**Title: HCN1 channels mediate mu opioid receptor long-term depression at insular
cortex inputs to the dorsal striatum**

Braulio Munoz¹, Brandon M. Fritz¹ and Brady K. Atwood^{1,2}.

¹Department of Pharmacology & Toxicology, Indiana University School of Medicine,
Indianapolis, IN, 46202, USA.

²Stark Neurosciences Research Institute, Indiana University School of Medicine,
Indianapolis, IN, 46202, USA.

Author Contact information

Corresponding author: Brady K. Atwood, PhD.

Indiana University School of Medicine

Department of Pharmacology & Toxicology

320 W 15th St.

NB 400C

Indianapolis, IN, USA 46202

Tel: 317-274-8917

Fax: 317-274-7714

Email: bkatwood@iu.edu

Abstract

Mu opioid receptors (MORs) are expressed in the dorsal striatum, a brain region that mediates goal-directed (via the dorsomedial striatum), and habitual (via the dorsolateral striatum, DLS) behaviors. Our previous work indicates that glutamate transmission is depressed when MORs are activated in the dorsal striatum, inducing MOR-mediated long-term synaptic depression (MOR-LTD) or short-term depression (MOR-STD), depending on the input. In the DLS, MOR-LTD is produced by MORs on anterior insular cortex (AIC) inputs and MOR-STD occurs at thalamic inputs, suggesting input-specific MOR plasticity mechanisms. Here, we evaluated the mechanisms of induction of MOR-LTD and MOR-STD in the DLS using pharmacology and optogenetics combined with patch clamp electrophysiology. We found that cAMP/PKA signaling and protein synthesis are necessary for MOR-LTD expression, similar to previous studies of endocannabinoid-mediated LTD (eCB-LTD) in DLS. However, MOR-LTD does not require mTOR signaling as eCB-LTD does. MOR-STD does not utilize these same mechanisms. We characterized the role of presynaptic HCN1 channels in MOR-LTD induction as HCN1 channels expressed in AIC are necessary for MOR-LTD expression in the DLS. These results suggest a mechanism in which MOR activation needs HCN1 to induce MOR-LTD, suggesting a new target for pharmacological modulation of synaptic plasticity, providing new opportunities to develop novel drugs to treat alcohol and opioid use disorders.

Keywords: synaptic plasticity, long-term depression, dorsal striatum, corticostriatal, mu opioid receptor, insular cortex, HCN1 channel.

Introduction

The opioid system is expressed throughout the brain(Le Merrer, Becker, Befort, & Kieffer, 2009) and promotes synaptic plasticity in many brain regions(Bao et al., 2007; Dacher & Nugent, 2011a, 2011b; Drake, Chavkin, & Milner, 2007; Iremonger & Bains, 2009), including the dorsal striatum (DS)(Atwood, Kupferschmidt, & Lovinger, 2014; Hawes, Salinas, Lovinger, & Blackwell, 2017; Lovinger, 2010; Munoz, Fritz, Yin, & Atwood, 2018; Muñoz, Haggerty, & Atwood, 2020). The DS is the primary input nucleus of the basal ganglia and can be subdivided in two structures: the dorsomedial striatum (DMS), which controls goal-directed learning, and the dorsolateral striatum (DLS), which regulates habit formation(Burton, Nakamura, & Roesch, 2015; Corbit & Janak, 2016; Hilário & Costa, 2008; Lovinger, 2010; O'Tousa & Grahame, 2014). Drug and alcohol use shift from being outcome-driven to becoming more stimulus-driven and compulsive in nature during addiction development(Corbit & Janak, 2016; Everitt & Robbins, 2016; Hopf & Lesscher, 2014; Shen et al., 2018). This transition from flexible goal-directed drug use to more habitual or compulsive use, is paralleled by a transition in the neural activity from the DMS to the DLS(Corbit & Janak, 2016; Renteria, Cazares, & Gremel, 2020; Shen et al., 2018), thereby shifting the balance of action control from DMS to DLS. Previous reports have demonstrated the role of glutamatergic synaptic plasticity in behavior, showing that the disruption of glutamatergic long-term synaptic depression (LTD) at orbitofrontal cortex-DMS synapses prevents habit learning(Gremel et al., 2016), and the disruption of glutamatergic LTD in DLS promotes habit learning(DePoy et al., 2013; C. Nazzaro et al., 2012). Therefore, changes in the expression of excitatory synaptic plasticity affect specific DS-related behaviors.

Our previous work established that mu opioid receptor (MOR) activation induces long-term synaptic depression (MOR-LTD) or short-term depression (MOR-STD) of glutamate release in both the DLS and DMS (Atwood, Kupferschmidt, et al., 2014; Munoz et al., 2018; Muñoz et al., 2020). We demonstrated that MOR-mediated synaptic inhibition is expressed at multiple DS synapses (cortical, thalamic, amygdala and cholinergic), but only a subset of corticostriatal synapses expressed MOR-LTD (Atwood, Kupferschmidt, et al., 2014) that was disrupted by prior *in vivo* ethanol exposure (Munoz et al., 2018; Muñoz et al., 2020). We also found that mice that were exposed to alcohol and the opioid oxycodone *in vivo* had ablated corticostriatal MOR-LTD when measured in brain slices (Atwood, Kupferschmidt, et al., 2014; Munoz et al., 2018; Muñoz et al., 2020). MOR plasticity at other DS synapses was unaffected by alcohol exposure (Munoz et al., 2018; Muñoz et al., 2020). Of particular interest, in DLS, alcohol and opioid-sensitive MOR-LTD of glutamatergic input is completely restricted to anterior insular cortex (AIC) inputs (Atwood, Kupferschmidt, et al., 2014; Munoz et al., 2018). Altogether, these data led us to hypothesize that MOR plasticity utilizes different mechanisms at distinct glutamatergic synapses in DS. As we previously demonstrated that MOR-LTD and endocannabinoid LTD (eCB-LTD) are mutually occlusive in DS (Atwood, Kupferschmidt, et al., 2014), we predicted that these two forms of LTD would utilize similar mechanisms. cAMP/PKA signaling mediates eCB-LTD in the nucleus accumbens, amygdala, cerebellum, and hippocampus (Atwood, Lovinger, & Mathur, 2014). In addition, local protein translation in the presynaptic region is enhanced by the activation of CB1 receptor, which is important for the induction of inhibitory LTD in the hippocampus (Younts et al., 2016) and likely also in the DS (Yin, Davis, Ronesi, &

Lovinger, 2006). eCB-LTD at GABAergic synapses in the hippocampus is also mediated by mTOR signaling (Younts et al., 2016). Therefore, we sought to test whether MOR-LTD in DLS, and specifically at AIC-DLS synapses is mediated by cAMP/PKA, protein translation and mammalian target of rapamycin (mTOR) signaling. As cAMP also regulates hyperpolarization-activated cyclic nucleotide-gated (HCN) channels, these channels can presynaptically regulate glutamate release, and the type 1 HCN (HCN1) channel may be localized to corticostriatal neurons, we also explored a potential role for HCN1 in mediating MOR-LTD in DLS (He, Chen, Li, & Hu, 2014; Z. Huang, Li, Aguado, Lujan, & Shah, 2017; Moosmang, Biel, Hofmann, & Ludwig, 1999; Postea & Biel, 2011).

In the current study, we probed glutamatergic synapses in the DLS using ex vivo mouse brain slice electrophysiology. We used broad electrical stimulation in C57BL/6J mice to test all glutamatergic inputs. We used an optogenetics approach to probe specific glutamate synapses in DLS, using Emx1-Ai32 mice to test all cortical inputs, VGluT2-Ai32 mice to test all thalamic inputs, and C57BL/6J mice stereotactically injected with an AAV-ChR2 vector into the AIC to specifically probe AIC-DLS synapses. In combination with these tools, we pharmacologically and genetically probed the mechanism involved in the expression of MOR-LTD in the DLS. We demonstrate that cAMP/PKA signaling and protein synthesis are necessary for MOR-LTD expression, but not for MOR-STD. We describe a novel role of presynaptic AIC HCN1 channels in the induction of MOR-LTD in DLS. We also report that mTOR signaling is required for eCB-LTD of glutamate transmission in the DLS, but this is not required for MOR-LTD indicating a divergence in the mechanisms of these two similar forms of synaptic plasticity.

Results

MOR activation express glutamatergic LTD in the DLS

We previously showed that the activation of MOR in DLS produces LTD of eEPSCs as well as oEPSCs(Munoz et al., 2018). Using three approaches, we confirmed that 5 min application of the MOR agonist DAMGO (0.3 μM) induces MOR-LTD. We found that DAMGO produced a persistent decrease in eEPSC amplitude in DLS MSNs (72 ± 5%; Figure 1A-D) in brain slices from C57Bl/6J mice. To generally probe corticostriatal synapses, we used 470 nm light delivered through the microscope objective to evoke oEPSCs in brain slices from Emx1-Ai32 mice (Figure 1E), where the activation of MOR again induced MOR-LTD (71 ± 5%; Figure 1F-H). Finally, to specifically stimulate the release of glutamate from AIC inputs to the DLS, we infused an AAV vector expressing ChR2 in the AIC (Figure 1I), as we previously did(Munoz et al., 2018). As shown previously, activation of MORs produced glutamatergic LTD of AIC-DLS transmission (66 ± 7%; Figure 1J-L). We also confirmed that thalamostriatal inputs expressed MOR-STD in the DLS using VGluT2-Ai32 transgenic mice (15-20 min: 70 ± 9%; 30-40 min: 93 ± 9%; Figure supplement 1A-B) as we previously showed(Atwood, Kupferschmidt, et al., 2014; Munoz et al., 2018). These data replicate our previous work that MORs mediated LTD of AIC-DLS synapses and STD of thalamostriatal synapses in the DLS. We previously showed that eCB-LTD and MOR-LTD were mutually occlusive in DLS(Atwood, Kupferschmidt, et al., 2014). In order to determine whether we could infer potential MOR-LTD mechanisms from what is known about eCB-LTD, we investigated whether eCB-LTD also occurred at AIC-DLS synapses. We used the same AAV approach described above (Figure supplement 2A) and tested the effects of stimulation-

induced LTD and responses to the CB1 receptor agonist WIN55,212-2 (1 μ M). We found that eCB-LTD ($63 \pm 7\%$; Figure 2B-D) and CB1 agonist-induced LTD ($68 \pm 8\%$; Figure 2B, C, E) both occur at AIC-DLS synapses. Next, we explored the signaling cascades that mediate MOR plasticity in DLS using what is known about eCB-LTD as a guide for our experiments.

Inhibition of presynaptic PKA blocks MOR-mediated LTD

The activation of opioid receptors produces an inhibition of adenylyl cyclase (AC) mediated by G α_i , decreasing cAMP levels, which leads to an attenuation of cAMP/protein kinase A (PKA) signaling (Atwood, Lovinger, et al., 2014). However, little is known about PKA's role in MOR-mediated LTD. Using KT5720 (1 μ M), a specific and cell permeable inhibitor of PKA, which brain slices were incubated in for 1h prior to recordings, we demonstrated that the inhibition of PKA blocks the maintenance of MOR-LTD of eEPSCs in DLS ($101 \pm 6\%$; Figure 3A-C). Corticostriatal MOR-LTD was also blocked by PKA inhibition ($90 \pm 5\%$; Figure 3G-I), as was MOR-LTD at AIC-DLS synapses ($96 \pm 4\%$; Figure 3M-O). We found that these effects were due to inhibiting presynaptic PKA activity as at least 30 min treatment with the membrane-impermeable PKA inhibitor PKI (1 μ M), intracellularly infused through the recording micropipette into the postsynaptic MSN, did not affect MOR-LTD in the DLS (eEPSCs: $84 \pm 2\%$; Figure 3D-F) (corticostriatal oEPSCs: $54 \pm 9\%$; Figure 3J-L) (AIC-DLS oEPSCs: $62 \pm 6\%$; Figure 3P-R). Furthermore, thalamostriatal inputs, that express MOR-mediated STD, were not affected by broad inhibition of PKA (15-20 min: $84 \pm 6\%$; 25-35 min: $107 \pm 10\%$) or just inhibition of postsynaptic PKA (15-20 min: $87 \pm 3\%$; 30-40 min: $111 \pm 9\%$;

Figure supplement 1C-D). As we expected, our data demonstrate the role of presynaptic PKA uniquely in MOR-LTD.

The activation of adenylyl cyclase disrupts MOR-mediated LTD

Demonstrating a role for PKA signaling in MOR-LTD implicates adenylyl cyclase (AC) signaling as mediating MOR-LTD. To more specifically test this, we persistently activated AC using 20 μ M forskolin throughout recording. In the presence of forskolin, DAMGO produced a transient increase of the EPSC amplitudes (Figure 4B, E and H), followed by a blockade MOR-LTD at all excitatory synapses ($105 \pm 4\%$; Figure 4A-C) and at corticostriatal ($102 \pm 3\%$; Figure 4D-F) and AIC inputs ($93 \pm 7\%$; Figure 4G-I). MOR-mediated STD at thalamic synapses was not affected by the application of forskolin (15-20 min: $83 \pm 6\%$; 25-35 min: $96 \pm 6\%$; Figure supplement 1E-F). Altogether, these findings indicate that cAMP/PKA signaling pathway is involved in the induction and maintenance of MOR-LTD in DLS.

Protein translation is required to produce MOR-mediated LTD

Next, we tested the role of protein translation in MOR-LTD. Using, bath application of 80 μ M cycloheximide (selective inhibitor of protein synthesis) during the entire recording, we found that the inhibition of protein translation blocked MOR-LTD generally ($104 \pm 9\%$; Figure 5A-C) and specifically at cortical ($105 \pm 9\%$; Figure 5D-F) and AIC inputs ($91 \pm 2\%$; Figure 5G-I). Similar to our experiments targeting cAMP/PKA signaling, thalamic inputs were unaffected by cycloheximide treatment (15-20 min: $73 \pm 6\%$; 30-40 min: $98 \pm 9\%$; Figure supplement 1G-H).

It had been reported that eCB protein translation-mediated-LTD involves mTOR signaling in the hippocampus. To test if mTOR is required for eCB-LTD also in the DLS, we used bath application of 100 nM Torin 2 (a selective mTOR inhibitor). We demonstrated that the inhibition of mTOR signaling disrupted eCB-LTD in the DLS (Control: $77 \pm 3\%$ vs Torin 2: $103 \pm 8\%$; Figure 6A-C). However, MOR-LTD does not require mTOR signaling in the DLS ($79 \pm 2\%$; Figure 6D-F). These data suggest that protein translation is necessary for MOR-mediated LTD, but mTOR signaling is not involved and indicate that eCB- and MOR-LTD, while qualitatively similar, do not utilize identical signaling mechanisms.

Calcium signaling is not required for MOR-mediated LTD

Our data to this point indicates that presynaptic cAMP/PKA and protein synthesis are critical for the induction and maintenance of MOR-LTD. However, calcium is vital to presynaptic neurotransmitter release and calcium-sensitive signaling mechanisms may be required for LTD induction (Atwood, Lovinger, et al., 2014). Therefore, we next tested whether Ca^{2+} is critical for MOR-LTD expression. First, we replaced Ca^{2+} for Sr^{2+} throughout the duration of the recording, to induce asynchronous glutamate release. We found that the replacement of Ca^{2+} did not impact MOR-LTD ($74 \pm 5\%$; Figure supplement 2). Thus, we used another approach where we prepared a DAMGO solution without Ca^{2+} in order to evaluate the role of Ca^{2+} in the induction of MOR-LTD. Our data show that Ca^{2+} is not necessary to induce MOR-LTD ($82 \pm 7\%$; Figure supplement 2).

Presynaptic HCN1 channels are indispensable for MOR-mediated LTD

Given that we identified cAMP signaling as important for MOR-LTD expression (Figure 4), we also explored the possibility of HCN1 channels as key factors for MOR-LTD. To

do so, we applied the general HCN blocker ZD7288 (25 μ M) to brain slices for at least 30 minutes before DAMGO application and during the entire recording. HCN inhibition blocked the expression of MOR-LTD ($101 \pm 7\%$; Figure 7A-C), suggesting a novel role for HCN channels in DLS glutamatergic synaptic plasticity. Next, we characterized the effects of HCN channels in glutamatergic DLS synaptic transmission, by bath applying 25 μ M ZD7288 for at least 15 minutes before recording sEPSCs. We found that the inhibition of HCN channels caused an enhancement in the frequency and a decrease in the amplitude of sEPSCs (Figure 7D-F), without affecting rise and decay times (Figure 6C, G and H). These data suggest, that HCN channels are present on both pre- and postsynaptic sites and they mediate MOR-LTD in the DLS.

To further characterize this new role of HCN channels and establish that presynaptic HCN1 channels are specifically involved in the mechanism of expression of MOR-LTD, we used HCN1^{flox} mice injected with AAV-cre vector into AIC to knock out the expression of HCN1 channel specifically in the AIC, and using AAV-GFP vector-infused mice as controls (Figure 7A, B). Using electrical stimulation and recording in the DLS, we found that the ablation of HCN1 in the AIC reduced the magnitude of MOR-LTD in AAV-cre-injected mice (Figure 7C-F), relative to AAV-GFP-injected mice (Figure 7C-G; cre: $90 \pm 4\%$ vs GFP: $76 \pm 3\%$; Figure 7E). Next, we tested if the deletion of presynaptic HCN1 channels produced changes in basal striatal glutamatergic transmission. We found that the lack of AIC HCN1 channels did not affect any synaptic parameters (Figure 7H-L). These data suggest that HCN1 channels specifically on AIC inputs are necessary for MOR-LTD expression in the DLS.

Discussion

The current study demonstrates that the mechanism of MOR-mediated LTD is similar to the mechanism of induction of the inhibition of glutamate of CB1 or eCB-LTD in that it involves cAMP-PKA-protein translation signaling and both occur at AIC-DLS synapses (Atwood, Lovinger, et al., 2014; Lonart et al., 2003; Mato, Lafourcade, Robbe, Bakiri, & Manzoni, 2008; Yasuda, Huang, & Tsumoto, 2008; Yin et al., 2006; Younts et al., 2016) (Figures. 2-5). However, we found notable differences such as a lack of dependence on mTOR signaling and calcium (Figure 6 and supplement 2) and a novel mechanism involving HCN1 signaling for MOR-LTD (Figures 7 and 8). Figure Supplement 3 summarizes the findings of the paper. Furthermore, the MOR-LTD mechanism is different than that of MOR-STD (Figure supplement 1), suggesting that MOR-mediated plasticity does not utilize a common pathway at all glutamatergic synapses.

Our previous report demonstrates that MORs expressed in AIC are the sole source of MOR-mediated LTD at glutamate inputs to the DLS(Munoz et al., 2018). However in the DMS, MORs on medial prefrontal cortex, anterior cingulate cortex, and basolateral amygdala inputs are capable of inducing LTD(Muñoz et al., 2020). MOR-LTD also occurs locally at cholinergic interneuron synapses that co-release glutamate on to MSNs(Munoz et al., 2018; Muñoz et al., 2020). However, alcohol only disrupts MOR-LTD at AIC inputs to DLS and at medial prefrontal and anterior cingulate cortex inputs to DMS leaving MOR plasticity at thalamostriatal, amygdalostriatal and cholinergic interneurons unaffected(Munoz et al., 2018; Muñoz et al., 2020). We also found that the opioid oxycodone disrupts MOR-LTD in DLS(Atwood, Kupferschmidt, et al., 2014), and given that this is mediated by AIC-DLS MORs(Munoz et al., 2018), it follows that alcohol

and opioids have similar actions on MOR plasticity at this synapse. Given that alcohol (and presumably oxycodone and other opioids) do not affect MOR plasticity at these other non-corticostriatal synapses, differential mechanisms that underlie MOR plasticity at each type of synapse may confer susceptibility to the deleterious effects of these drugs and could account for their mechanisms of action in the development of habitual or compulsive use of the drugs.

Previous work has shown that the inhibition of PKA disrupts eCB-LTD in the nucleus accumbens(Mato et al., 2008) and that presynaptic PKA is required to induce cannabinoid LTD in cerebellum(Lonart et al., 2003) and eCB synaptic plasticity in hippocampus(Yasuda et al., 2008). Moreover, previous reports had shown that the activation of AC by forskolin enhances corticostriatal EPSC amplitudes(Cho et al., 2008) and blocked delta opioid receptor-mediated LTD of inhibitory postsynaptic currents(Patton, Roberts, Lovinger, & Mathur, 2016). cAMP/PKA signaling has also been implicated in type 2/3 metabotropic glutamate receptor- and type 1b serotonin receptor-mediated LTD at various synapses in the brain(Atwood, Kupferschmidt, et al., 2014). Until now, the role of cAMP/PKA signaling pathway in MOR-mediated plasticity had not been addressed. Our results demonstrate that presynaptic PKA inhibition blocked AIC-DLS MOR-LTD (Figure 3), but not thalamostriatal MOR-STD (Figure supplement 1C-D). While the magnitudes and time courses differed slightly between measures of eEPSCs and oEPSCs, it appears that PKA is not required for MOR-mediated inhibition of glutamate release, but it is required for the maintenance of MOR-LTD. On the other hand, we found that AC activation disrupted both acute MOR-mediated inhibition (in fact turning it into a transient potentiation) and MOR-LTD (Figure

4), but had no effect on MOR-STD (Figure supplement 1E-F). Altogether these data suggest that cAMP/PKA pathway may be a general mechanism whereby GPCRs that signal through Gai may induce LTD, but a receptor's ability to couple to Gai does not necessarily mean that it will induce cAMP/PKA-dependent LTD upon activation.

Protein translation is necessary for striatal eCB-LTD(Yin et al., 2006) and is involved in GPCR-mediated LTD at some synapses, but not others(Atwood, Kupferschmidt, et al., 2014; Younts et al., 2016) suggesting that protein translation can be an important component of glutamatergic LTD, depending on the synapse. Indeed, local protein translation can take place in the presynaptic region, where the activation of CB1 receptors enhances protein translation, being critical for the induction and not the maintenance of inhibitory LTD(Younts et al., 2016). Here we demonstrated that protein translation is necessary to produce MOR-LTD (Figure 5), but not MOR-STD (Figure supplement 1G-H). Our data is consistent with the previous reports on protein translation-dependent eCB-LTD, and suggest that it is presynaptic protein translation that is involved in MOR-LTD expression, however more data is needed to determine this definitively.

Although we have previously reported that MOR- and eCB-LTD are mutually occlusive in the DLS(Atwood, Kupferschmidt, et al., 2014), and here we found that there were similarities between MOR-LTD and eCB-LTD mechanisms, we found some differences which make both types of LTD distinct. In the DLS, MOR-LTD of glutamate input is restricted to those from the AIC(Munoz et al., 2018), which differs from the more broadly expressed (and overlapping at AIC-DLS synapses, Figure supplement 2) eCB-LTD(Gremel et al., 2016; Wu et al., 2015), suggesting that these two forms of LTD may

occur in distinct synaptic environments that could utilize differing mechanisms. It is likely that even in synapses where both receptors are expressed (e.g. in AIC terminals) that each receptor shares partially overlapping, but not identical signaling pathways.

A clear example of distinct signaling mechanisms identified here is in the role of the mTOR pathway in eCB-LTD and MOR-LTD. The CB1 receptor can signal via mTOR in the hippocampus, where the increase of protein translation induced by CB1 receptor activation is mTOR-dependent producing inhibitory eCB-LTD (Younts et al., 2016). We predicted that we would see the same effect in the DLS for both eCB-LTD and for MOR-LTD. Partially correct in our prediction, we found that eCB-LTD is blocked after the inhibition of mTOR in the DLS (Figure 6A-C), but the mechanism of MOR-LTD did not utilize mTOR (Figure 6D-F). These novel data regarding eCB-mTOR signaling in DLS provide rationale for future experiments addressing the role of eCB-mTOR signaling in dorsal striatal- related behaviors.

Calcium signaling also plays an important role in the mechanism of synaptic plasticity (Frank, Pielage, & Davis, 2009; Tien & Kerschensteiner, 2018). Postsynaptic Ca^{2+} influx is needed to produce eCBs (Tanimura et al., 2010), which interacts with presynaptic mechanism to produce LTD (Adermark, 2011; Adermark, Talani, & Lovinger, 2009; Cristiano Nazzaro et al., 2012; Plotkin et al., 2013; Wang et al., 2006). Some forms of GPCR-mediated LTD, including eCB-LTD, also require presynaptic calcium signaling as well (Atwood, Lovinger, et al., 2014). However, we demonstrated that Ca^{2+} is not involved in MOR-LTD in DLS (Figure supplement 2), indicating another difference between the mechanisms of MOR-LTD and eCB-LTD.

315 Presynaptic cations such as Na⁺ and K⁺ are important for the propagation of action
316 potentials and subsequent glutamate release(Chen & Lui, 2021). HCNs channels are
317 permeable to Na⁺ and K⁺ and modulated by cAMP(Sartiani, Mannaioni, Masi, Novella
318 Romanelli, & Cerbai, 2017), and they can decrease neurotransmission by restricting
319 presynaptic Ca²⁺ influx(Zhuo Huang et al., 2011). However, while we did not find that
320 Ca²⁺ plays a role in MOR-LTD, we did identify a critical role for the HCN1 channel in
321 MOR-LTD expression (Figures 7 and 8). It has been reported that HCN1 channels can
322 be found at presynaptic sites in the globus pallidus, where the inhibition by ZD7288
323 produced an increase of miniature inhibitory postsynaptic current frequency, with no
324 changes in amplitude(Boyes, Bolam, Shigemoto, & Stanford, 2007). The block or
325 deletion of HCN1 causes an increase in the frequency of miniature EPSCs in entorhinal
326 cortical pyramidal neurons(Zhuo Huang et al., 2011). Here we showed that broad
327 blockade of HCN channels enhanced glutamate release, but decreased postsynaptic
328 responses to glutamate (Figure 7D-H), presumably by acting non-specifically at both
329 presynaptic and postsynaptic sites. However, when we deleted just HCN1 from AIC
330 inputs to DLS, we did not observe any changes in basal glutamate transmission (Figure
331 8H-L). The differences in effects of broad blockade and synapse-specific deletion may
332 be attributed to the overall contribution of HCN1s at AIC inputs relative to the population
333 of glutamate synapses. Genetic deletion did not completely ablate MOR-LTD
334 expression. This could be due to an incomplete AAV vector transduction (most
335 recordings had a complete loss of LTD) or perhaps that HCN1 contributes to LTD
336 differently at AIC inputs to different subtypes of MSNs. Therefore, in the future, it will
337 also be important to investigate the role of HCN1 in MOR-LTD at AIC inputs to D1- or

D2-expressing MSNs (direct or indirect pathway respectively). It has recently been reported that HCN1 contributes to alcohol preference(Salling & Harrison, 2020). Given that alcohol disrupts MOR-LTD in DLS, this mechanism of MOR-HCN1 signaling may be a target of alcohol's effects and contribute to alcohol-related, dorsal striatal-dependent behaviors such as habitual and compulsive alcohol drinking. As the opioid oxycodone also disrupts MOR-LTD it will be intriguing to determine the impact of both oxycodone and alcohol on AIC-DLS HCN1 function. HCN1 may therefore have therapeutic potential in alcohol and opioid use disorders.

In conclusion, these findings indicate that glutamatergic synapses in the DLS have distinct mechanisms of plasticity induction mediated by the same types of receptors. The specific signaling processes that result in these different forms of plasticity might be the answer to why different types of plasticity are differentially sensitive to drugs of abuse and alcohol(Atwood, Lovinger, et al., 2014; Munoz et al., 2018; Muñoz et al., 2020). However, further work is required to resolve what components of these mechanisms render some synapses susceptible to the deleterious effects of these drugs while others are resistant. It will also be important to decipher the behavioral relevance of AIC-DLS synapse HCN1 channels and eCB-mTOR signaling in the DLS.

Methods

Methods details

All experiments were performed similar to our previous dorsal striatal electrophysiological studies with some experiment-specific modifications (Atwood, Kupferschmidt, et al., 2014; Fritz, Munoz, & Atwood, 2019; Fritz, Munoz, Yin, Bauchle, & Atwood, 2018; Munoz et al., 2018). These methods are described in brief below.

Animals and materials

Animal care and experimental protocols for this study were approved by the Institutional Animal Care and Use Committee (IACUC) at the Indiana University School of Medicine and all guidelines for ethical protocols and care of experimental animals established by the NIH (National Institutes of Health, Maryland, USA) were followed. Male C57BL/6J mice were obtained from the Jackson Laboratory (JAX #000664, Bar Harbor, Maine, USA). Emx1Cre-Ai32, VGluT2Cre-Ai32, and HCN1fox mice were bred and genotyped in-house (Original stock strains: Ai32: JAX #012569; Emx1Cre: JAX #005628; VGluT2Cre: JAX #016963; HCN1f/f: JAX #028299). All mutant mice used in these studies were backcrossed to C57BL/6J mice for a minimum of 7 generations. The mice used in these studies were between PND 60-100 at the time of experimentation (with the exception of HCN1-flox AAV-cre-injected mice ~PND 98-154). Animals were group-housed in a standard 12-h light/dark cycle (lights on at 0800) at 50% humidity. Food and water were available *ad libitum*.

Reagents

We used the MOR agonist [D-Ala², NMe-Phe⁴, Gly-ol⁵]-enkephalin (DAMGO; H-2535,

Bachem), GABA_A receptor antagonist picrotoxin (PTX, P1675, Sigma-Aldrich), PKI (6221, Tocris), KT5720 (1288, Tocris), Forskolin (F3917, Sigma-Aldrich), Cycloheximide (0970, Tocris), ZD7288 (1000, Tocris), WIN55,212-2 (1038, Tocris), and Torin 2 (4248, Tocris). Other reagents used for making solutions were purchased from Sigma-Aldrich or Fisher Scientific.

Brain slice preparation.

Mice were euthanized via decapitation under deep isoflurane anesthesia, and the brain was quickly excised and placed in an ice-cold cutting solution containing (in mM): 194 sucrose, 30 NaCl, 4.5 KCl, 1 MgCl₂, 26 NaHCO₃, 1.2 NaH₂PO₄, 10 Glucose saturated with a mixture of 95% O₂ and 5% CO₂, and sliced to a thickness of 280 µm on a vibratome (Leica VT1200S, Germany). Slices were transferred to an artificial cerebrospinal fluid (aCSF) solution containing (in mM): 124 NaCl, 4.5 KCl, 1 MgCl₂, 26 NaHCO₃, 1.2 NaH₂PO₄, 10 Glucose, 2 CaCl₂ (310-320 mOsm) saturated with 95% O₂/5% CO₂ at 30°C for 1 hr before being moved to room temperature. When ready for recording, slices were transferred to a recording chamber continuously perfused with aCSF solution saturated with 95% O₂/5% CO₂.

Electrophysiology recordings

Whole-cell recordings of excitatory postsynaptic currents (EPSCs) in medium spiny neurons (MSNs) were carried out at 29–32 C° and aCSF was continuously perfused at a rate of 1-2 ml/min. Recordings were performed in the voltage clamp configuration using a Multiclamp 700B amplifier and a Digidata 1550B (Molecular Devices, San Jose, CA). Slices were visualized on an Olympus BX51WI microscope (Olympus Corporation of

America). MSNs were identified by their size, membrane resistance, and capacitance. Picrotoxin (50 μ M) was added to the aCSF for recordings to isolate excitatory transmission. Patch pipettes were prepared from filament-containing borosilicate micropipettes (World Precision Instruments) using a P-1000 micropipette puller (Sutter Instruments, Novato, CA), having a 2.0-3.5 M Ω resistance. The internal solution contained (in mM): 120 CsMeSO₃, 5 NaCl, 10 TEA-Cl, 10 HEPES, 5 lidocaine bromide, 1.1 EGTA, 0.3 Na-GTP and 4 Mg-ATP (pH 7.2 and 290-310 mOsm). MSNs were voltage clamped at -60 mV for the duration of the recordings. To evaluate the effect of Ca²⁺, aCSF Ca²⁺ was replaced with Sr²⁺ (2 mM) during the application of DAMGO (from 10 to 15 min after the start of recording) or through the entire recording (from 0 to 40 min). For electrically-evoked recordings, a twisted tungsten bipolar stimulating electrode (PlasticsONE, Roanoke, VA) was placed at the border of the white matter of the external capsule. Electrically-evoked excitatory postsynaptic currents (eEPSCs) were generated by a DS3 Isolated Current Stimulator (Digitimer, Ft. Lauderdale, FL) every 20 s and stimulus intensity was adjusted to produce stable eEPSCs of 200–600 pA in amplitude prior to the initiation of experimental recording. The spontaneous EPSC (sEPSC) recordings were made for 2 min and the analysis of frequency (Hz), decay constant (ms), rise constant (ms) and amplitude (pA) were measured to determine the effects of the deletion of HCN1 in AIC. The decay constant of sEPSCs was fitted as single exponential and both rise and decay-phase were fitted between 10 and 90% of the maximal amplitude. Data were acquired using Clampex 10.3 (Molecular Devices, San Jose, CA). Series resistance was monitored and only cells with a stable series resistance (less than 25 M Ω and that did not change more than 15% during recording) were included for data

analysis. Recordings were made 2–7 h after euthanasia.

Viral injections

C57BL/6J mice were anesthetized with isoflurane and stereotactically-injected with the adeno-associated viral (AAV) vector, AAV9.hSyn.ChR2(H134R)-eYFP (Addgene #26973) to drive ChR2 expression in AIC neurons. Bilateral injections were made into AIC: A/P: +2.4, M/L: ± 2.3 , D/V: -2.25 (50 nl/injection, 12.5 nl/min infusion rate).

To produce AIC projection neuron HCN1 knockout mice, HCN1-flox mice were anesthetized with isoflurane and stereotactically-injected with AAV9.hSyn.HI.eGFP.Cre.WPRE.SV40 (Addgene #105540) or AAV9.hSyn.eGFP.WPRE.bGH as control (Addgene #105539). Bilateral injections were made into AIC at coordinates: A/P: +2.4, M/L: ± 2.3 , D/V: -2.25 (75 nl/injection, 12.5 nl/min infusion rate). HCN1-flox mice were allowed to recover for at least 4 weeks to allow for adequate ablation of HCN1 channel expression before brain slices were made for electrophysiological recordings. Prior to recording, brain slices were imaged via an Olympus MVX10 microscope (Olympus Corporation of America) to verify GFP or GFP-tagged cre-recombinase expression in injected HCN1flox mice.

Optogenetic Recordings

AAV-ChR2 injection in C57BL/6J mice was performed to target ChR2 expression to inputs from AIC to DLS. Optically-evoked EPSCs (oEPSCs) in MSNs from Emx1-Ai32, VGlut2-Ai32 and C57BL/6J mice injected with AAV-ChR2 were produced in brain slices using 470-nm blue light (5-ms exposure time) delivered via field illumination through the microscope objective. Light intensity was adjusted to produce stable oEPSCs of 200–

600-pA amplitude prior to experimental recording. oEPSCs were evoked every 30s. AIC-DLS eCB-LTD was produced using optical high-frequency stimulation (oHFS) coupled with depolarization (4 blue light pulses of 50 Hz, 10 s inter-pulse interval). Prior to recording, brain slices were imaged via an Olympus MVX10 microscope (Olympus Corporation of America) to verify YFP-tagged ChR2, GFP-tagged cre-recombinase, or GFP expression in injected C57BL/6J mice.

Quantification and statistical analysis

Sample size

The target number of samples in each group for electrophysiological experiments was determined from our previously published studies (Atwood, Kupferschmidt, et al., 2014; Munoz et al., 2018; Muñoz et al., 2020).

Replication

All sample sizes indicated in figures for electrophysiological experiments represent biological replicates. 1 neuron was recorded per brain slice and all experiments involved recordings from at least 2 mice.

Data analyses

eEPSCs and oEPSCs were analyzed using Clampfit 10.1 (Molecular Devices, San Jose, CA) and sEPSCs with MiniAnalysis 6.0 (Synaptosoft Inc). Unless otherwise indicated, data are presented as the mean $\pm$ SEM. The analyses of normally distributed data were performed using two-tailed unpaired or two-tailed paired Student's t tests following an F test to confirm similar variances. Non-normally distributed data were

analyzed using two-tailed Welch's *t* test for unpaired data (Figure 7J-L). Data that were analyzed using this test are indicated in the figure legends. Statistical analyses were performed with Prism 9 (GraphPad, La Jolla, CA). The level of significance was set at *P* < 0.05 for all analyses. Representative traces are the average baseline EPSC (1-10 min) and average post-treatment EPSC of final 10 min of recording. For VGluT2-Ai32 recordings, representative traces are the average baseline EPSC (1-10 min) average of the peak of response after DAMGO (15-20 min) and average post-treatment EPSC of final 10 min of recording. Exclusion of individual data points was determined using a ROUT outlier calculator (Q=1%) included in the Prism 9 software package.

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480

Figures

505 oEPSC amplitudes (0–10 min baseline v. final 10 min of recording; paired t-test,
 506 $P = 0.0084$, $t_4 = 4.84$, $n = 5$ neurons from 2 mice). Data represent mean $\pm$ SEM.
 507 $**P < 0.01$.

508

Figure 2. Cannabinoid LTD occurs at AIC-DLS synapses. (A) Coronal brain slice showing the AAV-ChR2 infection of AIC and dorsal striatal terminal expression (bar scale = 1000 μm). (B) Representative oEPSC traces showing the effects of pairing optical high frequency stimulation with postsynaptic depression (oHFS) or WIN55,212-2 (1 μM, 10 min) application in brain slices of AAV-ChR2 C57Bl/6J mice. (C) Both oHFS and WIN55,212-2 induced AIC-DLS LTD (final 10 min of recording average: oHFS: 63 ± 7%; WIN55,212-2: 68 ± 8%). (D) oHFS significantly reduced oEPSC amplitudes (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.017$, $t_3 = 4.78$, $n = 4$ neurons from 2 mice). (E) WIN55,212-2 significantly reduced oEPSC amplitudes (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.0038$, $t_7 = 4.24$, $n = 8$ neurons from 4 mice). Data represent mean ± SEM. * $P < 0.05$, ** $P < 0.01$.

Figure 3. Inhibition of Presynaptic PKA blocks MOR-mediated LTD. (A) Representative eEPSC traces showing the effects of DAMGO (0.3 μ M, 5 min) application after the preincubation of PKA-selective inhibitor, KT5720 (1 μ M, $\geq$ 1 hr). (B-C) The preincubation of KT5720 blocked glutamatergic MOR-LTD ($101 \pm 6\%$), the eEPSC amplitudes did not change after DAMGO application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.9487$, $t_8 = 0.0664$, $n = 9$ neurons from 3 mice). (D) Representative eEPSC traces showing the effects of DAMGO (0.3 μ M, 5 min) application after the intracellular dialysis of PKI (1 μ M, $\geq$ 30 min). (E-F) The inhibition of postsynaptic PKA did not alter MOR-LTD ($84 \pm 2\%$). eEPSC amplitudes were reduced after DAMGO application (0–10 min baseline v. final 10 min of recording; paired t-test, $P < 0.0001$, $t_{11} = 6.374$, $n = 12$ neurons from 5 mice). (G) Representative oEPSC traces showing the effects of DAMGO (0.3 μ M, 5 min) application after the preincubation of PKA-selective inhibitor, KT5720 (1 μ M, $\geq$ 1 hr). (H-I) PKA inhibition blocked corticostriatal MOR-LTD ($90 \pm 5\%$), with no changes in oEPSC amplitudes after DAMGO application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.162$, $t_8 = 1.541$, $n = 9$ neurons from 4 mice). (J) Representative oEPSC traces showing the effects of DAMGO (0.3 μ M, 5 min) application after the intracellular dialysis of PKI (1 μ M, $\geq$ 30 min). (K-L) The inhibition of postsynaptic PKA did not alter MOR-LTD ($54 \pm 9\%$). oEPSC amplitudes were reduced after DAMGO application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.0014$, $t_6 = 5.588$, $n = 7$ neurons from 2 mice). (M) Representative AIC-DLS oEPSC traces showing the effects of DAMGO (0.3 μ M, 5 min) application after the preincubation of PKA-selective inhibitor, KT5720 (1 μ M, $\geq$ 1 hr). (N-O) KT5720 blocked corticostriatal MOR-LTD ($96 \pm 4\%$), with no changes in oEPSC amplitudes after DAMGO application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.2121$, $t_9 = 1.343$, $n = 10$ neurons from 3 mice). (P) Representative AIC-DLS oEPSC traces showing the effects of DAMGO (0.3 μ M, 5 min) application after the intracellular dialysis of PKI (1 μ M, $\geq$ 30 min). (Q-R) Postsynaptic PKA inhibition did not alter specific AIC MOR-mediated LTD ($62 \pm 6\%$). oEPSC amplitudes were reduced after DAMGO application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.0026$, $t_5 = 5.576$, $n = 6$ neurons from 3 mice). Data represent mean $\pm$ SEM. ns= no significant, ** $P < 0.01$, *** $P < 0.001$, **** $P < 0.0001$.

Figure 4. The activation of adenylyl cyclase disrupts MOR-mediated LTD. (A) Representative eEPSC traces showing the effects of 20 μ M forskolin before, during and after DAMGO (0.3 μ M, 5 min) application. (B-C) AC activation disrupted glutamatergic MOR-LTD ($105 \pm 4\%$), showing no changes in eEPSC amplitude after DAMGO bath application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.2681$, $t_{8} = 1.19$, $n = 9$ neurons from 4 mice). (D) Representative oEPSC traces showing the effects of 20 μ M forskolin before, during and after DAMGO (0.3 μ M, 5 min) application in brain slices from Emx1-Ai32 mice. (E-F) AC activation disrupted corticostriatal MOR-LTD ($102 \pm 3\%$), with no effects in oEPSC amplitude after DAMGO bath application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.4031$, $t_{5} = 0.913$, $n = 6$ neurons from 3 mice). (G) Representative AIC-DLS oEPSC traces showing the effects of 20 μ M forskolin before, during and after DAMGO (0.3 μ M, 5 min) application. (H-I) AC activation blocked specifically AIC MOR-LTD ($105 \pm 4\%$), without changes in oEPSC amplitude after DAMGO bath application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.379$, $t_{5} = 0.964$, $n = 6$ neurons from 2 mice). Data represent mean $\pm$ SEM. ns= no significant.

Figure 5. Protein translation is required for MOR-mediated LTD expression. (A) Representative eEPSC traces showing the effects of 80 μ M cycloheximide before, during and after DAMGO (0.3 μ M, 5 min) application. (B-C) Protein translation inhibition disrupted glutamatergic MOR-LTD ($104 \pm 9\%$), showing no changes in eEPSC amplitude after DAMGO bath application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.7006$, $t_{8} = 0.398$, $n = 9$ neurons from 4 mice). (D) Representative oEPSC traces showing the effects of 80 μ M cycloheximide before, during and after DAMGO (0.3 μ M, 5 min) application in brain slices from Emx1-Ai32 mice. (E-F) Protein translation inhibition blocked corticostriatal MOR-LTD ($105 \pm 9\%$) with no effects in oEPSC amplitude after DAMGO bath application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.4992$, $t_{7} = 0.713$, $n = 8$ neurons from 3 mice). (G) Representative AIC-DLS oEPSC traces showing the effects of 80 μ M cycloheximide before, during and after DAMGO (0.3 μ M, 5 min) application. (H-I) Protein translation is needed to induce MOR-LTD ($91 \pm 2\%$) from AIC inputs. DAMGO did not change in oEPSC amplitude (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.1208$, $t_{5} = 1.868$, $n = 6$ neurons from 2 mice). Data represent mean $\pm$ SEM. ns= no significant.

Figure 6. mTOR pathway is involved in eCB-LTD but not in MOR-LTD in the DLS. (A) Representative eEPSC traces before and after high-frequency stimulation (HFS) coupled with depolarization (4 pulses of 100 Hz, 10 s inter-pulse interval), showing the effects of torin 2 (100 nM) bath application in brain slices of C57BL/J. (B-C) mTOR inhibition blocked eCB-LTD in DLS (0–10 min baseline v. final 10 min of recording; Control: paired t-test, $P = 0.0002$, $t_{7} = 7.08$, $n = 8$ neurons from 3 mice; torin 2: paired t-test, $P = 0.9009$, $t_{7} = 0.129$, $n = 8$ neurons from 3 mice). (D) Representative eEPSC traces before and after DAMGO bath application (0.3 μ M, 5 min), showing the effects of torin 2 (100 nM) bath application in brain slices of C57BL/J. (E-F) mTOR inhibition did not affect the expression of MOR-LTD in DLS (0–10 min baseline v. final 10 min of recording; paired t-test, $P < 0.0001$, $t_{7} = 8.11$, $n = 8$ neurons from 3 mice). Data represent mean $\pm$ SEM. *** $P < 0.001$, **** $P < 0.0001$.

Figure 7. Inhibition of HCN channels disrupts MOR-mediated LTD. (A) Representative eEPSC traces showing the effects of 25 μM ZD7288 before, during and after DAMGO (0.3 μM, 5 min) application. (B-C) HCN channel inhibition disrupted glutamatergic MOR-LTD ($104 \pm 9\%$) in the DLS, showing no changes in eEPSC amplitude after DAMGO bath application (0–10 min baseline v. final 10 min of recording; paired t-test, $P = 0.8008$, $t_{14} = 0.2621$, $n = 8$ neurons from 5 mice). (D) Representative synaptic current traces from C57BL/6J mice showing the spontaneous excitatory postsynaptic currents (sEPSC) in absence (black trace) or presence of 25 μM ZD7288 (gray trace, 15 min) application. (E-H) HCN channel inhibition increased frequency (paired t-test, $P = 0.043$, $t_{14} = 2.223$, $n = 15$ neurons from 2 mice) and decreased amplitude (paired t-test, $P = 0.0005$, $t_{14} = 4.49$, $n = 15$ neurons from 2 mice), without changes in rise (paired t-test, $P = 0.542$, $t_{14} = 0.625$, $n = 15$ neurons from 2 mice) and decay times (paired t-test, $P = 0.324$, $t_{14} = 1.023$, $n = 15$ neurons from 2 mice) of the sEPSC events in DLS. Data represent mean $\pm$ SEM. ns= no significant, $*P < 0.05$, $***P < 0.001$.

Figure 8. Genetic ablation of HCN1 channels reduces MOR-mediated LTD expression. (A) Schematic figure of coronal brain slice showing injection paradigm enabling knock out of HCN1 channel expression in AIC in HCN1-flox mice. An AAV vector encoding for either cre-recombinase (AAV.hSyn.cre) or eGFP (AAV.hSyn.eGFP) was injected at least 4 weeks prior to recordings. (B) Coronal brain slice showing the AAV-cre or AAV-GFP infection of AIC and dorsal striatal terminal expression (bar scale = 1000 μ m). (C) Representative eEPSC traces showing the effects of DAMGO (0.3 μ M, 5 min) application in brain slices of AAV-injected HCN1-flox mice. (D-E) The conditional KO of HCN1 channel from AIC, blunted MOR-LTD in DLS (cre: $90 \pm 4\%$ vs GFP: $76 \pm 3\%$, Unpaired t-test, $P=0.017$, $t_{17}=2.64$, $n=10$ neurons from 5 cre-injected mice and $n=9$ neurons from 3 GFP-injected mice). (F) HCN1 deletion did not fully disrupt MOR-LTD, because there was still a small, but significant reduction of eEPSC amplitude after DAMGO application (0–10 min baseline v. final 10 min of recording;

638 paired t-test, $P=0.011$, $t_9=3.18$, $n=10$ neurons from 5 mice). **(G)** Control GFP-
639 injected mice showed normal decrease of eEPSC amplitude after DAMGO application
640 0–10 min baseline v. final 10 min of recording; paired t-test, $P<0.0001$, $t_8=7.51$,
641 $n=9$ neurons from 3 mice). **(H)** Representative synaptic current traces from HCN1-flox
642 mice showing the spontaneous excitatory postsynaptic currents (sEPSCs) in AAV-cre
643 (orange trace) and AAV-GFP (green trace) injected mice. **(I-L)** HCN1 channel ablation
644 from AIC synapses did not affect frequency (cre: 2.6 ± 0.4 Hz vs GFP: 2.7 ± 0.3 Hz,
645 Unpaired t-test, $P=0.886$, $t_{19}=0.146$, $n=10$ neurons from 3 cre-injected mice and $n=11$
646 neurons from 2 GFP-injected mice), amplitude (cre: 18.2 ± 1.2 pA vs GFP: 18.3 ± 0.6
647 pA, Unpaired t-test with Welch's correction, $P=0.957$, $t_{13}=0.055$, $n=10$ neurons from 3
648 cre-injected mice and $n=11$ neurons from 2 GFP-injected mice), rise time (cre: $1.23 \pm$
649 0.11 ms vs GFP: 0.98 ± 0.04 ms, Unpaired t-test with Welch's correction, $P=0.0501$,
650 $t_{11}=2.19$, $n=10$ neurons from 3 cre-injected mice and $n=11$ neurons from 2 GFP-
651 injected mice) and decay time (cre: 5.25 ± 0.43 ms vs GFP: 4.36 ± 0.14 ms, Unpaired t-
652 test with Welch's correction, $P=0.075$, $t_{11}=1.97$, $n=10$ neurons from 3 cre-injected mice
653 and $n=11$ neurons from 2 GFP-injected mice) of the sEPSC events in DLS. Data
654 represent mean $\pm$ SEM. ns= no significant, $*P<0.05$, $****P<0.0001$.

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