Chronic Ethanol and Withdrawal Differentially Modulate Pre- and Postsynaptic Function at Glutamatergic Synapses in Rat Basolateral Amygdala

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Läck AK, Diaz MR, Chappell A, DuBois DW, McCool BA. Chronic ethanol and withdrawal differentially modulate pre- and postsynaptic function at glutamatergic synapses in rat basolateral amygdala. J Neurophysiol 98: 3185–3196, 2007. First published September 26, 2007; doi:10.1152/jn.00189.2007. Withdrawal anxiety is a significant factor contributing to continued alcohol abuse in alcoholics. This anxiety is long-lasting, can manifest well after the overt physical symptoms of withdrawal, and is frequently associated with relapse in recovering alcoholics. The neurobiological mechanisms governing these withdrawal-associated increases in anxiety are currently unknown. The basolateral amygdala (BLA) is a major emotional center in the brain and regulates the expression of both learned fear and anxiety. Neurotransmitter system alterations within this brain region may therefore contribute to withdrawal-associated anxiety. Because evidence suggests that glutamate-gated neurotransmitter receptors are sensitive to acute ethanol exposure, we examined the effect of chronic intermittent ethanol (CIE) and withdrawal (WD) on glutamatergic synaptic transmission in the BLA. We found that slices prepared from CIE and WD animals had significantly increased contributions by synaptic NMDA receptors. In addition, CIE increased the amplitude of AMPA-receptor–mediated spontaneous excitatory postsynaptic currents (sEPSCs), whereas only WD altered the amplitude and kinetics of tetrodotoxin-resistant spontaneous events (mEPSCs). Similarly, the frequency of sEPSCs was increased in both CIE and WD neurons, although only WD increased the frequency of mEPSCs. These data suggest that CIE and WD differentially alter both pre- and postsynaptic properties of BLA glutamatergic synapses. Finally, we show that microinjection of the AMPA-receptor antagonist, CNQX (6-cyano-7-nitroquinoxaline-2,3-dione) into the BLA (Sajdyk and Shekhar 1997; Walker and Davis 1997a). Additionally, ethanol self-administration increases glucose utilization in the BLA (Porrino et al. 1998); moreover, modulation of BLA neuronal activity can modulate operant ethanol self-administration (Hyytia and Kianamia 2001). These data provide a clear rationale for examining the effects of chronic ethanol and withdrawal on BLA glutamatergic neurophysiology.

Given that classic fear-learning behaviors are associated with a N-methyl-D-aspartate (NMDA)–receptor-dependent increase in glutamatergic synaptic transmission within the BLA (McKernan and Shimnick-Gallagher 1997), we hypothesize that chronic ethanol-related increases in BLA NMDA-receptor function (Floyd et al. 2003) might mediate similar increases in glutamatergic synaptic transmission, ultimately driving increased anxiety during withdrawal. In rats, alcohol withdrawal leads to increased glutamate release in the hippocampus (Dahchour and De Witte 2003). Although they are distinctly different regions, NMDA-dependent changes in synaptic plasticity within the BLA and hippocampus are thought to rely on similar neurophysiological and cellular processes (Chapman et al. 2003). This suggests some parallels between these brain regions. Given that chronic ethanol facilitates NMDA-receptor expression and function in isolated BLA neurons (Floyd et al. 2003), it is reasonable to suggest that molecular mechanisms...
associated with the long-term anxiogenic effects of alcohol exposure are related to alterations in these receptor systems. Indeed, chronic ethanol exposure increases glutamate neurotransmission in the neighboring central nucleus of the amygdala (Roberto et al. 2004b). However, less is known about the effects of chronic ethanol and withdrawal in the BLA, an amygdala subdivision that controls information flow throughout the entire amygdala.

In the present study, we have used a chronic intermittent ethanol inhalation model to examine the effects of chronic ethanol exposure and withdrawal on glutamatergic synaptic transmission in the BLA. Repeated withdrawal from ethanol has been shown to enhance seizure severity in response to convulsants (Pinel 1980). Repeated cycles of exposure/withdrawal also appear to more dramatically alter several different neurotransmitters (Jarvis and Becker 1998; Kang et al. 1998). More important, recent findings suggest that the kindling-like effect of repeated ethanol exposure/withdrawal can occlude synaptic plasticity in the amygdala (Stephens et al. 2005). Because plastic synaptic processes in the BLA are induced by NMDA-type glutamate receptors and expressed as increased function of glutamatergic synapses (Huang and Kandel 1998), the current study therefore tested the hypothesis that repeated ethanol exposure/withdrawal cycles would enhance or alter similar glutamatergic mechanisms within the lateral/basolateral amygdala.

Methods

Animals

All animal procedures were performed in accordance with protocols approved by Wake Forest University School of Medicine Animal Care and Use Committee and were consistent with the National Institutes of Health animal care and use policy. Male Sprague–Dawley rats were obtained from Harlan (Indianapolis, IN) and were housed in an animal care facility at 23°C with a 12-h light/dark cycle and given food and water without restriction. Rats were weighed daily to ensure that ≥80% of their free-feeding weight was maintained during vapor chamber ethanol exposure.

Chronic ethanol exposure

Ethanol exposure was accomplished by ethanol inhalation using a method similar to that used in other studies (Becker and Hale 1993; Roberto et al. 2004b). All animals (100–120 g) were housed four to six animals per cage in large, standard polycarbonate cages (Allentown Caging Equipment, Allentown, NJ) containing wood-chip bedding. These home cages were placed within in large, custom-built Plexiglas chambers (Triad Plastics, Winston-Salem, NC) that were ventilated. These home cages were changed every 2–3 days. Tail blood was taken at the end of some exposure periods and then trunk blood was collected on the day of decapitation for subjects in the CIE group. Blood ethanol concentrations (BECs) were determined using a standard, commercially available alcohol dehydrogenase/NADH enzymatic assay (Diagnostic Chemicals, Oxford, CT). At 3–5 days of exposure, BECs from tail blood were 265 ± 12 mg/dL. At the time of decapitation (~30 min after removal from ethanol vapor), blood ethanol levels were 186 ± 18 mg/dL.

Behavioral assays

To confirm that the CIE exposure paradigm produced withdrawal anxiety, anxiety-like behaviors were assessed using a two-compartment light/dark box (McCool et al. 2003). Individuals were placed in the “light” side of a Plexiglas arena divided equally into “light” and “dark” sides by an opaque Plexiglas insert (Rat Truscan Activity Arena; Coulborn Instruments, Allentown, PA). The center of the animal ± 0.8 cm was followed for 300 s by two infrared sensor rings surrounding the entire apparatus, one in the floor plane and one located about 5 cm above the floor to measure rearing behavior. Data were collected and analyzed for general locomotor activity, time spent in the light and dark compartments, number of light–dark transitions, egress latency, reentry latency, and number of vertical beam breaks. Data were analyzed using one-way ANOVA across the different treatment groups. All variables are reported as the means ± SE.

Preparation of brain tissue

Animals were anesthetized with isoflurane and killed by decapitation. Coronal brain slices (400 μm) were prepared as described previously (Floyd et al. 2003). For in vitro slice preparations, 100 μM ketamine was added to a modified artificial cerebrospinal fluid (aCSF, in mM: 180 sucrose, 4.5 KCl, 1 MgCl₂·6H₂O, 26 NaHCO₃, 1.2 NaH₂PO₄, 10 glucose) during preparation of slices to minimize excitotoxicity. Slices were transferred and stored in 0.5 L of standard oxygenated aCSF solution (in mM: 126 NaCl, 3 KCl, 1.25 NaH₂PO₄, 2 MgSO₄, 26 NaHCO₃, 10 glucose, and 2 CaCl₂) at room temperature for ≥1 h and ≤6 h before performing electrophysiological analysis.

Electrophysiology

Methods for whole cell blind patch-clamp recordings from BLA neurons within slices were similar to those reported previously (Dubois et al. 2006). Electrodes were filled with an intracellular pipette solution containing (in mM): 122 CsOH, 17.5 CsCl, 10 HEPES, 1 EGTA, 5 NaCl, 0.1 CaCl₂, 4 Mg-ATP, and 0.3 Na-GTP, 2 QX-314 (Cl) (pH adjusted to 7.2 with gluconic acid, osmolality ranged from 280 to 290 mmol/kg with sucrose). Excitatory postsynaptic currents (EPSCs) were electrically evoked every 20 s by brief (0.2-ms) square-wave stimulations near the boundary between lateral/ basolateral amygdala (Fig. 1A) using platinum/iridium concentric bipolar stimulating electrodes (FHC, Bowdoinham, ME) with an inner pole diameter of 25 μm and resistance of 8–12 MΩ. Stimulation intensities ranged from 5 to 50 μA. This varied considerably across different experiments and was a function of the specific stimulating electrode used in that particular experiment. Unless stated otherwise, we used submaximal stimulations (just above threshold) that yielded consistent synaptic responses.

All glutamatergic events were pharmacologically isolated using 10 μM bicuculline to inhibit fast GABAergic transmission. Recordings were acquired with an Axoclamp 2B amplifier (Axon Instruments, Foster City, CA) and digitized with a Digidata 1200B (Axon Instruments). From a holding potential of ~60 mV, we included square-wave command hyperpolarizations (~5 mV) in every sweep to constantly monitor input resistance and capacitance. Neurons with high initial input resistance (>50 MΩ; presumptive interneurons;
Rainnie 1999) or whose input resistance increased >10% during data collection were excluded from the study.

NMDA SYNAPTIC RESPONSES. To record NMDA-mediated EPSCs, neurons were voltage-clamped at −60 mV in the presence of low extracellular magnesium (0.2 mM). For the ratio experiment, stimulus strength was adjusted until the amplitude of the compound (AMPA + NMDA) synaptic currents was approximately equivalent across all neurons; stimulus strength was held constant throughout the remainder of the experiment. After recording a stable baseline of the resulting compound AMPA/NMDA EPSC, the AMPA component was inhibited with bath application of 20 μM 6,7-dinitroquinoxaline-2,3-dione (DNQX). The relative contribution of each component within an individual neuron was calculated by dividing the amplitude of the remaining DNQX-resistant (NMDA-receptor–mediated) current by the amplitude of the DNQX-sensitive (AMPA-mediated) component. These ratio data are presented as means ± SE (averaged across cells) and were analyzed using one-way ANOVA across the different treatment groups.

For the NMDA input–output study, neurons from each treatment group were recorded at −60 mV holding potential and incubated with low extracellular Mg2⁺ and the AMPA-receptor antagonist DNQX (20 μM). Increasing stimulus intensities (10–50 μA) produced graded monosynaptic responses.

SPONTANEOUS GLUTAMERGIC SYNAPTIC EVENTS. Spontaneous excitatory post synaptic currents (sEPSCs) were acquired at 20 kHz and filtered at 2 kHz. For miniature (m)EPSCs, 1 μM tetrodotoxin (TTX) was bath applied for >5 min before recording spontaneous events. This concentration of TTX inhibited >99% of all electrically evoked EPSCs (not shown). TTX-sensitive and -resistant sEPSCs were detected and analyzed using Mini Analysis Program 6.0.3 (Synaptosoft, Fort Lee, NJ). Measures from individual cells were averaged across treatment groups (Van Sickle et al. 2004), reported as means ± SE, and analyzed using standard one-way ANOVA. In some cases, sEPSC and mEPSC data are expressed as cumulative probability distributions from individual cells representing the median across a treatment group. These data were analyzed using the Kolmogorov–Smirnov (KS) test for different population distributions. Significance was determined at a KS statistic of ≥0.09 rather than using the P value; this cutoff makes the less-conservative KS test more stringent.

NOISE ANALYSIS. Root-mean-square current noise (i_rms) was analyzed in the spontaneous event records. Noise was measured independently in four 50-ms epochs by comparing current amplitudes at each data point to the mean current amplitude across the epoch as previously described (Mitchechvili and Kapur 2006). Epochs containing obvious sEPSCs were avoided in the analysis. These data, reported as means ± SE, were statistically analyzed using standard one-way ANOVA.

PAIRED-PULSE RATIO. Paired-pulse (PP) facilitation was measured using pairs of electrical stimuli of equal intensity at 25-, 50-, or 250-ms interpulse intervals. Ratios of the amplitudes of the evoked EPSCs were calculated as the difference between the amplitude of the second event minus the amplitude of the first, with the result divided by the amplitude of the first synaptic response (Schulz et al. 1995). All values were expressed as means ± SE, and data were subjected to one-way ANOVA, with P < 0.05 considered statistically significant.

Microinjections

We accomplished microinjections into the BLA of male Sprague–Dawley rats according to previously published procedures (McCool and Chappell 2007). Briefly, we deeply anesthetized animals with pentobarbital (90 mg/kg) and affixed them to a stereotaxic instrument. Chronic guide cannulae were placed bilaterally into the dorsal aspect of the BLA and affixed to the skull with dental cement. We used sterile obturators to maintain the patency of the guide cannulae. During a 5-day recovery period, animals were handled extensively and habituated to the injection procedure and sound of the syringe pump. On test days, 0.5 μl drug in standard aCSF was directly infused into the BLA over a 1-min period. Injection cannulae were left in place for 1 min and animals were placed into the light/dark box 5 min after the microinjection. After the microinjections and behavioral measurements, guide cannulae placement was confirmed postmortem.

Two separate microinjection experiments were performed in the current study. In the first experiment, adult male Sprague–Dawley rats (n = 39; 303.9 ± 0.6 g at the time of surgery) were used to test the effects of muscimol microinjection into the BLA on light/dark box anxiety-like behavior. For these experiments, guide cannulae were implanted according to the following stereotaxic coordinates (millimeters relative to bregma; Paxinos and Watson 1997): −2.8 anterior/posterior; ±5.05 medial/lateral; and −6.20 dorsal/ventral (measured from the top of the brain). Sham, 88 nmol (10 μg), or 264 nmol (30 μg) muscimol (in aCSF) was microinjected into separate animals; individual animals were exposed to the light/dark box only once. Across two separate cohorts, we misplaced guide cannulae in only 3 of 39 animals used in the muscimol study; these are not included in the analysis (not shown; see text).

In the second experiment, male Sprague–Dawley rats (n = 19, 138.0 ± 2.6 g at the time of surgery) similar to those used for the electrophysiology experiments were used to test the effects of DNQX on withdrawal-related anxiety-like behavior expressed in the light/dark box. For these studies, animals were surgerized 1 h after the fifth ethanol intermittent inhalation. This time point was selected for two reasons. First, animals appeared behaviorally tolerant to the effects of the ethanol inhalation by the 5th day; this may help reduce interactions between the anesthetic and the ethanol during the surgery. Second, surgerizing animals before the ethanol exposure proved difficult because of changes in body weight and skull size during the long-term exposure. For the surgery, ethanol-exposed animals were deeply anesthetized with pentobarbital (50 mg/kg). In three separate cohorts (five to seven animals/cohort), guide cannulae were implanted according to the following stereotaxic coordinates (millimeters relative to bregma) that were empirically determined in pilot studies on juvenile animals: −2.7 anterior/posterior; +4.5 lateral on the right; −4.75 lateral on the left; −5.8 dorsal/ventral from the top of the brain. After the surgery, animals recovered for 10 h before being placed back into the inhalation chamber to complete the last 5 days of the chronic inhalation treatment. When not in the inhalation chamber, animals were handled during this period to habituate them to the injection procedure. After 24-h withdrawal following the 10th ethanol exposure, individual animals were micromanipulated with either sham or 100 pmol DNQX, and individuals were exposed to the light/dark box only once. In three separate cohorts, we placed guide cannulae outside the BLA in only 2 of 19 animals (see Fig. 6A).

RESULTS

Withdrawal from chronic ethanol inhalation increases anxiety-like behaviors

We initially assessed whether our treatment paradigm was relevant for the anxiety-related effects of chronic ethanol and withdrawal. Fifty-seven animals were divided equally into three groups (n = 19 each) with two groups receiving 10 days of 12 h/day ethanol vapor, whereas the third control group received room air in an identical chamber. We then measured anxiety-like behavior using the “light/dark” box assay in controls, immediately after the last ethanol exposure while individuals were still intoxicated (as evidenced by blood ethanol levels; “CIE” animals), or 24 h after withdrawal from ethanol vapors (“WD” animals). One CIE animal appeared sedated,
was clearly an outlier with regard to low locomotor activity, and was excluded from the behavioral study. Similarly, four animals from the withdrawn group appeared to freeze when initially placed on the light side of the behavioral apparatus (e.g., reduced locomotion with \( \geq 60 \) s to initially egress from the light side to the dark side) and were not analyzed further.

For the remaining animals, 24 h of withdrawal from 10 days of chronic intermittent ethanol exposure caused a significant decrease in time spent in the light side, a significant decrease in the number of light–dark transitions, and a significant increase in the reentry latency (time to reenter the light following first egress into the dark; Table 1). Egress latency (initial time to leave the light side and move into the dark side) was not significantly different between the treatment groups. The alterations in most “anxiety-related” variables are consistent with increased expression of anxiety-like behaviors in the WD animals. The total move time was not different between groups, indicating that the treatment did not affect the locomotor activity in the animals. Exploratory behaviors, represented by vertical-plane entries (“rears”), were significantly suppressed in both CIE and WD animals, again suggesting that the anxiety effects were specific to withdrawal.

Recent evidence suggests that anxiety-like behavior expressed in the light/dark box is dependent on BLA neurotransmitter systems (Bueno et al. 2005; McCool and Chappell 2007). To test this, we microinjected 0 (n = 12), 88 (10 \( \mu \)g; n = 14), or 264 nmol (30 \( \mu \)g; n = 13) muscimol into the BLA and measured light/dark box behavior. For several of the anxiety-related–dependent variables expressed in this assay, muscimol diminished anxiety-like behavior in a dose-dependent fashion. For example, 264 nmol muscimol increased the time spent in the light side of the apparatus (108 \( \pm 9 \) s; \( P < 0.05 \), one-way ANOVA) compared with sham (73 \( \pm 13 \) s) and 88 nmol muscimol (63 \( \pm 15 \) s). Likewise, the reentry latency (time to reenter the light side following egress into the dark) was significantly less in animals microinjected with 264 nmol muscimol (48 \( \pm 14 \) s; \( P < 0.05 \), one-way ANOVA) compared with either sham (140 \( \pm 33 \) s) or 88 nmol muscimol (111 \( \pm 22 \) s). However, the number of light–dark transitions and the initial egress latency (time to leave the light side after placement) were not significantly affected by muscimol. Likewise, locomotor-related behaviors were not altered by the test doses of muscimol examined here. The total distance moved during the assay was 1,343 \( \pm 42 \) cm in sham animals, 1,241 \( \pm 72 \) cm in animals microinjected with 88 nmol muscimol, and 1,377 \( \pm 38 \) with 264 nmol muscimol (\( P > 0.05 \), one-way ANOVA). Similarly, the total time spent moving was 227.3 \( \pm 5.0 \) s in sham, 209.7 \( \pm 6.6 \) s with 88 nmol muscimol, and 229.8 \( \pm 3.3 \) s with 264 nmol muscimol. Together, these data confirm that manipulation of BLA neural activity can alter anxiety-like behavior expressed in the light/dark box.

### Chronic ethanol and withdrawal increase the contribution of NMDA receptors at BLA glutamatergic synapses

Previous studies in our lab have demonstrated that chronic ethanol can lead to increased NMDA-receptor function measured in somatic compartments of acutely isolated BLA neurons (Floyd et al. 2003). To test whether these alterations are expressed within synaptic compartments, we investigated the effects of CIE and WD on NMDA-receptor function at glutamatergic synapses using whole cell in vitro slice electrophysiology. We first established a stable excitatory synaptic response using both low extracellular magnesium as well as the most minimal stimulation required to produce consistent monosynaptic responses. For these studies, response amplitudes in the low extracellular magnesium were 127 \( \pm 23 \) pA in CON neurons, 114 \( \pm 15 \) pA in CIE neurons, and 114 \( \pm 13 \) pA in WD neurons; these values were not statistically different from one another (\( P >> 0.05 \), one-way ANOVA). Within each neuron, we next perfused slices with the AMPA-receptor antagonist DNQX (20 \( \mu \)M; Fig. 1B) to pharmacologically isolate NMDA-receptor–mediated synaptic currents. When expressed as a ratio of NMDA-receptor–mediated current amplitude to the DNQX-sensitive AMPA-receptor–specific component, we found significant increases in the DNQX-insensitive NMDA-receptor–mediated current (Fig. 1C) in both CIE (\( P < 0.05 \), n = 6) and WD (\( P < 0.05 \), n = 7) neurons compared with control (n = 7). In a separate set of experiments, >95% of DNQX-insensitive synaptic current in BLA neurons was inhibited by 50 \( \mu \)M of the NMDA-receptor antagonist 2-amino-5-phosphonovaleric acid (APV; not shown). This inhibition did not differ between the three treatment groups.

We also examined the effects of CON, CIE, and WD treatments on NMDA-mediated synaptic currents by measuring responses to increasing stimulation intensities (input–output relationship). In all three treatment groups, local electrical stimuli elicited DNQX-insensitive synaptic currents whose amplitude increased in an intensity-dependent fashion (Fig. 1D). At the highest stimulation intensities, the amplitude of the DNQX-insensitive NMDA-mediated synaptic current was significantly greater in both CIE and WD neurons compared with control (\( P < 0.05 \), one-way ANOVA). Along with the ratio data, these input–output results clearly suggest that the contrib-

### Table 1. Increased anxiety-like behavior in the light/dark box following withdrawal from chronic intermittent ethanol

<table>
<thead>
<tr>
<th>Variable</th>
<th>CON</th>
<th>CIE</th>
<th>WD</th>
<th>Statistic</th>
</tr>
</thead>
<tbody>
<tr>
<td>Move time, s</td>
<td>234.1 ( \pm ) 2.5</td>
<td>231.8 ( \pm ) 4.9</td>
<td>221.2 ( \pm ) 4.3</td>
<td>ns</td>
</tr>
<tr>
<td>Light time, s</td>
<td>95.9 ( \pm ) 8.2</td>
<td>104.0 ( \pm ) 10.1</td>
<td>55.7 ( \pm ) 11.6</td>
<td>( P &lt; 0.01 )</td>
</tr>
<tr>
<td>Transitions</td>
<td>7.8 ( \pm ) 0.6</td>
<td>9.1 ( \pm ) 1.2</td>
<td>4.9 ( \pm ) 0.9</td>
<td>( P &lt; 0.01 )</td>
</tr>
<tr>
<td>Egress latency, s</td>
<td>21.5 ( \pm ) 3.0</td>
<td>22.4 ( \pm ) 4.9</td>
<td>16.4 ( \pm ) 3.2</td>
<td>ns</td>
</tr>
<tr>
<td>Reentry latency, s</td>
<td>72.2 ( \pm ) 15.4</td>
<td>65.5 ( \pm ) 19.6</td>
<td>171.4 ( \pm ) 30.8</td>
<td>( P &lt; 0.01 )</td>
</tr>
<tr>
<td>Vertical-plane entries</td>
<td>32.0 ( \pm ) 2.8</td>
<td>21.2 ( \pm ) 2.6</td>
<td>21.3 ( \pm ) 1.9</td>
<td>( P &lt; 0.01 )</td>
</tr>
</tbody>
</table>

Values are means \( \pm \) SE. *Anxiety-like behavior was assayed for 300 s immediately after 10 days’ exposure to room air (“CON,” n = 18), 10 days of intermittent ethanol vapor (“CIE,” n = 17), or 24-h withdrawal from ethanol vapor (“WD,” n = 14). See text for details. \( \dagger P < 0.05 \) versus CON and \( \ddagger P < 0.01 \) versus CON (one-way ANOVA)
Ethanol and withdrawal alter BLA glutamatergic transmission

 Contributions by synaptic NMDA receptors are increased by chronic ethanol exposure and are maintained during 24 h of withdrawal.

Chronic ethanol and withdrawal increase the amplitude and frequency of spontaneous EPSCs

Because the NMDA/AMPA ratio is also dependent on the relative contribution by AMPA receptors, a decrease in AMPA-receptor-mediated synaptic currents might increase the NMDA/AMPA ratio in slices prepared from CIE and WD animals. We attempted to test this by measuring the AMPA input–output relationship. However, under the current recording conditions (local stimulation and standard aCSF containing 10 μM bicuculline), AMPA-mediated synaptic responses had an extremely steep stimulus–response relationship with polysynaptic responses occurring at stimulation intensities >20 μA (responses >200 pA). This prevented a comparison of AMPA input–output across the treatment groups.

Alternatively, we examined spontaneous excitatory postsynaptic currents (sEPSCs; Fig. 2B). Under standard recording conditions (2 mM extracellular magnesium and −60-mV holding potential), spontaneous synaptic currents were entirely sensitive to the AMPA-receptor–specific antagonist GYKI 53655 (Fig. 2A). GYKI 53655 also inhibited electrically evoked responses by 96 ± 1% in CON neurons (n = 4), 98 ± 1% in CIE neurons (n = 6), and 96 ± 1% in WD neurons (n = 5; P > 0.05, one-way ANOVA). These data suggest that AMPA-type glutamate receptors mediate the spontaneous synaptic currents. With respect to the treatment groups, we found the sEPSC amplitude in CIE, but not WD, neurons was significantly increased relative to control (Fig. 2, B and C). However, the charge carried by these spontaneous events was not significantly different between the treatment groups (Fig. 2D).

Chronic ethanol and withdrawal decrease the PP ratio

The increased frequency of sEPSCs without substantial changes in amplitude might suggest increased presynaptic function following CIE and WD. To test this, we measured...
synaptic responses to two closely juxtaposed electrical stimuli (paired pulses; Fig. 3A). At short interpulse intervals, the ratio of the second synaptic response relative to the first is believed to be inversely related to the probability of release at synapses (Andreasen and Hablitz 1994; Katz et al. 1993). At interpulse intervals of 25 and 50 ms (Fig. 3A), CIE and WD both significantly decreased the PP ratios (Fig. 3B). In support of the sEPSC frequency data, the PP findings suggest that CIE and WD increase presynaptic glutamatergic function, potentially by increasing release probability at the terminal.

Withdrawal increases miniature EPSC amplitude, decay, and frequency

To confirm the possible alterations of presynaptic terminal function, we measured the effects of CIE and WD on the amplitude and frequency of TTX-resistant miniature EPSCs under standard recording conditions (2 mM extracellular magnesium and −60-mV holding potential; Fig. 4, A and B). WD robustly increased the frequency of mEPSCs (decreased interevent interval) compared with control (P < 0.01, one-way ANOVA) and CIE (P < 0.05; Fig. 4, E and F). Importantly, CIE did not alter mEPSC frequency, in contrast to the sEPSC and PP findings. Because TTX blocks action potentials, these data in the CIE neurons may imply that the presynaptic alterations evident in this treatment group are action potential dependent. Furthermore, comparisons between the sEPSC and mEPSC data suggest that CIE and WD may alter presynaptic glutamatergic function by distinct mechanisms.

In addition to these apparent presynaptic changes, WD also increased the amplitude of the mEPSCs (Fig. 4C). However, the charge carried by mEPSCs was not significantly altered by any treatment (Fig. 4D). Together these amplitude and charge data suggest some change in the kinetics of glutamatergic responses in WD neurons. We therefore examined the onset and decay kinetics of the miniature EPSCs (Fig. 5A). Average mEPSC traces were generated by aligning the rising phase of each event in each cell and scaling the amplitude. The resulting average mEPSC from each cell was fit to a first-order exponential equation. Rise times were not significantly different from each other (one-way ANOVA, Fig. 5B). However, the decay of mEPSCs was significantly faster in WD neurons compared with both CIE and CON (P < 0.05, one-way ANOVA; Fig. 5C). These data help explain the significant up-regulation of mEPSC amplitude in WD neurons in the absence of any substantial alterations in the amount of charge carried by events in this treatment group.
**BLA glutamatergic system and withdrawal-related anxiety-like behavior**

Given the focus of the current work on glutamatergic function in the BLA, we examined the effects of microinjection with the AMPA/kainate receptor antagonist DNQX (100 pmol) on anxiety-like behavior in withdrawn (WD) animals (Fig. 6A). Importantly, DNQX substantially diminished the anxiety-like behavior expressed by WD animals in the light/dark box. Both the time spent in the light side (Fig. 6B1) and the number of light-to-dark transitions (Fig. 6B2) were significantly increased (P < 0.05 and P < 0.01, respectively; t-test) by DNQX microinjection. Likewise, DNQX significantly (P < 0.01, t-test) decreased the reentry latency in WD animals (Fig. 6B3). These effects on anxiety-like behavior appear to be specific to delivery of the AMPA/kainate receptor antagonist DNQX into the BLA of WD animals can diminish anxiety-like behavior expressed in the light/dark box.

**DISCUSSION**

Our results suggest that the increased anxiety-like behavior observed during withdrawal may be due to increases in glutamatergic function seen during CIE and WD. This effect appears to have a presynaptic origin, as evidenced by the decreased PP ratio and increased frequency of mEPSCs during WD. We also found increases in the NMDA/AMPA ratio as well as increases in amplitude and frequency of spontaneous events during CIE and WD and during WD for mEPSCs; these latter data suggest that postsynaptic alterations may result from these treatments as well. Our NMDA findings parallel previous evidence showing an increase in the function and expression of NMDA receptors in isolated BLA neurons following chronic ethanol liquid diet exposure (Floyd et al. 2003). The most reasonable interpretation of the changes seen during withdrawal is that ethanol exposure increases presynaptic release of glutamate.

**FIG. 3.** Chronic ethanol and withdrawal decrease paired-pulse (PP) ratio in the BLA. A: sample traces of PP EPSCs at 50-ms interpulse interval in the presence of 10 μM bicuculline. Amplitude of the second synaptic response has been normalized across treatment groups to emphasize the relative differences between the first synaptic response and the second. PP ratio is significantly decreased at both the 25- and 50-ms intervals (*P < 0.05, **P < 0.01 vs. control, one-way ANOVA with Bonferroni’s posttest). These decreased ratios indicate that chronic ethanol and withdrawal may increase presynaptic release of glutamate.

**FIG. 4.** Withdrawal increases the frequency and amplitude of miniature (m)EPSCs. A: sample traces demonstrating the efficacy of 1 μM tetrodotoxin used in these studies. B: sample traces of mEPSCs in the presence of 10 μM bicuculline and 1 μM TTX (holding potential, −60 mV) are shown for each treatment group. C: mEPSC amplitude from WD neurons was significantly greater than that from CON (**P < 0.05) and CIE neurons (#P < 0.05; one-way ANOVA with Bonferroni’s posttest). D: charge carried by mEPSCs [amplitude (in pA) × decay time (in ms), expressed as femtocoulombs] was not significantly different from CON in the CIE and WD treatment groups. E: interevent interval of WD mEPSCs was significantly smaller than that of both control (**P < 0.01) and CIE neurons (#P < 0.05; one-way ANOVA with Bonferroni’s posttest). F: cumulative probability plot of interevent interval from individual CON, CIE, and WD neurons; each cell is representative of the median from that particular treatment group. Interevent interval distribution of the WD neuron (KS = 0.19, P < 0.0001), but not the CIE neuron (KS = 0.06, P > 0.05), was significantly different from that of the CON neuron.
CIE and WD include increased excitability of glutamatergic afferents in the BLA, increased presynaptic terminal function/number, and more modest postsynaptic increases in receptor function.

It should be noted that our findings must be interpreted in the context of the current exposure paradigm. Our intermittent exposure produces robust blood ethanol concentrations during the 10-day exposure period. Regardless, CIE rats with approximately 190 mg/dL blood-ethanol have little apparent motor incoordination, at least as it is represented in the light/dark box (Table 1). Nonetheless, this same exposure paradigm produces elevated anxiety-like behavior during subsequent withdrawal. Chronic human alcoholics can also have remarkable functional tolerance despite blood levels that would produce coma and death in naıve individuals (Davis and Lipson 1986; Hammond et al. 1973). It is clear then that the intermittent exposure pattern and severity of exposure used in the current study may model this type of profound “tolerance-dependence.”

**Chronic ethanol facilitates synaptic-NMDA and -AMPA receptors in the basolateral amygdala**

Our previous work has shown that chronic exposure to an ethanol liquid diet increased NMDA-receptor function in the somatic compartments of acutely isolated BLA neurons (Floyd et al. 2003). Here we show that chronic intermittent ethanol inhalation also significantly increased NMDA-receptor function, exemplified by the NMDA/AMPA ratio, at BLA glutamatergic synapses. This is similar to findings in the central nucleus after chronic ethanol (Roberto et al. 2006). Together, these results strongly support the hypothesis that CIE up-regulates postsynaptic NMDA-receptor number or function in the amygdala. The amygdala therefore joins a growing list of brain regions responding to chronic ethanol by increased expression/function of NMDA receptors.

Chronic ethanol also increases AMPA-receptor protein in primary cortical cultures (Chandler et al. 1999) as well as AMPA-receptor–dependent calcium signaling in cerebellar Purkinje neurons (Netzeband et al. 1999). However, we found no evidence that CIE produces any alteration in AMPA-receptor postsynaptic function. Preliminary evidence (J. Weiner, personal communication) suggests that acute ethanol robustly inhibits BLA NMDA-mediated synaptic responses, whereas AMPA-mediated currents are relatively insensitive. This differential acute sensitivity might explain the robust up-regulation of NMDA synaptic currents to chronic ethanol, whereas AMPA-mediated events are more modestly affected.

Chronic ethanol facilitates TTX-sensitive presynaptic function in the basolateral amygdala

Our paired-pulse data, specifically the decrease in facilitation, clearly indicate increased presynaptic function with CIE. This is supported by an increased frequency of spontaneous events in these neurons. Our data are further supported by increased presynaptic glutamate release in both the central nucleus of the amygdala using a similar exposure paradigm (Roberto et al. 2004b) as well as in the hippocampus during ethanol consumption (Sabria et al. 2003). However, the increase in BLA sEPSC frequency was entirely TTX sensitive; CIE did not alter the frequency of miniature, TTX-resistant EPSCs. These data may imply that CIE facilitates action-potential–dependent mechanisms within the BLA glutamatergic afferents themselves. In support of this, chronic ethanol can alter a number of action-potential–related processes (Scott and Edwards 1981), including decreased calcium-activated potassium channels (Pietrzykowski et al. 2004) and increased TTX-sensitive voltage-gated sodium channels (Brodie and Sampson 1990) or L-type calcium channels (Watson and Little 1999). In fact, dihydropyridine-sensitive voltage-gated calcium channels appear to be recruited to BLA glutamatergic synapses following fear learning (Shinnick-Gallagher et al. 2003). Changes within the context of action-potential–related, TTX-sensitive cellular processes could therefore lead to increased neurotransmitter release at BLA glutamatergic synapses.

Finally, it is worth noting that slices prepared from CIE animals were incubated at room temperature in aCSF for as
long as 6 h during the data collection. Because we did not include ethanol in the slice incubation media, it is possible that CIE slices were experiencing “acute in vitro” withdrawal during this period. Our data do not support this hypothesis. First, all CIE animals were killed within 30 min after removal from the inhalation chamber. Because we know precisely when each data file was collected, we performed a correlation analysis between the various sEPSC and mEPSC parameters and the time elapsed between collection of these data and preparation of the slices. For all CIE neurons, there was no significant correlation between these event-related dependent variables and the time the slice had been stored in vitro. This argues against a significant contribution by “acute in vitro” withdrawal to CIE-related alterations in glutamatergic transmission.

Withdrawal maintains postsynaptic changes seen during chronic exposure

After 24 h of withdrawal from chronic ethanol, we saw an increase in NMDA/AMPA ratio and the NMDA input–output relationship in BLA neurons. Given that sEPSC data suggest that AMPA-receptor function is not down-regulated during WD, the NMDA data indicate that the increased receptor function seen with CIE is maintained for at least 24 h after the exposure. There are conflicting reports concerning the effects of ethanol withdrawal, relative to chronic ethanol, on NMDA receptors in other brain regions. NMDA-receptor function appears to increase during ethanol withdrawal in the mouse hippocampus (Whittington et al. 1995), although in the central amygdala, withdrawal did not alter NMDA-receptor synaptic function (Roberto et al. 2006). These data indicate that WD-related changes in NMDA-receptor function are clearly brain region dependent, even within the amygdala.

Withdrawal also induced postsynaptic facilitation of AMPA-mediated glutamatergic spontaneous EPSCs in the BLA. Specifically, we found a large increase in the amplitude of TTX-resistant spontaneous EPSCs. This effect appears to share similarities with other drugs of abuse. For example, the AMPA-receptor subunit GluR1 has been shown to be up-regulated after 1 and 30 days of withdrawal from cocaine in the BLA of rats (Lu et al. 2005). Similarly, TTX-resistant EPSCs from WD neurons decayed faster than those measured in both control and CIE. Increased AMPA-mediated EPSC decay is associated with increased delivery of GluR1 homomeric channels to glutamatergic synapses in response to sensory stimulation of barrel cortex (Clem and Barth 2006) and pharmacologic blockade of AMPA receptors in hippocampal neuron cultures (Thiagarajan et al. 2005). The faster decay (Atassi and Glavinovic 1999; Hiraseawa et al. 2001; Veruki et al. 2003), along with increased amplitude (Carroll et al. 1998; Thio et al. 1992), may indicate postsynaptic alterations within the AMPA-receptor complex expressed by WD neurons. One possible mechanism explaining increased EPSC amplitude/altered kinetics following WD would be increased delivery of specific AMPA-receptor subtypes to BLA glutamatergic synapses. However, this remains to be directly investigated. Regardless, the increase in mEPSC amplitude was not associated with substantial increases in charge carried by individual events. The contribu-
tion of postsynaptic alterations to BLA-dependent, WD-related behaviors may therefore be subtle.

Withdrawal produces presynaptic, terminal-specific alterations

There is a general lack of data on the effects of ethanol withdrawal on presynaptic mechanisms. In the central nucleus of the amygdala, glutamate release remains elevated during both chronic ethanol exposure (Roberto et al. 2004b). The results of our mEPSC and paired-pulse data in the BLA complement these findings and suggest that WD specifically alters presynaptic terminal function. Potential mechanisms include altered release machinery (Bacci et al. 2001; Capogna et al. 1997; Herreros et al. 1995; Pang et al. 2006), increased resting calcium levels in the terminal (Cummings et al. 1996; Levesque and Atchison 1988; Li et al. 1998; Nishimura et al. 1990), or increased numbers of synapses (Lauri et al. 2003) following WD. Chronic blockade of glutamate receptors in hippocampal cultures selectively increases AMPA-mediated mEPSC frequency without altering synapse number (Bacci et al. 2001). Conversely, withdrawal from repeated cocaine exposures increases the number of glutamate-containing terminals in the shell of the nucleus accumbens (Kozell and Meshul 2004). Ultimately, the mechanisms regulating increased mEPSC frequency during WD from CIE are likely to be both brain region and treatment specific.

Behavioral implications

Given the association between the BLA glutamatergic system and anxiety (Sajdyk and Shekhar 1997), increases in glutamatergic function like those observed in the current work may contribute to the anxiety-like behavior evident during withdrawal from chronic ethanol. Additional studies that follow a withdrawal time course would be necessary to more robustly illustrate this. However, we have used juvenile animals to facilitate the whole cell electrophysiology recordings, and this creates confounds between developmental issues and a protracted time course. Studies in adult animals would perhaps be more appropriate in this respect but are also substantially more challenging.

Regardless, our microinjection results with DNQX clearly suggest that BLA mechanisms associated with AMPA/kainate-type glutamate receptors help regulate the expression of anxiety-like behavior during withdrawal. These findings parallel studies showing that amygdala AMPA receptors are important for the expression of learned fear-potentiated startle (Walker and Davis 1997b) and learned avoidance tasks (Mesches et al. 1996). Importantly, the expression of learned fear is coincident with increased AMPA-receptor-mediated synaptic function at BLA glutamatergic synapses (McKernan and Shinnick-Gallagher 1997) and with increased delivery of AMPA-receptor subunits to the cell surface (Yeh et al. 2006). The coincidental increase in BLA glutamatergic synaptic function during withdrawal, the increase in withdrawal-related anxiety-like behavior, and the apparent contribution of BLA AMPA-type glutamate receptors to withdrawal-associated anxiety-like behavior together suggest an intimate relationship between these parameters.

Paradoxically, we found that glutamate function was elevated during the CIE exposure, despite an absence of elevated anxiety-like behavior measured in the light/dark box. One potential explanation is that the BLA contributes minimally to CIE-related behaviors expressed in the light/dark box. Most evidence does not support this hypothesis. Several recent studies (de la Mora et al. 2005; McCool and Chappell 2007; Perez de la Mora et al. 2006; Salome et al. 2006) as well as the muscimol study reported in the current work have demonstrated the contribution of BLA-dependent processes to light/dark box anxiety-like behavior in naïve animals. Therefore compensatory changes in other BLA/downstream neurotransmitter systems or continued sensitivity of these systems to acute ethanol are more likely to explain the paradoxical increase in BLA glutamatergic function in the absence of significant increases in anxiety-like behavior. This latter “continued sensitivity” hypothesis is particularly relevant because anxiety-like behaviors in CIE animals were measured while the individuals were still intoxicated (based on BEC at decapitation).

The potential neurotransmitter systems contributing to CIE-related behaviors are quite numerous. Within the amygdala fear/anxiety circuit, recent reports suggest that GABAergic function is increased (Roberto et al. 2004a), whereas glutamatergic function is decreased (Roberto et al. 2004b), in the neighboring central amygdala in ethanol-dependent rats. Because the projections from BLA to central amygdala are critical for the expression of learned fear (Davis 2006), it is possible these central amygdala alterations might offset increased glutamatergic function in the BLA during CIE treatment, ultimately masking anxiety-related behavioral manifestations in these animals. Within the BLA itself, alcohol has well-known effects on numerous neurotransmitter systems. For example, acute and chronic ethanol alter GABAergic function in BLA neurons (Floyd et al. 2004; McCool et al. 2003; Zhu and Lovenger 2006). In addition, BLA serotonergic (Gonzalez et al. 1996), dopaminergic (de la Mora et al. 2005), noradrenergic (Schroeder et al. 2003), and various neuropeptide systems (Rupniak et al. 2003; Sajdyk et al. 1999a,b; Wunderlich et al. 2002) all regulate anxiety-like behavior. Disruption or alteration of any of these systems might also contribute to anxiety-related behavior observed in intoxicated rats exposed to chronic ethanol.

In conclusion, we have shown that chronic intermittent ethanol exposure and withdrawal lead to significant increases in glutamatergic synaptic transmission in the basolateral amygdala. Our working hypothesis for the sequence of events that result from this treatment is that chronic ethanol produces both increased postsynaptic NMDA-receptor function as well as increased glutamatergic afferent excitability. At some point during the withdrawal process, these alterations trigger significant changes in presynaptic terminal function and more modest alterations of postsynaptic AMPA-receptor function. This increased glutamatergic system in the basolateral amygdala was not observed behaviorally until withdrawal because we hypothesize that the additional ethanol-sensitive systems may be engaged while the animal is still intoxicated.

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