

The reinforcing effects of ethanol within the nucleus accumbens shell involve activation of local GABA and serotonin receptors

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Abstract

Ethanol is reinforcing within the nucleus accumbens shell (NACsh), but the underlying mechanisms remain unclear. Ethanol can potentiate the function of the GABA_A, GABA_B, and 5-HT₃ receptors. Therefore, the current study tested the hypothesis that activation of these receptors would be involved in the reinforcing effects of ethanol in the NACsh. An intracranial self-administration (ICSA) procedure was used to assess the reinforcing effects of ethanol *in the NACsh of alcohol preferring (P) rats*. The ICSA consisted of 7 sessions: 4 sessions to establish 150 mg% ethanol self-infusion into the NACsh; sessions 5 and 6 with co-infusion of ethanol plus one concentration of the GABA_A antagonist bicuculline (10 or 100 μM), the GABA_B antagonist SCH 50911 (50, 75 or 100 μM), or the 5-HT₃ receptor antagonist zacopride (10 or 100 μM); and session 7 with 150 mg% ethanol alone. All groups self-infused ethanol into the NACsh and readily discriminated the active from inactive lever during the acquisition sessions. Co-infusion of 100 μM, but not 10 μM, bicuculline or zacopride significantly decreased active responses during sessions 5 and 6. Co-infusion of 75 μM, but not 50 or 100 μM, SCH 50911 significantly attenuated responses for ethanol. Overall, the results suggest that the reinforcing effects of ethanol in the NACsh may be modulated by activation of local GABA_A, GABA_B and 5-HT₃ receptors.

Key words: Intracranial self-administration, nucleus accumbens, ethanol, GABA_A receptor, GABA_B receptor, 5-HT₃ receptor

Introduction

Identification of brain reward mechanisms underlying the reinforcing effects of ethanol is critical to the development of effective therapies for combating alcohol abuse and alcoholism. A recent intracranial self-administration (ICSA) study demonstrated that ethanol can be readily self-infused into the nucleus accumbens (NAC) of the rat; the self-infusion was sub-region dependent with self-infusion into the NAC shell (NACsh) but not NAC core (NACcr), and genetically influenced with alcohol preferring (P) rats being more sensitive than stock Wistar rats to the rewarding effects of ethanol (Engleman et al. 2009). These findings suggest that the NACsh is one neuro-anatomical substrate supporting the reinforcing and rewarding effects of ethanol. However, neurochemical mechanisms underlying these effects remain unknown and elucidation of such mechanisms are important.

The NAC is a heterogeneous nucleus and approximately 90% of its neurons are GABAergic medium spiny neurons (MSNs); it receives synaptic inputs from various brain regions (Meredith 1999; Sesack and Grace 2010). GABA synapses provide a major inhibitory regulation of MSN activity. These GABAergic inputs originate mainly from GABA neurons in the VTA (Van Bockstaele and Pickel 1995) and ventral pallidum (Groenewegen et al. 1999), as well as intrinsic axonal collaterals from local MSNs (Meredith 1999). Moderate expression of both GABA_A and GABA_B receptors were found in the NAC (Bowery et al. 1987; Schwarzer et al. 2001). In addition, the NAC receives modulatory serotonin (5-HT) afferents from the dorsal raphe nucleus (Van Bockstaele and Pickel 1993). The presence of 5-HT receptors has been demonstrated in the NAC (Meredith 1999), including the serotonin-3 (5-HT₃) receptor (Ge et al. 1997). The interactions between GABA/5-HT and their respective receptors play an important role in regulating the MSN activity and other neurotransmitter systems converging on the NAC

(Meredith 1999). It has been proposed that the rewarding information is encoded in the activity of MSNs in the NAC and forwarded to downstream structures including the ventral pallidum (Carlezon and Thomas 2009).

Substantial evidence suggests that GABA neurotransmission is involved in the effects of ethanol (Koob 2004). Electrophysiological studies indicate that acute ethanol can potentiate GABA_A- and GABA_B- receptor mediated inhibitory currents (Allan et al. 1987; Federici et al. 2009; Grobin et al. 1998). Systemic administration of GABA_A receptor antagonists or an inverse benzodiazepine agonist reduced voluntary alcohol drinking and the motivation in rats to respond for ethanol self-administration in an oral operant setting (McBride et al. 1988; Petry 1997; Rassnick et al. 1993; Samson et al. 1987). GABA_A receptors within the mesolimbic system appeared to mediate such effects (Hodge et al. 1995; Hyytia and Koob 1995). Similarly, pharmacological manipulations of GABA_B receptor activity can regulate the reinforcing and rewarding effects of ethanol (Maccioni and Colombo 2009; Vlachou and Markou 2010). In addition, acute ethanol can facilitate 5-HT₃ receptor-mediated ion currents (Lovinger and White 1991). The 5-HT₃ receptor has been implicated in the effects of drugs of abuse (Engleman et al. 2008). Systemic administration of 5-HT₃ receptor antagonists suppressed voluntary ethanol consumption in rats under 24-hr free-choice conditions (Knapp and Pohorecky 1992; McKinzie et al. 1998). Nonetheless, the involvement of these receptors in the local reinforcing effects of ethanol within the NACsh has not been examined. Given the above-mentioned findings, it is possible that activation of these receptors may be involved in mediating the self-infusion of ethanol into the NACsh. The current study was designed to test this hypothesis by examining local self-infusion of ethanol in the presence of antagonists for these receptors.

Materials and methods

Animals: Experimentally-naïve adult female alcohol preferring (P) rats (body weight 250 to 300 g, Indiana University) were housed in-pairs in a reverse 12-hr light-dark cycle room (light off at 10:00 am) with temperature and humidity maintained constant for at least 2 weeks before the commencement of experiment. Food and water were available *ad libitum* except during the ICSA test sessions. Female rats were used in the present study due to their ability to maintain their head size better than male rats for more accurate stereotaxic placements (Ding et al. 2009; Rodd-Henricks et al. 2000a). The estrous cycle was not monitored in the present study. Experiments were performed during the dark phase. Protocols used were approved by the Institutional Animal Care and Use Committee of Indiana University School of Medicine. All experiments were performed in accordance with the principles outlined in the Guide for the Care and Use of Laboratory Animals (National Research Council, 2011).

Chemical agents: The artificial cerebrospinal fluid (aCSF) consisted of 120 mM NaCl, 4.8 mM KCl, 1.2 mM KH_2PO_4 , 1.2 mM MgSO_4 , 25 mM NaHCO_3 , 1.2 mM CaCl_2 , 10 mM d-glucose, pH 7.2-7.4. Ethyl alcohol (190 proof) was obtained from Decon Laboratories, Inc, (King of Prussia, PA). The GABA_A receptor antagonist bicuculline, the GABA_B receptor antagonist SCH 50911 (Bolser et al. 1995), and the 5-HT₃ receptor antagonist zacopride were obtained from Tocris (Ellisville, MO). All chemicals were dissolved in the aCSF solution to the desired concentrations prior to use.

Stereotaxic surgery procedures: Rats were implanted unilaterally with a 22-gauge cannula (Plastics One Inc., Roanoke, VA) aimed at the right NACsh (AP + 1.7 mm, ML + 2.4 mm, DV – 7.0 mm), as described previously (Engleman et al. 2009). Stylets were inserted into cannulae when no experiments were being conducted. Rats were individually housed after

surgery and were allowed to recover from surgery for at least 7 days prior to tests. Rats were habituated and handled on a daily basis during the 7-day period.

Intracranial self-administration (ICSA) procedure: The ICSA tests followed procedures previously described (Ding et al. 2009; Engleman et al. 2009). Briefly, rats were placed into standard two-lever operant chambers on test days. One active lever was connected to an isolated pulse stimulator (A-M Systems, Inc, Carlsborg, WA) controlled by an operant control system Graphic State 3.02 (Coulbourn Instruments, Allentown, PA). The pulse stimulator was connected to two electrodes that were immersed in a drug-filled tank equipped with a 28-gauge injection cannula inserted into the NACsh. Each response on the active lever (FR1 reinforcement schedule) produced a pulse infusion of 100 nl of solution into the NACsh during a 5-sec period. Each infusion was followed by a 5-sec timeout period. During both the infusion and timeout periods, responses on the active lever were recorded but did not produce further infusions. The responses on the inactive lever were recorded but did not result in any infusions. The assignment of active and inactive lever was counterbalanced among rats.

There were a total of 7 sessions conducted every other day with the duration of each session being 4 hr. The first four sessions were acquisition sessions with 150 mg% ethanol available for self-infusion for each rat. *This concentration of ethanol is physiologically relevant and can be readily achieved with free-choice voluntary drinking by P rats (Murphy et al., 1986). In addition, this concentration of ethanol was demonstrated to be the optimal concentration for self-infusion into the NACsh by P rats (Engleman et al. 2009).* Sessions 5 and 6 were co-infusion sessions, during which 150 mg% ethanol plus one concentration of the GABA_A antagonist bicuculline (10 or 100 μ M), the GABA_B receptor antagonist SCH 50911 (50, 75 or 100 μ M), or the 5-HT₃ receptor antagonist zacopride (10 or 100 μ M) was self-infused. Session 7 was a

session with the return of self-infusion of only 150 mg% ethanol. This procedure has been carried out in the past to examine receptor mechanisms involved in ethanol ICSCA within the posterior VTA (Ding et al. 2012; Rodd-Henricks et al. 2003). *In addition, the Engleman et al. study (2009) showed that replacement of ethanol with aCSF during sessions 5 & 6 resulted in extinction of lever responding. Therefore, such a group was not included in the current study.*

Histology: At the end of the experiments, rats were euthanized and bromophenol blue was injected into the NACsh, as described previously (Engleman et al. 2009). Brains were quickly removed and frozen at -20 °C. Brain sections (40 µm thick) were sliced in a cryostat microtome and stained with cresyl violet for the verification of the placement of injection sites according to the rat brain atlas of Paxinos & Watson (Paxinos and Watson 1998).

Statistical analysis: For each group, lever discrimination was determined by ‘lever (active vs inactive) x session’ mixed ANOVA with repeated measures on session; t-tests were used to compare responses between active and inactive lever following a significant main effect ($p < 0.05$). For determination of drug effect, paired t-tests were used to compare the responses on the active lever and number of infusions during sessions 5 and 6 to average responses on the active lever and number of infusions during acquisition sessions 3 and 4.

Results

Figure 1 depicts the representative non-overlapping placements of injection sites within the NACsh. A majority of injection sites were located in the medial and ventral portions of the NACsh. Only rats with correct placements in the NACsh were included in analysis. *There were six rats with injection sites ventral to the NACsh (Fig. 1). These rats displayed low responses on active and inactive levers (8 ± 2 vs 5 ± 1 responses/session, respectively), as well as low*

infusions per session (5 ±1). These results, along with the previous study (Engleman et al., 2009), suggest that adjacent areas outside the NACsh do not support the self-infusion of ethanol.

Lever responses for the bicuculline co-infusion groups are shown in Fig. 2. The ‘lever x session’ mixed ANOVAs indicated that the control group (Fig. 2, upper panel) demonstrated a significant effect of lever ($F_{1,12} = 20.3, p = 0.001$) and interaction ($F_{6,72} = 2.8, p = 0.02$), but no effect of session ($F_{6,72} = 1.3, p = 0.28$). Lever discrimination was observed during sessions 3 through 7. The 10 μM bicuculline group demonstrated significant effects of session ($F_{6,60} = 5.3, p < 0.001$), lever ($F_{1,10} = 21.2, p = 0.001$) and interaction ($F_{6,60} = 2.5, p = 0.03$). Lever discrimination was observed during each session. Microinjection of 10 μM bicuculline did not significantly reduce responses on the active lever (Fig. 2, middle panel, *e.g., responses in session 6 vs average during sessions 3 & 4: 36 ± 6 vs $42 \pm 5, t_5 = 0.9, p = 0.39$*) or number of infusions (Table 1) during either treatment session. The significant effects of session and interaction resulted mainly from the significant difference in responses on the active lever between session one and the other sessions. The 100 μM bicuculline group (Fig. 2, bottom panel) demonstrated significant effects of session ($F_{6,60} = 11.7, p < 0.001$), lever ($F_{1,10} = 21.2, p = 0.001$), and interaction ($F_{6,60} = 9.0, p < 0.001$). Lever discrimination was evident during sessions 1 to 4, and session 7. Co-infusion of 100 μM bicuculline with ethanol significantly depressed responses on the active lever (*e.g., responses in session 6 vs average during sessions 3 & 4: 13 ± 3 vs $35 \pm 2, t_5 = 7.4, p = 0.001$*) and number of infusions (Table 1) during both treatment sessions. Responses on the active lever during session 7 returned to levels observed during the acquisition sessions.

Lever responses in the SCH 50911 co-infusion groups are shown in Fig. 3. The 50 μM SCH 50911 group (Fig. 3, upper panel) demonstrated significant effects of session ($F_{6,60} = 4.7, p$

= 0.001), lever ($F_{1, 10} = 13.8$, $p = 0.004$) and interaction ($F_{6, 60} = 2.5$, $p = 0.03$). Lever discrimination was observed during sessions 3 to 5, and 7. SCH 50911 at 50 μM slightly reduced active responses during session 6. The 75 μM SCH 50911-treated group demonstrated significant effects of session ($F_{6, 60} = 3.8$, $p = 0.003$), lever ($F_{1, 10} = 8.3$, $p = 0.02$) and interaction ($F_{6, 60} = 5.3$, $p < 0.001$). Lever discrimination was observed during sessions 3, 4, and 7. Co-infusion of 75 μM SCH 50911 with ethanol significantly depressed responses on the active lever during both co-infusion sessions (Fig. 3 middle panel, *e.g.*, *responses during session 6 vs average during sessions 3 & 4: 13 ± 2 vs 33 ± 4 , $t_5 = 5.3$, $p = 0.003$*). The number of infusions was also reduced (Table 1). The 100 μM SCH 50911 group (Fig. 3, bottom panel) demonstrated significant effects of session ($F_{6, 60} = 2.9$, $p = 0.02$), lever ($F_{1, 10} = 18.9$, $p = 0.001$), but no significant effect of interaction ($F_{6, 60} = 0.8$, $p = 0.79$). Lever discrimination was observed during sessions 3, 4 and 7. Co-infusion of 100 μM SCH 50911 did not significantly alter responses on the active lever during either co-infusion session (*e.g.*, *responses during session 6 vs average during sessions 3 & 4: 36 ± 3 vs 49 ± 10 , $t_5 = 1.2$, $p = 0.29$*). However, co-infusion of 100 μM SCH 50911 disrupted lever discrimination during both treatment sessions, as a result of a trend to increase responses on the inactive lever (Fig. 3, bottom panel).

Lever responses in the zacopride co-infusion groups are shown in Fig. 4. The 10 μM zacopride group (Fig. 4, upper panel) demonstrated significant effects of lever ($F_{1, 12} = 15.58$, $p = 0.002$), but no significant effect of session ($F_{6, 72} = 1.1$, $p = 0.35$) or interaction ($F_{6, 72} = 0.15$, $p = 0.99$). Lever discrimination was observed during sessions 1 to 4, and session 6. Co-infusion of 10 μM zacopride did not significantly alter responses on the active lever (*e.g.*, *responses during session 6 vs average during sessions 3 & 4: 27 ± 5 vs 32 ± 2 , $t_6 = 1.0$, $p = 0.34$*) or number of infusions (Table 1) in either co-infusion session. The 100 μM zacopride group demonstrated

significant effects of session ($F_{6,60} = 5.8, p < 0.001$), lever ($F_{1,10} = 32.3, p < 0.001$) and interaction ($F_{6,60} = 6.3, p < 0.001$). Lever discrimination was observed during sessions 2, 3, 4 and 7. Co-infusion of 100 μ M zacopride with ethanol significantly decreased responses on the active lever (Fig. 4, bottom panel, *e.g., responses during session 6 vs average during sessions 3 & 4: 16 ± 2 vs $34 \pm 3, t_5 = 7.1, p = 0.001$*) and number of infusions (Table 1) during both co-infusion sessions. Responses on the active lever returned to baseline levels during session 7.

Discussion

The major findings of the current study are that co-infusion of antagonists for the GABA_A, GABA_B or 5-HT₃ receptor with ethanol attenuated the responses on the active lever and the self-infusion of ethanol into the NACsh, suggesting that the reinforcing effects of ethanol in the NACsh may be mediated, at least in part, by activating local GABA_A, GABA_B and 5-HT₃ receptors. *None of the antagonists, at any concentration, appeared to reduce responses on the inactive lever. These results suggest that a general effect of motor impairment cannot account for the reduced responses on the active lever.* In addition, the highest concentration of the GABA_B receptor antagonist disrupted lever discrimination mainly by increasing the responses on the inactive lever, which suggests a non-specific effect of the antagonist at this concentration.

The finding with bicuculline is in agreement with previous studies demonstrating that both systemic administration (McBride et al. 1988; Petry 1997; Samson et al. 1987) and *local microinjection of GABA_A receptor antagonists into the NACsh (Hyytia and Koob 1995) impaired oral ethanol self-administration in the rat. Another study (Hodge et al. 1995) also indicated that microinjection of bicuculline into the NAC reduced oral ethanol self-administration, although a detailed histological map of injection sites was not presented. In addition, bicuculline did not*

significantly alter responses on the inactive lever (Fig. 2), suggesting the effects of bicuculline may not involve alteration of locomotor activity. Moreover, a previous study (Znamensky et al. 2001) demonstrated that bilateral microinjection of bicuculline (up to 1.6 mM) into the NACsh did not significantly alter food intake in the rat, providing support that bicuculline is not impairing general motor activity. Taken together, these findings indicate the important role of NACsh GABA_A receptors in modulating the local positive motivational properties of ethanol (Koob 2004). A recent study using a similar ICSS procedure indicated that co-infusion of bicuculline with ethanol did not appear to alter ethanol self-infusion directly into the posterior VTA (Ding et al. 2011). These results suggest that local GABA_A receptors may be differentially involved in ethanol self-infusion between the posterior VTA and NACsh.

In addition, these findings are in line with *in vitro* electrophysiological studies demonstrating that ethanol can potentiate GABA_A receptor-mediated currents (Allan et al. 1987). In the NAC, ethanol may potentiate GABA_A receptor function through both pre- and post-synaptic mechanisms (Weiner and Valenzuela 2006). In a NAC slice preparation, application of ethanol produced a dose-dependent enhancement of GABA_A receptor-mediated currents in a subpopulation of NAC neurons; the optimal concentration of ethanol (44 mM or ~ 200 mg%) was comparable to that used in the current study (150 mg%) and increased GABA_A current amplitude by approximately 40% in about 45% of neurons examined (Nie et al. 2000). Furthermore, ethanol was shown to increase presynaptic GABA release in the NAC slice (Weiner and Valenzuela 2006). Therefore, the net effect of ethanol through the GABA_A receptor on these MSNs may be an enhanced inhibition of their activity and reduce GABA release in the ventral pallidum. In line with this idea, previous findings indicated that ethanol administration decreased spike activity in approximately 30% of NAC neurons, and reduced extracellular GABA levels in

the ventral pallidum (Janak et al. 1999; Kemppainen et al. 2010). *It should be noted that the exact location of ethanol-sensitive GABA_A receptors within the NAC sub-regions remains unknown in these electrophysiological studies due to the lack of detailed histological presentation of electrode placements.* Furthermore, GABA_A receptor subunits were also found in non-GABAergic interneurons and axonal processes converging on the NAC (Schwarzer et al. 2001). Activation of these receptors can modulate release of acetylcholine, GABA and/or dopamine (Aono et al. 2008; Rada et al. 1993; Yoshida et al. 1997), and differentially alter the activity of MSNs. For example, activation of these receptors on GABAergic terminals may inhibit GABA release, resulting in attenuated inhibition of post-synaptic MSNs. Indeed, ethanol was shown to decrease GABA_A-mediated currents and increase the excitability of a subpopulation of NAC neurons (Janak et al. 1999; Nie et al. 2000). Moreover, the sensitivity of GABA_A receptors to ethanol enhancement depends on the subunit compositions of the GABA_A receptor (Kumar et al. 2009), which are heterogeneously expressed within the NAC (Schwarzer et al. 2001). Therefore, the action of ethanol on MSNs depends on micro-circuits within the NAC that may differ in the relative location of GABA_A receptors and their subunit composition. Taken together, it is possible that multiplex mechanisms may contribute to the self-infusion of ethanol and the effects of bicuculline on ethanol self-infusion into the NACsh.

Similar to GABA_A receptors, activation of local GABA_B receptors appeared to mediate the reinforcing effects of ethanol within the NACsh, as indicated by the results that 75 μ M SCH 50911 attenuated ethanol self-infusion. This finding is consistent with *in vitro* evidence that the GABA_B receptor-mediated inhibitory postsynaptic potential is sensitive to ethanol potentiation (Federici et al. 2009). The highest concentration of SCH 50911 disrupted lever discrimination between the active and inactive levers due in part to increased responding on the inactive lever

(Fig. 3, bottom panel). These results suggest possible non-specific actions of SCH 50911. The mechanism underlying these effects are unknown, but may be due to different subtypes of GABA_B receptors, GABA_B receptors being located on both pre- and post-synaptic membrane, and multiple actions of SCH 50911 occurring at the highest concentration. For example, a high concentration of SCH 50911 was shown to act as an inverse agonist for the GABA_B receptor to increase cAMP production in a cell culture (Grunewald et al. 2002).

In addition to the GABA receptors, activation of local 5-HT₃ receptors also appeared to be involved in mediating the reinforcing effects of ethanol, as suggested by the findings that co-infusion of zacopride with ethanol dampened ethanol self-infusion into the NACsh. *A previous study (Herges & Taylor 2000) showed that antagonism of 5-HT₃ receptors within the NAC with ondansetron did not significantly alter basal locomotor activity. In addition, zacopride did not appear to alter responses on the inactive lever in the current study (Fig. 4). All these findings suggest that the effects of zacopride may not involve impairment of locomotor activity.* This finding is in line with electrophysiological findings that ethanol can potentiate 5-HT₃ receptor mediated currents *in vitro* (Lovinger and White 1991; Machu and Harris 1994). Our results provide mechanistic evidence supporting the involvement of activation of NACsh 5-HT₃ receptors in mediating systemic ethanol self-administration in rodents (Engleman et al. 2008; Knapp and Pohorecky 1992; McKinzie et al. 2000; Rodd-Henricks et al. 2000b). Interestingly, local 5-HT₃ receptors within the VTA also seem to be involved in the reinforcing and rewarding effects of ethanol (Rodd-Henricks et al. 2003; Rodd et al. 2010). These studies suggest a close relationship between ethanol self-administration and 5-HT₃ receptors within the mesolimbic system.

Autoradiographic studies indicated that low to moderate densities of 5-HT₃ receptor

binding sites were present in the NAC (Barnes et al. 1990; Ge et al. 1997; Kilpatrick et al. 1987). However, the relative cellular location and distribution in *each NAC sub-region* of these receptors remains unclear. *In situ* hybridization studies (Morales et al. 2004; Tecott et al. 1993) did not detect 5-HT₃ receptor mRNA in the NAC, which agrees with an immunocytochemistry study showing that very low levels of 5-HT₃ receptor-expressing cells were detected in the NAC (Morales et al. 1998). These studies suggest that these receptors may be preferentially localized on synaptic terminals as opposed to NAC MSNs. One likely possibility is on dopaminergic terminals. Microdialysis studies demonstrated that local perfusion of a 5-HT₃ receptor agonist could increase basal and ethanol-evoked dopamine release in the NAC, whereas antagonizing local 5-HT₃ receptors could attenuate ethanol-induced dopamine release in the NAC (Campbell and McBride 1995). In addition, systemic administration of 5-HT₃ receptor antagonists was shown to attenuate ethanol-induced dopamine release in the NAC (Carboni et al. 1989; Wozniak et al. 1990). These findings suggest that 5-HT₃ receptors on dopaminergic terminals may contribute to ethanol self-infusion and the effects of zacopride observed in the current study.

In summary, the current study demonstrated that antagonism of local GABA_A, GABA_B or 5-HT₃ receptors attenuated ethanol self-infusion into the NACsh, suggesting that the local reinforcing effects of ethanol in the NACsh may be mediated by activation of these receptors. These results are consistent with previous findings and suggest these receptors within the NACsh may contribute to the global reinforcing and rewarding effects of ethanol. *On the other hand, it remains unknown whether there are possible interactions among these different receptors and how they interact in mediating ethanol reinforcement, which warrants future studies.*

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Declaration of conflicting interests

The authors declare that there is no conflict of interest.

Figure legend

Figure 1: Representative placements of microinjection sites in the nucleus accumbens shell for the self-infusion of ethanol. For clarity purpose, the overlapping sites are not included. Circles represent sites within the nucleus accumbens shell and squares represent sites ventral to the nucleus accumbens shell.

Figure 2: Effects of co-infusion of aCSF, 10 or 100 μ M bicuculline (a GABA_A receptor antagonist) with 150 mg% ethanol on lever responses for the intracranial self-infusion of ethanol into the nucleus accumbens shell (n = 6-7 / group). * p < 0.05, significantly higher responses than those on the inactive lever. # p < 0.05, significantly lower responses than the average responses on the active lever during acquisition sessions 3 and 4.

Figure 3: Effects of co-infusion of 50, 75 or 100 μ M SCH 50911 (a GABA_B receptor antagonist) with 150 mg% ethanol on lever responses for the intracranial self-infusion of ethanol into the nucleus accumbens shell (n = 6 / group). * p < 0.05, significantly higher responses than those on the inactive lever. # p < 0.05, significantly lower responses than the average responses on the active lever during acquisition sessions 3 and 4.

Figure 4: Effects of co-infusion of 10 or 100 μ M zacopride (a 5-HT₃ receptor antagonist) with 150 mg% ethanol on lever responses for the intracranial self-infusion of ethanol into the nucleus accumbens shell (n = 6-7 / group). * p < 0.05, significantly higher responses than those on the inactive lever. # p < 0.05, significantly lower responses than the average responses on the active lever during acquisition sessions 3 and 4.

Table 1: Effects of co-infusion of an antagonist for the GABA_A, GABA_B or 5HT₃ receptor with 150 mg% ethanol on number of infusions into the NACsh

Group	Sessions 3 & 4	Session 6	t value	p value
150 mg% EtOH + aCSF	24 ± 3	25 ± 3	0.3	0.76
150 mg% ethanol + 10 μM bicuculline	23 ± 4	22 ± 4	0.4	0.72
150 mg% ethanol + 100 μM bicuculline	17 ± 1	8 ± 2	5.6	0.002
150 mg% ethanol + 50 μM SCH 50911	21 ± 4	19 ± 3	1.2	0.29
150 mg% ethanol + 75 μM SCH 50911	18 ± 1	8 ± 1	8.0	< 0.001
150 mg% ethanol + 100 μM SCH 50911	22 ± 2	32 ± 5	1.6	0.16
150 mg% ethanol + 10 μM zacopride	19 ± 2	14 ± 3	1.4	0.22
150 mg% ethanol + 100 μM zacopride	17 ± 1	10 ± 1	17.0	< 0.001

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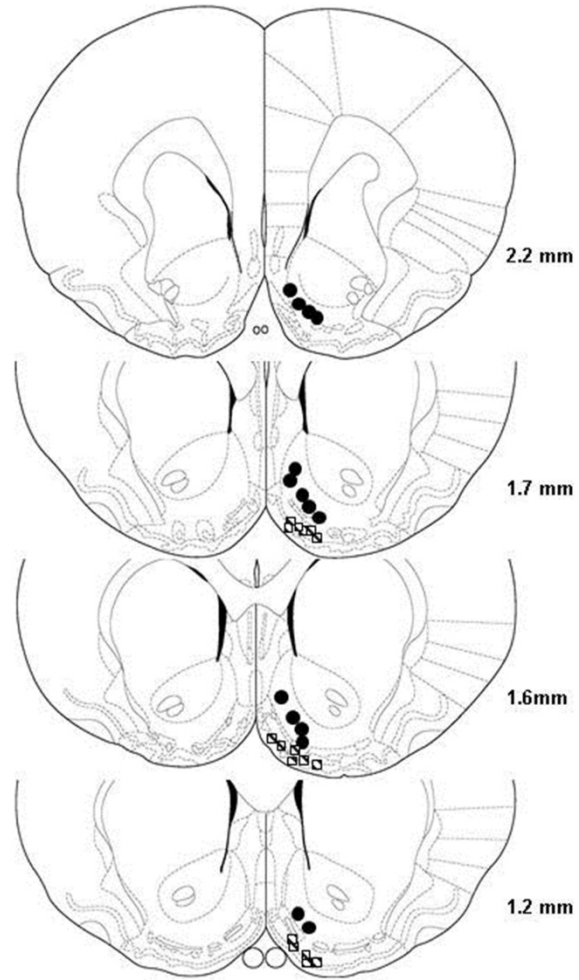


Figure 1

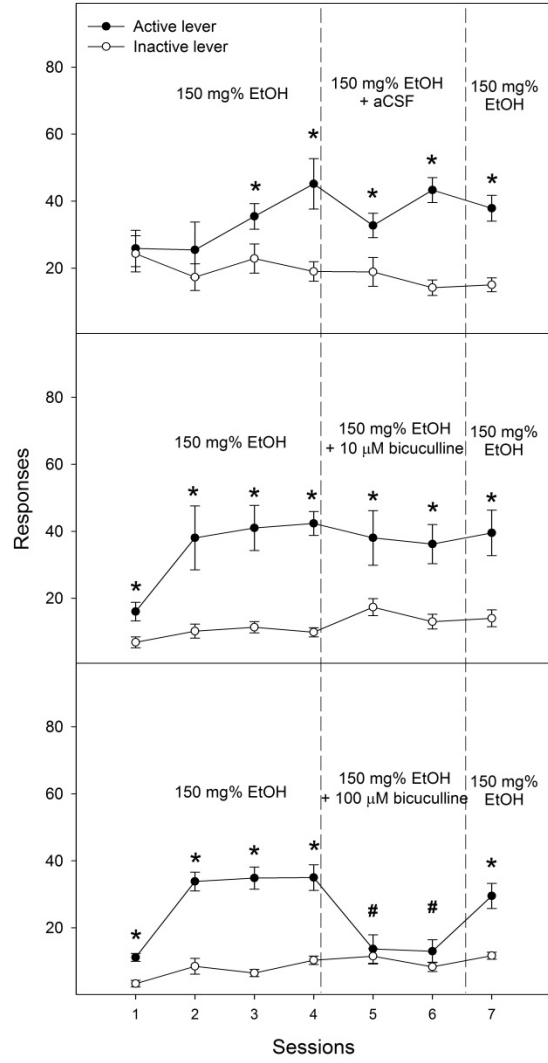


Figure 2

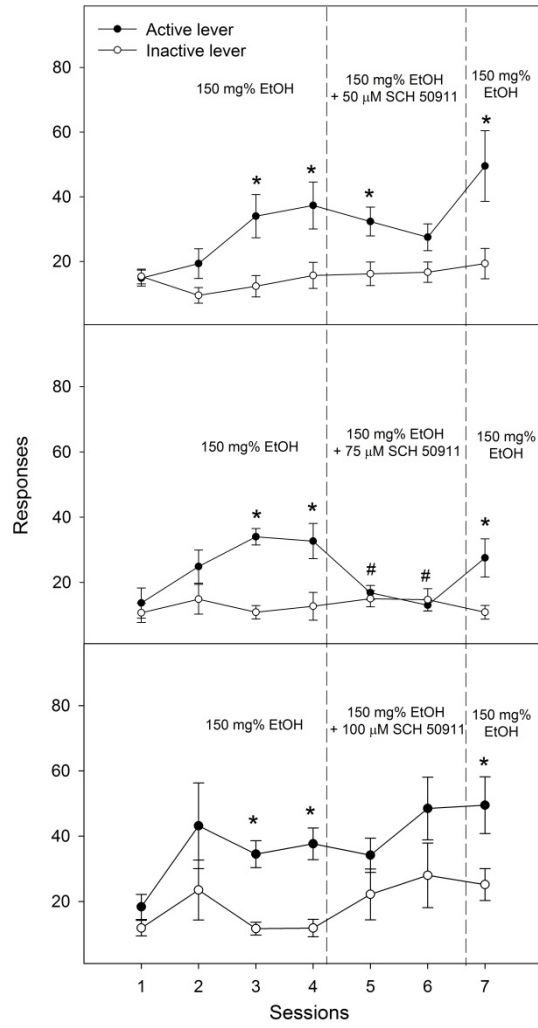


Figure 3

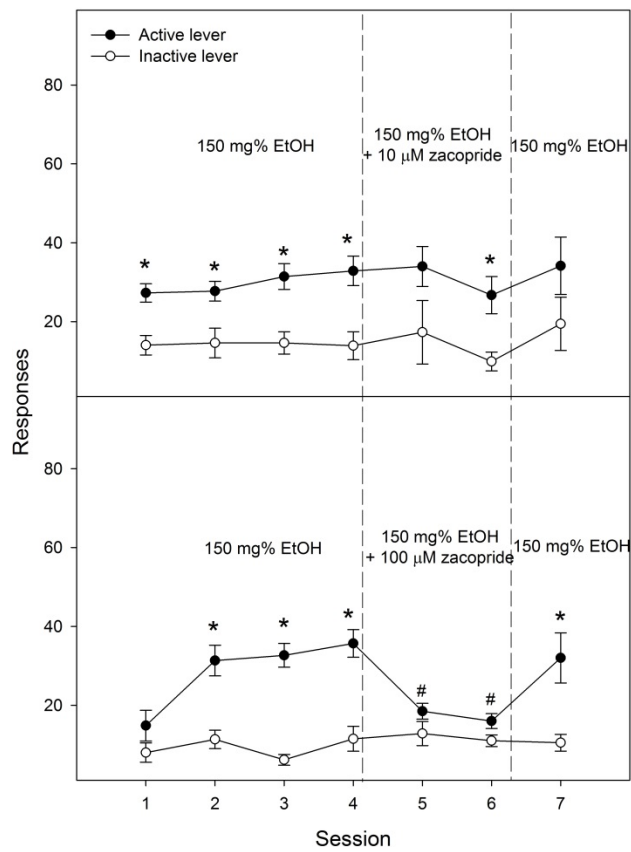


Figure 4