Dopamine Regulates Aversive Contextual Learning and Associated In Vivo Synaptic Plasticity in the Hippocampus

Highlights

- Molecular approaches verified dopamine innervation of the hippocampus
- Inhibitory avoidance (IA) learning induces ex vivo and in vivo LTP in the CA1
- D1-like dopamine receptor inhibition prevents IA induction of LTP
- Dopamine activation enhances and inhibition prevents long-term retention of IA

Authors

John I. Broussard, Kechun Yang, Amber T. Levine, ..., Fu-Ming Zhou, Mariella De Biasi, John A. Dani

Correspondence

johndani@upenn.edu

In Brief

The role of dopamine in signaling rewards is well known, but here Broussard et al. show that dopamine D1-like receptor activity in the hippocampus is necessary for retention of aversive memories. The results indicate dopaminergic innervation and function within the hippocampus underlying long-term synaptic potentiation associated with aversive memory retention.
Dopamine Regulates Aversive Contextual Learning and Associated In Vivo Synaptic Plasticity in the Hippocampus

John I. Broussard,1 Kechun Yang,2 Amber T. Levine,1 Theodoros Tsetsenis,2 Daniel Jenson,1 Fei Cao,1 Isabella Garcia,3 Benjamin R. Arenkiel,3,4 Fu-Ming Zhou,5 Mariella De Biasi,6 and John A. Dani2,*

1Department of Neuroscience, Baylor College of Medicine, Texas Children’s Hospital, Houston, TX 77030, USA
2Department of Neuroscience, Mahoney Institute for Neurosciences, Perelman School for Medicine, Philadelphia, PA 19104, USA
3Program in Developmental Biology
4Jan and Dan Duncan Neurological Research Institute
Baylor College of Medicine, Texas Children’s Hospital, Houston, TX 77030, USA
5Department of Pharmacology, University of Tennessee, Memphis, TN 38163 USA
6Department of Psychiatry, Mahoney Institute for Neurosciences, Perelman School for Medicine, Philadelphia, PA 19104, USA
*Correspondence: johndani@upenn.edu
http://dx.doi.org/10.1016/j.celrep.2016.01.070
This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

SUMMARY

Dopamine release during reward-driven behaviors influences synaptic plasticity. However, dopamine innervation and release in the hippocampus and its role during aversive behaviors are controversial. Here, we show that in vivo hippocampal synaptic plasticity in the CA3-CA1 circuit underlies contextual learning during inhibitory avoidance (IA) training. Immunohistochemistry and molecular techniques verified sparse dopaminergic innervation of the hippocampus from the midbrain. The long-term synaptic potentiation (LTP) underlying the learning of IA was assessed with a D1-like dopamine receptor agonist or antagonist in ex vivo hippocampal slices and in vivo in freely moving mice. Inhibition of D1-like dopamine receptors impaired memory of the IA task and prevented the training-induced enhancement of both ex vivo and in vivo LTP induction. The results indicate that dopamine-receptor signaling during an aversive contextual task regulates aversive memory retention and regulates associated synaptic mechanisms in the hippocampus that likely underlie learning.

INTRODUCTION

Dopamine (DA) neurons arising from the ventral tegmental area (VTA) and substantia nigra pars compacta (SNc) contribute during the formation of rewarded behaviors (Bayer and Glimcher, 2005; Schultz, 1986, 1998). DA neurons fire phasic bursts in response to unpredicted reward, and their phasic firing begins to track neutral stimuli that predict those rewards (Hollerman and Schultz, 1998). This firing characteristic of DA neurons suggests that they are highly effective at pairing neutral stimuli to unconditioned stimuli, and this property provided evidence that DA signals are a neural substrate of reward prediction (Berridge and Robinson, 1998; Dayan and Balleine, 2002; Montague et al., 2004; Schultz et al., 1997).

Recent studies indicate that DA neurons have a more heterogeneous response profile (Henny et al., 2012). For example, dorsal VTA neurons are typically inhibited by footshocks, but ventral VTA neurons may be phasically excited by noxious stimuli (Brischoux et al., 2009; Bromberg-Martin et al., 2010; Zweifel et al., 2011). Aversive events have been shown to increase the firing rate of a subset of VTA DA neurons and, as a result, increase DA release in target areas, such as the striatum or medial prefrontal cortex (Budygin et al., 2012; Dong et al., 2010; Lammel et al., 2011). Recent studies indicate that a DA neuron’s response to negative or positive stimuli is largely dependent upon the neuron’s presynaptic inputs, and the dopaminergic signal influences separate brain regions depending on valence (Lammel et al., 2011, 2012, 2014). These findings suggest that DA signaling may encode beyond prediction errors and contribute to synaptic plasticity required for updating memory of environmental salience, and that these DA signals act upon specific neural targets.

One such potential target is the hippocampus. Earlier evidence indicated that dopaminergic projections originating primarily from the midbrain (including the VTA, substantia nigra, and retrorubral field) project directly to the hippocampus (Gasbarri et al., 1994a, 1994b, 1996). Quantitative real-time PCR confirmed that D1 and D5 receptors are found in the hippocampus, including the CA1 (Mu et al., 2011). Physiological evidence demonstrated that D1 and D5 receptors are important for controlling spike timing dependent plasticity within the hippocampus (Yang and Dani, 2014). Also, there is functional evidence that indicates that drugs of abuse, such as nicotine (Tang and Dani, 2009; Zhang et al., 2010) and methylphenidate (Jenson et al., 2015), recruit dopaminergic neurotransmission to influence synaptic plasticity within the hippocampus. Dopaminergic neurotransmission in the hippocampus during specific time
points in the retention interval also seems to be important for successful consolidation of long-term memories (Rossato et al., 2009). Despite this evidence, there is currently some controversy about the source of dopaminergic innervation in the hippocampus (Smith and Greene, 2012), suggesting that significant dopaminergic neurotransmission is the result of extra-synaptic volume transmission from other sources (Agnati et al., 1995; Borgkvist et al., 2012), such as the locus coeruleus (Walling et al., 2012).

In this study, we used virally introduced molecular markers to label dopaminergic synaptic terminals arising from the midbrain to determine whether there is significant innervation from midbrain DA areas. Then, we tested whether neurotransmission at D1-like receptors was important for the acquisition of aversive memories in an inhibitory avoidance (IA) task. After finding a sparse direct projection of midbrain DA neurons to the hippocampus and evidence of dopaminergic influence in the retention of IA learning, we tested the effects of D1-like receptors on measures of synaptic plasticity. We found that dopaminergic activity regulated IA-induced CA1 LTP measured from single pyramidal neurons in ex vivo slices or from in vivo field excitatory postsynaptic potentials (fEPSPs) in freely moving mice.

RESULTS

Evidence for Innervation of the Hippocampus by Midbrain Dopamine Neurons

Dopamine transporters (DATs) have been shown to be located on dopamine (DA) fibers and terminals (Nirenb erg et al., 1996; Shimada et al., 1991, 1992). To examine whether there is a direct dopaminergic projection to the hippocampus, we injected a recombinant adeno-associated virus (Grimm et al., 2008; Tsai et al., 2009) (AAV-EF1a-DIO-synaptophysin:GFP) containing a double-floxed inverted open reading frame encoding synaptophysin-GFP into the midbrain DA area of adult Slc6a3ires-cre/+ knockin and Slc6a3+/+ (wild-type [WT]) mice (Figures 1A and 1B). This AAV vector facilitates the expression of GFP specifically in the synaptic terminals of neurons containing DATs, exclusively in Cre-expressing cells. Because synaptophysin is a synaptic vesicle protein, this procedure concentrated the GFP fluorophore into DA terminals of neurons from the ventral tegmental area (VTA) and the substantia nigra (SN). By concentration of the fluorophore into the synaptic terminals, we increased the likelihood of finding DA-positive innervation.

As positive controls, 2 weeks after injection, we inspected the brains of injected mice and found prominent labeling of projections and cell bodies in the VTA (Figure 1C) and labeled terminals in the ventral striatum (Figure 1D) and medial prefrontal cortex (Figure 1E), major known targets for innervation by midbrain DA neurons. GFP-positive terminals were also found throughout the CA1 region of Cre-positive mice (Figures 1F and 1H). This expression was significantly greater relative to Cre-negative mice (Figure 1G): 0.83% ± 0.03%, n = 5 for Slc6a3ires-cre/+; 0.09% ± 0.004%, n = 3 for WT (Figure 1I). We inspected the locus coeruleus and found punctate terminals, but no cell bodies labeled (Figure 1J), verifying that the labeling arose from the midbrain not the locus coeruleus. To test the specificity of Slc6a3ires-cre/+ expression, we checked the cerebellum of injected Slc6a3ires-cre/+ mice and found virtually no green fluorescence in the cerebellum (0.003%, data not shown). Likewise, there was no GFP fluorescence in the hypothalamus, which is near to the midbrain DA areas (Figures S1A and S1B). Thus, the injections were restricted to the midbrain DA areas, and there the synaptophysin-GFP was found to co-localize with the catecholamine synthesis enzyme, tyrosine hydroxylase (TH) (Figure S1C).

To verify that the synaptophysin signal we observed in the hippocampal CA1 is indeed dopaminergic terminals, we injected the midbrains of a second group of Slc6a3ires-cre/+ mice with Synaptophysin-Ruby-Red (AAV-EF1a-DIO-synaptophysin:RFP) and immunostained with an antibody against TH (Figure 2A). We observed nearly complete co-localization between the Synaptophysin-Ruby-Red and TH (Figures 2A and 2B), confirming that the synaptic terminals are dopaminergic and originate from the midbrain: Ruby-Red/TH double-positive terminals, 93% ± 2%, n = 6.

Dopamine-Regulated Acquisition of Inhibitory Avoidance Memories

To test the hypothesis that DA regulates acquisition of IA memory, we injected groups of animals with SCH 23390 (SCH), a D1/D5 receptor antagonist, prior to footshock training (Figure 3A). During IA training, the mice received a footshock upon entering the dark side of the training chamber from the lighted side of the chamber. The latency to enter the darkened side was examined at three SCH doses (0.2, 0.1, and 0.05 mg/kg, i.p.). Only the higher doses of SCH slowed the approach latency during training (Figure S2): p < 0.01, n = 4–8. Therefore, we used the lowest dose of SCH (0.05 mg/kg) that did not slow the initial approach of mice to the dark side of the light/dark chamber (Figure S2).

24 hr after training (i.e., footshock), control mice injected with saline significantly delayed entering the dark (previously shocked) side (Figure 3B, black bars). On the contrary, the latency to approach the dark (shocked) side was not delayed in SCH-treated mice at three different shock intensities (Figure 3B, red bars). This dose of SCH (0.05 mg/kg, i.p.) also did not impair short-term memory (STM) because approach latency remained similar to controls at 1 and 3 hr after the shock training (Figure 3C). In summary, the D1/D5 antagonist, at a dose that did not influence approach latency during training and did not impair short-term memory, inhibited long-term memory retention in this IA task.

To test whether the affected dopaminergic receptors resided within the hippocampus, we implanted bilateral cannulas directly above the CA1 and infused SCH (1 μl, 1 mg/ml concentration given at 0.5 μl/min) 15 min prior to IA training. When tested 24 hr later, direct hippocampal infusions of the D1-like antagonist significantly reduced approach latency compared to saline controls (Figure 3D): p < 0.05, unpaired t test, n = 7, 9. Injecting a higher dose (0.2 mg/kg, i.p.) of SCH 23390 after IA training did not influence long-term memory retention 24 hr later, as indicated by an approach latency significantly higher after testing (Figure 3E): training versus testing p < 0.01, n = 10. Acting opposite to the D1-like antagonist, the D1-like agonist, SKF 81297...
(0.9 mg/kg), enhanced IA retention 72 hr after a moderate (0.4 mA) footshock (Figure 3F): p < 0.05, n = 10, 32. This result suggests that elevated DA D1R activity during training enhances retention of a more difficult training task, which was IA memory at a longer retention interval (i.e., 72 hr) to a moderate footshock (0.4 mA).

In order to test whether β-adrenergic (norepinephrine, NE) neurotransmission was important for learning the IA task, we injected two doses (10 mg/kg and 20 mg/kg, i.p.) of the β2-adrenergic antagonist, Timolol (Tim), prior to IA training. The approach latency was significantly greater when tested after 24 hr retention, indicating that mice still learned to avoid the dark side (shocked side) of the chamber even when β2-adrenergic receptors were inhibited (Figure 3G): p < 0.01, paired t tests, n = 10, 10.

Histological Analysis of Synaptophysin-GFP Labeled Terminals

(A) Didactic vector map of the AAV-EF1a-DIO-synaptophysin:GFP virus constructed for use in the viral tracing experiments. This DJ8 vector specifically expressed synaptophysin, which was primarily targeted in the fiber terminals of infected neurons.

(B) AAV-EF1α-DIO-synaptophysin:GFP was injected into the midbrain dopamine area (VTA/SNc) of Slc6a3ires-cre/+ mice and WT mice served as controls.

(C) Confocal image (10×) taken from the VTA of a Slc6a3ires-cre/+ mouse injected with Synaptophysin-GFP virus. The light blue is DAPI (Vector Laboratories), and green fluorescence indicates the reporter from Synaptophysin-GFP. Cell bodies and processes were labeled in the VTA.

(D) As a positive control, a confocal image is shown of the dense innervation of the ventral striatum, which receives innervation from VTA DA neurons.

(E) A second positive control showing green DA-terminal puncta in the medial prefrontal cortex (PFC).

(F) Image from Slc6a3ires-cre/+ mouse indicating direct dopaminergic projections from the midbrain DA area revealed as green puncta in the CA1.

(G) The CA1 of a Slc6a3ires-cre/+ (WT littermate) shows practically no green fluorescence.

(H) Confocal image (40×) of DAT terminals projecting directly to cell bodies in the PCL of the hippocampal CA1.

(I) Quantification of the number of green pixels from Slc6a3ires-cre/+ mice compared to WT littermates: 0.83% ± 0.03%, n = 5 for Slc6a3ires-cre/+; 0.09% ± 0.004%, n = 3 for WT.

(J) Locus coeruleus image illustrating Synaptophysin-GFP (DAT)-labeled terminals, but no cell somas, indicating that these are not DAT positive cell bodies.

SO, stratum oriens; PCL, pyramidal cell layer; SR, stratum radiatum. Scale bars, 100 μm (C, D, and E); 50 μm (F, G, and J); and 25 μm (H).

Inhibitory Avoidance Training Increased the AMPA/NMDA Ratio in the CA1

Learning the context of IA is expected to engage synaptic plasticity mechanisms in the CA1 region of the hippocampus (Whitlock et al., 2006). To gain insight into whether LTP occurred in the CA1 region following IA training, we cut ex vivo slices and determined the AMPA/NMDA ratio at the CA3-CA1 synapses when measured at 1.5 hr after the footshock, as compared to walk-through controls that were not shocked (Figures 4B and 4C): 0.59 ± 0.06 in control versus 0.93 ± 0.06 after IA, n = 12, 6, p < 0.01 unpaired t test. The ratio returned to baseline 24 hr after the footshock (Figures 4D and 4E): 0.66 ± 0.10 in control versus 0.59 ± 0.10 after IA, n = 6, 6,
Inhibitory Avoidance Training Increased the Slope of CA1 fEPSPs In Vivo

The ex vivo slice studies (Figure 4) suggest that IA-induced long-term synaptic potentiation (LTP) in the CA1 region. Therefore, we measured in vivo synaptic strength along the Schaffer collateral input to CA1 in freely behaving mice to determine whether there was synaptic potentiation associated with the IA learning task in real time. We monitored synaptic transmission from the stratum radiatum of the CA1 by stimulating the contralateral Schaffer collateral axons (indicated in Figure 5A; also see Figure S3) before and after IA training (Figure 5B). The day before training (−1 day) we recorded baseline fEPSP measurements from freely moving mice that were well habituated to the recording box (Figure 5C). These baseline recordings of synaptic responses were required to remain stable over the 30-min period on the day before (−1 day) and the day of IA training (0 day) to justify continuation of the multi-day in vivo recording paradigm (Figure 5C). The baseline recordings on 0 day were taken again for 30 min immediately prior to IA training, and we required continued stability from −1 day of the fEPSP slope in both the control and IA groups: p > 0.05, n = 4 for control, n = 9 for IA. Walk-through controls that were exposed to the IA apparatus but not shocked demonstrated no significant change in the slope of the fEPSP of the CA3-CA1 circuit (Figure 5C, white circles): F(41,123) = 1.165; p > 0.05. IA training significantly increased the slope of the fEPSPs (Figure 5C, black circles): F(41, 328) = 5.065, p < 0.01. Example fEPSPs are shown (Figure 5B) at the times indicated in the recording time course (Figures 5C, 1, and 2). In both groups synaptic transmission returned to baseline levels the following day (1 day): controls, F(17, 51) = 0.50; IA, F(17, 136) = 0.45; both p > 0.05, n = 4, 9.

Learning-Induced In Vivo LTP Required Activity of D1-like DA Receptors

We recorded fEPSPs from the stratum radiatum of the CA1 while the contralateral Schaffer collaterals were stimulated (Figure 6A). As always, a long-term stable baseline was required on −1 day and 0 day before the injections and before the continuation of recording on 0 day. As in Figure 5 (without the inhibitors), IA training increased the slope of the fEPSP that persisted after inhibition of β2-adrenergic receptors with Tim (10 mg/kg, i.p., Figures 6B and 6C, gray traces and circles): F(41,164) = 1.98; p < 0.01, n = 5. However, following inhibition of D1-like receptors with SCH (0.05 mg/kg, i.p.), IA training did not significantly change the slope of the fEPSP (Figures 6B and 6C, red traces and circles): F(41, 246) = 1.35; p > 0.05. As in previous experiments (Figure 2), SCH significantly impaired footshock memory retention of IA by the implanted mice (Figure S4, red bar): 61.2 ± 31.0 s approach latency, n = 7, p = 0.12, paired t test. After Tim injection, implanted mice retained the IA memory (Figure S4, gray bar): 131.2 ± 20.1 s approach latency, n = 5, p < 0.05, paired t test. Separate in vivo fEPSP recordings showed that neither drug in the absence of training changed basal synaptic transmission (Figure 6C): F(41, 246) = 1.03, p > 0.05, n = 7 for SCH alone; F(41,164) = 0.94, p > 0.05, n = 5 for Tim alone. These results suggest that DA neurotransmission influences learning and the associated synaptic plasticity along the CA3-CA1 pathway after aversive conditioning.

Inhibiting D1-like Dopamine Receptors Prevented IA Learning and Synaptic Plasticity

After demonstrating that the IA learning-induced increase in the AMPA/NMDA ratio were specific in the hippocampus to the CA3-CA1 circuit, we tested the hypothesis that DA D1-like activity is necessary for increases in the AMPA/NMDA ratio. The D1-like receptor inhibitor, SCH, at doses that inhibited learning (0.05 mg/kg; Figures 3B and S4) also inhibited IA-induced increases in the AMPA/NMDA ratio (Figures 7A–7C). In contrast, the β2-adrenergic receptor antagonist, Tim (10 mg/kg), did not block learning-induced changes in the AMPA/NMDA ratio (Figures 7D–7F). These results are consistent with D1-like receptor
activation significantly influencing CA1-CA3 IA-induced synaptic potentiation.

**DISCUSSION**

Fluorescent markers associated with DATs and TH indicated that dopaminergic fibers and terminals project sparsely from the midbrain DA neurons to the dorsal CA1 (Figures 1 and 2). A low dose of a D1-like receptor antagonist did not impair the approach time (latency) for IA training and did not impair short-term memory (Figure 3C) but did prevent the long-term retention (24 hr) of the fear memory (Figure 3B). Training in the IA task produced significant increases in the AMPA/NMDA ratio of the CA1 synapses (Figure 4C) and significantly increased the slope of the fEPSP in the CA1 of freely moving mice (Figure 5C). D1-like receptor inhibition blocked learning-induced increases in the slope of the fEPSP in the CA3-CA1 circuit (Figure 6) and blocked the learning-induced enhancement in the AMPA/NMDA ratio in the CA1 (Figure 7C). Taken together, these results suggest that D1-like receptor activity in the hippocampus contributes to the synaptic plasticity associated with acquisition of contextual memory required for long-term retention of aversive memories.

**Anatomical Evidence of a Direct Midbrain Dopaminergic Projection to the Hippocampus**

There is known noradrenergic innervation of the hippocampus, and D1-like receptor activity can arise from DA released from noradrenergic innervation. Much of the DA receptor activity may arise from neurons originating in the locus coeruleus (Smith and Greene, 2012). Our results also support a sparse direct projection from midbrain DA neurons to the hippocampus as indicated by GFP- and Ruby-Red-synaptophysin staining of DAT-expressing neurons (Figures 1 and 2). Injections were localized to the midbrain and no fluorescence was found in the hypothalamus or from the cell bodies in the locus coeruleus (Figure 1J), which precludes noradrenergic neurons as the source of the labeling that we observed in the hippocampus. Furthermore, fluorescent labeling of terminals in the CA1 resulting from viral
infection of the VTA/SNc region highly co-localized with TH, identifying these terminals as dopaminergic (Figure 2). These data support previous evidence arising from retrograde tracers injected in the hippocampus that indicated dopaminergic innervation of the hippocampus from the VTA/SNc (Broussard et al., 2012; Gasbarri et al., 1994b, 1997). Our results also confirm recent studies using viral expression of channelrhodopsin in Slc6a3ires-cre/+ mice that labeled relatively sparse dopaminergic axons in the CA1 region (McNamara et al., 2014; Rosen et al., 2015).

Despite the sparse innervation of the hippocampus by midbrain dopaminergic axons, several lines of evidence indicate a significant functional role for this dopaminergic innervation. It was shown that nicotine can cause in vivo hippocampal long-term synaptic plasticity, and this effect required local activation of DA receptors (Tang and Dani, 2009). Inactivation of the VTA with local infusion of TTX inhibited this effect, demonstrating that the required dopamine signal originated, at least in part, from the midbrain (Tang and Dani, 2009). In more recent studies, researchers expressed channelrhodopsin-2 specifically in dopaminergic neurons in the VTA/SNc. Optical stimulation of these dopaminergic fibers in the VTA or locally in the hippocampus in mice exploring novel environments enhanced hippocampal reactivation and improved spatial learning and memory (McNamara et al., 2014). In addition, optogenetic release of dopamine exclusively from the VTA was shown to cause a bidirectional, activity-dependent modulation of Schaffer collateral synapses in hippocampal slices (Rosen et al., 2015). Our data are consistent with those findings, indicating the existence of anatomical connectivity between midbrain dopamine centers and the hippocampus.

Although DA neurons have been traditionally viewed to reinforce rewarding behaviors and/or to signal the expectation of reward, studies also have indicated that DA can influence the persistence of aversive learning (Moncada et al., 2011; Ortiz et al., 2010; Rossato et al., 2009). A population of ventral VTA neurons located within the paranigral nucleus (PN) has been shown to respond specifically to aversive stimuli within seconds (Brischoux et al., 2009), resulting in increased DA in target areas (Dong et al., 2010; Horvitz, 2000; Kienast et al., 2008). Anatomically, this subpopulation of DA neurons projects to the medial prefrontal cortex and the medial shell of the nucleus accumbens (Lammel et al., 2011, 2012). Interestingly, in an older retrograde labeling study, it was shown that a population of TH-positive neurons within the PN also connects to the hippocampus (Gasbarri et al., 1994b).

**D1-like Receptor Activity Enhanced Memory and Plasticity in CA3-CA1 Synapses**

Our data support that the synaptic plasticity associated with aversive learning is dependent upon the temporal coincidence of glutamatergic and dopaminergic neurotransmission. D1-like

**Figure 4. IA Training Increased the AMPA/NMDA Current Ratio in CA1 Pyramidal Neurons, but Not in Dentate Gyrus Granule Cells**

(A) Diagram illustrating the whole-cell recording from CA1 pyramidal neurons in hippocampal slices. The stimulating electrode (S) was placed on the Schaffer collateral path and the recording electrode (R) onto a CA1 pyramidal neuron. 

(B and D) Representative traces of AMPA (gray traces) and NMDA (black traces)-receptor-mediated whole-cell currents recorded from CA1 pyramidal neurons from control (unshocked, left) and IA (shocked, right) mice, which were decapitated either 1.5 hr after training (B), or after testing (D). 

(C and E) The average of AMPA/NMDA ratios from CA1 pyramidal neurons are plotted. IA training significantly increased the AMPA/NMDA ratio in slices prepared from animals decapitated 1.5 hr after training: control versus IA, 0.59 ± 0.06 versus 0.93 ± 0.06, n = 12, 6, p < 0.01, but not in slices prepared from animals after testing: control versus IA, 0.66 ± 0.10 versus 0.59 ± 0.10, n = 6, 6, p > 0.05. 

(F) Diagram illustrating the whole-cell recording from dentate gyrus granule cells. The stimulating electrode (S) was placed on the medial perforant path (MPP) and the recording electrode (R) onto a DG granule cell. 

(G and I) Representative traces of AMPA (gray) and NMDA (black)-receptor-mediated whole-cell currents recorded in granule cells from control (unshocked, left) and IA (shocked, right) mice, which were sacrificed either 1.5 hr after training (G) or after testing (I). 

(H and J) IA training had no effect on the AMPA/NMDA ratio measured from dentate gyrus granule cells at these time points: 1.5 hr, control versus IA, 0.77 ± 0.06 versus 0.63 ± 0.07, n = 10, 11, p > 0.05; post test, control versus IA, 0.70 ± 0.10 versus 0.57 ± 0.06, n = 5, 7, p > 0.05.
receptor inhibition after the behavioral training did not prevent remembering the training that occurred just minutes before the D1-like receptor inhibition. The 2-min window after training was sufficient to allow DA to bind to D1 receptors to initiate the mechanisms of LTP and memory retention. Activity initiated by the D1-like receptors continued despite the later arrival of the D1-like antagonist. Thus, synaptic potentiation and learning proceeded owing to the original initiation of these memory-related processes before the antagonist arrived 2 min after the training period.

An interesting concept was that D1-like receptor activity in the CA1 was necessary for retention of long-term, but not short-term IA memory. In the 1- to 3-hr time window, mice still avoided the footshocked side even after D1-like receptors were inhibited during training. Similarly, DA was important for long-term retention at 6 hr, but not at 20 min in a different behavioral task (O’Carroll et al., 2006), and a D1-like antagonist applied after training did not affect memory consolidation (Rossato et al., 2009).

Shorter-term memory will likely involve different mechanisms from longer-term memory that is consolidated over time, often requiring periods of sleep. The hippocampal synaptic potentiation indicated in vivo by the fEPSPs or ex vivo by the AMPA/NMDA ratios correlated with the long-term memory tested 24–72 hr later. In both cases (i.e., the in vivo fEPSP and ex vivo AMPA/NMDA ratio), the indication of LTP was expressed as a relatively global feature of the CA1 area we were studying.

That is, different measures that in one case averaged a broad area (in vivo fEPSP) or in the other case picked individual CA1 pyramidal neurons (ex vivo AMPA/NMDA ratio) both indicated the presence of LTP. These results suggest that to retain memory for 24–72 hr a high percentage of a general area of the CA1 region was contributing: nearly every single CA1 pyramidal neuron we studied had an elevated AMPA/NMDA ratio. Because a large area and high percentage of neurons were contributing, we were able to measure those diverse indications of LTP with our techniques. Even under those successful circumstances, however, the in vivo fEPSP rose slowly over approximately 30 min (see Figure 5C, black data), suggesting an imperfect match between the learning and presence of LTP as measured by the in vivo fEPSP.

When D1Rs were inhibited, in vivo the global fEPSP did not change during IA training and the ex vivo slices did not show LTP, and the memory was not observed at 24–72 hr. All of these data to this point are consistent and easily understandable. However, at early times after IA training with D1-like receptors inhibited, short-term memory was observed without global indications of LTP by fEPSPs or AMPA/NMDA ratios. That result may be because without D1-like activity a high percentage of the dorsal CA1 is not contributing to the memory process needed to retain information for days. However, smaller components of the dorsal hippocampus (or other areas of the hippocampus or brain) could be contributing to the short-term memory seen at 1–3 hr. Consistent with this speculation, for example, hippocampal dopaminergic activity was shown to influence long-term but not short-term memory retention in appetitive tasks (Bethus et al., 2010). In the present experiments, DA was shown to be important during the encoding phase of a long-term memory in an aversive paradigm. Thus, DA neurotransmission within the CA1 may serve to associate neutral spatial contexts with...
unexpected unconditioned stimuli. This conclusion is consistent with the hypothesis that DA signals under these circumstances contribute to the updating of the rodent’s internal perception (or map) of environmental saliency (Bethus et al., 2010; McNamara with the hypothesis that DA signals under these circumstances stabilize learning-induced synaptic change (Frey and Morris, 1997, 1998). Our data indicate that aversive IA training produces broad synaptic plasticity in the CA3-CA1 synapses that is strongly influenced by D1-like neurotransmission. Thus, DA signaling within the hippocampus serves to consolidate the association of a neutral context with aversive, unconditioned stimuli by contributing to the plasticity of the CA3-CA1 circuitry that likely contributes to memory retention (McHugh et al., 2007; Rumpel et al., 2005).

**Conclusions**

In summary, DA D1-like receptor activity modulates and regulates hippocampal CA1 synaptic plasticity, associated learning, and long-term memory arising from an aversive contextual task. Although DA signaling from noradrenergic innervation may contribute, dopaminergic innervation of the CA1 arises from the VTA/SNc and $\beta_2$ adrenergic signaling did not significantly influence memory retention examined in our paradigms. These results support the hypothesis that aversive events recruit subpopulations of DA neurons (Brischoux et al., 2009; Henny et al., 2012; Lammel et al., 2011, 2012, 2014) to contribute to learning and memory.

**EXPERIMENTAL PROCEDURES**

The methodology and reagents for immunohistochemistry, ex vivo electrophysiology, surgery and in vivo electrophysiology, and statistical analysis are described in the Supplemental Experimental Procedures.

**Animals**

Wild-type C57BL/6J mice 2–4 months old (Jackson Laboratory) had free access to food and water and were housed in accordance with the guidelines specified by the Institutional Animal Care and Use Committee at Baylor College of Medicine or University of Pennsylvania and the Public Health Service Policy on Humane Care and Use of Laboratory Animals. All animals were maintained on a reverse light cycle with the lights off from 6 a.m. to 6 p.m., and all studies were conducted during the dark phase. For anatomy experiments (Figures 1 and 2), Slc6a3ires-cre/+ mice were obtained from Jackson Laboratory (stock #: 006660, which are commonly referred to as DATires-cre mice).

**Inhibitory Avoidance Behavioral Task**

Mice were handled in the experiment room for 3–4 days and habituated to needle injections with saline before the start of the experiments, and all experiments were conducted during the dark cycle. Prior to IA training, all animals were injected i.p. with either saline, SCH 23390, SKF 82197, or Tim. Training consisted of 40 s acclimatizing to the light chamber, and then the trap door to the dark chamber was opened. Once mice walked into the dark chamber, the trap door closed and the mice were given a single footshock (2 s) followed by 2 min within the dark chamber before being returned to their home cages. Footshock intensity was 0.8 mA unless otherwise noted. After IA training, animals were tested for avoidance retention at 1, 3, 24, or 72 hr or sacrificed for in vitro neurophysiological assays. IA retention was assayed after training by replacing mice into the light side of the chamber and measuring the latency before the animal returned to the dark context side where the shock had been administered. The footshock was not re-administered during the retention assay and measurements.
were terminated at a ceiling delay interval of 180 s. Statistical analysis was determined using repeated-measures ANOVA with the criterion significance set at $p < 0.05$.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Supplemental Experimental Procedures and five figures and can be found with this article online at http://dx.doi.org/10.1016/j.celrep.2016.01.070.

**AUTHOR CONTRIBUTIONS**

J.I.B. led the wet lab effort and did the in vivo electrophysiology assisted by D.J. K.Y. did the ex vivo electrophysiology. A.T.L. did much of the IA behavior. I.G. and B.R.A. devised and assisted J.I.B. with the vectors and GFP studies. F.C. did viral injections and aspects of histology and imaging. F.-M.Z. helped to devise studies on dopamine innervation of the hippocampus. J.A.D. originated, planned, and oversaw the experiments with M.D.B.'s assistance. J.A.D. and J.I.B. did much of the writing. T.T. performed the TH co-localization studies of Figure 2 and assisted with the writing.

**ACKNOWLEDGMENTS**

This work was supported by grants from the NIH (NS21229 and DA09411 to J.A.D., DA021194 and MH067119 to F.-M.Z., and DA017173 to M.D.B.). We wish to thank Andrea Stout for help with the Metamorph software. J.A.D., DA021194 and MH067119 to F.-M.Z., and DA017173 to M.D.B.). We wish to thank Andrea Stout for help with the Metamorph software.

Received: July 2, 2015
Revised: November 4, 2015
Accepted: January 22, 2016
Published: February 18, 2016

**REFERENCES**


Yang, K., and Dani, J.A. (2014). Dopamine D1 and D5 receptors modulate spike timing-dependent plasticity at medial perforant path to dentate granule cell synapses. J. Neurosci. 34, 15888–15897.
