An Initial, Three-Day-Long Treatment with Alcohol Induces a Long-Lasting Phenomenon of Selective Tolerance in the Activity of the Rat Hypothalamic–Pituitary–Adrenal Axis

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We determined whether an initial alcohol challenge induced long-lasting changes in the activity of the hypothalamic–pituitary–adrenal (HPA) axis. Adult male rats received intragastric injections of the vehicle or a moderately intoxicating dose of alcohol (3.0 gm/kg) daily for 3 d. When animals were acutely challenged with alcohol 3–12 d later, their ACTH and corticosterone responses were significantly blunted, compared with that of vehicle-pretreated rats. In contrast, exposure to mild electric foot shocks induced a pattern of ACTH secretion that was comparable in animals administered alcohol or the vehicle previously, indicating a lack of cross-tolerance. No significant differences were observed in pituitary responsiveness to corticotropin-releasing factor or vasopressin in rats pretreated with the vehicle or alcohol. The influence of the initial drug treatment was not mimicked by exposure to foot shocks, nor was it prevented by administering a potent corticotropin-releasing factor antagonist to block the elevations in plasma ACTH and corticosterone induced by this initial treatment. Finally, we found that rats injected initially with the vehicle and challenged subsequently with alcohol exhibited the expected increased neuronal activation (measured by the upregulation of steady-state mRNA and protein levels of immediate early genes) in the paraventricular nucleus of their hypothalamus. In contrast, this response was markedly decreased in animals exposed previously to the drug.

To our knowledge, this is the first report that exposure to a stress (i.e., alcohol), although not immediately altering the response of the HPA axis to this particular signal, induces a selective tolerance that is both slow to develop and long-lasting. The primary mechanism mediating the ability of an initial drug treatment to decrease subsequent responses of the HPA axis to a second drug challenge seems to be the inability of hypothalamic neurons to respond adequately to this second challenge.

Key words: alcohol; ACTH; corticosterone; c-fos; NGFI-B; PVN; rat

It is well known that an initial exposure of the hypothalamic–pituitary–adrenal (HPA) axis to a stress can alter the response of this axis to the same, as well as other, noxious signals. Depending on the nature, intensity, and duration of the initial stimulus, tolerance (habituation) or sensitization develops. Tolerance refers to the situation in which repeated exposure to the same stimulus elicits a diminishing effect on ACTH secretion; sensitization, on the other hand, is observed when repeated treatments elicit a stronger effect on ACTH release than was initially observed.

Hormones of the HPA axis influence many bodily functions. Glucocorticoids (GC), for example, promote the availability of energy substrates (Dallman et al., 1993) and modulate vascular tone (see Munck and Guyre, 1986). After antigenic challenges and once appropriate defense mechanisms have become activated, GC serve to restrain the activity of the immune system and thus prevent it from overshooting (Munck and Guyre, 1986). Corticotropin-releasing factor (CRF), on the other hand, regulates behavior (Koob, 1990) and sympathetic activity (Brown and Fisher, 1990), as well as appetite and gastrointestinal functions (Taché et al., 1993), immune responses, and reproductive and cardiovascular parameters (see Munck et al., 1984; McEwen, 1992; Rivest and Rivier, 1995). Alcohol administration represents a powerful stimulant of the HPA axis, resulting in increased circulating levels of ACTH and GC, as well as activation of the hypothalamic neurons containing the two peptides that are most important for ACTH release, namely, CRF and vasopressin (VP) (Vale et al., 1981; Rivier et al., 1982; Antoni, 1993). It is therefore obvious that this drug, particularly if it is administered repeatedly, will threaten homeostasis via a variety of mechanisms. In addition, an emerging hypothesis is that hormones of the HPA axis can promote drug use (Cador et al., 1992; Deroche et al., 1993; Piazza et al., 1993; Fahlke et al., 1995; Lamblin and De Witte, 1996; Shaham et al., 1996). Consequently, sensitization to alcohol may reinforce its consumption, whereas, conversely, animals rendered tolerant may attempt to restore normal concentrations of ACTH, corticosterone, and CRF by consuming ever-increasing doses of the drug.

We (Rivier et al., 1984; Lee and Rivier, 1993; Rivier, 1996) and others (Spencer and McEwen, 1990) have reported previously that after an initial rise in plasma ACTH and corticosterone levels, continuous exposure to alcohol resulted in a blunting of these responses to the drug itself as well as to other stimuli. What we do not know is whether an initial treatment with alcohol induces long-lasting changes in the activity of the HPA axis that persist after exposure to the drug has ceased. We therefore tested the hypothesis that a first exposure to alcohol would also alter the response of the HPA axis to subsequent challenges. Unexpect-

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edly, we observed a significant blunting of the ACTH and corticosterone responses to a second alcohol challenge, but not to other stimuli, administered 3–12 d after the initial drug treatment. We then determined whether this phenomenon was because of reduced pituitary responsiveness to trophic factors, whether the initial alcohol-induced rise in plasma ACTH and corticosterone levels participated in this altered response, and whether a blunted neuronal response of hypothalamic neurons responsible for ACTH release might account for the decreased effectiveness of the second alcohol challenge.

Many investigators have reported an adaptation of the HPA axis to chronic stress (Dallman, 1993). Most studies have focused on animals that were exposed to a particular stressor immediately after a period of repeated or continuous exposure to this same or another signal (Kant et al., 1985; Akana et al., 1992; Ottenweller et al., 1992; vanRaaij et al., 1997), including work showing adaptation of the HPA axis of rats chronically exposed to alcohol and to the drug itself but not to other stresses (Spencer and McEwen, 1990). To our knowledge, the only report of a long-lasting change to the drug itself but not to other stresses is that of Kant et al. (1985) who showed that the pituitary response to repeated exposure to alcohol was blunted. We then determined whether this phenomenon was because of a change in the sensitivity of the pituitary gland to the ACTH liberating factor.

The experiment was performed 7 d after the end of the initial intragastric treatment. Initially alcohol-induced rises in plasma ACTH and corticosterone were significant (p < 0.05) at all times indicated. However, alcohol-induced changes in hormone levels were significant (p < 0.01) at all times indicated.

### Table 1. Weights

<table>
<thead>
<tr>
<th>Initial treatment</th>
<th>At arrival</th>
<th>At end of initial intragastric treatment</th>
<th>At time of second challenge*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vehicle</td>
<td>181 ± 7</td>
<td>240 ± 22</td>
<td>285 ± 20</td>
</tr>
<tr>
<td>Alcohol</td>
<td>184 ± 9</td>
<td>238 ± 19</td>
<td>282 ± 21</td>
</tr>
</tbody>
</table>

n = 5–7 rats per group.

*The experiment was performed 7 d after the end of the initial intragastric treatment.

### Materials and Methods

#### Animals

Adult Sprague Dawley male rats were maintained in groups (three to four rats per cage) on a 12 hr light/dark cycle (lights on at 6:00 A.M.). Food and water were available ad libitum.

#### Cannulae

Under halothane anesthesia, indwelling intragastric cannulae were inserted at least 1 week before experimentation. Preliminary experiments indicated that it was not necessary to fast the animals before surgery (Ogilvie et al., 1997a). Cannulae were constructed of polyethylene tubing (PE60). The end to be inserted into the stomach was expanded twice to make two bubbled portions, positioned ~5 mm apart. To place the intragastric cannula, we cut the abdominal wall on the midline and pulled the stomach through this opening. A purse string suture was placed in the nonglandular fundus, after which a hole was opened in the middle of the suture using a pair of spreader forceps. The cannula was passed into the stomach between the two bubbled portions of PE tubing to retain the tip in the stomach lumen. With the aid of a trocar, we passed the free end of the cannula through the body wall and under the skin so that it exited at the nape of the neck, where it was capped. After intragastric cannulation, rats were housed individually to prevent chewing of the exteriorized cannula. In experiments requiring blood sampling, intravenous cannulae were inserted 48 hr earlier (Rivier, 1993).

#### Treatments

Alcohol. For the daily alcohol injections, the intragastric cannulae were extended with a PE50 tubing connected to a syringe. Alcohol was diluted with saline to <20% v/v. An equal volume of saline was administered to control rats. During treatment, the rats were awake and freely moving in their home cage. Because the animals were not fasted, alcohol was injected 4–5 hr after lights were turned on, a time when most of the food consumed during the previous night had left the stomach. This treatment schedule also ensured that blood alcohol levels (BALs) had returned to baseline before the animals started feeding again, and indeed alcohol-treated rats maintained normal weight gains (Table 1). Gross examination of the stomach lining failed to indicate any tissue damage, and

![Figure 1](image-url). Effects of the intragastric (ig) injection of the vehicle (○) or alcohol (●, 3.0 gm/kg), administered daily for 3 d, on plasma ACTH and corticosterone levels over a 60 min time course. Each point represents the mean ± SEM of five to six intact male rats. For the sake of clarity, no statistical levels of significance are indicated on the figure. However, alcohol-induced changes in hormone levels were significant (p < 0.01) at all times indicated.
indeed alcohol has been administered through gavage over prolonged periods of time by other investigators (e.g., see Maier et al., 1995). On the day of an experiment, animals were removed to a soundproof room and housed in opaque buckets with cannulae connected such that animals could be injected and bled without being handled. Rats were left undisturbed for 3 hr so that hormone levels would return to basal levels by the time of injection. All experiments began between 11:00 and 12:00 A.M.

**Peptides.** The potent CRF antagonist astressin (Gulyas et al., 1995) was administered in some experiments to determine whether blockade of the ACTH and corticosterone responses to the initial alcohol exposure would interfere with the long-term influence of the drug. In other experiments, pituitary responsiveness was determined by administering CRF or VP intravenously at concentrations chosen to provide dose-related increases in plasma ACTH levels (Lee and Rivier, 1995). Peptides were synthesized by solid-phase methodology and generously provided by Dr. Jean Rivier (The Salk Institute, La Jolla, CA). They were diluted in 0.04 M phosphate buffer, pH 7.4, containing 0.1% bovine serum albumin and 0.01% ascorbic acid. This vehicle was injected in control rats in the corresponding experiments.

**Mild electric foot shocks.** These shocks (1 mA; 1 sec in duration; 2 shocks/min for 30 min) were delivered to the paws of the rats as described previously (Rivier and Vale, 1988).

All protocols were approved by the Salk Institute Institutional Animal Care and Use Committee.

**Blood alcohol levels (BALs)**

Samples for the measurement of alcohol in whole blood (100 μl) were immediately diluted in 6.25% trichloroacetic acid (900 μl) in plastic vials with screw tops (Sarstedt, Nümbrecht, Germany). Samples were stored at 4°C until assayed. BALs were measured using a kit purchased from Sigma (St. Louis, MO; 333A), optimized for use with small samples. The coefficient of variation of this method, determined with a serum pool provided by the manufacturer, never exceeded 8%.

**cRNA probe synthesis and preparation**

The pBluescript SK-1 vector (Stratagene, La Jolla, CA) containing rat NGFI-B cDNA (provided by Dr. J. Milbrandt, Washington University, St. Louis, MO) or c-fos (provided by Dr. I. Verma, The Salk Institute, La Jolla, CA) was linearized with BamHI or SmaI, respectively. Radioactive cRNA copies were synthesized by incubation of 250 ng of linearized plasmid in 6 mM MgCl₂; 36 mM Tris, pH 7.5; 2 mM spermidine; 8 mM dithiothreitol; 25 mM ATP, GTP, CTP, and α-35S-UTP; 1 U of RNAse (Promega, Madison, WI); and 10 U of T₃ (for NGFI-B) or T₇ (for c-fos)
for 60 min at 37°C. Unincorporated nucleotides were removed using Quick-Spin columns (Boehringer Mannheim, Indianapolis, IN). A sense probe was used as a control for nonspecific signal in some adjacent sections for in situ hybridization.

**In situ hybridization histochemistry**

Four to six rats per group were deeply anesthetized with 35% chloral hydrate, a drug that does not increase immediate early gene RNA levels or ACTH concentrations. The animals were then perfused transcardially with saline followed by 4% paraformaldehyde and 0.1 M borate buffer, pH 9.5. The brains were removed, post-fixed in 4% paraformaldehyde for 4–5 d, and then placed overnight in 10% sucrose, 4% paraformaldehyde, and 0.1 M borate buffer. They were cut into 30 μm coronal slices obtained at 120 μm intervals throughout the hypothalamus and stored at −20°C in a cryoprotectant solution (50% 0.1 M PBS, 30% ethylene glycol, and 20% glycerol) until histochemical analysis.

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Hybridization histochemical localization of each transcript was performed using 35S-labeled cRNA probes. Protocols for riboprobe synthesis, hybridization, and autoradiographic localization of mRNA signals were adapted from Simmons et al. (1989).Brains from the same experiments were always processed and analyzed in the same hybridization experiment. All solutions were treated with diethylpyrocarbonate (Dep. C) and sterilized to prevent RNA degradation. Sections mounted onto gelatin- and poly-L-lysine-coated slides were desiccated under vacuum overnight, fixed in 4% paraformaldehyde for 30 min, and digested by proteinase K (10 μg/ml in 50 mM Tris HCl, pH 7.5, and 5 mM EDTA at 37°C for 25 min). Thereafter, brain sections were rinsed in sterile Dep. C water followed by a solution of 0.1 M triethanolamine (TEA), pH 8.0, acetylated in 0.25% acetic anhydride in 0.1 M TEA, and dehydrated via graded concentrations of alcohol (50, 70, 95, and 100%). After vacuum drying for a minimum of 2 hr, 90 μl of hybridization mixture (10^7 cpm/ml) was spotted on each slide, sealed under a coverslip, and incubated at 60°C overnight in a slide warmer. Coverslips were then removed, and the slides were rinsed in 4 × SSC at room temperature. Sections were digested by RNase A (20 μg/ml; 37°C; 30 min), rinsed in descending concentrations of SSC (2×, 1×, and 0.5×), washed in 0.1× SSC for 30 min at 65°C (1× SSC, 0.15 M NaCl, and 15 mM trisodium citrate buffer, pH 7.0), and dehydrated via graded concentrations of alcohol. After being dried under the vacuum, sections were exposed at 4°C to x-ray film (Eastman Kodak, Rochester, NY) for 15 hr, defatted in xylene, and dipped in NTB2 nuclear emulsion (Kodak; diluted 1:1 with distilled water). Slides were exposed for 6 d, developed in D19 developer (Kodak) for 3.5 min at 15°C and fixed in rapid fixer (Kodak) for 6 min. Thereafter, tissues were rinsed in running distilled water, counterstained with thionin (0.25%), dehydrated via graded concentrations of alcohol, cleared in xylene, and coverslipped with a mixture of distrene, tricresyl phosphate, and xylene (DPX).

**Quantitative analysis of in situ hybridization results**

Semi-quantitative densitometric analyses of hybridization signals for RNAs of interest were performed on nuclear emulsion-dipped slides. Brain paste standards containing serial dilutions of 35S-UTP, used for quantification of mRNA signals, were prepared concurrently to ensure that optical density was found within the linear range of the standard curve. In addition, analyses with emulsion-coated slides were performed with two to three different exposure times to confirm that signals were not saturated. Densitometric analyses of autoradiographic signals were...
done over the confines of cells within the paraventricular nucleus (PVN), using a Leitz optical system coupled to a Macintosh II computer and Image software (version 1.60; W. Rasband, National Institutes of Health). For statistical analysis, the paraventricular (pPVN) and magnocellular divisions of the PVN (mPVN) were first delineated in each section under dark-field illumination at the rostral and caudal level, and optical density was measured under dark-field illumination as described previously (Ogilvie et al., 1997a,b). Although it is possible to exclude magnocellular neurons scattered in the pPVN region, we did not use this method. It thus remains possible that counts throughout the medial pPVN included some magnocellular neurons. Gray level measurements (optical density) were taken under dark-field illumination of hybridized sections over the medial pPVN, as defined by redirected sampling from the corresponding Nissl-stained sections under bright-field images. Data were expressed in gray scale values of 1–256. All gray level measurements were corrected for background. Signals were measured in both sides of the brain, and mean values for all animals (four to six per group) were determined for each rat in three to four sections throughout the PVN.

**Immunohistochemistry**

Series of brain sections were washed in 0.05 M KPBS and incubated 10 min in 0.3% H₂O₂. Sections were then incubated at 4°C for 48 hr with primary antiserum applied at a 1:20,000 dilution. Fos immunoreactivity was localized using polyclonal antisera raised in rabbits against a synthetic N-terminal fragment (residues 4–17) of human c-fos (Oncogene Sciences). Next, the sections were incubated with a biotinylated anti-rabbit IgG (1:1500 dilution; Vector Laboratories, Burlingame, CA) and subsequently incubated at room temperature with a conventional nickel-enhanced avidin–biotin immunoperoxidase complex (Vetastain ABC Elite reagents). After the reaction, the sections were mounted onto gelatin–chrome alum-coated slides and coverslipped with DPX.

**RIAs**

Plasma ACTH levels were measured in duplicate using a two-site immunoradiometric assay (Allegro kit; Nichols Institute, San Juan Capistrano, CA), which we have validated for rat ACTH (Rivier and Shen, 1994). Total corticosterone was measured by double-antibody RIA. The primary antibody was anti-corticosterone-3-BSA (377; G. Niswender, Fort Collins, CO) diluted to give a final concentration of 1:30,000. The sensitivity and coefficient of variation of this assay were 4 ng/ml (0.022 ng/tube) and <15%, respectively.

**Statistical analysis**

Results were analyzed by one- or two-way ANOVA. Student–Newman–Keuls tests were then used to determine differences between treatments. Levels of NGFI-B or c-fos mRNA in the pPVN and mPVN were subjected to a two-way ANOVA, with time after injection of alcohol and pretreatment as the variables. When differences were indicated by the ANOVA procedure, the least squares means test was used for post hoc analysis.

**RESULTS**

**Effect of an initial exposure to alcohol on the hormonal response to a second alcohol challenge delivered 3–7 d later**

The first set of experiments was designed to investigate the influence of single or repeated alcohol injections on the ACTH and corticosterone responses to a second alcohol challenge. The vehicle or alcohol (3.0 gm/kg) was injected intragastrically either once or daily for 3 consecutive days. The ACTH and corticosterone responses were comparable whether alcohol was administered once or daily for 3 d (Fig. 1). Three to seven days after either the single injection or the last of the three injections, the animals received an acute intragastric injection of the vehicle or alcohol (3.0 gm/kg), yielding four experimental groups: vehicle/vehicle; vehicle/alcohol; alcohol/vehicle; and alcohol/alcohol. Serial blood samples were obtained over a 120 min period after the second (acute) treatment. Plasma ACTH and corticosterone levels of the vehicle/vehicle and alcohol/vehicle groups remained at baseline concentrations (data not shown). A single initial alcohol injection did not significantly alter the course of ACTH and corticosterone release induced by the second drug challenge administered 3–7 d later (data not shown). In contrast, rats initially injected with alcohol daily for 3 d showed significantly (p < 0.01) blunted neuroendocrine responses when rechallenged with the drug 3–7 d later (Fig. 2). Interestingly, the patterns of ACTH and corticosterone secretion after the third daily alcohol injection were comparable in vehicle- and alcohol-pretreated rats, indicating that tolerance had not yet developed. There was no obvious difference between the magnitude of this blunting at the various test times (i.e., 3 or 7 d after the initial alcohol treatment). BALs were statistically comparable after all alcohol injections (Fig. 3).

In a second set of experiments, we determined whether a smaller dose of alcohol [1.5 gm/kg, intragastrically (i.g.), daily for 3 d] would also blunt the response of the HPA axis to the second
alcohol challenge (using our standard dose of 3.0 gm/kg) and found that such was not the case (data not shown).

**Effect of an initial exposure to alcohol on the hormonal response to mild electric foot shocks delivered 7 d later**

Rats were injected with the vehicle or alcohol (3.0 gm/kg) daily for 3 consecutive days and submitted to either an alcohol challenge (3.0 gm/kg) or mild electric foot shocks (30 min) 7 d later. The initial alcohol treatment induced the expected blunted ACTH response after rechallenge with the drug but did not significantly alter the response to the shocks (Fig. 4). It should be mentioned that although in the experiment illustrated here the ACTH response to shocks was higher than that to alcohol, a subsequent experiment, conducted with shocks of lower intensity, indicated a similar lack of cross-tolerance to foot shocks in alcohol-pretreated rats. These results suggest that there was no ceiling effect that might have masked the development of tolerance to the shocks.

**Effect of an initial exposure to mild electric foot shocks on the hormonal response to an alcohol challenge delivered 12 d later**

This experiment was performed to determine whether previous exposure to foot shocks would mimic the effect of the initial

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![Figure 7](image-url)

*Figure 7.* Effects of three consecutive daily intragastric injections of the vehicle or alcohol (3.0 gm/kg) on steady-state mRNA levels of the immediate early gene NGFI-B in the PVN of rats injected with alcohol (3.0 gm/kg) 7 d later. Dark-field photomicrographs illustrating NGFI-B mRNA levels in the PVN of rats injected with alcohol 30, 60, or 120 min earlier are shown. 3v, Third ventricle. Magnification, 340×.*
alcohol treatments. The animals were either injected with the vehicle or alcohol (3.0 gm/kg, i.g.) as described for Figures 1 and 2 or exposed to mild electric foot shocks (30 min) daily for 3 d. The 30 min shock paradigm we chose induces a pattern of ACTH levels that resembles the hormone profile induced by the intragastric injection of EtOH at 3.0 gm/kg (C. Rivier, unpublished observations). Five, 7, 9, or 12 d after the last drug treatment or shock session, the animals were injected with the vehicle or alcohol (3.0 gm/kg, i.g.). Animals originally administered alcohol and later rechallenged with the drug showed the expected blunting of their ACTH release (Fig. 5). Because results were similar at the 5–12 d time points, Figure 5 only shows data obtained on the 12th day. In contrast, rats originally exposed to shocks and subsequently injected with alcohol exhibited a pattern of ACTH release that was comparable with that of control animals.

Changes in pituitary responsiveness to CRF or VP in rats exposed previously to alcohol

The observation that an initial alcohol challenge did not significantly alter the subsequent ACTH response to electroshocks suggested that pituitary responsiveness to CRF was probably not significantly diminished. Nevertheless, a dose–response curve to CRF was generated in both vehicle-treated rats and rats pretreated with alcohol 7 d earlier. It indicated that, as expected, the ACTH response to the intravenous injection of CRF at 0.2, 1, or 5 μg/kg, measured 10 and 30 min after peptide administration, was statistically comparable in both experiment groups (data not shown). We had shown previously that rats exposed to an alcohol diet for 7–10 d displayed a significantly decreased ACTH secretory pattern when injected with VP at the end of the diet treatment (Lee and Rivier, 1995). We therefore also investigated the possibility that pituitary responsiveness to VP might similarly be decreased in the present paradigm. However, there was no significant difference (p > 0.05) in baseline ACTH levels in rats pretreated with the vehicle (6.23 ± 0.82 pg/ml ACTH) or alcohol at 3.0 gm/kg 7 d earlier (6.30 ± 0.74 pg/ml ACTH) or in the cumulative ACTH responses to the injection of VP at 0.5, 1.5, or 4.5 μg/kg, measured at the 10 and 20 min time points (vehicle-pretreated, 190 ± 21, 499 ± 45, and 1130 ± 120 pg/ml ACTH, respectively; alcohol-pretreated, 211 ± 22, 543 ± 49, and 1079 ± 115 pg/ml ACTH, respectively).
Effects of preventing the ACTH and corticosterone response to the initial alcohol treatment on hormonal responses to the second alcohol challenge

In this experiment, we investigated the contribution of increases in plasma ACTH and corticosterone release induced by the initial alcohol challenge on the subsequent blunting of the HPA axis of rats rechallenged with the drug. A first series of preliminary experiments was performed to determine whether the potent CRF antagonist astressin would be able to prevent the ACTH and corticosterone responses during the initial alcohol treatments. We found that the subcutaneous injection of astressin at 10 mg/kg 60 min before alcohol injection achieved this aim and maintained plasma ACTH and corticosterone levels below 80 pg/ml and 30 ng/ml, respectively. This regimen was therefore used in the experiment illustrated in Figure 6, which compares the ACTH responses to an acute alcohol challenge (3.0 gm/kg) delivered 7 d after a series of three daily alcohol treatments (3.0 gm/kg, i.g.) in rats pretreated with the vehicle or astressin. Vehicle pretreatment, whether or not preceded by astressin, led to the expected rise in plasma ACTH levels when the animals were acutely challenged with alcohol (3.0 gm/kg) 7 d later. Alcohol pretreatment induced the expected blunting of the ACTH response to a second drug challenge, an effect that was not significantly reversed or altered by astressin at the time of the initial alcohol injections. These results indicate that blunting the ACTH and corticosterone release during the initial alcohol challenge did not prevent the development of neuroendocrine tolerance that characterizes our model.

Changes in the appearance of immediate early genes, taken as an index of neuronal activation, during the second alcohol challenge

The protocol was similar to that used for Figures 1 and 2, and rats were pretreated with the vehicle or alcohol (3.0 gm/kg) 7 d before a vehicle or alcohol rechallenge. The appearance of the immediate early genes c-fos and NGFI-B was taken as an index of hypothalamic neuronal activation in response to the second challenges. A 30, 60, 120, and 180 min time course was chosen because it allowed us to detect potential changes both in peak hypothalamic responses as well as in the pattern of these responses. NGFI-B mRNA signals were not detected in the PVN after vehicle injection (data not shown). On the other hand, a significant upregulation of NGFI-B transcripts was observed after alcohol treatment, but this response was markedly blunted in alcohol-pretreated rats (Figs. 7, 8). Similar results were obtained with c-fos mRNA levels in the PVN of the hypothalamus (Fig. 9). Because of the possibility that alcohol might differentially alter gene transcription and translation, in particular by influencing mRNA stability, we then decided to confirm results obtained by also measuring protein levels, an approach that for technical
reasons was only possible for c-fos. We observed that although in vehicle-pretreated rats alcohol significantly increased levels of the c-fos protein measured 60 or 180 min later, this change was markedly blunted in animals pretreated with the drug (Fig. 10).

DISCUSSION
We show here that one daily intragastric injection of alcohol for 3 d produced a significant and long-lasting decrease in the ability of the HPA axis to respond to a second alcohol challenge. We further demonstrate that repeated treatment was necessary because rats injected with alcohol only once showed no change in their subsequent ACTH response. Finally, we found that this phenomenon of selective tolerance did not take place immediately, because ACTH release was undiminished on the third day of the three daily alcohol injections. This latter observation suggests that we are not dealing with a phenomenon of behavioral habituation similar to that described in rats repeatedly exposed to foot shocks or forced swimming (Kant et al., 1985).

The blunted ACTH and corticosterone release that we observed was unexpected in view of several findings showing that previous activation of the HPA axis usually results in increased responsiveness to subsequent challenges. For example, repeated exposure to a particular stressor tends to facilitate the ACTH and corticosterone response to a subsequent novel acute stressor (Hennessy, 1991; Marti et al., 1994), a phenomenon that has been attributed to increased CRF expression in the hypothalamus (Makino et al., 1995) and to changes in the sensitivity of the HPA axis to steroid feedback (Akana et al., 1992). Also, a single intraperitoneal injection of IL-1 is reported to prolong the ACTH response to a second challenge, whether it is immune (IL-1 injection) or nonimmune (exposure to foot shocks) (Schmidt et al., 1995). In this model, increased median eminence levels of VP have been held responsible for the changes in HPA axis activity (Dijken et al., 1993). Finally, there is the well known example of the permanent increase in HPA axis function in rats exposed to maternal deprivation during neonatal development (Stanton et al., 1988; Rosenfeld et al., 1991; Suchecki et al., 1993; Viau et al., 1993), a response in part caused by an upregulation of CRF in nerve terminals (Ladd et al., 1996). It therefore seems that, in the present model, alcohol exerts an influence that is unique in its ability to dampen further responses of the HPA axis.

We had observed previously that rats exposed to various noxious stimuli on the last day of a 7 to 10 d alcohol diet released less ACTH and corticosterone than did control animals, but this phenomenon was not caused by decreased pituitary responsiveness to CRF (Lee and Rivier, 1995). The present finding that rats exposed previously to alcohol can adequately release ACTH when challenged with electroshocks several days later suggests that, in these animals also, CRF remained a potent secretagogue of ACTH release. On the other hand, although animals fed the alcohol diet responded to VP with significantly blunted ACTH secretion (Lee and Rivier, 1995), such was not the case in the paradigm discussed here. We therefore hypothesized that in the model of selective tolerance, an initial exposure to alcohol prevented hypothalamic neurons from responding to a second drug challenge. To investigate this possibility, we measured the appearance of the immediate early genes NGFI-B and c-fos in the PVN, which is the site of CRF perikarya (Swanson et al., 1983). We show here that, indeed, a subsequent acute injection of alcohol resulted in the appearance of lower NGFI-B and c-fos mRNA levels in rats previously administered this drug, compared with animals administered the vehicle. Although additional studies are necessary to demonstrate unambiguously whether this change took place in CRF and/or VP cell bodies, the fact that these immediate early genes were primarily found in the pPVN suggests that it probably corresponded to a dampening in the activity of CRF perikarya. Our results therefore indicate that the neuroendocrine hyporesponsiveness we observed may arise from a relatively slow developing, although long-term, effect of the drug on afferent pathways to the PVN and/or on the ability of CRF neurons to mount an adequate response to the second alcohol challenge.

We also considered the potential influence of another mechanism, namely, the effect of increased hormone levels during the initial drug challenge. We show here that when we blocked the ACTH and corticosterone response to the three repeated intragastric alcohol injections with a potent CRF antagonist, we did not alter the ability of the initial course of alcohol treatments to induce a phenomenon of neuroendocrine tolerance to the drug. These results agree with the finding that foot shocks, which also elevate ACTH and corticosterone levels, were not able to duplicate the long-term inhibitory influence of alcohol. They also suggest that widely operative mechanisms controlling PVN responses to stressors, such as those responsible for the ability of the hippocampus to regulate the negative feedback of glucocorticoids (Jacobson and Sapolsky, 1991; Feldman and Weidenfeld, 1993), are probably not of primary importance. It must be noted that alcohol and shocks, although they both stimulate the pituitary–adrenal axis, also exert influences on the brain that are distinct from each other and that may account for their differential influence in our paradigm. One feature that distinguishes these two stresses is that even a short course of alcohol treatment can induce withdrawal (Buck and Harris, 1991), which results in a course of brain activation (Matsumoto et al., 1993) most probably not present in shocked rats. A detailed study of the response of the hypothalamus during and after exposure to alcohol or shocks will be necessary to probe this hypothesis. It is also possible that alcohol pretreatment decreases the brain levels of a neurotransmitter that participates in the ability of the drug, but not of shocks, to release ACTH. One such candidate is serotonin, a stimulator of CRF-dependent pathways (Gartsdine and Cowen, 1990; Fuller, 1992; Pan and Gilbert, 1992; Calogero et al., 1993), which is inhibited by repeated or prolonged drug treatment (Car-michael and Israel, 1975; Woods and Druse, 1996) but which does not seem to be involved in physicoemotional stresses (Harbuz et al., 1993).

To our knowledge, this is the first report that any stimulus, given for a relatively short time, causes a long-lasting selective tolerance with regard to neuroendocrine responses. Alcoholism is a disease that affects between 8 and 10% of the population. If we accept the concept that some of the individuals who abuse alcohol do so in part because of the changes the drug induces in their CNS, it seems reasonable to propose that if such changes cannot be achieved with a given dose of alcohol, these individuals may consume more of the drug in an attempt to regain the wanted changes. A drug exerts its reinforcing effects via a number of distinct actions, including the ability to stimulate brain circuits involved in the reward system and to induce symptoms of withdrawal that can only be averted by resuming drug consumption (Nestler et al., 1993; Altman et al., 1996; Koob, 1996). We suggest that our studies may provide the basis for a testable, neuroendocrine-based hypothesis of vulnerability to alcohol abuse.
REFERENCES


Schmidt ED, Janssen AWJW, Wouterlood FG, Tilders FJH (1995) Interleukin-1-induced long-lasting changes in hypothalamic corticotropin-
Viau V, Sharma S, Plotsky PM, Meaney MJ (1993) Increased plasma ACTH responses to stress in nonhandled compared with handled rats require basal levels of corticosterone and are associated with increased levels of ACTH secretagogues in the median eminence. J Neurosci 13:1097–1105.