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Bidirectional and long-lasting control of alcohol-seeking behavior by corticostriatal LTP and LTD

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Abstract

Addiction is proposed to arise from alterations in synaptic strength via mechanisms of long-term potentiation (LTP) and depression (LTD). However, the causality between these synaptic processes and addictive behaviors is difficult to demonstrate. Here we report that LTP/LTD induction altered operant alcohol self-administration, a motivated drug-seeking behavior. We first induced LTP by pairing presynaptic glutamatergic stimulation with optogenetic postsynaptic depolarization in the dorsomedial striatum, a brain region known to control goal-directed behavior. Blockade of this LTP by NMDA receptor inhibition unmasked an endocannabinoid-dependent LTD. *In vivo* application of the LTP-inducing protocol caused a long-lasting increase in alcohol-seeking behavior, while the LTD protocol decreased this behavior. We further identified that optogenetic LTP/LTD induction at cortical inputs onto striatal dopamine D1 receptor-expressing neurons controlled these behavioral changes. Our results demonstrate a causal link between synaptic plasticity and alcohol-seeking behavior, and that modulation of this plasticity may inspire a therapeutic strategy for addiction.

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Data availability

The data that supports the findings of this study are available from the corresponding author upon reasonable request.

Author Contributions

J.W. conceived, designed, and supervised all experiments in the study. J.W. and T.M. wrote a draft of the manuscript. J.W., T.M., and Y.C. revised the manuscript. T.M. also designed and performed electrophysiology experiments and analyzed the data. Y.C. and E.R.H. designed and performed the behavior experiments and analyzed the data. X.W. conducted behavior experiments. J.Y.J. supervised the collection and analyses of biochemical experiments. J.L. and X.G. performed and analyzed the biochemical experiments. C.H. conducted the electrophysiology experiment for virus validation and analyzed the data. Y.C. and X.Y.W. conducted confocal imaging experiments and analyzed the data. All authors discussed the results and commented on the manuscript.

Competing interests

The authors declare no competing financial interests.

Drug addiction is a mental illness that is viewed as a transition from recreational use to compulsive drug-seeking and -taking¹⁻³. This behavioral transition is proposed to be controlled by “drug-evoked plasticity”¹⁻⁴. However, exactly how synaptic plasticity controls the adaptive changes in drug-seeking behavior remains unclear. The dorsomedial striatum (DMS), a brain region crucially involved in drug and alcohol addiction, receives glutamatergic inputs from several brain areas^{2, 5-7}. In these neural circuits, the medial prefrontal cortex (mPFC) afferent into the DMS is essential for the control of goal-directed behaviors^{2, 5-7}, and this connection is linked to drug or alcohol addiction^{2, 5, 8}. For example, exposure to drugs of abuse or alcohol potentiates AMPA receptor (AMPA)- and NMDA receptor (NMDAR)-mediated glutamatergic transmission in the DMS⁸⁻¹¹, while pharmacological inhibition of striatal glutamatergic transmission transiently suppresses operant alcohol self-administration and cocaine relapse^{9, 10, 12, 13}. Although these studies indicate that drugs and alcohol evoke corticostriatal plasticity, which may, in turn, contribute to drug-seeking and -taking behaviors, there has been no direct demonstration that synaptic plasticity drives addictive behaviors.

The DMS contains two types of medium spiny neurons (MSNs): D1-MSNs express dopamine D1 receptors (D1Rs) and D2-MSNs contain D2Rs¹⁴. Both neuronal types receive mPFC inputs¹². While synaptic plasticity, including long-term potentiation (LTP) and depression (LTD), was observed in both D1- and D2-MSNs¹⁵, drug- or alcohol-induced plasticity was found predominantly in striatal D1-MSNs^{12, 16, 17}. Mimicking alcohol-evoked plasticity by inducing LTP, or reversing this plasticity by inducing LTD, will provide a new understanding of how this plasticity controls alcohol-seeking behavior. LTP/LTD induction at specific neuronal circuits requires simultaneous control of both pre- and post-synaptic neurons, which can be achieved using a recently developed dual-channel optogenetic technique^{18, 19}.

In this study, we paired optogenetic postsynaptic depolarization (oPSD) with presynaptic glutamatergic stimulation; this greatly enhanced NMDAR-mediated transmission and induced a reliable NMDAR-dependent LTP, as well as an endocannabinoid (eCB)-dependent LTD. Importantly, *in vivo* optogenetic delivery of the LTP protocol to the corticostriatal synapses within the DMS produced a long-lasting increase in operant self-administration of alcohol. Conversely, delivery of the LTD protocol led to a long-lasting decrease in this behavior. Furthermore, we discovered that the *in vivo* LTP and LTD protocols preferentially induced plasticity in D1-MSNs and that selective induction of LTP/LTD in this neuronal type produced the corresponding changes in alcohol-seeking behavior. These findings demonstrate a causal link between DMS corticostriatal synaptic plasticity and alcohol-seeking behavior and indicate that the reversal of drug-evoked synaptic plasticity may provide a novel therapeutic strategy for the treatment of alcohol use disorder.

RESULTS

oPSD facilitates induction of NMDAR-dependent LTP and eCB-LTD in the DMS

In dorsostriatal slices, LTD is the easiest form of synaptic plasticity to observe, while LTP is more difficult to detect^{6, 20}. A D2R antagonist was thus included in the recording solution in order to prevent LTD¹⁵ and favor the induction of LTP. Field excitatory postsynaptic

potentials/population spikes (fEPSP/PS) were evoked by electrical stimulation within the DMS (Fig. 1a), but these were not potentiated by electrical high-frequency stimulation (eHFS) (Fig. 1b and Supplementary Fig. 1a). This observation is consistent with previous reports^{6, 21}. Previous studies also suggested that sufficient postsynaptic depolarization was necessary for reliable LTP induction²⁰. Therefore, we examined whether postsynaptic depolarization by somatic current injection (iPSD) facilitated LTP induction. Using whole-cell recording, we discovered that paired eHFS and iPSD produced little LTP (Supplementary Fig. 1b). This is consistent with the notion that current injection-elicited action potentials do not back-propagate to the distal dendrites of striatal neurons²⁰; insufficient depolarization of this region means that no LTP is generated. In contrast, optogenetics can be used to depolarize any process of postsynaptic neurons, with no limitation of their distance to the soma, and in a non-invasive manner²². We observed that optogenetic postsynaptic depolarization (oPSD) induced a higher distal dendritic calcium transient than iPSD (Supplementary Fig. 2), suggesting that oPSD produced more effective depolarization of this region.

Next, we assessed whether oPSD facilitates LTP induction. An adeno-associated virus (AAV) expressing a channelrhodopsin, C1V1²³, was infused into the DMS, resulting in C1V1 expression in the soma and distal dendrites (Fig. 1c). We found that pairing of eHFS with oPSD of DMS neurons induced a robust and reliable LTP, whereas oPSD alone did not (Fig. 1d). Furthermore, paired presynaptic stimulation and oPSD enhanced synaptic NMDAR activity and consequently Ca²⁺ influx through this channel (Supplementary Fig. 3). LTP was blocked by bath application of an NMDAR antagonist, APV (Fig. 1e). In addition, it was reported that dorsostriatal LTP induction also depended on D1R activation^{6, 15}. We found that optogenetic LTP was inhibited by a D1R antagonist, SCH 23390, and was facilitated by D1R activation (Supplementary Fig. 4). Collectively, these results suggest that paired presynaptic stimulation and oPSD produced effective distal dendrite depolarization and induced robust NMDAR-dependent LTP that is strongly regulated by D1R signaling.

Surprisingly, after blockade of optogenetic LTP by APV, LTD was observed (Fig. 1e). This LTD was completely abolished by bath application of an eCB CB1 receptor (CB1R) antagonist, AM251 (Fig. 1f). This was consistent with previous reports indicating that LTD in the dorsal striatum was mediated by the CB1R^{24, 25}. Since this eCB-LTD only emerged after LTP was blocked, we reasoned that LTP and LTD were induced simultaneously, and that LTD was masked by LTP. To assess this possibility, we bath-applied AM251 throughout the recording period; this produced a significantly greater magnitude of LTP, as compared with that recorded in the absence of AM251 (Fig. 1g). Collectively, these results suggest that both NMDAR-dependent LTP and eCB-LTD were induced simultaneously and that the LTP masked the LTD.

oPSD facilitates corticostriatal LTP in the DMS

Corticostriatal plasticity is critical for drug-seeking behaviors^{2, 5-7}. We, therefore, examined whether oPSD facilitated LTP induction at specific corticostriatal afferents in the DMS. We expressed two channelrhodopsins simultaneously in order to selective stimulation of cortical inputs and oPSD of DMS neurons: Chronos¹⁸ was expressed in the mPFC and Chrimson¹⁸

was expressed in the DMS (Fig. 2a,b). Chronos-expressing mPFC neurons and their projections to the DMS were able to follow high-frequency (up to 50 Hz) light stimulation (Supplementary Fig. 5). We thus used light stimulation of 50 Hz for 2 sec (oHFS), paired with oPSD of DMS neurons, to induce LTP. While oHFS of corticostriatal fibers or oPSD alone caused little potentiation (Fig. 2c,d), a robust LTP was observed following paired oHFS and oPSD (Fig. 2c). This corticostriatal LTP was abolished by APV or MK801, as expected (Fig. 2e and Supplementary Fig. 6a). Surprisingly, no LTD was observed after LTP was blocked (Fig. 2e and Supplementary Fig. 6b); this contrasted with the results produced by pairing eHFS and oPSD (Fig. 1e). However, whole-cell recording detected a robust LTD in specific D1-MSNs (Supplementary Fig. 6c). This was consistent with a recent report showing that selective optogenetic stimulation of cortical inputs induced LTD only in D1-MSNs²⁶. Collectively, these data suggest that paired oHFS and oPSD induces corticostriatal LTP, as well as LTD in D1-MSNs.

LTP stimulation has been demonstrated to induce expression of immediate early genes, contributing to drug addiction^{27, 28}. We analyzed DMS slices and found that paired oHFS and oPSD, but not oHFS or oPSD alone, significantly increased mRNA levels of *Npas4* (*neuronal PAS domain protein 4*) gene, which encodes the Npase4 protein (Fig. 2f). This immediate early gene is associated with synaptic plasticity and positive valence experience^{29, 30}.

***In vivo* optogenetic induction of corticostriatal LTP in the DMS produces a long-lasting increase in operant alcohol self-administration in rats**

Our *ex vivo* findings revealed that paired oHFS and oPSD elicited LTP in the DMS. We thus asked whether *in vivo* delivery of this LTP-inducing protocol (oHFS+oPSD) altered alcohol-seeking behavior. To test this possibility, rats were trained to self-administer alcohol in operant chambers. Chronos and Chrimson were expressed as described above, and optical fibers were implanted into the DMS (Fig. 3a). We found that *in vivo* delivery of this optogenetic protocol produced significant increases in active lever presses, alcohol deliveries, and alcohol intake at 30 min, 2 days, and 4 days (Fig. 3b and Supplementary Fig. 7a). This increased alcohol intake resulted in elevated blood alcohol concentrations (Supplementary Fig. 7b,c). In contrast, oHFS or oPSD alone did not alter alcohol-seeking behavior (Supplementary Fig. 7d,e). Together, these data suggest that *in vivo* delivery of this LTP-inducing protocol is sufficient to cause long-lasting enhancement of alcohol-seeking behavior. However, systemic administration of antagonists of NMDARs (MK801) or D1Rs (SCH 23390) blocked this effect of the LTP-inducing protocol on alcohol-seeking behavior (Supplementary Fig. 7f,g). Note that administration of SCH 23390 alone did not affect this behavior (Supplementary Fig. 7h). These results suggest that both NMDARs and D1Rs are required for the enhancement of alcohol-seeking behavior by *in vivo* LTP induction.

Next, we asked whether the LTP-inducing protocol specifically enhanced alcohol-seeking behavior. Another cohort of rats was trained to self-administer sucrose prior to receiving the same LTP-inducing protocol as the alcohol group. We found that the LTP protocol did not alter the active lever presses, sucrose deliveries, or sucrose intake (Fig. 3c and Supplementary Fig. 7i). We then asked why the same LTP-inducing protocol specifically

promoted alcohol-seeking, but not sucrose-seeking, in rats. On day 2 post-LTP induction, both AMPAR-mediated excitatory postsynaptic currents (EPSCs) and the AMPAR/NMDAR ratio were increased in the alcohol group (Fig. 3d,e), but not in the sucrose group (Fig. 3f,g). Interestingly, prior to *in vivo* LTP induction, operant alcohol or sucrose self-administration had increased the AMPAR/NMDAR ratio, as compared with the water group without operant training (Supplementary Fig. 8a). The increase was slightly lower in the alcohol-treated rats than in the sucrose controls, but the difference was not statistically significant ($Q = 2.86$, $P = 0.051$, SNK test). These data suggest that while operant-training produced plasticity in alcohol and sucrose groups, *in vivo* delivery of the LTP-inducing protocol caused further long-lasting synaptic potentiation selectively in the alcohol group. This difference may reflect the distinct effects of alcohol and sucrose on NMDAR activity^{6, 31–33}. To investigate this, we measured NMDAR activity in the DMS of rats that self-administered alcohol or sucrose. We found that the amplitude of NMDAR-mediated EPSCs was significantly higher in the alcohol group than in the sucrose group (Supplementary Fig. 8b,c). Furthermore, LTP was induced in DMS slices from the alcohol-treated rats, but not in those from the sucrose-drinking animals (Supplementary Fig. 8d), suggesting that alcohol-mediated facilitation of NMDAR activity^{34, 35} promotes subsequent *ex vivo* and *in vivo* induction of LTP. In addition, the rectification index of AMPAR-EPSCs was significantly enhanced following *in vivo* LTP induction (Supplementary Fig. 9), suggesting that this plasticity is mediated by an increase in calcium-permeable AMPARs.

Collectively, these results suggest that induction of corticostriatal LTP in the DMS produces a long-lasting and specific increase in operant alcohol self-administration in rats.

***In vivo* optogenetic delivery of an LTD-inducing protocol in the DMS produces a long-lasting decrease in alcohol-seeking behavior in rats**

Having observed the link between LTP and alcohol-seeking behavior, we reasoned that reversal of alcohol-induced potentiation of corticostriatal inputs by LTD should reduce alcohol-seeking behavior. To induce LTD *in vivo*, we systemically administered a cocktail of MK801 and a D2R antagonist, raclopride, 30 min before delivering oHFS and oPSD (Fig. 4a). We used these two categories of antagonists because they were employed in the *ex vivo* LTD experiments described above (Fig. 1e and Supplementary Fig. 6c). Thirty minutes after oHFS and oPSD delivery, significant reductions in active lever presses, alcohol deliveries, and alcohol intake were observed; these reductions were maintained for at least 7 days (Fig. 4b and Supplementary Fig. 10a). Note that neither the cocktail (MK801+raclopride) plus oHFS (Supplementary Fig. 10b–d) nor the cocktail alone (Supplementary Fig. 10e–g) affected alcohol-seeking behavior. Together with our finding that oHFS+oPSD+MK801 induced no changes in alcohol consumption (Supplementary Fig. 7g), these data suggest that the *in vivo* LTD-inducing protocol (oHFS+oPSD+MK801+raclopride) produces a long-lasting reduction of alcohol-seeking behavior and that this induction requires D2R blockade.

Since the LTD induction is eCB-dependent (Fig. 1e), we examined whether blockade of CB1Rs attenuated the effect of the LTD-inducing protocol on alcohol-seeking behavior. We found that systemic administration of additional AM251 completely abolished the LTD-induced reduction of active lever presses, alcohol deliveries, and alcohol intake (Fig. 4c and

Supplementary Fig. 10h–i). In contrast, AM251 itself did not alter alcohol-seeking behavior (Supplementary Fig. 10j–l). Lastly, our *ex vivo* results further ascertained that the paired-pulse ratio was significantly increased (Fig. 4d) and that the frequency of spontaneous miniature EPSCs was decreased (Fig. 4e) 2 days after delivery of the LTD protocol, confirming a presynaptically expressed striatal eCB-LTD. These data indicate that delivery of the *in vivo* LTD-inducing protocol at the corticostriatal synapses within the DMS produced a long-lasting suppression of alcohol-seeking behavior.

***In vivo* deliveries of LTP and LTD protocols cause plasticity preferentially in DMS D1-MSNs**

The DMS contains D1- and D2-MSNs, which have been reported to exert opposite effects on drug and alcohol drinking behaviors^{16, 36}. We thus explored how LTP or LTD induction altered glutamatergic transmission in these two neuronal types.

First, to examine *ex vivo* LTP induction in D1- and D2-MSNs, we infused AAV-DIO-ChR2-mCherry into the DMS of *Drd1a*- and *Drd2-Cre* transgenic mice, to enable selective depolarization of D1- or D2-MSNs. Paired eHFS and oPSD induced significant LTP, which did not differ between D1- and D2-MSNs (Fig. 5a,b). This promoted us to explore how synaptic transmission changed in these two neuronal populations following *in vivo* LTP induction. To achieve this, we infused Chronos into the mPFC and Chrimson into the DMS of adult rats, as described above (Fig. 3a). D1-MSNs were labeled by retrograde beads infused into the substantia nigra pars reticulata (SNr) (Fig. 5c, left), whereas D2-MSNs were labeled by infusion of AAV-D2SP-eYFP (Fig. 5c, right). Two days after *in vivo* delivery of the LTP-inducing protocol, the AMPAR-EPSC amplitude and the AMPAR/NMDAR ratio were increased in D1-MSNs, but not in D2-MSNs (Fig. 5d,e), as compared with slices from control rats that were not exposed to light stimulation. This cell type-specific LTP induction is likely attributable to the higher GluN2B/NMDA ratio in D1-MSNs than in D2-MSNs (Fig. 5f), as alcohol-mediated enhancement of GluN2B promotes LTP induction^{9, 37}. Collectively, these data indicate that the *in vivo* LTP protocol potentiated synaptic transmission selectively in D1-MSNs.

We next investigated whether LTD was also preferentially induced in D1-MSNs *ex vivo* and *in vivo*. To induce oPSD selectively in rat D1-MSNs, we infused a retrograde AAV encoding Cre (AAV5-Cre) into the SNr and a Cre-inducible AAV expressing Chrimson (AAV-Flex-Chrimson-tdTomato)¹⁸ into the DMS. We found that in DMS slices from alcohol-naïve rats, a protocol (eHFS+oPSD+MK801+raclopride) that was similar to that used to successfully induce LTD (Fig. 1e) caused a robust LTD in D1-MSNs; this LTD was abolished by AM251 (Fig. 6a,b). To induce oPSD specifically in D2-MSNs, we infused AAV-D2SP-ChR2³⁸ into the DMS of alcohol-naïve rats. We found that the same protocol of eHFS+oPSD+MK801+raclopride did not produce any LTD in the D2-MSNs (Fig. 6a,c), which was consistent with previous reports^{21, 39}.

Lastly, to ascertain whether *in vivo* LTD induction caused glutamatergic depression in D1-MSNs, we measured corticostriatal EPSCs in DMS slices prepared two days after *in vivo* delivery of the LTD-inducing protocol. We found that the LTD protocol reduced the release probability, as indicated by the increased paired-pulse ratio, and reduced the mEPSC frequency in D1-, but not D2-, MSNs, as compared to neurons from control animals without

LTD induction (Fig. 6d–g). These results indicate that *in vivo* LTD induction leads to long-lasting depression of corticostriatal inputs selectively onto DMS D1-MSNs.

Taken together, our results suggest that *in vivo* delivery of optogenetic LTP and LTD protocols preferentially induced plasticity in DMS D1-MSNs.

Selective LTP and LTD induction in DMS D1-MSNs produces long-lasting changes in controls alcohol-seeking behavior

Finally, we examined whether *in vivo* induction of corticostriatal LTP or LTD directly in DMS D1-MSNs altered alcohol-seeking behavior. For oHFS, we infused AAV-Chronos into the mPFC; for oPSD of D1-MSNs, we infused the retrograde AAV5-Cre into the SNr and AAV-Flex-Chrimson into the DMS (Fig. 7a). These infusions led to Chronos expression at the mPFC inputs and selective Chrimson expression in DMS D1-MSNs (Fig. 7b). *In vivo* LTP induction produced significant increases in active lever presses, alcohol deliveries, and alcohol intake at 30 min; this effect persisted for at least 2 days (Fig. 7c and Supplementary Fig. 11a). However, D1-MSN oPSD alone did not alter alcohol-seeking behavior (Fig. 7d and Supplementary Fig. 11b,c). These data suggest that *in vivo* corticostriatal LTP in DMS D1-MSNs caused a long-lasting potentiation of alcohol-seeking behavior.

In contrast, we found that delivery of the *in vivo* LTD-inducing protocol to the mPFC input onto D1-MSNs caused sustained decreases in active lever presses, alcohol deliveries, and alcohol intake at 30 min and 2 days (Fig. 7e and Supplementary Fig. 11d). This behavioral effect was abolished by systemic administration of AM251 (Fig. 7f and Supplementary Fig. 11e,f), which confirmed that eCB signaling regulated this inhibition of alcohol-seeking behavior. These data demonstrate that eCB-LTD in D1-MSNs is required for the long-lasting decrease in alcohol-seeking behavior.

DISCUSSION

In this study, we provide evidence to suggest that alcohol intake induces glutamatergic plasticity, which can be further potentiated by *in vivo* LTP induction, and that this causes long-lasting enhancement of alcohol-seeking behavior (Fig. 8a). In contrast, *in vivo* LTD induction suppresses this plasticity and produces a long-lasting reduction of this behavior. We report that pairing HFS of corticostriatal afferents with oPSD of DMS neurons induces a robust NMDAR-dependent LTP, which masks an eCB-LTD (Fig. 8b). Furthermore, we discovered that LTP and LTD in D1-MSNs contributed to the alteration of alcohol-seeking behavior (Fig. 8b,c). These results provide a direct causal link between long-term synaptic plasticity within a given neural circuit (mPFC → DMS D1-MSNs) and alcohol-seeking behavior. Our findings also demonstrate that induction of D1-MSN LTD might be a potential therapeutic strategy for alcohol use disorder.

oPSD facilitates LTP and LTD induction in the dorsal striatum

It has long been known that dorsostriatal LTP induction proves difficult, possibly due to insufficient depolarization of striatal neurons^{20, 40}. In this study, oPSD was used to strongly depolarize the distal dendrites of these neurons, thus enhancing NMDAR channel opening and calcium influx, which is required for LTP induction^{20, 40} (Fig. 8b). Interestingly,

blockade of LTP by NMDAR antagonists leads to LTD; this is consistent with a study showing that LTP blockade by memantine shifted LTP to LTD⁴¹. However, APV or MK801 was found to shift the plasticity in the current research, but not in the study by Mancini et al.⁴¹. This discrepancy may reflect the fact that eCB-LTD was induced mainly by oPSD in the presence of a D2R antagonist in the current study, and by D2R activation in the previous research. In addition, we report that dopamine D1R signaling plays a critical modulatory role in optogenetic LTP. The observations that blockade of LTP unmasks eCB-LTD and that blockade of eCB-LTD enhances the LTP magnitude suggest that paired HFS and oPSD simultaneously induces NMDAR-dependent LTP and eCB-LTD.

The current oPSD also facilitated LTP in the specific corticostriatal input when we used dual-channel optogenetics. The precise control of both pre- and post-synaptic activity allowed us to reliably induce LTP for the first time. LTP induction is known to activate the expression of immediate early genes such as *Npas4*, which has recently been identified as an important factor in brain plasticity³⁰. Expression of the *Npas4* gene requires Ca²⁺ influx and is associated with drug addiction²⁹. Indeed, the mRNA level *Npas4* gene is increased only after paired oHFS and oPSD, suggesting that oPSD predominantly facilitates LTP at corticostriatal synapses.

Optogenetic LTP induction promotes, and LTD induction suppresses, alcohol-seeking behavior

The corticostriatal circuit is believed to control goal-directed behaviors, including drug-seeking behavior^{2, 5-7}. *In vivo* optogenetic induction of corticostriatal LTP enhances alcohol-seeking behavior, suggesting a link between this plasticity and the behavior. The selective effects of LTP on operant self-administration of alcohol versus sucrose may reflect the distinct activities of these two chemicals on the rats. Operant training with alcohol, but not with sucrose, enhanced NMDAR activity and facilitated subsequent *in vivo* and *ex vivo* induction of LTP. This finding is consistent with previous reports indicating that *ex vivo* or *in vivo* alcohol exposure caused long-term facilitation of NMDAR activity^{6, 10, 31, 32}, which is required for LTP induction in the dorsal striatum⁶ and for operant alcohol self-administration¹⁰. Operant alcohol self-administration induced a smaller, but not significant, increase in AMPAR/NMDAR ratio than did operant sucrose training. This ratio difference might be attributable to the higher NMDAR activity in alcohol-treated rats than in sucrose controls.

How drug (e.g., cocaine)-evoked plasticity affects subsequent LTP induction is likely to depend on the degree of saturation of the plasticity. Our study reveals that operant alcohol self-administration using the FR3 schedule induced glutamatergic plasticity (increased AMPAR/NMDAR ratio). This plasticity was not saturated because the AMPAR/NMDAR ratio was further potentiated by *in vivo* LTP induction, and LTP was induced in slices from alcohol-drinking animals. It is known that 1-2 day(s) withdrawal from cocaine exposure induces silent synapses that contain NMDARs but not AMPARs^{13, 42-44}; these can mature over time (e.g., at 45 days)^{13, 44, 45} and potentially contribute to subsequent LTP occlusion^{46, 47}. These studies suggest that short-term withdrawal from drug exposure induces unsaturated plasticity. Since we induced LTP 24 hours after the last alcohol

exposure, it is not surprising that no occlusion was observed; this is consistent with previous reports^{9, 37}.

Pharmacological inhibition of alcohol-evoked glutamatergic strengthening in the DMS attenuates alcohol consumption⁹. However, this inhibition is transient and disappears as the inhibitory compounds are metabolized. Furthermore, structural plasticity, such as an increased density of mushroom spines, has been observed following alcohol consumption⁴⁸. In this study, *in vivo* eCB-LTD induction elicited a long-lasting decrease in alcohol-seeking behavior, indicating that this plasticity mediates more sustained behavior changes⁴⁹.

LTP and LTD in D1-MSNs affect alcohol-seeking behavior

While the present study and others^{15, 20} report that LTP can be induced in both D1- and D2-MSNs in slices from alcohol-naïve animals, *in vivo* delivery of our LTP-inducing protocol selectively causes long-lasting potentiation of corticostriatal transmission in D1-MSNs of alcohol-drinking rats. This selectivity may be attributed to the fact that alcohol consumption potentiates NMDAR activity in D1-, but not D2-, MSNs¹⁶. The current study further identified that alcohol consumption specifically potentiated GluN2B-containing NMDAR activity at the mPFC input onto D1-MSNs. Alcohol-mediated potentiation of GluN2B-NMDARs was reported to facilitate LTP induction^{9, 37}. Our *in vivo* LTD-inducing protocol also caused LTD in D1-, but not D2-, MSNs because we included a D2R antagonist, which blocks LTD induction in D2-MSNs^{15, 21} (Fig. 8c,d). Our findings are in agreement with a recent report showing that eCB-LTD was induced at corticostriatal inputs to D1-, but not D2-, MSNs²⁶. Given that D1-MSNs positively control alcohol consumption¹⁶, it is not surprising that induction of D1-MSN LTP produces long-lasting enhancement of operant alcohol self-administration, while LTD induction in this neuronal type reduces the same behavior. The induction of LTP/LTD by inducing oPSD selectively in D1-MSNs confirmed that synaptic plasticity in this neuronal type is sufficient to control alcohol-seeking behavior in a bi-directional manner. Therefore, blockade of striatal LTP induction and promotion of eCB-LTD in D1-MSNs may inspire a therapeutic strategy to cause a long-lasting reduction of alcohol-seeking behavior. Although optogenetic intervention cannot be immediately translated to human use, deep brain stimulation (DBS) is an FDA-approved treatment that has the potential to cause LTP⁴⁶, and probably LTD. Thus, we believe that the combined use of DBS and antagonists of the NMDAR (e.g., memantine) and D2R may provide novel clinical treatments for alcohol use disorder.

In summary, we have demonstrated that optogenetic induction of bidirectional long-term synaptic plasticity at corticostriatal afferents within the DMS produced long-lasting increases or decreases in alcohol-seeking behavior. Importantly, we show that the plasticity of DMS D1-MSNs controls alcohol-seeking behavior. Our research establishes a causal link between corticostriatal synaptic potentiation and alcohol-seeking behavior and provides an evidence base for therapeutic strategies to reduce excessive alcohol consumption.

ONLINE METHODS

Reagents

AAV5-CaMKIIa-C1V1(E122T/E162T)-eYFP (3×10^{12} vg/ml), AAV8-Syn-Chronos-GFP (5.6×10^{12} vg/ml), AAV8-Syn-Chrimson-tdTomato (5.5×10^{12} vg/ml), AAV8-Syn-Flex-Chrimson-tdTomato (4.1×10^{12} vg/ml) and AAV-EF1a-DIO-ChR2-mCherry (2×10^{12} vg/ml) were purchased from the University of North Carolina Vector Core. AAV5-CAG-GCaMP6s (2.2×10^{13} vg/ml) and AAV5-CMV-Cre-eGFP (4.9×10^{12} vg/ml) were obtained from the University of Pennsylvania Vector Core. AAV8-D2SP-eYFP (2.5×10^{12} vg/ml) and AAV8-D2SP-eChR2 (H134R)-eYFP (2.5×10^{12} vg/ml) were purchased from Gene Vector and Virus Core of Stanford University School of Medicine. NBQX and APV were purchased from R&D systems. Tetrodotoxin (TTX) was obtained from Tocris. Alexa Fluor 594 was purchased from Invitrogen. MK801, sulpiride, raclopride and the other reagents were obtained from Sigma.

Animals

Male Long-Evans rats (3 months old, Harlan Laboratories) and *Drd1a-Cre* (D1-Cre) and *Drd2-Cre* (D2-Cre) transgenic mice (C57BL/6 background, 3 months old, Mutant Mouse Regional Resource Centers) were used. Both rats (2/cages) and mice (5/cage) were group-housed. All animals were kept in a temperature- and humidity-controlled environment with a light: dark cycle of 12 h (lights on at 7:00 a.m.), and food and water available *ad libitum*. All behavior experiments were conducted in their light cycle, and animals had no history prior to the behavior reported in this paper. All animal care and experimental procedures were approved by the Texas A&M University Institutional Animal Care and Use Committee and were conducted in agreement with the National Research Council *Guide for the Care and Use of Laboratory Animals*.

Stereotaxic virus infusion

The stereotaxic viral infusion was performed as described previously⁸. Depending on experimental design, viruses or beads were infused into the mPFC (AP: +3.2 and +2.6 mm, ML: ± 0.65 mm, DV: -4.0 mm from Bregma), the DMS (AP1: +1.2 mm, ML1: ± 1.9 mm, DV1: -4.7 mm; AP2: +0.36 mm, ML2: ± 2.3 mm, DV2: -4.7 mm), and the SNr (AP1: -4.92 mm, ML1: ± 2.3 mm, DV1: -8.3 mm; AP2: -5.5 mm, ML2: ± 2.0 mm, DV2: -8.6 mm) for rats. For mice, the viruses were infused into the DMS (AP1: +1.18 mm, ML1: ± 1.3 mm, DV1: -2.9 mm; AP2: +0.38 mm, ML2: ± 1.55 mm, DV2: -2.9 mm from Bregma). 0.5-1 μ l of the virus was infused bilaterally at a rate of 0.08 μ l/min. At the end of the infusion, the injectors remained at the site for 10 min to allow for virus diffusion. Animals infused for electrophysiology were maintained in their home cages for 6-8 weeks before recordings. For animals infused with viruses for behavioral experiments, we started training them to self-administer alcohol or sucrose one week after surgery.

Slice preparation

The procedure has been described previously^{8, 50, 51}. Briefly, coronal sections of the striatum (250 μ m in thickness) were cut in an ice-cold solution containing the following (in mM): 40

NaCl, 143.5 sucrose, 4 KCl, 1.25 NaH₂PO₄, 26 NaHCO₃, 0.5 CaCl₂, 7 MgCl₂, 10 glucose, 1 sodium ascorbate, and 3 sodium pyruvate, saturated with 95% O₂ and 5% CO₂. Slices were then incubated in a 1:1 mixture of cutting solution and external solution at 32°C for 45 min. The external solution was composed of the following (in mM): 125 NaCl, 4.5 KCl, 2 CaCl₂, 1 MgCl₂, 1.25 NaH₂PO₄, 25 NaHCO₃, 15 sucrose and 15 glucose, saturated with 95% O₂ and 5% CO₂. Slices were maintained in external solution at room temperature until use.

Field potential recording

For LTP experiments, extracellular field recordings were conducted as previously described⁸. Specifically, the recording used a patch pipette filled with 1 M NaCl and was placed within the DMS. DMS slices were visualized under an epifluorescent microscope (Examiner A1, Zeiss, Germany). Bipolar stimulating electrodes were positioned 100–150 μm away from the recording electrode. Field excitatory postsynaptic potentials/population spikes (fEPSP/PS)⁵² were evoked by electrical stimuli through stimulating electrodes at 0.05 Hz. Picrotoxin (100 μM) was bath applied to block GABAergic transmission. The dopamine D2R antagonist, sulpiride (20 μM), was included externally for all experiments conducted in Figures 1, 5a, and Supplementary Figures 1b, 4. Optical stimuli (2 ms, 405 nm) were delivered through the objective lens to elicit fEPSP/PS. fEPSP/PS were measured using a MultiClamp 700B amplifier with Clampex 10.4 software (Molecular Devices). After a stable baseline had been established for 10 min, high-frequency stimulation (HFS) was delivered through the stimulating electrodes or objective lens to induce LTP. HFS consists of 4 trains of stimuli repeated at an interval of 20 sec. Each train contains 100 pulses at 100 Hz (electrically HFS, eHFS) or 50 Hz (optogenetic HFS, oHFS). For pairing experiments, optogenetic postsynaptic depolarization (oPSD) was induced by light stimulation (590 nm, 1 sec for eHFS or 2 sec for oHFS) of DMS neurons through the objective lens.

Whole-cell recording

In Supplementary Figures 1, 2, and 5b, we used a K⁺-based intracellular solution, containing (in mM): 123 potassium gluconate, 10 HEPES, 0.2 EGTA, 8 NaCl, 2 MgATP, 0.3 NaGTP (pH 7.3). All other experiments utilized the Cs-based solution, containing (in mM): 119 CsMeSO₄, 8 TEA.Cl, 15 HEPES, 0.6 EGTA (10 BAPTA for Supplementary Figures 3d,e), 0.3 Na₃GTP, 4 MgATP, 5 QX-314.Cl, 7 phosphocreatine, 0.05 Alexa Fluor 594 (Supplementary Figures 3a–e), and 0.1 spermine (Supplementary Figure 9). The pH was adjusted to 7.3 with CsOH. Neurons were clamped at –70 mV.

For measuring NMDAR-EPSPs in distal dendrites, Alexa Fluor 594 was infused through patch-pipettes into the recorded neurons to label their dendrites. Under the guidance of fluorescence, the stimulating electrodes were positioned close to the Alexa Fluor-labeled dendrites and were 100–150 μm away from the soma. AMPA receptor (AMPA)-mediated EPSPs were recorded in 1.0 mM extracellular Mg²⁺. NBQX (10 μM) was then bath applied to block AMPAR-EPSPs. Next, simultaneous presynaptic electrical stimulation and oPSD of striatal neurons induced a response that was mediated by a C1V1-induced depolarization (V_{C1V1}) plus an NMDAR response (EPSP_{NMDA}). Lastly, the EPSP_{NMDA} component was blocked by bath application of APV (50 μM), and V_{C1V1} was isolated. The optogenetic-mediated EPSP_{NMDA} was calculated by digital subtraction of V_{C1V1} from $V_{C1V1} +$

EPSP_{NMDA}. The input-output relationships for AMPAR-mediated excitatory postsynaptic currents (EPSCs) were measured at 5 different stimulating laser powers. For measurement of the AMPAR/NMDAR ratio, the peak currents of AMPAR-mediated EPSCs were measured at a holding potential of -70 mV and the NMDAR-mediated EPSCs were estimated as the EPSCs at $+40$ mV at 30 ms after the peak AMPAR-EPSCs, when the contribution of the AMPAR component was minimal. The AMPA/NMDA ratio was calculated by dividing the NMDAR-EPSC by AMPAR-EPSC. To measure the GluN2B/NMDA ratio, NMDAR-EPSCs were recorded in the absence and presence of Ro 25-6981, and GluN2B-EPSCs were calculated by subtraction of these two responses. For measuring mEPSCs, we added TTX (1 μ M) to the external solution to suppress action potential-driven release. The paired-pulse ratio (PPR) was calculated by dividing the second light-evoked EPSC by the first with 100-ms intervals between the two. To measure above synaptic transmission in specific D1- and D2-MSNs (Fig. 5, 6), we first patched bead-positive (D1-MSNs) or eYFP-positive neurons (D2-MSNs) within a DMS area containing strong green mPFC fibers (expressing Chronos-GFP) and red Chrimson-positive neurons. We then sequentially delivered 405-nm light to stimulate the mPFC inputs and 590-nm light to induce Chrimson-mediated oPSD. The synaptic inputs and oPSD were distinguished using 2- and 100-ms light stimulation, since the prolonged light stimulation increased the duration of oPSD, but not of synaptic transmission⁵³. Only those neurons that received mPFC inputs and exhibited oPSD were selected for further experiments. At the end of the recording, NBQX was applied to confirm synaptic transmission induced by 405-nm light stimulation. To measure AMPAR rectification, AMPAR-EPSCs were recorded at three holding potentials of -70 , 0 , and $+40$ mV in the presence of APV (50 μ M). Rectification index of the AMPAR-EPSC was calculated by plotting the EPSC magnitude at these potentials, and using the slope of the lines connecting the data between -70 - 0 mV and between 0 - 40 mV to calculate the ratio.

Calcium image

An AAV-C1V1²³ and an AAV-GCaMP6s⁵⁴ were infused into the DMS. Whole-cell recordings were made in C1V1-expressing neurons. The GCaMP6s measures the calcium signal that is induced by current injection (iPSD), or optogenetic depolarization (oPSD) (Supplementary Figure 2). In supplementary Figures 3f–g, fluorescent Ca^{2+} signals were elicited by eHFS or eHFS+oPSD without whole-cell recording. The distal dendrite (~ 120 μ m from the soma) was chosen for analysis. Ca^{2+} signals were acquired and analyzed with the Zen program (Zeiss) and Origin software (Origin Lab Corporation, MA), and calculated as previously described⁵⁵. The fluorescence signals were quantified by measuring the mean pixel intensities of the circular regions of interest (ROI). Fluorescence intensity is expressed as $\Delta F/F$ values vs. time, where F is the baseline fluorescence and ΔF is the baseline-subtracted fluorescence.

Operant self-administration of alcohol

After one week recovery from viral infusions, Long-Evans rats were trained to self-administer a 20% alcohol solution in operant self-administration chambers as described⁵⁶. Each chamber contains two levers; an active lever, in which presses result in a delivery of 0.1 ml of the alcohol solution, and an inactive lever, in which presses are recorded, but no programmed events occur. After 48-h of exposure to 20% alcohol in the home cage, and one

overnight session in the chamber in which pressing the active lever delivers 0.1 ml of water in a fixed ratio 1 (FR1), operant sessions were conducted 5 days per week for two weeks in a FR1 schedule with an active lever press resulting in the delivery of 20% alcohol with sessions shortened from 3 h to 30 min. Following the first two weeks, operant sessions were run three days per week for one week, and the schedule requirement was increased to FR3. After one week of FR3 training, rats underwent surgery for optical fiber implantation. FR3 training was resumed one week after the surgery. Once a stable baseline of active lever presses was achieved, animals underwent *in vivo* LTP and LTD induction. Following the induction, some rats were continuously monitored with their operant behavior for 7–9 days, while other rats were euthanized at day 2 post-induction for electrophysiology recordings. To test drugs' effect without LTP/LTD induction, we systematically administered them 30 min before the operant behavior test. Simultaneously, we also measured inactive lever presses before and after treatment (Supplementary Figure 12).

Operant self-administration of sucrose

After a one-week recovery from viral infusions, Long-Evans rats were trained to self-administer a 2% sucrose in operant chambers using the same procedure as the alcohol group described above. Optical fiber implantation was also conducted in an identical manner to the alcohol group.

Optical fiber implantation

One-week following operant training with the FR3 schedule, animals were anesthetized with isoflurane and mounted in a stereotaxic frame. An incision was made, bilateral optical fiber implants (300-nm core fiber secured to a 2.5-mm ceramic ferrule with 5-mm fiber extending past the end of the ferrule) were lowered into the DMS (AP: +0.36 mm; ML: \pm 2.3 mm; DV: –4.6 mm from Bregma). Implants were secured to the skull with metal screws and dental cement (Henry Schein) and covered with denture acrylic (Lang Dental). The incision was closed around the head cap and the skin vet-bonded to the head cap. Rats were monitored for one week or until they resumed normal activity.

***In vivo* LTP and LTD induction and operant testing**

Once a stable baseline of active lever presses was attained after optical fiber implantation, an LTP/LTD-inducing protocol was delivered 30 minutes before operant testing sessions in a neutral Plexiglass chamber, with no visual cues. LTP-induction consisting of paired oHFS +oPSD used the following protocol: 100 pulses at 50 Hz of 473-nm light (2 ms) with or without constant 590-nm light for 2 sec, repeated 4 times with 20-sec intervals. The protocol was repeated three times with 5-min intervals. LTD induction employed the following protocol: animals were injected with a cocktail of MK-801 (0.1 mg/kg) and raclopride (0.01 mg/kg) 15 minutes before delivery of oHFS and oPSD. The complete LTP/LTD-inducing procedure was performed once and 30 min later animals were allowed to press levers for alcohol in a 30-min session. Operant sessions were repeated every 48 or 72 h until active lever presses returned to their levels prior to the induction.

Measurement of blood alcohol concentration (BAC)

To measure blood alcohol concentration, we used the same procedure as in Figure 3b to train two groups of rats. One week prior to LTP induction, we collected blood samples from the one side of the lateral saphenous vein⁵⁷ in both groups to measure baseline BAC. Thirty minutes after LTP induction, we collected blood samples from the other side of the lateral saphenous in one group of rats. Two days after the LTP induction, we collected blood samples from the other group of rats. BAC was measured using gas chromatography as previously described⁵⁸.

RNA extraction and quantitative PCR (qPCR) analysis

The rats were infused with AAV-Chronos-GFP in the mPFC and AAV-Chrimson-tdTomato in the DMS. Coronal striatal sections (250 μ m) were prepared as described in Slice Preparation section above. A slice was placed in a recording chamber and perfused with the external solution at a flow rate of 3 mL/min. Optical stimuli were delivered through the objective lens to fluorescent DMS areas, which contained both GFP-expressing mPFC axons and tdTomato-expressing neurons using one of the following stimulation protocols: oHFS, oPSD, or oHFS+oPSD. All protocols were repeated three times with 5-min intervals, which is the same as the *in vivo* LTP-inducing protocol. Thirty minutes after completing the optogenetic stimulation, the DMS tissues from experimental and control groups were collected on ice. The RNA isolation, reverse transcription, and the qRT-PCR analyses were performed as described previously⁵⁹. The mRNA level of *Npas4* was normalized against the GAPDH mRNA level in the same sample and presented as fold changes over baseline using the delta-delta CT method. The following primers were used: *Npas4*, Forward: 5'-GAACCTCAAGGAAGTCTGC-3', reverse: 5'-GTGCCTCCAGCAAAGAAGAC-3'; GAPDH, Forward: 5'-TGCCACTCAGAAGACTGTGG-3', reverse: 5'-TTCAGCTCTGGGATGACCTT-3'. For each experimental condition, two slices per rat were treated, and the averaged mRNA values were used.

Histology

Rats with viral and beads infusion were perfused intracardially with 4% paraformaldehyde (PFA) in phosphate-buffered saline (PBS). The brains were taken out and put into 4% PFA/PBS solution for post-fixation overnight at 4°C, followed by dehydration in 30% sucrose solution and cryostat frozen sectioning. The brains were cut into 50- μ m coronal sections. A confocal laser-scanning microscope (Fluorview-1200, Olympus) was used to image these sections with a 470-nm laser to excite eYFP and GFP and with a 593-nm laser to excite Alexa Fluor 594 and tdTomato. All images were processed using Imaris 8.3.1 (Bitplane, Zurich, Switzerland).

Data acquisition and statistics

In electrophysiology experiments, we used 184 rats and 10 D1- and D2-Cre mice, with 10 rats excluded before data collection due to virus expression in the incorrect place or expression that was too weak. In behavioral tests, we used 156 rats, among which 28 were excluded due to lack of alcohol responding in the operant setting (10 active lever presses/session), 6 were removed from data analysis due to death during surgery, and 21 were

removed from the last experiments due to head cap loss. In the imaging experiments, we used 11 rats.

All data are expressed as mean \pm SEM. Each experiment was replicated in 3–7 animals. The data collection was randomized. Data were obtained and analyzed by experimenters who did not know the types of treatments of the animals except Figures 3d–g, 4d, 4e, 5d, 5e, 6d–g. No data points were excluded unless specified, and the only exclusion standard was the health condition of the animal. Data from the repeated experiments for the same sub-study were pooled together for statistics. The sample size for each experiment was determined to be either at least 3 animals or 10 neurons. If animals in Figures 3d–g, 4d, 4e, 5d, 5e, 6d–g were successfully induced *in vivo* changes, we measured responses *ex vivo* in enough neurons to evaluate the effect of light stimulation. The sample size was presented as “n = x, y”, where “x” refers to the number of slices or neurons, and “y” refers to the number of animals. In electrophysiological experiments, 1–4 recordings were performed using slices from a single animal except for Figures 3d, 3e, 4d, 4e. Slice or neuron-based statistics were performed and reported for electrophysiology and animal-based statistics for behavioral data. Normal distribution was assumed and tested. Variance was estimated for most major results, and no significant difference was found between control and manipulation groups. Statistical significance was assessed in electrophysiological studies using the unpaired or paired *t* test, or two-way RM ANOVA followed by Student-Newman-Keuls (SNK) method. Behavioral studies were analyzed using the paired *t* test and one-way RM ANOVA followed by the SNK method. Two-tail tests were performed for all studies. Statistical significance was set at $P < 0.05$.

Life Sciences Reporting Summary

Further information on experimental design is available in the Life Sciences Reporting Summary.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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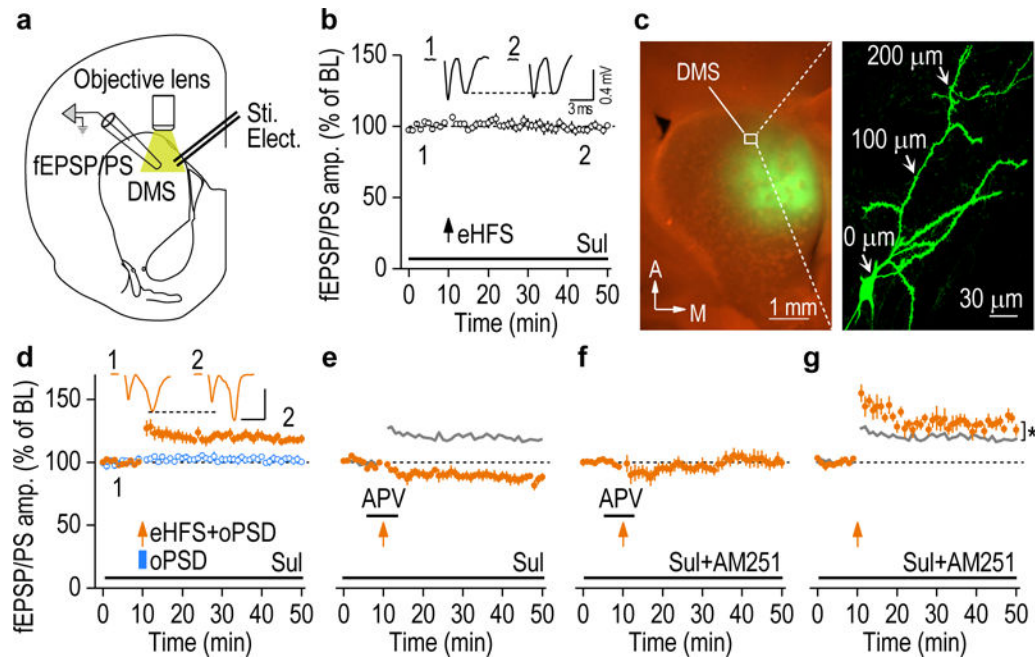


Figure 1.

oPSD facilitated induction of NMDAR-dependent LTP and eCB-LTD in DMS slices. (a) Schematic showing the bipolar stimulating electrodes (Sti. Elect.) used to evoke fEPSP/PS and the objective lens for optogenetic depolarization. (b) Presynaptic eHFS did not potentiate fEPSP/PS ($98.59 \pm 2.39\%$ of baseline [BL], $t_{(7)} = 0.59$, $P = 0.57$; $n = 8$ slices, 6 rats). Inset, sample fEPSP/PS traces at time 1 and 2. Sulpiride (Sul; $20 \mu\text{M}$) was bath-applied to prevent LTD and favor LTP induction in this and following recordings as indicated. (c) Representative fluorescent images showing C1V1-eYFP expression in the DMS (left) and in the full-length dendrites of a DMS neuron (right). The section was counter-stained with NeuroTrace (red). (d) Pairing of presynaptic eHFS and oPSD (1 sec), but not oPSD alone, induced robust LTP. eHFS+oPSD: $118.32 \pm 2.96\%$ of BL, $t_{(9)} = -6.18$, $P = 0.00016$; $n = 10$ slices, 6 rats; oPSD: $101.59 \pm 2.26\%$ of BL, $t_{(6)} = -0.70$, $P = 0.51$; $n = 7$ slices, 3 rats. Scale bars: 3 ms, 0.4 mV. (e) Optogenetic induction of LTP was abolished by APV ($50 \mu\text{M}$), leading to LTD ($86.75 \pm 3.06\%$ of BL, $t_{(6)} = 4.33$, $P = 0.0049$; $n = 7$ slices, 6 rats). The grey line is the control LTP from d for reference. (f) LTD was completely abolished by the CB1R antagonist, AM251 ($3 \mu\text{M}$) ($101.32 \pm 4.71\%$ of BL, $t_{(8)} = -0.28$, $P = 0.79$, $n = 9$ slices, 5 rats). (g) AM251 facilitated LTP induction ($130.73 \pm 2.89\%$ of BL, $t_{(5)} = -10.62$, $P = 0.00013$; compared with the control LTP: $t_{(14)} = -2.79$, $*P = 0.015$; $n = 6$ slices, 3 rats, unpaired t test). Two-sided paired t test for b and d-g, unless otherwise stated. Data are presented as mean \pm s.e.m.

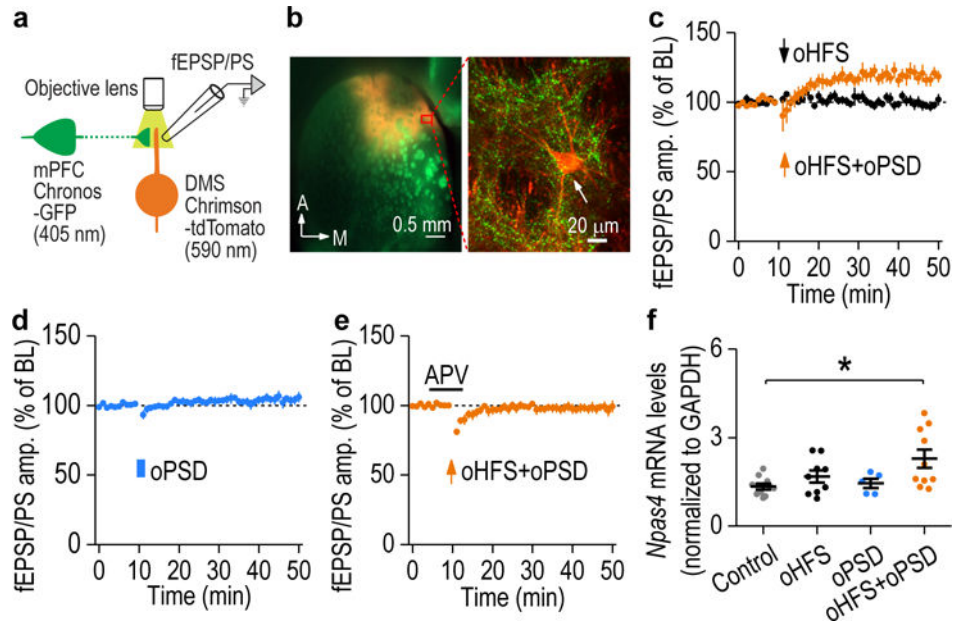


Figure 2.

oPSD facilitated corticostriatal LTP induction in the DMS. **(a)** Schematic illustration of selective pre- and post-synaptic stimulation of DMS corticostriatal synapses using dual-channel optogenetics. AAV-Chronos-GFP was infused into the mPFC and AAV-Chrimson-tdTomato into the DMS of rats. Chronos and Chrimson were activated by 405- and 590-nm light, respectively. **(b)** Confocal fluorescent images showing Chronos-GFP-expressing mPFC fibers (green) and Chrimson-tdTomato-expressing neurons (red) in the DMS. Images shown in b is representative of 3 experiments of 3 rats. **(c)** Pairing oHFS with oPSD produced robust LTP in the DMS ($119.06 \pm 4.69\%$ of baseline [BL], $t_{(6)} = -4.07$, $P = 0.0066$; $n = 7$ slices, 4 rats). oHFS alone did not alter fEPSP/PS ($99.60 \pm 3.12\%$ of BL, $t_{(7)} = 0.13$, $P = 0.90$; $n = 8$ slices, 5 rats). **(d)** oPSD alone did not induce LTP ($104.55 \pm 3.15\%$ of BL, $t_{(6)} = -1.44$, $P = 0.20$; $n = 7$ slices, 3 rats). **(e)** Dual-channel optogenetic induction of LTP was blocked by APV ($98.06 \pm 3.27\%$ of BL; $t_{(8)} = 0.59$, $P = 0.57$; $n = 9$ slices, 5 rats). **(f)** *Npas4* mRNA levels were significantly increased following paired oHFS+oPSD, but not after oHFS or oPSD only. $F_{(3,30)} = 3.86$, $P = 0.019$; $*P < 0.05$; $n = 10$ (Control), 9 (oHFS), 5 (oPSD), and 10 (oHFS+oPSD) rats. Two-sided paired *t* test for c-e; one-way ANOVA followed by SNK test for f. Data are presented as mean \pm s.e.m.

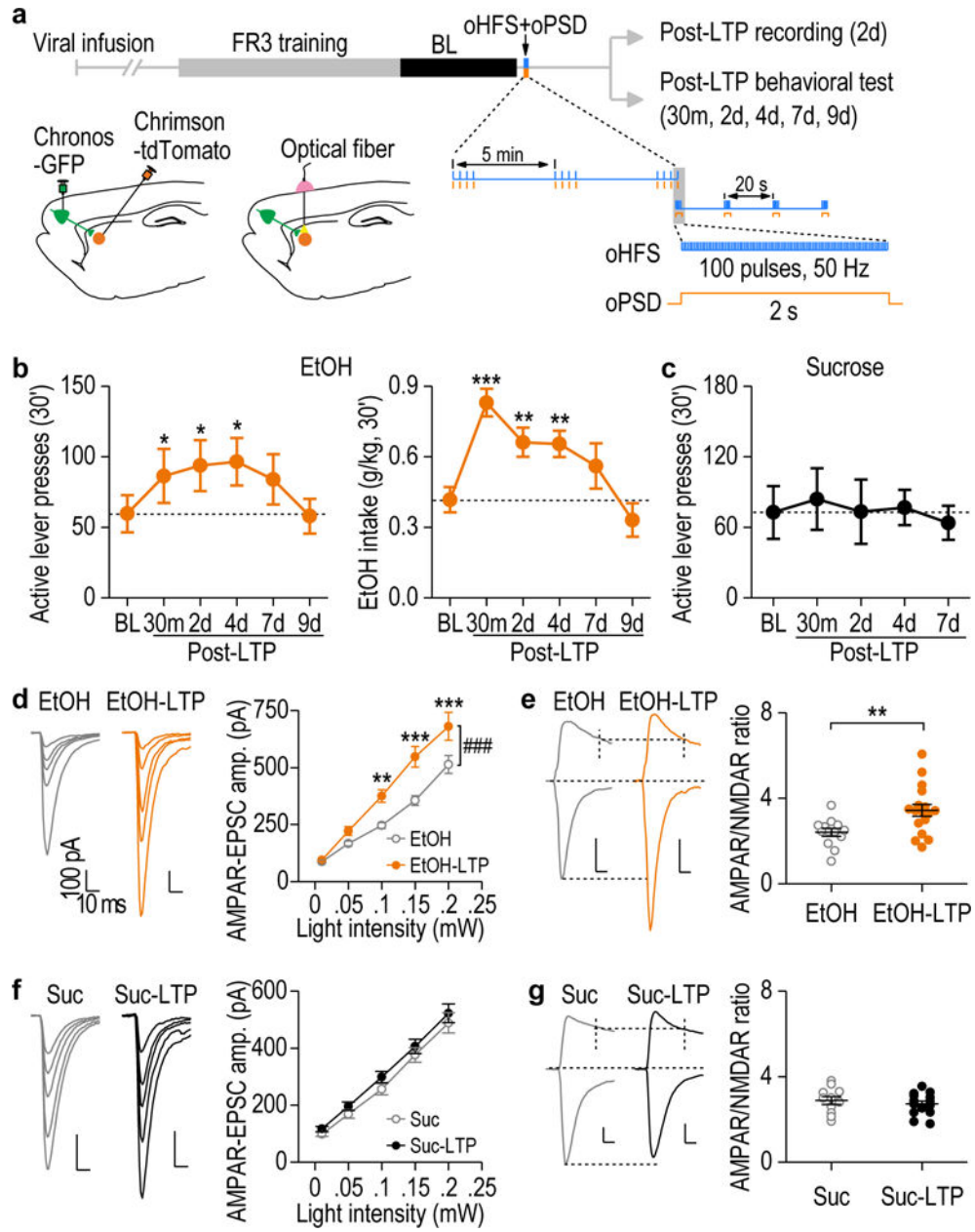


Figure 3. *In vivo* optogenetic LTP induction in the DMS produced a long-lasting increase in operant alcohol self-administration. **(a)** Schematic of the training procedure and *in vivo* LTP induction protocol. **(b)** *In vivo* LTP induction produced long-lasting increases in active lever presses (left, $F_{(5,54)} = 3.26$, $P = 0.012$) and alcohol (EtOH) intake (right, $F_{(5,49)} = 10.91$, $P < 0.0001$). * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ vs. baseline (BL); $n = 14$ rats. **(c)** *In vivo* LTP induction did not alter active lever presses for sucrose. $F_{(4,18)} = 1.10$, $P = 0.39$; $n = 6$ rats. **(d)** *In vivo* LTP induction caused potentiation of AMPAR-EPSCs on day 2 in alcohol-administered rats. Left and middle: Sample traces of AMPAR-EPSCs. Right: Input-output curves for AMPAR-EPSCs with (EtOH-LTP) and without (EtOH) *in vivo* LTP induction. $F_{(1,107)} = 15.62$, $P = 0.0005$; $n = 16$ neurons, 5 rats (EtOH) and 13 neurons, 3 rats (EtOH-

LTP). (e) Representative traces and averaged data showing a significant increase in the AMPAR/NMDAR ratio by LTP induction. $t_{(28)} = -2.88$, $P = 0.0075$; $n = 13$ neurons, 5 rats (EtOH) and 17 neurons, 3 rats (EtOH-LTP). (f) *In vivo* LTP induction did not change AMPAR-EPSCs from the sucrose-administered rats. $F_{(1,99)} = 1.30$, $P = 0.27$; $n = 11$ neurons, 4 rats (Suc) and 16 neurons, 5 rats (Suc-LTP). (g) AMPAR/NMDAR ratio did not change in the sucrose group after *in vivo* LTP induction. $t_{(23)} = 0.73$, $P = 0.47$; $n = 11$ neurons, 5 rats (Suc) and 14 neurons, 5 rats (Suc-LTP). Scale bars: 10 ms, 100 pA for d-g. One-way RM ANOVA for b, c; two-way RM ANOVA for d, f; Two-sided unpaired t test for e, g. Data are presented as mean \pm s.e.m.

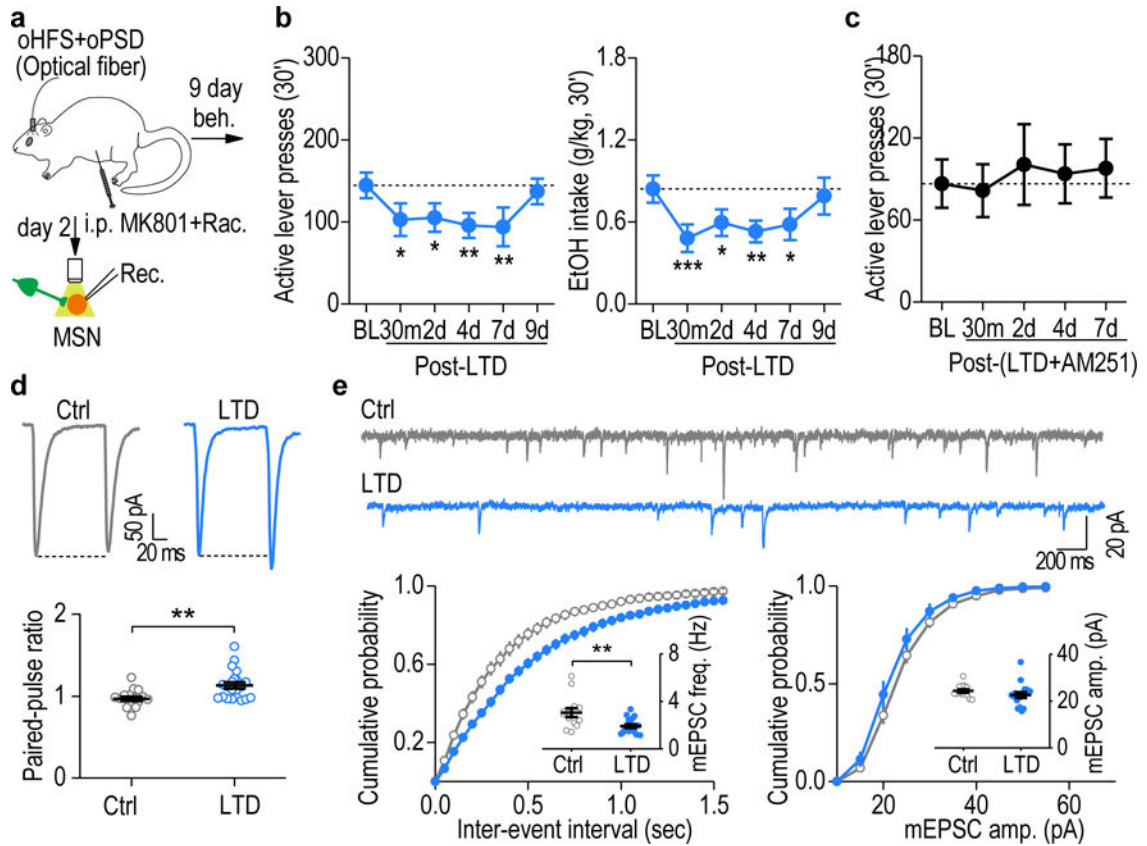


Figure 4.

In vivo LTD induction caused a long-lasting reduction of alcohol-seeking behavior in an eCB-dependent manner. (a) Schematic of the *in vivo* LTD-inducing protocol (oHFS+oPSD +MK801+raclopride) and *ex vivo* LTD measurement. MK801 (0.1 mg/kg) and raclopride (0.01 mg/kg) were systemically administered 15 min before the optogenetic stimulation. (b) Delivery of the *in vivo* LTD-inducing protocol in the DMS produced a long-lasting decrease in active lever presses (left, $F_{(5,38)} = 3.89$, $P = 0.006$) and alcohol intake (right, $F_{(5,35)} = 5.17$, $P = 0.0012$). * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ vs. baseline (BL); $n = 9$ rats. (c) Delivery of the *in vivo* LTD-inducing protocol in the presence of AM251 failed to alter active lever presses. $F_{(4,31)} = 0.40$, $P = 0.81$; $n = 9$ rats. (d,e) Delivery of the *in vivo* LTD-inducing protocol produced a long-lasting depression of glutamatergic transmission in the DMS on day 2 post-stimulation. (d) Top, sample traces showing paired-pulse ratios (100-ms inter-stimulus interval) measured in fluorescent neurons from LTD-induced rats and their controls (without light stimulation). Bottom, averaged data showing an increased paired-pulse ratio after LTD induction. $t_{(36)} = -3.31$, $P = 0.0021$; $n = 17$ neurons, 3 rats (Ctrl) and 21 neurons, 5 rats (LTD). (e) Top, representative traces of mEPSCs in fluorescent neurons in LTD-induced and control rats. Bottom, cumulative distributions of inter-event intervals and amplitudes of mEPSCs. Inset, reduced frequency (left), but not amplitude (right), of mEPSCs after *in vivo* LTD induction. $t_{(28)} = 2.97$, $P = 0.006$ for frequency; $t_{(28)} = 1.11$, $P = 0.28$ for amplitude; $n = 13$ neurons, 3 rats (Ctrl) and 17 neurons, 3 rats (LTD). One-way RM ANOVA for b, c; Two-sided unpaired t test for d, e. Data are presented as mean \pm s.e.m.

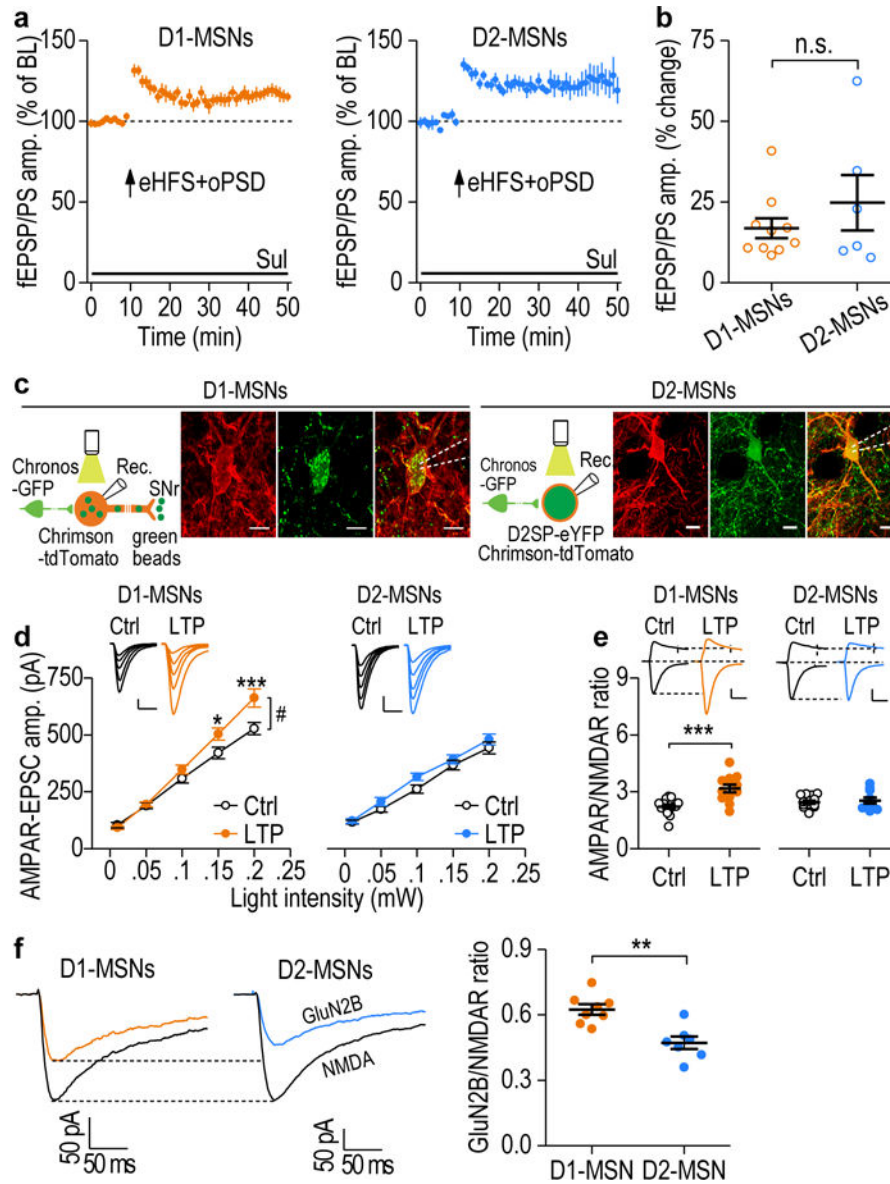


Figure 5.

Corticostriatal LTP was preferentially induced in DMS D1-MSNs. **(a)** AAV-DIO-ChR2-mCherry was infused into the DMS of D1-Cre and D2-Cre mice. Pairing of eHFS and oPSD induced LTP in both D1-MSNs ($116.88 \pm 3.07\%$ of baseline [BL], $t_9 = -5.49$, $P = 0.00038$; $n = 10$ slices, 5 mice) and D2-MSNs ($124.79 \pm 8.58\%$ of BL, $t_5 = -2.89$, $P = 0.034$; $n = 6$ slices, 5 mice). **(b)** There was no significant difference (n.s.) in D1- and D2-MSN LTP. $t_{(14)} = -1.04$, $P = 0.32$. **(c)** Left, experimental design and sample images of retrograde bead labeling of a D1-MSN. AAV-Chronos-GFP, AAV-Chrimson-tdTomato, and green beads were infused into the mPFC, DMS, and SNr, respectively. Right, experimental design and sample images of labeling of a D2-MSN with D2SP-eYFP. AAV-D2SP-eYFP was infused into the DMS. Scale bar: $10 \mu\text{m}$. **(d)** Left, *In vivo* LTP induction resulted in higher AMPAR-EPSC amplitudes in D1- (left), but not D2- (right), MSNs from LTP-induced (LTP) rats, as compared with rats that were not exposed to light stimulation (Ctrl). Both groups of rats

were trained to self-administer alcohol. D1-MSNs: $F_{(1,95)} = 4.72$, $P = 0.04$; $n = 13$ neurons, 4 rats (Ctrl) and 13 neurons, 5 rats (LTP). D2-MSNs: $F_{(1,90)} = 2.11$, $P = 0.16$; $n = 14$ neurons, 4 rats (Ctrl) and 11 neurons, 5 rats (LTP). Scale bars: 20 ms, 100 pA. (e) Left, the AMPAR/NMDAR ratio increased in D1- (Left), but not D2- (Right), MSNs after *in vivo* LTP induction. D1-MSNs: $t_{(25)} = -4.43$, $P = 0.00016$; $n = 15$ neurons, 5 rats (Ctrl) and 12 neurons, 5 rats (LTP). D2-MSNs: $t_{(22)} = -0.52$, $P = 0.61$; $n = 14$ neurons, 5 rats (Ctrl) and 10 neurons, 5 rats (LTP). Scale bars: 20 ms, 100 pA. (f) Operant alcohol self-administration resulted in a higher GluN2B/NMDA ratio in D1-MSNs than in D2-MSNs. Left and middle, sample trace of NMDAR-EPSCs in the absence or presence of Ro 25-6981 (0.5 μM). Right, summarized data of the ratios in D1-MSNs and D2-MSNs. $t_{(13)} = 4.16$, $P = 0.0011$; $n = 8$ neurons, 5 rats (D1-MSNs), $n = 7$ neurons, 5 rats (D2-MSNs). Two-sided paired t test for a; Two-sided unpaired t test for b, e, f; two-way RM ANOVA for d. Data are presented as mean \pm s.e.m.

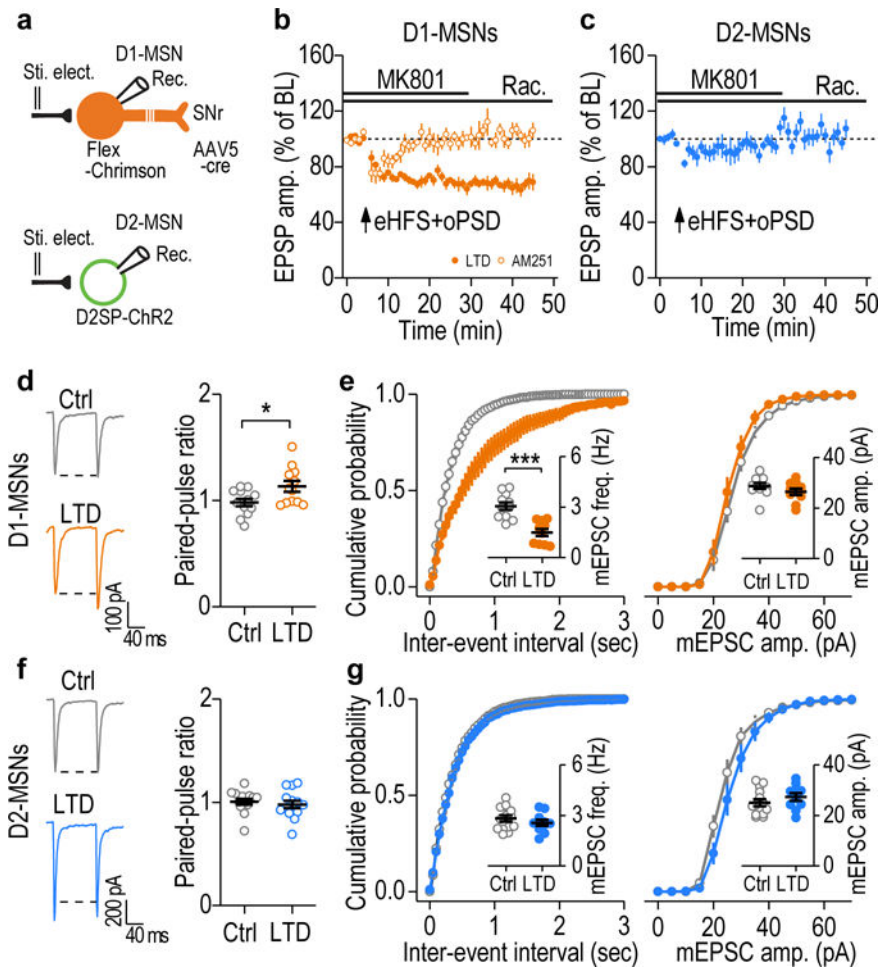


Figure 6.

Corticostriatal LTD was preferentially induced in DMS D1-MSNs. (a) Schematic of viral infusion and whole-cell recordings of rat D1- or D2-MSNs. (b) Paired eHFS and oPSD in the presence of NMDAR and D2R antagonists caused a robust LTD in DMS D1-MSNs ($66.99 \pm 4.56\%$ of baseline [BL], $t_7 = 7.23$, $P = 0.00017$; $n = 8$ neurons, 5 rats), which was completely abolished by AM251 ($3 \mu\text{M}$) ($102.82 \pm 4.28\%$ of BL, $t_6 = -0.66$, $P = 0.53$; $n = 7$ neurons, 3 rats). (c) Paired eHFS and oPSD in the presence of NMDAR and D2R antagonists did not induce LTD in D2-MSNs ($100.08 \pm 5.15\%$ of BL, $t_7 = -0.02$, $P = 0.99$; $n = 8$ neurons, 3 rats). (d) Sample traces and averaged data showing an increased paired-pulse ratio in D1-MSNs 2 days after *in vivo* LTD induction. $t_{22} = -2.45$, $P = 0.023$; $n = 12$ neurons, 4 rats (Ctrl) and 12 neurons, 3 rats (LTD). (e) *In vivo* optogenetic LTD induction reduced the mEPSC frequency (left), but not the mEPSC amplitude (right) of DMS D1-MSNs. $t_{19} = 5.00$, $P < 0.0001$ for frequency; $t_{19} = 1.30$, $P = 0.21$ for amplitude; $n = 11$ neurons, 4 rats (Ctrl) and 10 neurons, 3 rats (LTD). (f) *In vivo* LTD induction did not change paired-pulse ratios in D2-MSNs. $t_{26} = 0.59$, $P = 0.56$; $n = 14$ neurons, 4 rats (Ctrl) and 14 neurons, 3 rats (LTD). (g) *In vivo* LTD induction did not alter the mEPSC frequency (left, inset) or the mEPSC amplitude (right) in DMS D2-MSNs. $t_{22} = 0.95$, $P = 0.35$ for frequency; $t_{22} = -1.11$, $P = 0.28$ for amplitude; $n = 14$ neurons, 4 rats (Ctrl) and 10

neurons, 3 rats (LTD). Two-sided paired *t* test for b, c; Two-sided unpaired *t* test for d-g.
Data are presented as mean \pm s.e.m.

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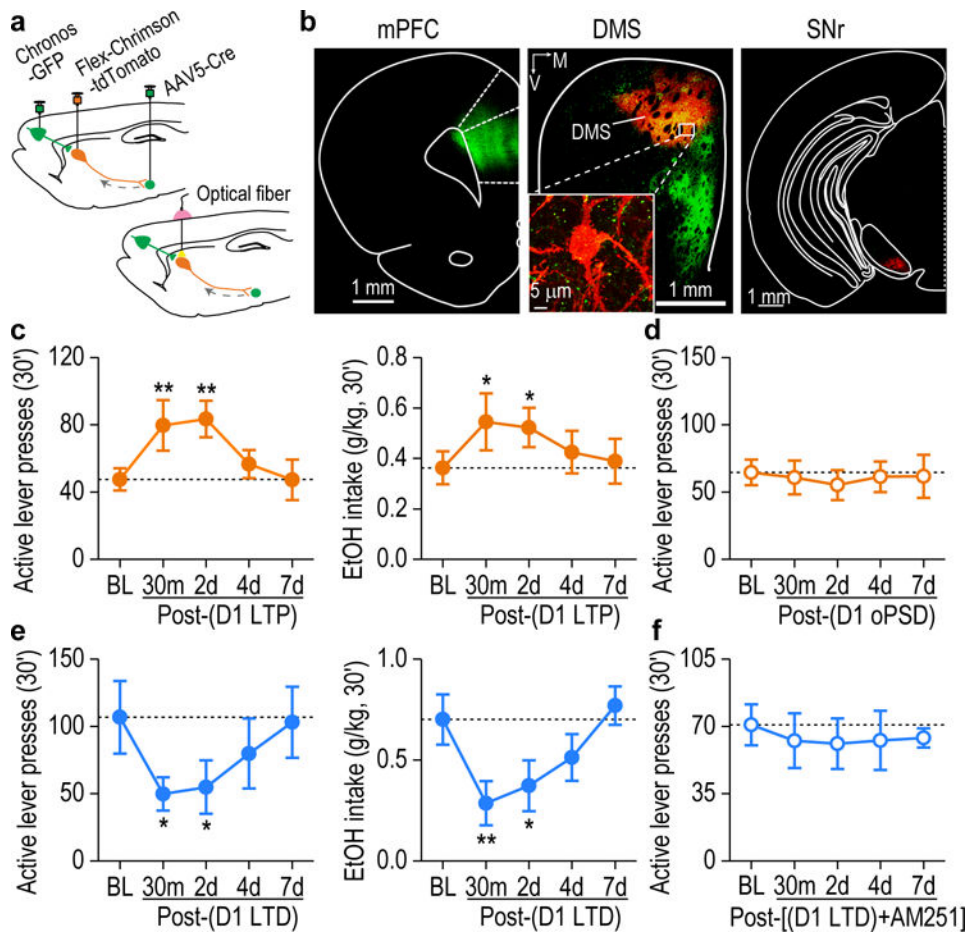


Figure 7. Selective *in vivo* LTP or LTD induction in D1-MSNs produced a long-lasting control of alcohol-seeking behavior. **(a)** Schematic showing viral infusion (top) and optical fiber implantation (bottom). **(b)** Representative fluorescent images showing that mPFC (left) and SNr infusions produced Chronos-expressing fibers (green) and Chrimson-tdTomato expression (red) in DMS D1-MSNs (middle), which project to the SNr (right). Images shown in b is representative of 3 experiments of 3 rats. **(c)** Paired oHFS of mPFC inputs and oPSD of D1-MSNs induced long-lasting increases in active lever presses (left, $F_{(4,24)} = 6.02$, $P = 0.0017$) and alcohol intake (right, $F_{(4,24)} = 3.74$, $P = 0.017$). * $P < 0.05$, ** $P < 0.01$ vs. baseline (BL); $n = 8$ rats. **(d)** *In vivo* delivery of oPSD alone in DMS D1-MSNs did not alter active lever presses for alcohol. $F_{(4,18)} = 0.36$, $P = 0.83$; $n = 6$ rats. **(e)** Delivery of the *in vivo* LTD-inducing protocol to mPFC inputs onto DMS D1-MSNs produced a long-lasting attenuation in active lever presses (left, $F_{(4,23)} = 4.07$, $P < 0.0001$) and alcohol intake (right, $F_{(4,15)} = 6.67$, $P = 0.0027$). * $P < 0.05$, ** $P < 0.01$ vs. BL; $n = 7$ rats for active lever presses and $n = 6$ rats for alcohol intake. **(f)** *In vivo* delivery of the LTD-inducing protocol in the presence of AM251 (oHFS+oPSD+MK801+raclopride+AM251) failed to alter active lever presses for alcohol. $F_{(4,22)} = 0.12$, $P = 0.97$, $n = 7$ rats. One-way RM ANOVA for c-f. Data are presented as mean \pm s.e.m.

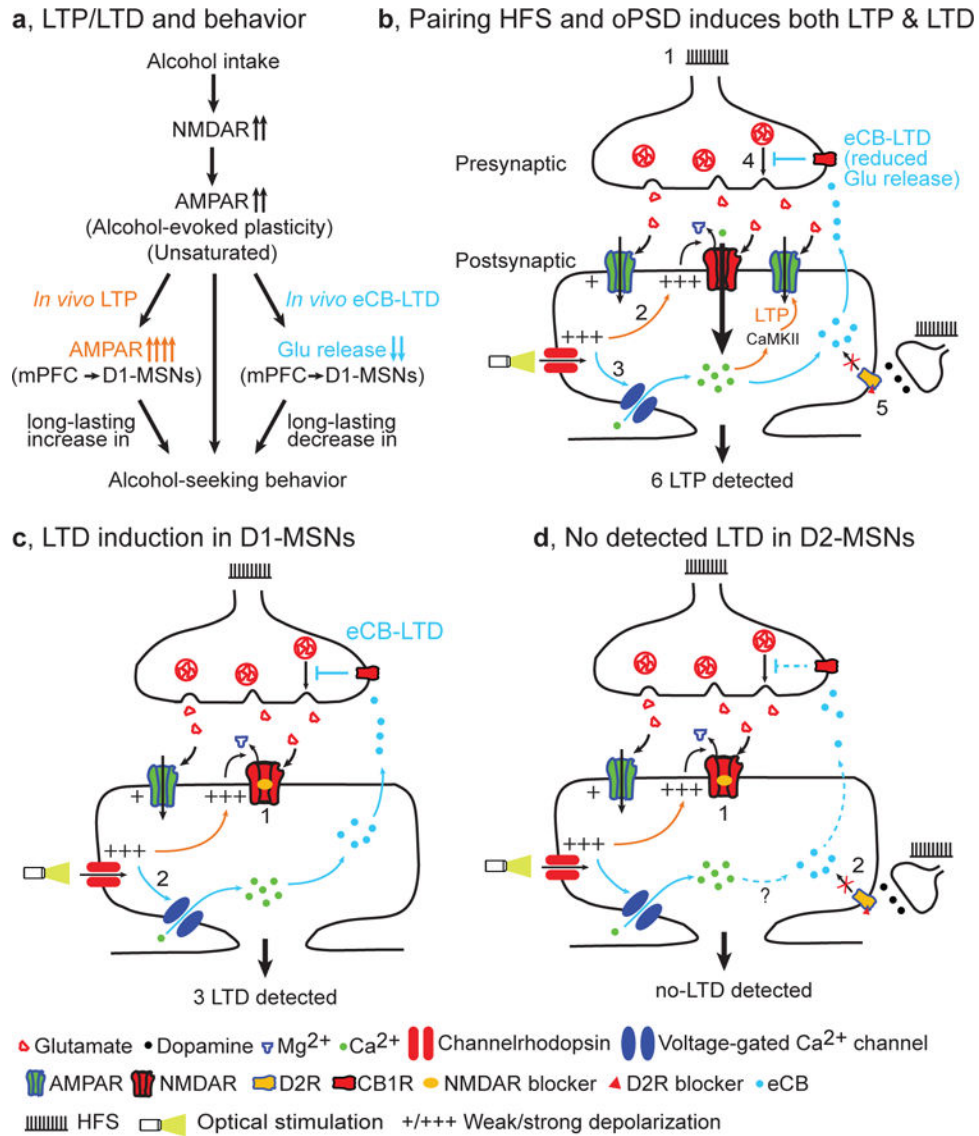


Figure 8. Model of bidirectional and long-lasting control of alcohol-seeking behavior by corticostriatal plasticity. (a) Alcohol intake facilitates NMDAR activity, leading to potentiation of AMPAR activity (unsaturated alcohol-evoked plasticity). This is further potentiated by *in vivo* LTP induction at mPFC inputs to DMS D1-MSNs, producing a long-lasting enhancement of alcohol-seeking behavior. Conversely, *in vivo* eCB-LTD induction at the same synapses elicited a long-lasting suppression of this behavior. (b) Paired HFS and oPSD induces both LTP and LTD, but only LTP is detected. 1, HFS causes presynaptic release of glutamate, which activates AMPARs. The resultant weak membrane depolarization is insufficient to remove the Mg²⁺ blockade of NMDARs and thus fails to induce LTP. 2, Optical stimulation of channelrhodopsin expressed on postsynaptic neurons causes strong membrane depolarization (oPSD). This is sufficient to remove Mg²⁺ blockade of NMDARs, leading to greater Ca²⁺ influx, activation of Ca²⁺/calmodulin-dependent protein kinase II (CaMKII) signaling pathways, and consequently AMPAR insertion (LTP induction). 3, oPSD also

opens voltage-gated Ca^{2+} channels, causing Ca^{2+} influx and production of endocannabinoids; these are retrogradely released into the synaptic cleft, where they activate presynaptic CB1Rs. 4, CB1R activation reduces glutamate release (eCB-LTD induction). 5, D2R antagonists are used to blocking D2R-mediated eCB-LTD, which may occur following eHFS *ex vivo* or oHFS *in vivo*. 6, Since the magnitude of LTP (~31%) is greater than that of LTD (~13%), only LTP is detected. (c) LTD induction and detection in D1-MSNs. Since LTP induction is blocked by NMDAR antagonists (1) and eCB-LTD is induced (2), only eCB-LTD is detected (3). (d) No LTD is detected in D2-MSNs because LTP is blocked by NMDAR antagonists (1) and eCB-LTD is blocked by D2R antagonists (2).