations of plasma prolactin levels and mitogenic responses and lymphocyte subset populations in submaxillary lymph nodes and spleen. Animals received the ethanol liquid diet starting on day 35 of life, the diet being similar to that given to controls except for that maltose was isocalorically replaced by ethanol. Ethanol provided 36% of the total caloric content. Every week, the discontinuous ethanol group received the ethanol diet for 3 days and the control liquid diet for the remaining 4 days. After 4 weeks, rats were killed at 6 time intervals, beginning at 0900 h. A significant decrease of splenic cells’ response to concanavalin A, and of lymph node and splenic cells’ response to lipopolysaccharide was found in rats under the discontinuous ethanol regime, as compared to control or ethanol-chronic rats. Under discontinuous ethanol feeding mean values of lymph node and splenic CD8⁺ and CD4⁺-CD8⁺ cells decreased, whereas those of lymph node and splenic T cells, and splenic B cells, augmented. In rats chronically fed with ethanol, splenic mean levels of CD8⁺ and CD4⁺-CD8⁺ cells augmented. Both modalities of ethanol administration disrupted the 24 h variation in immune function seen in controls. Mean plasma prolactin levels
increased by 3.6 and 8.5 fold in rats chronically or discontinuously fed with alcohol, respectively. The immune parameters examined in an additional group of rats fed regular chow and water ad libitum did not differ significantly from control liquid diet. The results support the view that the discontinuous drinking of a moderate amount of ethanol can be more harmful for the immune system than a continuous ethanol intake, presumably by inducing a greater stress as indicated by the augmented plasma prolactin levels observed.

**Key Words:** Binge drinking – Lymph nodes - Spleen - Mitogenic responses - Lymphocyte subsets - Circadian rhythms – Prolactin - Stress
Introduction

Excessive alcohol consumption continues to be a major public health problem in adolescent and young adults (Stolle et al., 2009). High levels of ethanol consumption followed by repeated episodes of withdrawal, as typifies by binge drinking, are common in these age groups and appears to be a particularly harmful way to drink (Duka et al., 2004; Stephens et al., 2005). Even in young social drinkers, with relatively low overall levels of alcohol consumption, both cognitive deficits and mood changes are seen in those with a history of discontinuous drinking (Townshend and Duka, 2005).

There is considerable evidence indicating that ethanol consumption alters immune system function and leads to increased susceptibility to infections and neoplastic diseases (Nath and Szabo, 2009; Lau et al., 2009; Nava-Aguilera et al., 2009; Szabo and Mandrekar, 2009). As indicated by studies in peripubertal rats, chronic administration of ethanol also resulted in significant changes of the circadian organization of the immune response (Jiménez et al., 2005).

In view of the evidence that a significant proportion of the adolescent and young adult populations tend to consume alcohol in a discontinuous pattern at weekends, studies on the effects of alcohol exposure on the adolescent immune system using the chronic alcohol feeding models would probably fail in giving information on the consequence of ethanol drinking behavior. Moreover, none is known as to whether the discontinuous consumption of ethanol affects the circadian organization of the immune response. This prompted us to undertake the present study whose objective was to
compare the effect of the discontinuous feeding (3 days/week) of a liquid diet containing a moderate amount of ethanol with that of a continuous ethanol administration or a control diet, taking as end points the 24-h variations of mitogenic responses and lymphocyte subset populations in lymph nodes and spleen of peripubertal rats. Plasma prolactin levels were also measured as an index of the stress produced by ethanol intake.

**Materials and methods**

*Animals and experimental design*

Five week-old, peripubertal, male Wistar rats were kept under standard conditions of controlled light (12:12 h light/dark schedule) and temperature (22 ± 2 C). Prior to treatment, animals were randomly assigned to one of the following three treatment groups (with an n of 48 animals per group, chosen based on a power analysis performed using preliminary data): (a) control liquid diet; (b) discontinuous ethanol diet; (c) chronic ethanol diet. An additional group of 48 rats fed with regular chow and water *ad libitum* was also studied.

A liquid diet mode of ethanol administration was employed (Dodd and Shorey-Kutschke, 1987; Lieber and DeCarli, 1994). Animals received the liquid diet for 4 weeks, starting on day 35 of life. The diet contained an aqueous suspension of pulverized casein, l-methionine, vitamin mixture, mineral mixture, sucrose, xanthum gum, choline bitartrate, Celufil cellulose, corn oil and maltose. Percent composition of the diet was 35% fat, 18% protein and 47% carbohydrates. The ethanol-fed group received a similar diet except for that maltose was replaced by 96% ethanol. Final ethanol concentration was 6.2% (wt./vol)
and ethanol replacement was isocaloric providing about 36% of the total caloric content of the diet. To mask the ethanol taste, saccharin was added to the experimental ethanol diet; animals under control liquid diet also received saccharin. For the discontinuous ethanol group, the rats received the ethanol diet 3 days/week, the remaining 4 days receiving the control liquid diet.

Rats were caged in groups of 4 animals/cage and had access to the liquid diet ad libitum. An additional group of rats fed with regular chow and water ad libitum was also studied. Control and experimental liquid diets were freshly made each day. Daily consumption of liquid diet (mL/rat, average of 4 rats) was 36.9 ± 7.3 (control), 34.8 ± 7.3 (discontinuous ethanol); 30.0 ± 8.1 (chronic ethanol). Rats were under the liquid diet without ethanol since 5 days before the study to allow the animals to become accustomed to the new diet.

The care and use as well as all procedures involving animals were approved by the Institutional Animal Care Committee, Faculty of Medicine, Complutense University, Madrid. The study was in accordance with the guidelines of the Institutional Care and Use Committee of the National Institute on Drug Abuse, National Institutes of Health and the Guide for the Care and Use of Laboratory Animals (Institute of Laboratory Animal Resources, 1996)

After 4 weeks of treatment groups of 8 rats were killed by decapitation at six different time intervals, every 4 h, throughout a 24-h cycle starting at 0900 h. At night intervals animals were killed under red dim light. Trunk blood was collected and plasma samples were obtained by centrifugation of blood at 1,500 g for 15 min and were stored
at -20 °C until further analysis. The submaxillary lymph nodes and spleen were removed aseptically, weighed and placed in Petri disks containing balanced salt solution, the cells being gently teased apart. After removing the clumps by centrifugation, the cells were suspended in sterile supplemented medium (RPMI 1640), containing 10% heat-inactivated, fetal bovine serum, 20 mM L-glutamine, 0.02 mM 2-mercaptoethanol and gentamicin (50 mg/ml), and were counted.

**Mitogen assays**

Mitogen assays were performed as described in detail elsewhere (Esquivino et al., 1996). Thymidine [methyl-\(^3\)H] (specific activity 20 Ci/mmol) was purchased from NEN Research Products, Boston, MA, USA. Lymph nodes or splenic cells were used at a final number of cells/well (0.1 ml) of 5 x 10^5. Control and experimental cultures were run in triplicate. Mitogens were added to the cultures at final supramaximal concentrations of 5 μg/ml. The cultures were incubated in a humidified 37°C incubator in an atmosphere of 5% CO₂. After 48 h incubation, \(^3\)H-thymidine (0.2 μCi) was added to each well in a volume of 0.02 ml. Cells were harvested 5 h later using an automated sample harvester, and the filters were counted in a liquid scintillation spectrometer. The proliferation index was estimated as the ratio between stimulation in the presence of mitogens/controls. Results were expressed as proliferation index/number of cells.

**Lymphocyte subsets**

The relative size distributions of lymph cells in the lymph nodes and the spleen were determined by FACS analysis, as previously described (Castrillon et al., 2000). For these studies, we used the following monoclonal antibodies: Anti-rat LCA (OX-33) for B
lymphocytes (Serotec, Oxford, UK), Anti-rat TCR alpha/beta (R7.3) for T lymphocytes (Serotec, Oxford, UK), Anti-rat CD4 (OX-35) which recognize a rat T helper cell differentiation antigen (Pharmingen, San Diego, CA, USA), and Anti-rat CD8a (OX-8) which recognize the reactive antigen expressed on rat T cytotoxic/suppressor cells (Pharmingen, San Diego, CA, USA). Lymphocytes isolated as indicated above, were washed in cold PBS with 0.02% sodium azide and then incubated (3 x 10^5 cells/tube) with appropriate primary antibodies for 30 min at 4ºC. Following two washes, the cells were incubated with 1 ml of PBS-BSA 1%, during 5 min at 4ºC, washed three times, resuspended in 1% paraformaldehyde in PBS. Fluorescence intensity was analyzed by fluorescence activated cell sorting (FACStarplus; Beckton Dickinson, Mountain View, CA).

**Prolactin assay**

Plasma prolactin levels were measured by a homologous-specific double antibody radioimmunoassay (RIA), using materials kindly supplied by the NIDDK’s National Hormone and Pituitary Program and by Dr. A.Parlow (Harbor UCLA Medical Center, 1000 West Carson Street, Torrance, CA, USA). The intra- and inter-assay coefficients of variation were 6 and 8%, respectively. Sensitivity of the RIA was 50 pg/mL using the NIDDK rat prolactin standard.

**Statistical analysis**

Statistical analysis of results was performed by a factorial analysis of variance (ANOVA). Generally, the analysis included assessment of the group effect (i.e. the occurrence of differences in mean values between discontinuous ethanol, chronic ethanol and control groups), of time-of-day effects (the occurrence of daily changes) and of the
interactions, from which inference about differences in timing and amplitude could be obtained. One-way ANOVA was then applied to show which time points were significantly different within each experimental group to define the existence of peaks. Post-hoc Bonferroni’s multiple comparisons tests were employed. $P$ values lower than 0.05 were considered evidence for statistical significance.

**Results**

Figure 1 depicts the changes in mitogenic responses to Con A and LPS of cells derived from lymph nodes and spleen of discontinuous ethanol-fed rats, chronic ethanol-fed rats and controls along the 24 h span. A factorial ANOVA indicated a significant decrease in splenic cells’ response to Con A, and in lymph node and splenic cells’ response to LPS in the discontinuous ethanol group ($p<0.001$, Bonferroni’s test). Significant interactions “treatment x time of day” were found in a factorial ANOVA for every parameter examined indicating that both modalities of ethanol administration disrupted the 24 h changes observed in control rats ($p<0.001$, Fig. 1).

Figure 2 shows the percentage of CD4$^+$, CD8$^+$ and CD4$^+$-CD8$^+$ cells, and the ratio between CD4$^+$ and CD8$^+$ cells, in the three groups of animals examined. The discontinuous ethanol administration brought about a significant decrease of lymph node and splenic CD8$^+$ and CD4$^+$-CD8$^+$ cells ($p<0.0001$, Bonferroni’s test). Consequently, lymph node and splenic CD4$^+$/CD8$^+$ ratios were higher in rats receiving ethanol in a discontinuous regime ($p<0.001$). After chronic ethanol administration, splenic mean levels of CD8$^+$ and CD4$^+$-CD8$^+$ cells augmented ($p<0.01$ as compared to controls, Bonferroni’s test). Significant
interactions “treatment x time of day” were detected for all the parameters tested and in both ethanol-administered groups, except for lymph node and splenic CD4+ cells (p<0.001, Fig. 2).

Figure 3 shows the changes in lymph node and splenic T and B cells along the 24 h span. Analyzed by a factorial ANOVA, mean values of lymph node and splenic T cells, and of splenic B cells in the discontinuous ethanol group, were significantly higher, and B/T ratios significantly lower, than those of ethanol-chronic and control rats (p<0.001, Bonferroni’s test). Significant interactions “treatment x time of day” were observed in the factorial ANOVA for every parameter tested except for the lymph node T cell population, i.e. both modalities of ethanol administration disrupted the 24 h changes observed in control rats (p<0.001, Fig. 3). As shown in Fig. 4, the total number of cells in lymph nodes and spleen did not differ significantly among groups, nor as a function of time of day.

Plasma prolactin levels in the 3 groups of animals examined are depicted in Fig. 5. Peak circulating levels occurred at the second part of scotophase in the 3 groups of animals. Mean values of prolactin (ng/mL) were 2.21 ± 0.23, 7.97 ± 0.87 and 18.9 ± 1.65 in control rats and rats chronically or discontinuously fed with ethanol, respectively, differences among the 3 groups being significant (p<0.001, factorial ANOVA). When analyzed within individual groups, prolactin levels did not correlate with any of the immune parameters over the time course studied.

An additional group of rats fed regular chow and water ad libitum was compared to rats receiving the control liquid diet as far as the different immune parameters tested.
Neither the mitogenic responses (Fig. 6) nor lymphocyte subset populations (results not shown) differed significantly between both groups.

Discussion

Foregoing results are compatible with the view that the discontinuous drinking of a moderate amount of ethanol can be more harmful for the immune system than a continuous administration of a similar amount of ethanol. Rats receiving a discontinuous ethanol diet exhibited an impaired mitogenic response of splenic cells to Con A, and of lymph node and splenic cells to LPS, as compared to chronic ethanol feeding or to control rats. These findings coexisted with a decreased number of lymph node and splenic CD8+ and CD4+-CD8+ cells and augmented number of lymph node T cells and splenic T and B cells, in the discontinuous ethanol group. Both modalities of ethanol administration disrupted the 24 h changes in most immune parameters examined.

Clinical and experimental evidence demonstrate that acute, moderate and chronic, excessive ethanol use result in various abnormalities in the functions of the immune system [see for ref. (Szabo and Mandrekar, 2009)]. This effect is observed both on innate as well as on adaptive immunity. Ethanol feeding of rodents is associated with suppression of T-cell proliferation and B-cell antibody production to T-cell-dependent antigens, decreased natural killer cell activity, and increased susceptibility to incidence of infection (Goral et al., 2008). Other changes in the immune system include loss of lymphoid cells from spleen, thymus, and mesenteric lymph nodes; impaired lymphocyte proliferation in response to various stimuli and diminished leukocyte mobilization and phagocytosis.
Most of these studies were performed at single time points in the 24 cycle (usually in the morning). Therefore, the effect of ethanol on the circadian organization of the immune response remaining largely undefined. This can be of particular importance in view of the significant effect of chronic ethanol administration on various circadian rhythms (sleep, motor activity and food intake) (Rosenwasser, 2001). In addition, circadian rhythms modulate body’s response to ethanol (Chen et al., 2004; Arjona et al., 2004; Arjona and Sarkar, 2005,2006; Spanagel et al., 2005a,b; Zghoul et al., 2007). We previously reported that chronic ethanol feeding resembling that employed herein in the ethanol-chronic group brought about significant modifications in the 24 h pattern of several immune parameters in the thymus and spleen of peripubertal rats (Jiménez et al., 2005). In the present study both ways of ethanol administration brought about circadian disruption of most immune parameters examined.

Binge drinking models were achieved in rodents by repeated alcohol administration for 3 to 4 consecutive days using gastric gavage (Nagy, 2008). The amounts of ethanol administered by this procedure are very high; in contrast, when complete liquid diets containing adjusted nutritional components and ethanol with approximately 36% of
calories from ethanol are used (Lieber and DeCarli, 1994), the amounts of ethanol administered are moderate and consequently the immunological sequels are more subtle. Since a moderate discontinuous way of drinking is probably the most frequent way of ethanol drinking in adolescent and young adults, we employed a liquid ethanol diet to examine its effect on the 24-h organization of the immune response. Our results demonstrate that under a discontinuous ethanol regime the mitogenic response to Con A or LPS is heavily impaired, concomitantly with an important decrease in lymph node and splenic CD8+ and CD4+-CD8+ cell population. We previously reported that a chronic ethanol feeding resembling that employed herein the mitogenic response of splenic cells to mitogens remained unaffected, while mean values of splenic CD8+ and CD4+-CD8+ number augmented (Jiménez et al., 2005). The present data in the ethanol-chronic group as far as splenic immune parameters were essentially similar to those reported earlier.

Although no mechanistic conclusions can be derived from the present observational study, some considerations on this aspect deserve comment. In the present experiments, we employed the Lieber–DeCarli liquid diet in a discontinuous way to minimize the stress caused by the intragastric administration of a high amount of ethanol in the binge drinking models. However, the possibility that the periodic ethanol feeding and withdrawal could be stressors was supported by the observation that mean plasma prolactin levels increased by 3.6-fold in rats chronically fed with alcohol and by 8.5-fold in rats under discontinuous ethanol administration. Indeed, prolactin secretion is increased by stress (Neill, 1970; Gala, 1990; Franci et al., 1992), e.g, ether stress in male rats (Deis et al., 1989). A role for the stress response has been proposed in cell loss and changes in
immune cell functions in ethanol-fed rodents (Jerrells et al., 1990). Since adrenalectomy in rats partially abolishes the effects of ethanol on thymic and splenic atrophy and suppression of lymphocyte proliferation of T-cell mitogens, corticosteroids have been implicated in this phenomenon (Padgett et al., 2000).

During the diurnal cycle plasma corticosterone starts to increase at approximately light/dark transition and attains maximal levels during the first part of scotophase (Haus, 2007; Claustrat et al., 2008). In the case of prolactin an increase during the scotophase has also been described in several studies (Haus, 2007; Claustrat et al., 2008), including the present one. In spite of the apparently similar sensitivity of the two endocrine systems to stress, no consistent correlations were found between the activation of hypothalamic-pituitary-adrenal axis and prolactin secretory responses (Courvoisier et al., 1996; Márquez et al., 2002). This lack of consistent correlations suggests that stress-related factors underlying activation of the two endocrine systems are different. Further studies are needed to determine whether the changes reported herein in the extent of prolactin release are associated with different levels of adrenal stimulation.

Acknowledgements

This work was supported by grants from Ministerio de Interior, Plan Nacional sobre Drogas, Spain (PR201/02-11474), Programa de Creación y Consolidación de Grupos de Investigación, Universidad Complutense de Madrid (GR58/08), and Agencia Nacional de Promoción Científica y Tecnológica, Argentina (PICT 2007-01045). DPC is a Research
Career Awardee from the Argentine Research Council and Emeritus Professor, University of Buenos Aires.
References


Figure 1.
Effect of a continuous or discontinuous drinking of ethanol on mitogenic responses in cells derived from submaxillary lymph nodes and spleen of peripubertal rats. Groups of 8 rats subjected to a discontinuous ethanol liquid diet, a chronic ethanol liquid diet or to a control liquid diet as described in Methods were killed by decapitation at 6 different time intervals throughout a 24 h cycle. Bar indicates scotophase duration. Shown are the means ± SEM. Letters indicate the existence of significant differences between time points within each group after a Bonferroni’s multiple comparisons test, as follows: a p< 0.01 vs. all time points; b p< 0.02 vs. 0900, 2100, 0100 and 0500 h; c p< 0.02 vs. 0500 h; d p< 0.02 vs. 1300 and 2100 h; e p< 0.01 vs. 1700, 2100 and 0500 h; f p< 0.01 vs. 0900 and 1700 h; g p< 0.02 vs. 1300 h. For further statistical analysis, see text.
Figure 2.
Effect of a continuous or discontinuous drinking of ethanol on 24-h changes of CD4\(^+\), CD8\(^+\) and CD4\(^+\)-CD8\(^+\) cells, and CD4\(^+\)/CD8\(^+\) ratio, in submaxillary lymph nodes and spleen of peripubertal rats. For experimental details see legend to Fig. 1. Shown are the means ± SEM. Letters indicate the existence of significant differences between time points within each group after a Bonferroni’s multiple comparisons test, as follows: \(^a\) p < 0.01 vs. 0100, 1300 and 2100 h; \(^b\) p < 0.03 vs. 1300, 2100, 0100 and 0500 h; \(^c\) p < 0.02 vs. 2100 h; \(^d\) p < 0.01 vs. all time points; \(^e\) p < 0.01 vs. 0900, 1300, 1700 and 0500 h; \(^f\) p < 0.01 vs. 0900, 1300 and 1700 h; \(^g\) p < 0.01 vs. 0900, 2100, 0100 and 0500 h; \(^h\) p < 0.01 vs. 2100, 0100 and 0500 h; \(^i\) p < 0.02 vs. 0100 and 1300 h; \(^j\) p < 0.001, 0900 and 2100 h. For further statistical analysis, see text.
**Figure 3.**
Effect of a continuous or discontinuous drinking of ethanol on 24-h changes of T and B lymphocytes, and T/B ratio, in submaxillary lymph nodes and spleen of peripubertal rats. For experimental details see legend to Fig. 1. Shown are the means ± SEM. Letters indicate the existence of significant differences between time points within each group after a Bonferroni’s multiple comparisons test, as follows: a $p < 0.02$ vs. 1300 h; b $p < 0.01$ vs. 1700 and 0100 h; c $p < 0.02$ vs. 0900, 0100 and 0500 h; d $p < 0.01$ vs. all time points; e $p < 0.01$ vs. 0900, 1300, 1700 and 0500 h; f $p < 0.01$ vs. 1300, 2100 and 0100 h; g $p < 0.01$ vs. 0900, 1700, 2100 and 0500 h. For further statistical analysis, see text.
Figure 4.
Effect of a continuous or discontinuous drinking of ethanol on 24-h changes of total number of viable cells recovered from submaxillary lymph nodes and spleen of peripubertal rats. For experimental details see legend to Fig. 1. Shown are the means ± SEM. In a factorial ANOVA there were no significant differences among groups, nor as a function of time of day.
Figure 5.
Effect of a continuous or discontinuous drinking of ethanol on plasma prolactin levels in peripubertal rats. For experimental details see legend to Fig. 1. Shown are the means ± SEM. Letters indicate the existence of significant differences between time points within each group after a Bonferroni’s multiple comparisons test, as follows: \(^a\) p< 0.01 vs. 0900, 1300 and 1700 h; \(^b\) p< 0.01 vs. 0900, 2100 and 0500 h; \(^c\) p< 0.01 vs. 0900, 1700 and 0100 h; \(^d\) p< 0.05 vs. 0900 h.
Figure 6.
Mitogenic responses in cells derived from submaxillary lymph nodes and spleen of peripubertal rats receiving regular rat chow and water *ad libitum* as compared to control liquid diet animals. Groups of 8 rats fed with regular rat chow and water were killed by decapitation at 6 different time intervals throughout a 24 h cycle. Control liquid diet data are from Fig. 1. Bar indicates scotophase duration. Shown are the means ± SEM. Letters indicate the existence of significant differences between time points within each group after a Bonferroni’s multiple comparisons test, as follows: a \( p < 0.01 \) vs. all time points; b \( p < 0.02 \) vs. 0900 and 0500 h; c \( p < 0.01 \) vs. 0900 and 1700 h; d \( p < 0.02 \) vs. 0900 and 2100 h. Differences between groups were not significant when analyzed as main factors in a factorial ANOVA.