

Ethanol Enhances Glutamate Transmission by Retrograde Dopamine Signaling in a Postsynaptic Neuron/Synaptic Bouton Preparation From the Ventral Tegmental Area

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It is well documented that somatodendritically released dopamine is important in the excitability and synaptic transmission of midbrain dopaminergic neurons. Recently we showed that in midbrain slices, acute ethanol exposure facilitates glutamatergic transmission onto dopaminergic neurons in the ventral tegmental area (VTA). The VTA is a brain region critical to the rewarding effects of abused drugs, including ethanol. We hypothesized that ethanol facilitation might result from an increase in somatodendritically released dopamine, which acts retrogradely on dopamine D₁ receptors on glutamate-releasing axons and consequently leads to an increase in glutamate release onto dopaminergic neurons. To further test this hypothesis and to examine whether ethanol facilitation can occur at the single-cell level, VTA neurons were freshly isolated from rat brains using an enzyme-free procedure. These isolated neurons retain functional synaptic terminals, including those that release glutamate. Spontaneous excitatory postsynaptic currents (sEPSCs) mediated by glutamate α -amino-3-hydroxy-5-methylisoxazole-4-propionic acid receptors were recorded from these freshly isolated putative dopaminergic neurons. We found that acute application of clinically relevant concentrations of ethanol (10–80 mM) significantly facilitated the frequency of sEPSCs but not their mean amplitude. Ethanol facilitation was mimicked by the D₁ agonist SKF 38393 and by the dopamine uptake blocker GBR 12935 but was blocked by the D₁ antagonist SKF 83566, and by depleting dopamine stores with reserpine, as well as by chelating postsynaptic calcium with BAPTA. Furthermore, the sodium channel blocker tetrodotoxin eliminated the facilitation of sEPSCs induced by ethanol but not by SKF 38393. These results constitute the first evidence from single isolated cells of ethanol facilitation of glutamate transmission to dopaminergic neurons in the VTA. In addition, we show that ethanol facilitation has a postsynaptic origin and a presynaptic locus. Furthermore, ethanol stimulation of a single dopaminergic neuron is capable of eliciting the release of somatodendritic dopamine, which is sufficient to influence glutamatergic transmission at individual synapses.

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INTRODUCTION

In the ventral tegmental area (VTA) and substantia nigra, dopaminergic (DAergic) neurons can release dopamine somatodendritically (Bjorklund and Lindvall, 1975; Geffen *et al*, 1976; Cheramy *et al*, 1981). Dopamine acts on two families of receptors, D₁-like (consisting of dopamine D₁ and D₅ receptors) and D₂-like (D₂, D₃, and D₄ receptors). D₁ and D₂ receptors (D₁R and D₂R) exert opposite effects on the same signaling pathway, with D₁Rs positively and D₂Rs negatively affecting adenylyl cyclase. It has been well

documented that somatodendritic dopamine is important in the excitability of DAergic neurons through the activation of D₂ autoreceptors in the postsynaptic membrane of the DAergic neurons (Aghajanian and Bunney, 1977; Cragg and Greenfield, 1997; Tepper *et al*, 1997). Nevertheless, somatodendritic dopamine can also function as a rapid, retrograde signal that activates dopamine receptors on synaptic terminals and regulates synaptic transmission (Abarca *et al*, 1995; Koga and Momiyama, 2000).

Excitatory synaptic input mediated by glutamate is a key component in the regulation of DAergic cell excitability (Overton and Clark, 1997). D₁Rs are situated on glutamatergic synaptic terminals on VTA DAergic neurons but not on the postsynaptic membrane of the DAergic neurons (Lu and Hagg, 1997). A previous *in vivo* study demonstrated that stimulation of the D₁Rs increases glutamate release in the VTA (Kalivas and Duffy, 1995). However, it is unknown whether dopamine released from the somatodendritic area

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of an isolated cell can stimulate the D₁Rs on the glutamatergic synapses on the same DAergic neurons.

The mechanisms underlying alcohol addiction are unclear, but it has been well documented that the glutamatergic and DAergic systems may be critical. Ethanol suppresses glutamatergic transmission in several brain regions (Siggins *et al*, 2005). Nevertheless, in a recent study in midbrain slices, we found that acute ethanol facilitates glutamatergic transmission in the VTA, a key region in the brain reward system (Xiao *et al*, 2008). We hypothesize that ethanol facilitation might be mediated by somatodendritically released dopamine, which retrogradely activates the D₁Rs on the glutamate-releasing terminals on the DAergic neurons to increase glutamate release. However, it is unknown whether ethanol facilitation can occur at the single cell level.

To determine whether ethanol facilitation can occur at the single cell level, we took advantage of a highly reduced preparation consisting of an isolated postsynaptic neuron from the VTA and attached glutamatergic presynaptic terminals. We examined the effects of ethanol on spontaneous excitatory postsynaptic currents (sEPSCs) mediated by glutamate α -amino-3-hydroxy-5-methylisoxazole-4-propionic acid (AMPA) receptors and demonstrated that ethanol at pharmacologically relevant concentrations (10–80 mM) significantly increased sEPSC frequency in the single cell preparation.

MATERIALS AND METHODS

Slice Preparation and Mechanical Dissociation

The midbrain slices were prepared as described previously by Ye *et al* (2004, 2006). Sprague–Dawley rats (aged 10–20 postnatal days) were anesthetized and then killed by decapitation. Coronal midbrain slices (300 μ m thick) were cut using a VF-200 slicer (Precisionary Instruments Inc., Greenville, NC). Slices were prepared in an ice-cold glycerol-based artificial cerebrospinal fluid (GACSF) containing 250 mM glycerol, 1.6 mM KCl, 1.2 mM NaH₂PO₄, 1.2 mM MgCl₂, 2.4 mM CaCl₂, 25 mM NaHCO₃, and 11 mM glucose, and saturated with 95%O₂/5%CO₂ (carbogen; Ye *et al*, 2006). Slices (two per animal) were allowed to recover for at least 1 h in a holding chamber at room temperature (22–24°C) in carbogen-saturated regular artificial cerebrospinal fluid, which has the same composition as GACSF, except that glycerol was replaced by 125 mM NaCl.

Neurons with functional presynaptic terminals attached were obtained by mechanical dissociation (Figure 1a and b), as previously described (Akaike and Moorhouse, 2003), with some modifications (Ye *et al*, 2004, 2006). Briefly, a midbrain slice was transferred to a 35 mm culture dish (Falcon, Rutherford, NJ, USA) and held down with a flat U-shape wire. The dish was filled with standard external solution containing 140 mM NaCl, 5 mM KCl, 2 mM CaCl₂, 1 mM MgCl₂, 10 mM HEPES, and 10 mM glucose (320 mOsm, pH adjusted to 7.3 with Tris base). Under an inverted microscope (Nikon Diaphot, Tokyo, Japan), VTA was identified. A heavily fire-polished pipette, lightly touched the surface of the VTA, and vibrated horizontally at 15–20 Hz for 2–5 min using a homemade device. Finally, the slice was removed. After 20 min, isolated neurons that

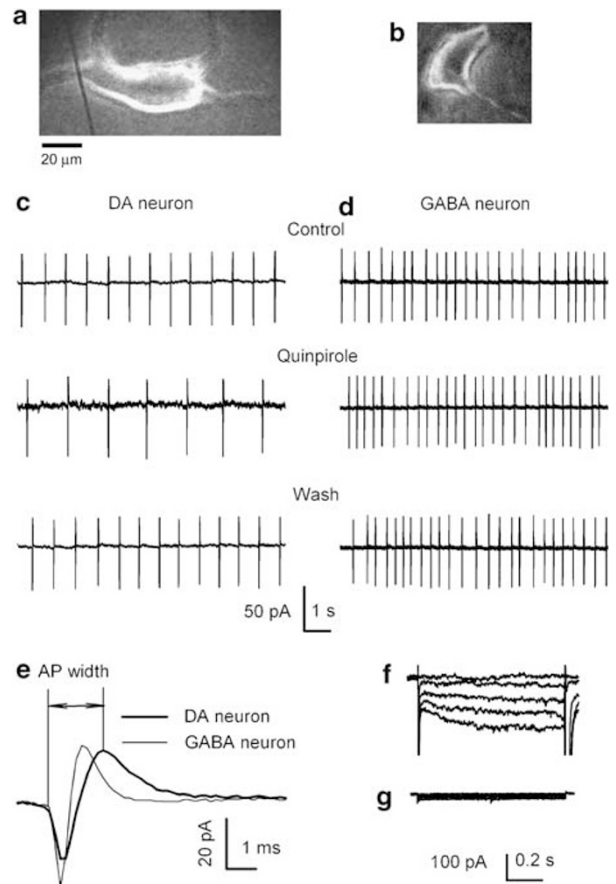


Figure 1 Identification of DAergic neurons. Photomicrograph of a mechanically dissociated putative DAergic neuron (a) and a putative GABAergic neuron (b). The much reduced dendritic arbors of such cells facilitate space clamp. Spontaneous firings recorded with loose-patch cell-attached mode showing the inhibitory response of the DAergic neuron (c) and the no response of the GABAergic neuron (d) to the D_{2/3} receptor antagonist, quinpirole. (e) Comparison of single spontaneous action potentials recorded from a putative DAergic neuron (thicker line, with a half-width of 1.7 ms) and a putative GABAergic neuron (thinner line, with a half-width of 1.1 ms). Under voltage clamp, the currents induced in such a DAergic cell by a series of hyperpolarizing voltage pulses (the 10 mV steps to -110 mV) show the prominent I_h of the DAergic (f) but not in the GABAergic (g) neuron.

adhered to the bottom of the dish were used for electrophysiological recordings. These mechanically dissociated neurons often preserved some functional nerve terminals, including those that release glutamate (Yamanaka *et al*, 2002; Akaike and Moorhouse, 2003; Ye *et al*, 2004; Zhu and Lovinger, 2005; Jang *et al*, 2006).

Electrophysiological Recordings

Electrical signals were obtained in whole-cell and loose-patch cell-attached patch clamp configurations with an Axon 200B amplifier (Molecular Devices Co., Union City, CA, USA), a Digidata 1320A A/D converter (Molecular Devices Co.), and pCLAMP 9.2 software (Molecular Devices Co.). Data were filtered at 1 kHz and sampled at 5 kHz. The patch electrodes had a resistance of 2–5 M Ω when filled with the pipette solution containing: 135 mM CsF, 5 mM KCl, 2 mM MgCl₂, 10 mM HEPES, 2 mM Mg ATP, and 0.2 mM

GTP. The pH of the pipette solution was adjusted to 7.2 with Tris base. All sEPSCs were recorded in the presence of 10 μ M bicuculline at a holding potential (V_H) of -65 mV.

Chemicals and Applications

The chemicals, including bicuculline, 6,7-dinitroquinoxaline-2,3-dione (DNQX), reserpine, tetrodotoxin (TTX), [D-Ala², N-Me-Phe⁴, Gly⁵-ol] enkephalin (DAMGO) were obtained from Sigma-Aldrich Chemical Company (St Louis, MO, USA). (–)-Quinpirole hydrochloride, GBR 12935, N-allyl-(\pm)-SKF 38393, and (\pm)-SKF 83566 were from Tocris Bioscience (Ellisville, MO, USA). Ethanol (95%, prepared from grain) was from Pharmco (Brookfield, CT, USA) and stored in glass bottles. Drugs were added to the superfusate applied to the cell using a fast perfusion system (Y tube). Solutions in the vicinity of a neuron can be completely exchanged within 40 ms without damaging the seal (Zhou *et al*, 2006).

Data Analysis

sEPSCs and miniature excitatory postsynaptic currents (mEPSCs) were counted and analyzed with Clampfit 9.2 (Molecular Devices Co.); sEPSCs and mEPSCs were screened automatically (6 pA amplitude threshold), checked visually, and accepted or rejected according to their rise and decay times. The frequency and amplitude of all events, during and after drug application, were normalized to the mean of the values observed during the initial control period. Cumulative probability plots of the incidences of various interevent intervals and amplitudes, recorded under different conditions from the same neuron, were compared using the Kolmogorov–Smirnov (K–S) test. For other plots, data obtained over a 0.5–2 min period at the peak of a drug response were normalized to the average values of frequency (or amplitude) of the sEPSC (or mEPSC) recorded during the initial control period (1–4 min). Data are expressed as means (\pm SEM). The statistical significance of drug effects was assessed by a paired two-tailed *t*-test on normalized data. For effects of multiple drug concentrations an appropriate one-way analysis of variance (ANOVA) was used. Statistical analyses were performed with SigmaStat (SPSS, Chicago, IL). Values of $p < 0.05$ were considered significant.

RESULTS

Identification of VTA DAergic Neurons

DAergic neurons in the VTA were identified on the basis of their well-established pharmacological and electrophysiological properties (Lacey *et al*, 1989; Zhou *et al*, 2006; Xiao *et al*, 2007, 2008). Figure 1 shows sample traces of spontaneous discharges of a putative DAergic neuron (Figure 1c) and a putative GABAergic neuron (Figure 1d) recorded in the cell-attached mode. Although the dopamine D₂/D₃ receptor agonist quinpirole inhibited ongoing discharges of the putative DAergic neurons, the μ -opioid receptor agonist Tyr-D-Ala-Gly-N-Me-Phe-Gly-ol enkephalin (DAMGO) had no effect (data not shown). In contrast, quinpirole had no effect on the ongoing discharges of the

GABAergic neuron (Figure 1d). The spontaneous pacemaker (1–5 Hz) firing with spikes exhibiting action potential widths of ≥ 1.2 ms (Figure 1e, measured in cell-attached mode from the initial inward current to the peak of the outward current) is a marker of tyrosine hydroxylase expressing principle cells (Ford *et al*, 2006; Xiao *et al*, 2007, 2008). In addition, DAergic neurons exhibited a prominent hyperpolarization-activated inward current (I_h) in response to a series of voltage steps from -70 to -110 mV (with decrement of 10 mV) when recorded under whole-cell voltage clamp conditions (Figure 1f), whereas VTA GABAergic neurons have no prominent I_h (Figure 1g; Jones and Kauer, 1999). These are characteristic electrophysiological properties of DAergic neurons. The following experiments were carried out on putative DAergic neurons identified according to the aforementioned characteristics.

The identification of VTA DAergic neurons by their responses to quinpirole and the expression of I_h has been challenged: I_h is present in $\sim 60\%$ of VTA DAergic neurons (Margolis *et al*, 2006). Nevertheless, as our result relies on somatodendritically released dopamine, it is highly unlikely that the recordings are from nonDAergic neurons, because of the fact that there is no DAergic axonal projection in the VTA (Tepper *et al*, 1987; Adell and Artigas, 2004).

Ethanol Raises sEPSC Frequency in Mechanically Dissociated DAergic Neurons

In a recent study in midbrain slices, we found that clinically relevant concentrations of ethanol (10–80 mM) increased glutamatergic transmission (Xiao *et al*, 2008). To determine whether this effect could be observed in isolated neurons, we repeated the experiments in mechanically dissociated neurons ('nerve-bouton' preparations). Such preparations combine several advantages: good space clamp, preservation of some functioning synaptic terminals, including those that release glutamate as well as better control of the surrounding solution (Akaike and Moorhouse, 2003; Ye *et al*, 2004; Zhu and Lovinger, 2005). sEPSCs were recorded in the presence of the GABA_A receptor antagonist bicuculline (10 μ M) at a holding potential of -65 mV. In addition, the use of the CsF-based internal solution in these experiments suppressed GABA_A receptor-mediated postsynaptic currents (Bormann *et al*, 1987). Under these conditions, all the spontaneous synaptic events were reversibly abolished by the antagonist of AMPA/kainate receptor DNQX (10 μ M), indicating that they are AMPA/kainate receptor-mediated sEPSCs (Figure 2a and b).

After obtaining stable control sEPSCs, 40 mM ethanol was applied to the cell using a fast perfusion system (Y tube). In keeping with the results from brain slices, 40 mM ethanol robustly increased the frequency of sEPSCs in the dissociated neurons. This effect, which was reversible upon washout of ethanol, is illustrated in Figure 2a–c (normalized increase by $91 \pm 16\%$; from 1.5 ± 0.4 Hz in control to 2.6 ± 0.5 Hz in ethanol; $n = 5$, $p < 0.001$, by paired *t*-test). It is further illustrated in Figure 2d by the large increase in the incidence of shorter intervals between successive sEPSCs. Additional tests with a range of ethanol concentration indicate that the effect of ethanol on sEPSC frequency was dependent on its concentrations. Specifically, 10, 20, and 80 mM increased the frequency by $29 \pm 10\%$ ($n = 4$,

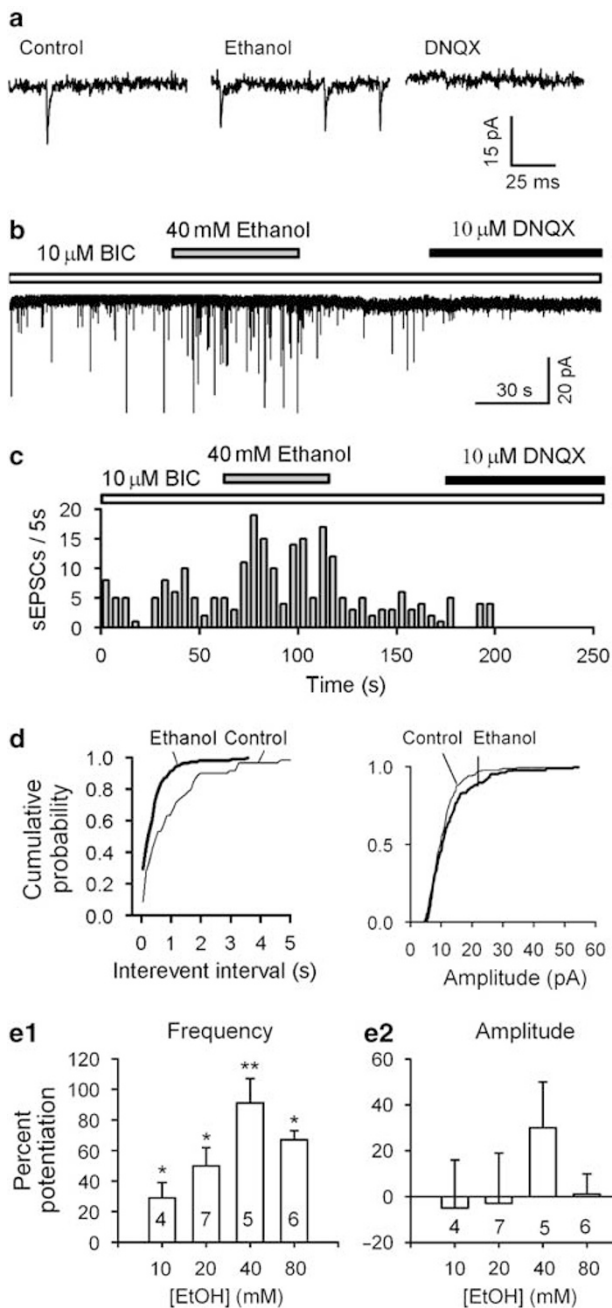


Figure 2 Ethanol enhances frequency of spontaneous excitatory postsynaptic currents (sEPSCs) recorded from mechanically dissociated DAergic neurons. sEPSCs were recorded in the presence of bicuculline (10 μ M) at a V_H of -65 mV. The frequency of sEPSCs was increased by 40 mM ethanol and the sEPSCs were eliminated by 10 μ M DNQX (a–c). (c) Time course of ethanol-induced enhancement of sEPSC frequency in one experiment. (d) Cumulative probability of sEPSC interevent intervals (left panel) and current amplitudes (right panel) of the same data. (e) Dose-dependent potentiation of sEPSC frequency (e₁), but unaltered amplitude (e₂; mean \pm SEM, number of cells in each group is indicated). * $p < 0.05$, ** $p < 0.01$, paired *t*-test for ethanol application vs pre-ethanol control.

$p = 0.02$), $50 \pm 12\%$ ($n = 7$, $p < 0.05$), and $67 \pm 6\%$ ($n = 6$, $p = 0.04$), respectively. In contrast, ethanol did not significantly change sEPSC amplitude. 10, 20, 40, and 80 mM ethanol altered the amplitude of sEPSCs by -5 ± 21 , -3 ± 22 , 30 ± 22 , and $1 \pm 9\%$, respectively. This finding is

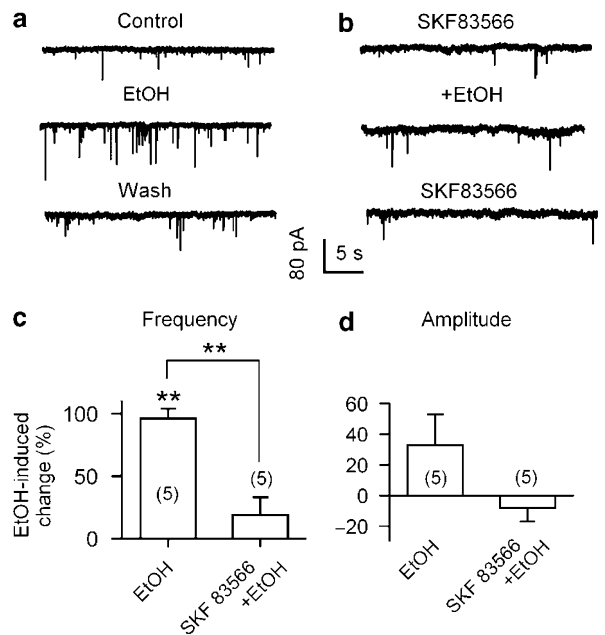


Figure 3 Effect of the D₁R antagonist on ethanol facilitation of sEPSCs. (a) Current traces show that ethanol enhancement of sEPSCs was eliminated by 10 μ M SKF 83566 (D₁R antagonist; b). Summary of ethanol-induced changes (%) in sEPSC frequency (c) and amplitude (d) in the absence (EtOH) and the presence of SKF 83566 (SKF 83566 + EtOH, data from six cells). ** $p < 0.01$, paired *t*-test for ethanol application vs pre-ethanol control.

the first electrophysiological demonstration of the potentiating effect of acute ethanol on glutamatergic transmission at the single cell level.

Effects of Dopamine D₁-Like Antagonist on Ethanol Facilitation of sEPSCs

Systemic administration of ethanol increases somatodendritic dopamine release in the VTA *in vivo* (Campbell *et al.*, 1996). In the VTA, D₁Rs are expressed on glutamatergic axons but not on the soma of DAergic neurons (Lu and Hagg, 1997). Microdialysis data have shown that D₁-like agonists increase the extracellular levels of glutamate in the VTA (Kalivas and Duffy, 1995). To test for the possible involvement of D₁Rs in ethanol-induced enhancement of glutamatergic transmission, we compared the effect of ethanol on sEPSC frequency in the absence and the presence of the D₁ antagonist SKF 83566 (10 μ M). The application of 40 mM ethanol robustly increased sEPSC frequency (Figure 3a). Though ineffective when applied alone, SKF 83566 (10 μ M) suppressed ethanol (40 mM)-induced increase in sEPSC frequency (Figure 3b). Specifically, ethanol (40 mM) alone increased sEPSC frequency by $96 \pm 8\%$ ($n = 5$, $p < 0.01$) and by $19 \pm 11\%$ ($n = 5$, $p = 0.274$) in the absence and the presence of SKF 83566 (10 μ M), respectively (Figure 3c). Ethanol (40 mM) has no significant effect on sEPSC amplitude both in the absence and the presence of SKF 83566 (10 μ M), by $23 \pm 17\%$ ($p = 0.41$, $n = 5$) and $-7 \pm 7\%$ ($p = 0.26$, $n = 5$), respectively (Figure 3d). These results suggest that presynaptic dopamine D₁-like receptors

are essential in ethanol facilitation of glutamatergic transmission.

Dopamine D₁-Like Agonist Increases the Frequency of sEPSCs in Dissociated DAergic Neurons

To further establish the involvement of D₁Rs, we tested the effects of the D₁ agonist SKF 38393 on sEPSCs in dissociated putative DAergic neurons. SKF 38393 (10 μM) significantly increased sEPSC frequency by 106 ± 19% (control, 1.0 ± 0.1 Hz; SKF 38393, 2.0 ± 0.4 Hz; *n* = 5, *p* = 0.01; Figure 4A and B), and shortened intervals between successive sEPSCs (K-S test, *p* < 0.01; Figure 4C). However, there was no significant change in the corresponding plot of sEPSC

amplitudes (by -8 ± 6%, *n* = 5, *p* > 0.3; Figure 4D). The facilitation effect was observed to peak within 30 s. Subsequently the frequency of sEPSCs recovered to or slightly lower than the base line level (Figure 4B). Similar phenomenon was observed in five other neurons tested (see also Figure 4F). We next tested the effect of SKF 38393 on mEPSCs in the presence of TTX (1 μM). SKF 38393 (10 μM) caused a significant increase of mEPSC frequency by 156 ± 63% (basal, 0.8 ± 0.3 Hz, SKF 38393, 2.1 ± 0.3 Hz, *p* < 0.01; *n* = 6), without changing the amplitude (by 13 ± 19%, basal, 13 ± 2 pA, SKF 38393, 13 ± 1 pA, *p* = 0.85; *n* = 6). Similar to that for sEPSCs, the response of mEPSC to SKF 38393 desensitized quickly in the continuous presence of SKF 38393 (Figure 4F). These results provide unique

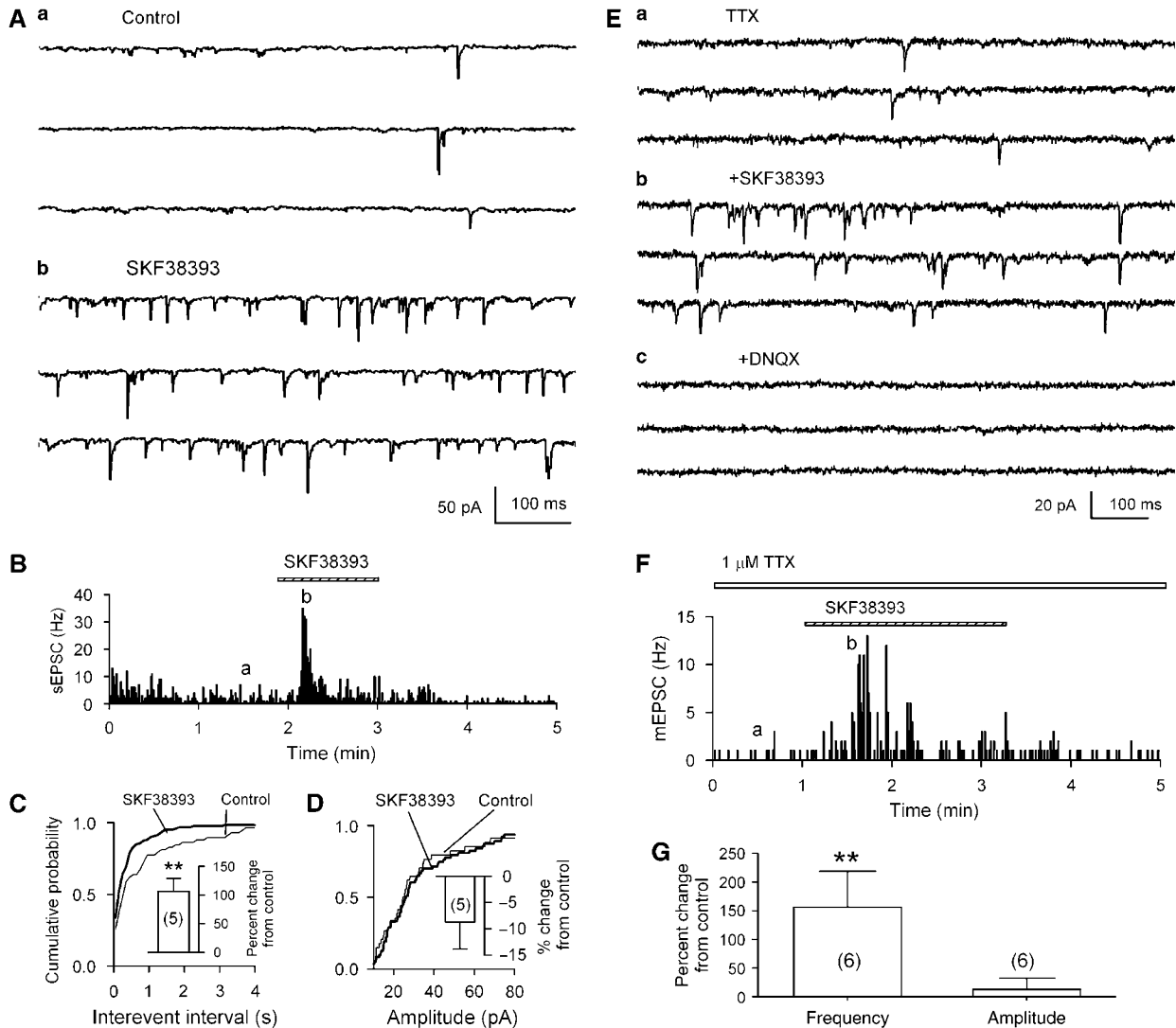


Figure 4 Effect of the D₁R agonist on sEPSCs and mEPSCs in isolated neurons. (A) The D₁R agonist SKF 38393 (10 μM) increased sEPSC frequency in a putative DAergic neuron in the VTA. (B) Time course of changes induced by SKF 38393. The traces in (A) were taken at times indicated by (a) and (b). The change in frequency is also demonstrated by the much increased incidence of shorter intervals between sEPSCs. A representative cumulative probability plot of intervent interval (C), but there was no associated change in sEPSC amplitudes (D). The inset shows mean ± SEM of % changes in sEPSC frequency (C) and amplitude (D) induced by SKF 38393 (10 μM; *n* = 5). ***p* < 0.01, paired *t*-test for SKF 38393 application vs pre-SKF 38393. (E) SKF 38393 (10 μM) increased mEPSC frequency in the presence of TTX (1 μM) in a putative DAergic neuron in the VTA. (F) Time course of changes induced by SKF 38393. The traces in (E) were taken at times indicated by (a) and (b). (G) Summary of results (mean ± SEM of % changes) from six cells showing that SKF 38393 (10 μM) significantly increased mEPSC frequency (left column) but not the amplitude (right column). ***p* < 0.01, paired *t*-test for SKF 38393 application vs pre-SKF 38393.

electrophysiological evidence indicating that functional D₁Rs exist on glutamatergic terminals, which make synapses on the DAergic neurons in the VTA, and that activation of these presynaptic D₁Rs facilitates glutamatergic neurotransmission.

Our results might appear inconsistent with the study in brain slices by Schilström *et al* (2006) reporting that application of the D₁ agonist SKF 81297 (10 μM) caused a slight but significant reduction in mEPSC frequency. However, our hypothesis is that the slightly reduction in mEPSC frequency observed by these investigators might be a consequence of D₁R desensitization (Memo *et al*, 1982; Balmforth *et al*, 1990; Cameron and Williams, 1993), because our data show that even in the isolated cells, in which D₁ agonist application is much more rapid than that in brain slices, D₁R activation induced facilitation of EPSCs desensitized very rapidly. In addition, in the experiments of Schilström *et al* (2006), the effect of SKF 81297 on mEPSCs was determined by the data collected after 10 min of drug application. Thus, the short increase in mEPSC frequency induced by the D₁ agonist, if it exists, might have been overlooked in their study. The latter possibility is supported by our recent study showing that SKF 38393 did increase sEPSC frequency in brain slices (Xiao *et al*, 2008).

Ethanol does not Alter mEPSCs in Mechanically Dissociated DAergic Neurons

Next, we recorded mEPSCs in the presence of the sodium channel blocker tetrodotoxin (TTX, 1 μM) and bicuculline (10 μM). In nerve-bouton preparations, TTX (1 μM) sharply decreased sEPSC frequency to $33 \pm 10\%$ of control (from 3.3 ± 1.1 Hz in control to 1.0 ± 0.5 Hz in TTX; $n = 5$, $p < 0.01$) and moderately reduced the amplitude to $73 \pm 5\%$ of control (from 16 ± 4 pA in control to 12 ± 3 pA in TTX; $n = 5$, $p > 0.05$).

In the presence of TTX (1 μM), the mEPSCs recorded from four nerve-bouton preparations were not sensitive to ethanol, consistent with the observations in brain slices (Xiao *et al*, 2008). Ethanol (40 mM) increased the frequency of mEPSCs by $11 \pm 7\%$ ($p = 0.33$, $n = 4$) and the amplitude by $-7 \pm 8\%$ ($p = 0.19$; $n = 4$; Figure 5A–D).

The effects of TTX suggest that ethanol-induced increase in glutamatergic transmission is not only because of an increase in dopamine released from the postsynaptic cell, as that should not be blocked by TTX in a well-clamped cell. Ethanol might additionally act on a presynaptic site other than that directly affected by the D₁R. It appears that ethanol augments the D₁R-mediated enhancement of glutamate release in a way that is dependent on the presynaptic TTX-sensitive Na⁺ channels. To test this possibility, we examined ethanol-induced augmentation of the SKF 38393-induced enhancement of sEPSCs in the absence and presence of TTX. As expected, in the absence of TTX, the co-application of ethanol (20 mM) and SKF 38393 (10 nM) increased sEPSC frequency by $181.8 \pm 14.6\%$ ($n = 4$), which is significantly larger than that induced by SKF 38393 (10 nM) alone (by $50.9 \pm 16.0\%$, $n = 4$, $p < 0.01$). Conversely, in the presence of TTX, the co-application of SKF 38393 (10 nM) and ethanol (20 mM) increased mEPSC

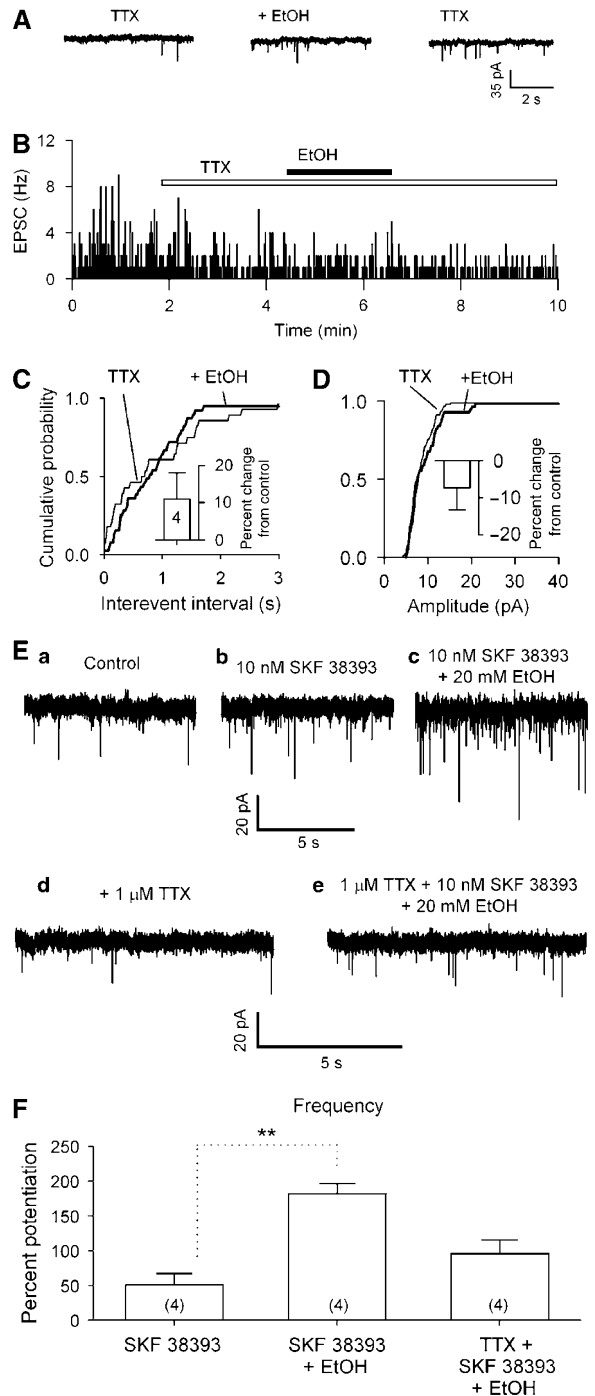


Figure 5 Ethanol has no effect on mEPSCs. The mEPSCs were recorded in VTA DAergic neurons at a V_H of -65 mV, in the presence of bicuculline (10 μM) and tetrodotoxin (TTX, 1 μM). (A) Application of 40 mM ethanol did not change the incidence of mEPSCs. (B) Time course of ethanol-induced changes of sEPSC frequency in one experiment. (C, D) Representative cumulative probability plots of intervals between mEPSCs (C) and their amplitudes (D) remained unchanged. Inset: mean data (\pm SEM, from four neurons) confirm absence of significant effect of ethanol on frequency and amplitude of mEPSCs. (E) Ethanol (20 mM) increased the enhancement of sEPSC frequency induced by 20 nM SKF 38393 (c) but not in the presence (e) of TTX (1 μM). (F) Mean data (\pm SEM, from four neurons) confirm ethanol (20 mM) increased 20 nM SKF 38393-induced enhancement of sEPSC frequency in the absence but not in the presence of TTX (1 μM). ** $p < 0.01$, t-test for SKF 38393 application vs SKF 38393 + ethanol.

frequency by $95.8 \pm 19.4\%$ ($n = 4$), which is not significantly different from that induced by SKF 38393 alone ($p = 0.1$, $n = 4$; Figure 5E and F).

Somatodendritic Dopamine Release in the VTA is Essential for the Action of Ethanol

To determine whether dopamine release is necessary for the action of ethanol, we applied ethanol to nerve-bouton preparations in which dopamine was depleted by reserpine. Because reserpine was dissolved in acetic acid (final concentration 1:5000, v/v), control tests were carried out on neurons isolated from slices pretreated for > 90 min with this concentration of the vehicle alone. After the treatment with acetic acid, the frequency and amplitude of the sEPSCs were not significantly changed. Ethanol (40 mM) increased sEPSC frequency by $77 \pm 6\%$ (baseline, 1.0 ± 0.2 Hz; ethanol, 1.8 ± 0.4 Hz, $n = 4$, $p < 0.05$, Figure 6A and E2). By contrast, in neurons isolated from slices pretreated for similar periods with reserpine (10 μ M), sEPSC frequency was much lower (see below), and also did not respond to 40 mM ethanol: by $39 \pm 19\%$ of control (baseline, 0.2 ± 0.0 Hz; ethanol 0.3 ± 0.1 Hz, $n = 4$, $p = 0.233$, Figure 6B and E). These data indicate that the release of dopamine is essential for ethanol-induced facilitation of sEPSCs.

Somatodendritic dopamine release depends on membrane depolarization (Llinas *et al*, 1984; Grace, 1990). It is augmented following potassium (Kalivas and Duffy, 1991) or veratridine depolarization (Irvani *et al*, 1996). Therefore we tested whether depolarizing step applied to isolated dopamine neurons can increase sEPSC frequency. A 4 s depolarizing step from -60 to 0 mV increased sEPSC frequency in 7 out of 13 cells tested (by $52 \pm 18\%$, prepulse, 1.9 ± 0.3 Hz; postpulse, 2.9 ± 0.4 Hz, $n = 7$, $p < 0.05$; Figure 6C). These responses varied widely and may have been complicated by depolarization induced the release of other neurotransmitters/neuromodulators, such as endocannabinoids, which inhibited presynaptic glutamate release in the VTA (Melis *et al*, 2004).

The release of dopamine in the VTA is Ca^{2+} dependent (Kalivas and Duffy, 1991; Campbell *et al*, 1996; Reith *et al*, 1997), and thus can be prevented or reduced by chelating postsynaptic Ca^{2+} . To further ensure a role for somatodendritic dopamine in ethanol facilitation of sEPSCs, we tested the effect of ethanol (40 mM) on cells recorded with pipette solution containing the Ca^{2+} chelator BAPTA (20 mM). Under these experimental conditions, ethanol (40 mM) changed neither sEPSC frequency (by $-6 \pm 8\%$, baseline, 1.8 ± 0.4 Hz; ethanol, 1.8 ± 0.5 Hz, $n = 3$, $p = 0.66$; Figure 6D and E), nor sEPSCs amplitude (baseline, 34 ± 4 pA; ethanol, 38 ± 3 pA, $n = 3$, $p = 0.29$). These data indicate that somatodendritic release of dopamine is necessary for ethanol-induced facilitation of sEPSCs.

Blockade of Dopamine Reuptake Raises sEPSC Frequency in Mechanically Dissociated DAergic Neurons

Previous *in vivo* evidence indicates that acute administration of ethanol (by intraperitoneal injection) elevates extracellular dopamine levels in the VTA (Campbell *et al*, 1996; Yan *et al*, 2005). To elicit a similar rise in local

dopamine levels in single neurons, we applied the selective blocker of dopamine transport GBR 12935. Application of GBR 12935 (1 μ M) remarkably increased sEPSC frequency by $157 \pm 31\%$ (control, 0.7 ± 0.2 Hz; GBR 12935, 1.5 ± 0.3 Hz; $n = 4$, $p = 0.01$, Figure 7b). GBR 12935 had no effect on the mean amplitude of sEPSCs (control, 19 ± 2 pA; GBR 12935, 21 ± 2 pA; $n = 4$, $p = 0.01$, Figure 7c). The effect of GBR 12935 depended on its concentrations. On average, 0.02, 0.2, and 1 μ M GBR 12935 changed sEPSC frequency by $66 \pm 20\%$ ($n = 5$), $85 \pm 18\%$ ($n = 3$), and $157 \pm 31\%$ ($n = 4$), respectively (one-way ANOVA, $F = 5.16$, $p = 0.018$). This finding shows that a rise in local dopamine levels also mimics the action of ethanol.

Our next step was to investigate the mechanism of GBR 12935-induced facilitation of sEPSC frequency. Application of SKF 83566 (10 μ M) alone had no effect on sEPSC frequency (by $3 \pm 2\%$ of control. Baseline, 0.9 ± 0.1 Hz; SKF 83566, 0.9 ± 0.1 Hz; $n = 4$, $p = 0.10$, not illustrated). However, SKF 83566 abolished the enhancement of sEPSC frequency induced by GBR 12935 (1 μ M). In the presence of SKF 83566 (10 μ M), the sEPSC frequency changed by $2 \pm 4\%$ (GBR 12935, 1.5 ± 0.3 Hz; GBR 12935 + SKF 83566, 0.9 ± 0.2 Hz; $n = 4$, $p = 0.01$; Figure 7b). These results further indicate that D_1 Rs are essential in dopamine facilitation of glutamatergic transmission to the DAergic neurons in the VTA.

Finally, we reexamined the effects of GBR 12935 (1 μ M) and SKF 38393 (10 μ M) on cells recorded with pipette solution containing 20 mM BAPTA. Under these experimental conditions, the sEPSCs were virtually identical in the absence and presence of 1 μ M GBR 12935 (frequency: control, 0.490 ± 0.129 Hz; GBR 12935, 0.487 ± 0.131 Hz, $n = 4$, $p = 0.81$; amplitude: control, 27.7 ± 5.2 pA; GBR 12935 28.5 ± 5.7 pA, $n = 4$, $p = 0.21$). Conversely, SKF 38393 increased sEPSC frequency by $101.4 \pm 39.6\%$ (control, 0.41 ± 0.12 Hz, SKF 38393, 0.76 ± 0.17 Hz; $n = 4$; $p = 0.032$), without changing their mean amplitude (control, 30.9 ± 2.6 pA; SKF 38393, 31.3 ± 3.8 pA; $n = 4$, $p = 0.94$; Figure 7d and e). These results indicate that the dopamine reuptake blocker works only if there is ambient dopamine.

DISCUSSION

The current study using isolated neurons provides solid evidence that: (1) ethanol at clinically relevant concentrations (10–80 mM) facilitates glutamatergic transmission to VTA DAergic neurons, (2) although ethanol facilitation has a presynaptic locus, it is mediated not by the direct activation of the presynaptic D_1 Rs, but by the retrograde messenger dopamine, which is released from somatodendritic area and travels backward across synapses to facilitate glutamatergic neurotransmission by the activation of the D_1 Rs on the glutamatergic terminals, and (3) ethanol facilitation of glutamate transmission to VTA DAergic neurons can occur in a preparation containing only a postsynaptic neuron and attached glutamatergic synapses.

Of note, in keeping with our recent result from brain slices, the single cell data show that ethanol enhances glutamatergic EPSCs at concentrations close to the blood alcohol levels observed in rats after voluntary ethanol consumption (Robinson *et al*, 2000; Doyon *et al*, 2003,

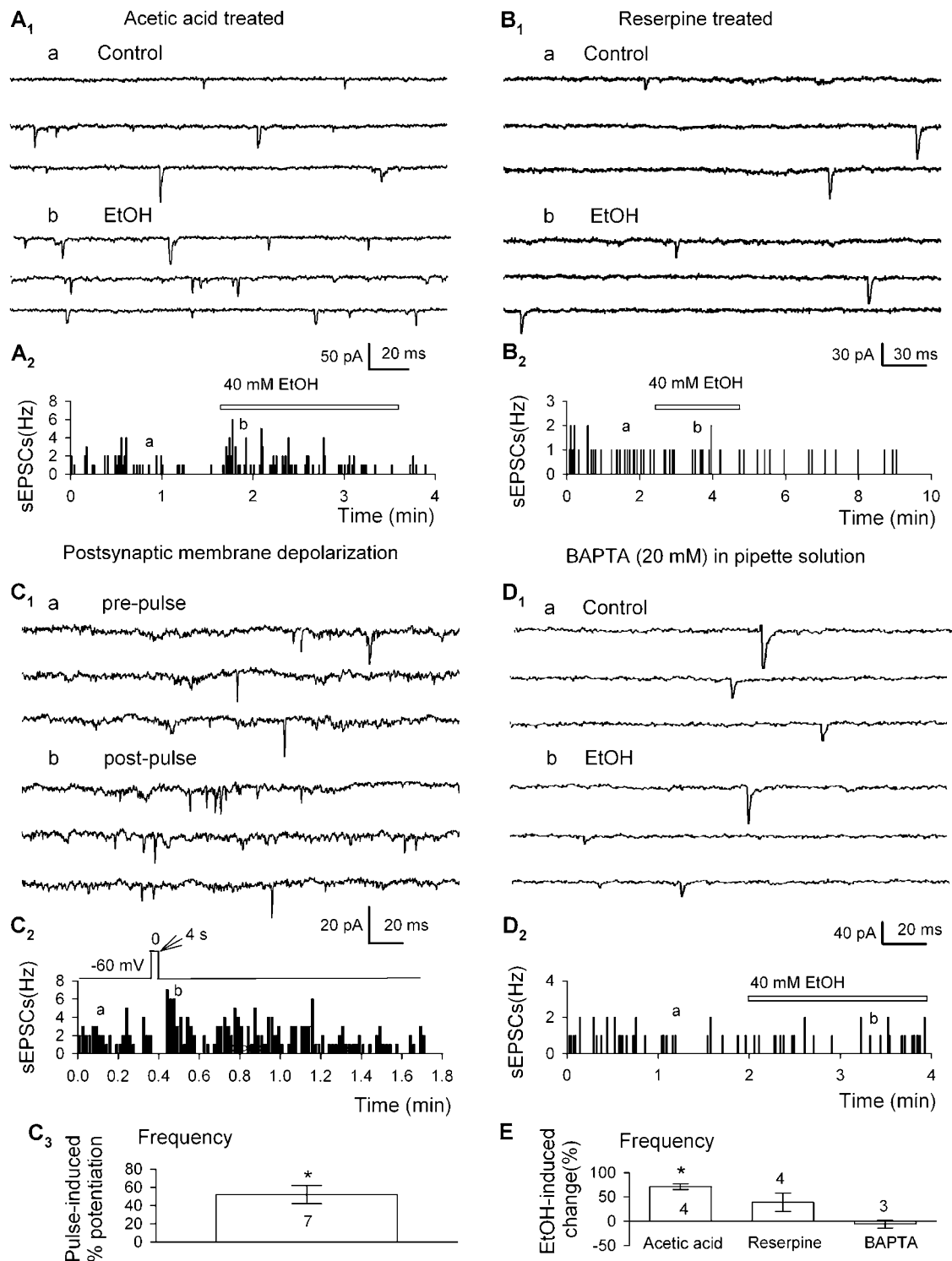


Figure 6 The release of dopamine is necessary for ethanol's facilitation of sEPSCs. (A₁, B₁) Current traces show that ethanol enhancement of sEPSCs was observed in neurons from slices pretreated with acetic acid (A₁), but not from reserpine (B₁). (A₂, B₂) Time course of ethanol-induced change of sEPSC frequency in one experiment. The traces in A₁ and B₁ were taken at times indicated by (a) and (b). (C₁) Current traces show that a 4 s depolarization pulse (from -60 to 0 mV) increased sEPSC frequency. (C₂) Time course of depolarization-induced changes of sEPSC frequency in one experiment. (C₃) Mean data (\pm SEM, from seven neurons) show depolarization induced potentiation of sEPSC frequency. * $p < 0.05$, paired t -test for postpulse vs prepulse control. (D₁) Current traces show that 40 mM ethanol did not change the sEPSCs when the pipette solution contained 20 mM BAPTA. (D₂) Time course of 40 mM ethanol-induced changes of sEPSC frequency in one experiment in which BAPTA was in the pipette solution. (E) Mean data (\pm SEM) show changes in sEPSC frequency induced by 40 mM ethanol under different experimental conditions. * $p < 0.05$, paired t -test for ethanol application vs pre-ethanol control, or as indicated, nonpaired t -test for reserpine group vs reserpine + ethanol group.

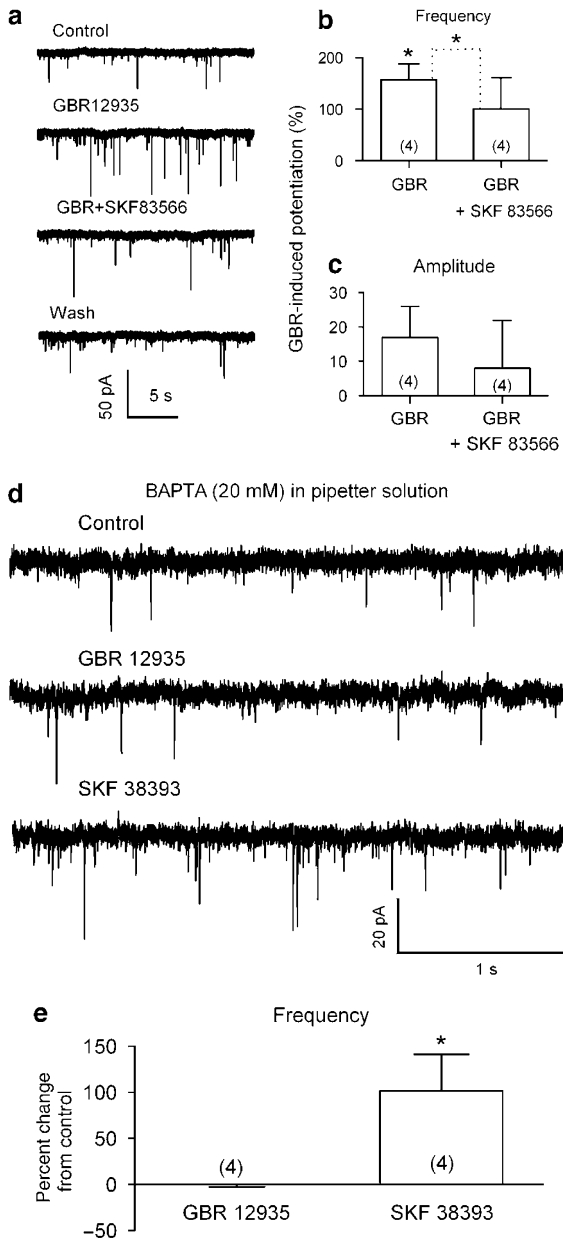


Figure 7 Ethanol-induced increase in sEPSC frequency is mimicked by blocking dopamine transport. (a) Traces illustrating increase in sEPSC frequency produced by GBR 12935 (1 μ M, a dopamine transport blocker), which is blocked by the D₁R antagonist SKF 83566 (10 μ M, $n = 4$). (b) Summary of GBR 12935-induced changes in sEPSC frequency, which was blocked by SKF 83566 (data from four cells). * $p < 0.05$, paired t -test for GBR 12935 application vs pre-GBR 12935, or as indicated for GBR 12935 group vs GBR 12935 + SKF 83566. (c) No change in amplitude was seen in GBR 12935 either in the absence or the presence of 10 μ M SKF 83566. (d) Traces illustrating increase in sEPSC frequency produced by 10 μ M SKF 38393 but not by 1 μ M GBR 12935 when 20 mM BAPTA was in the pipette solution. (e) Mean data (\pm SEM, from four neurons) show that SKF 38393, but not GBR 12935, increased sEPSC frequency when pipette solution contained 20 mM BAPTA. * $p < 0.05$, paired t -test for SKF 38393 application vs pre-SKF 38393.

2005). As glutamatergic transmission is an important mediator of behavioral changes caused by ethanol (Eckardt *et al*, 1998), we propose that this somatodendritic dopamine-mediated increased glutamate release in the VTA could be an important aspect of alcohol addiction.

Ethanol Facilitates Glutamatergic Transmission in Single VTA DAergic Neurons

In contrast to the suppression effects ethanol has on the neurons in many of the brain areas, previous studies have shown that ethanol can excite the VTA DAergic neurons (Gessa *et al*, 1985; Brodie *et al*, 1990). The mechanism underlying this excitatory effect of ethanol is not well understood, but it may contribute to the addictive property of ethanol and thus is a rapidly growing area of research in the neuroscience community. Ethanol may excite the DAergic neurons directly (Brodie *et al*, 1999b), by an enhancement of the hyperpolarization-activated current (Okamoto *et al*, 2006), and/or by an inhibition of the M channel (Koyama *et al*, 2007). Nevertheless, recent studies from this and other laboratories have shown that ethanol may excite VTA DAergic neurons indirectly, by an inhibition of the GABAergic transmission (Stobbs *et al*, 2004; Xiao *et al*, 2007; Xiao and Ye, 2008; but see Theile *et al* (2008) who reported that ethanol increases GABAergic transmission onto VTA DAergic neurons), and by an enhancement of the glutamatergic transmission to the VTA DAergic neurons (see below for more; Xiao *et al*, 2008).

Glutamate is the most abundant excitatory neurotransmitter in the brain, mediating as much as 70% of synaptic transmission within the CNS. Glutamate is important in drug addiction and alcoholism (Gass and Olive, 2008). Most previous studies found that ethanol inhibits NMDA and nonNMDA glutamate receptors (Lovinger *et al*, 1989; Weitauf and Woodward, 2008), as well as glutamate release (Siggins *et al*, 2005). In a recent study in midbrain slices, however, we demonstrated that acute ethanol facilitates glutamatergic transmission to VTA DAergic neurons (Xiao *et al*, 2008). In the current investigation in isolated neurons, we offer further support for ethanol potentiation of glutamate transmission to VTA DAergic neurons. This is the first electrophysiological evidence demonstrating a potentiating effect of acute ethanol on glutamatergic transmission in single cells.

Note that the effect of ethanol on sEPSC frequency is larger in isolated neurons than in slices. For example, 40 mM ethanol increased sEPSC frequency by $\geq 77\%$ in single cells, whereas the same concentration of ethanol increased sEPSC frequency by 50% in brain slices. We hypothesize that this results from a more efficient space clamping of isolated neurons and much faster and more complete exchange of the surrounding solution. Ethanol application is much more rapid in the isolated neurons, and there may be less time for the development of tolerance. In addition, a much greater role of reuptake in dopamine clearance in slices than in isolated neurons may also contribute to the difference (Brodie *et al*, 1999a).

It has been well documented that glutamatergic afferents of the VTA DAergic neurons comprise abundant neurons within a large formation extending from the prefrontal cortex to the caudal brainstem (Geisler *et al*, 2007). In addition, there are glutamatergic neurons in the VTA (Yamaguchi *et al*, 2007). Furthermore, DAergic neurons release glutamate as a cotransmitter with dopamine (Trudeau, 2004; Descarries *et al*, 2007). Future studies should identify the source of the glutamatergic afferents of

the VTA DAergic neurons, which is responsible for ethanol facilitation.

Ethanol Acts Presynaptically to Enhance Glutamatergic Transmission to Single VTA DAergic Neurons

In keeping with our recent observation in brain slices (Xiao *et al*, 2008), in the current investigation in isolated neurons, we showed that ethanol increased the frequency of sEPSCs but had no effect on the amplitude of mEPSCs and of sEPSCs, indicating that ethanol acts on the presynaptic site to increase glutamate release. A possible explanation for the unusual positive action on glutamate release—which is in contrast to the more common depressant actions (Siggins *et al*, 2005)—would be an indirect mechanism, based on the special organization of the VTA, whose DAergic neurons receive dopamine (released from the somatodendritic area) and glutamate (released from glutamate-releasing terminals) of particular importance is the presence of D₁Rs on the glutamatergic axons.

Of note, a recent study shows that the DAergic cells in the substantia nigra make chemical DAergic synapses on one another (Vandecasteele *et al*, 2008). Perhaps VTA DAergic cells do as well. That is, the dopamine, which activates the presynaptic D₁Rs could be released from the somatodendritic areas of the adjacent DAergic neuron, instead of the same DAergic neuron. However, ethanol facilitation was blocked by chelating postsynaptic calcium with BAPTA, indicating that the dopamine released from adjacent neuron, if it exists, may not have major function in ethanol facilitation of glutamate transmission in the VTA.

Ethanol Enhances Glutamate Release in Single Cells by D₁R Activation

This notion is supported by the following evidence. The ethanol-induced increase in sEPSC frequency was eliminated by a D₁ antagonist. Moreover, like ethanol, a D₁ agonist increased the frequency of sEPSCs and mEPSCs without affecting their amplitude; a similar effect was produced by an increase in endogenous dopamine, induced by a blocker of dopamine reuptake. All these changes were abolished by the D₁ antagonist.

Our data also suggest a crucial involvement of somatodendritic released dopamine, because ethanol's action was suppressed when endogenous monoamines (including dopamine) were depleted by reserpine, or when the postsynaptic Ca²⁺ was chelated by BAPTA (20 mM). In addition, these data indicate that blocking dopamine reuptake may not be the major player in ethanol facilitation of sEPSCs, although previous *in vivo* studies suggested that ethanol may raise extracellular dopamine by blocking dopamine reuptake (Campbell *et al*, 1996; Yan *et al*, 2005).

A rise in local dopamine levels is also suggested by the bell-shaped dose-dependence of ethanol's action, both in our experiments (both in slices and in single cells) and in the *in vivo* experiments of Rodd *et al* (2004). This may result from opposite effects of dopamine (the nonselective endogenous agonist) acting on different receptors. At low concentration, dopamine appears to bind predominantly to D₁Rs and at higher concentrations to D₂Rs. Thus, Trantham-Davidson *et al* (2004) found that low

concentrations of dopamine (<0.5 μM) enhanced IPSCs by activating mainly D₁Rs; whereas higher concentrations of dopamine (>1 μM) reduced IPSCs by predominantly activating D₂Rs. Similarly, Alberto *et al* (2006) found that a low dose of dopamine (1 μM) increased mEPSC frequency by activating D₁Rs; whereas higher doses of dopamine (10–100 μM) decreased mEPSC frequency by activating D₂Rs. In comparable experiments on VTA and globus pallidus, Koga and Momiyama (2000) and Hernandez *et al* (2007) found that dopamine or a D₂ agonist inhibits EPSCs in DAergic neurons through presynaptic D₂Rs (in VTA, with a high IC₅₀ of 16 μM for dopamine). That the bell-shaped concentration dependence of ethanol's action is resulting from opposite presynaptic actions by D₁Rs and then D₂Rs is strongly supported by our finding in slices that the facilitation of sEPSCs by 80 mM ethanol was greatly potentiated when D₂Rs were blocked (Xiao *et al*, 2008).

Dopamine Reuptake Blocker (GBR 12935) Enhances Glutamate Release in Isolated VTA DAergic Neurons

The application of GBR 12935 facilitates sEPSC frequency in single isolated DAergic neurons. This observation has specific significance to the mechanism underlying somatodendritic release of dopamine.

At the axon terminal, dopamine is stored in vesicles and released exocytotically. In the dendrite/soma, traditional secretory vesicles are scarce. Because synaptic sites and classic vesicular storage are uncommon in somatodendritic areas, some investigators have questioned whether somatodendritic dopamine is released exocytotically. Hallmarks of exocytotic release include sensitivity to Ca²⁺ concentration, Na⁺ channel blockade, and neuronal vesicular monoamine transporter inhibition. Several studies have shown that somatodendritic dopamine release obeys these ionic requirements (Geffen *et al*, 1976; Cheramy *et al*, 1981; Kalivas and Duffy, 1991; Rice *et al*, 1994). However, nonexocytotic mechanisms of somatodendritic dopamine release have also been suggested, including reversal of the dopamine transporter (DAT; Nirenberg *et al*, 1996; Falkenburger *et al*, 2001). Nevertheless, a recent study found that dopamine could be measured in VTA slices from DAT knockout mice, and that the mechanisms governing electrically evoked somatodendritic dopamine release mechanisms are similar between DAT knockout and C57BL/6 mice (John and Jones, 2006). These data support an exocytotic mechanism for dopamine release in the VTA. This is also supported by a recent study suggesting that somatodendritic dopamine transmission is more analogous to classical synaptic transmission than previously believed (Beckstead *et al*, 2004). In isolated neurons, we found that GBR 12935 significantly increased sEPSC frequency. Higher concentrations of GBR 12935 may have effects other than blocking dopamine reuptake. However, this is unlikely as the facilitation of sEPSCs induced by GBR 12935 was blocked by a D₁R antagonist. Thus, our results using GBR 12935 provide strong support for the exocytotic mechanism of somatodendritic dopamine release in the VTA.

In conclusion, this study provides evidence from single isolated VTA DAergic neurons that ethanol facilitation results from increased somatodendritic dopamine release, which retrogradely activates the D₁Rs on glutamate-

releasing terminals. These results demonstrate that ethanol stimulation of a single cell is capable of eliciting the release of dopamine, which is sufficient to influence glutamatergic transmission at individual synapses. Given that excitatory afferents regulate the activity of DAergic neurons, the release of dopamine is essential for the development of addiction (Kalivas and Duffy, 1995), and ethanol facilitation of glutamate release might contribute to its reward property. Our findings indicate that somatodendritic dopamine release is crucial in modulating the activity of DAergic neurons and in the rewarding effect of ethanol, and thus a key target of ethanol. Furthermore, the current study also demonstrates a powerful tool for pharmacological and physiological studies of dopamine signaling in the VTA.

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DISCLOSURE/CONFLICT OF INTEREST

The authors declare no conflict of interest.

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