

Hippocampus-driven feed-forward inhibition of the prefrontal cortex mediates relapse of extinguished fear

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The medial prefrontal cortex (mPFC) has been implicated in the extinction of emotional memories, including conditioned fear. We found that ventral hippocampal (vHPC) projections to the infralimbic (IL) cortex recruited parvalbumin-expressing interneurons to counter the expression of extinguished fear and promote fear relapse. Whole-cell recordings *ex vivo* revealed that optogenetic activation of vHPC input to amygdala-projecting pyramidal neurons in the IL was dominated by feed-forward inhibition. Selectively silencing parvalbumin-expressing, but not somatostatin-expressing, interneurons in the IL eliminated vHPC-mediated inhibition. In behaving rats, pharmacogenetic activation of vHPC→IL projections impaired extinction recall, whereas silencing IL projectors diminished fear renewal. Intra-IL infusion of GABA receptor agonists or antagonists, respectively, reproduced these effects. Together, our findings describe a previously unknown circuit mechanism for the contextual control of fear, and indicate that vHPC-mediated inhibition of IL is an essential neural substrate for fear relapse.

Extinction learning is essential for cognitive-behavioral therapies in patients with trauma-related and stressor-related disorders, including post-traumatic stress disorder^{1,2}. Despite the success of these approaches, extinction training does not erase traumatic memories, but instead only dampens the expression of those memories. Consequently, work in both rats and humans has revealed that extinguished fear ‘relapses’ under a number of circumstances^{3,4}. For example, extinction training only transiently suppresses learned fear responses, such as freezing behavior, in rats conditioned to fear an auditory conditioned stimulus (CS)⁵. Notably, fear to an extinguished CS is ‘renewed’ if it is encountered outside of the extinction context⁶, a phenomenon that reveals that extinction memories are context dependent. In humans undergoing exposure therapy, the context-dependence of extinction leads to fear relapse outside of the clinic^{3,4}. For this reason, understanding the neural circuits underlying the relapse of extinguished fear is critical for developing effective neurobehavioral interventions for trauma-related and stressor-related disorders.

Considerable evidence suggests that encoding and consolidation of extinction memories require the mPFC^{7–9}. After extinction is learned, projections from the IL cortex to the amygdala inhibit the production of conditioned fear responses, including freezing behavior^{10,11}. The recruitment of this prefrontal-amygdala circuit is context dependent: Fos expression in the IL is maximal in the extinction context, but is reduced in other contexts (where fear relapses)^{12,13}. This contextual control of prefrontal-amygdala inhibition appears to depend, at least in part, on the hippocampus¹⁴. Specifically, lesions or pharmacological inactivation of the hippocampus lead to a context-independent expression of extinction

and prevent fear renewal^{15–17}. These findings indicate that the hippocampus drives fear renewal when an extinguished CS is encountered outside of the extinction context. Indeed, the renewal of fear recruits vHPC neurons projecting to the amygdala^{13,18} and engages ‘fear’ neurons in the amygdala that receive hippocampal input^{19,20}. Moreover, optogenetic silencing of vHPC neurons projecting to the central nucleus of the amygdala, but not the basolateral amygdala, prevents renewal²¹.

Although renewal engages neural circuits involved in fear expression, prevailing models of relapse phenomena, including renewal, posit that a loss of inhibitory control is necessary for the return of conditional responding to an extinguished CS^{5,14}. Notably, we recently observed that renewal increased Fos expression in vHPC neurons projecting to the mPFC²², suggesting that vHPC→mPFC projections might mediate the loss of inhibitory control hypothesized to yield fear relapse. Here we characterized the physiology of these projections and explored their contribution to the contextual control of extinguished fear memories.

Results

Characterization of vHPC→mPFC projections *ex vivo*. Hippocampal projections to the mPFC have been identified²³, but the precise projection patterns, local circuits that are driven by these inputs and functional role of these projections are not fully understood. To determine the nature of hippocampal projections to the mPFC, we expressed channelrhodopsin (ChR2) using viral-mediated transduction of neurons in the area CA1 of the vHPC (Fig. 1a,e), the primary projection site from the hippocampus to the mPFC²⁴. *Ex vivo* whole-cell recordings from neurons in the mPFC

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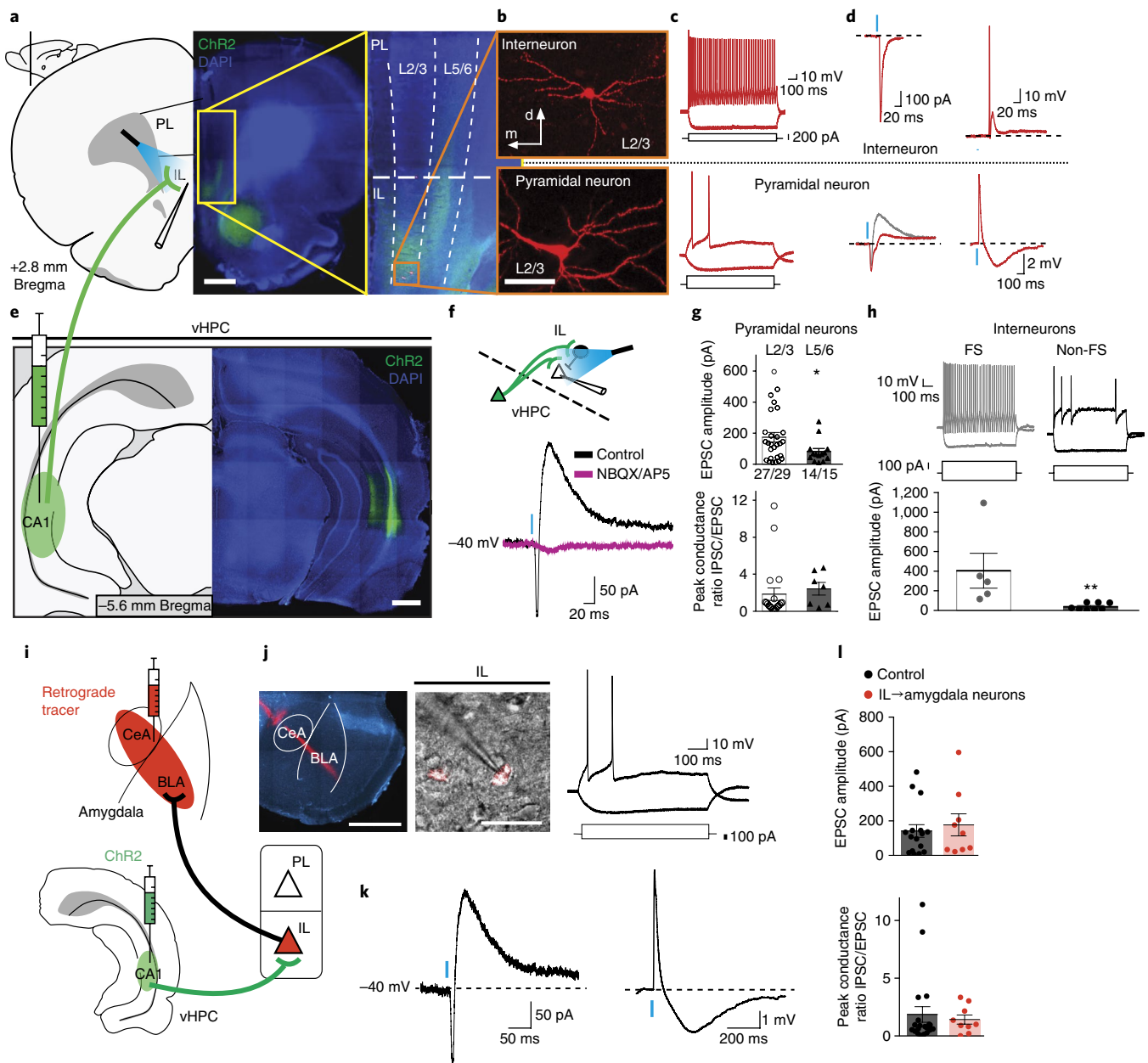


Fig. 1 | vHPC projection to the mPFC is dominated by strong local feed-forward inhibition mediated by FS interneurons in the IL. **a, e**, Coronal images of terminal labeling in the IL following injection of ChR2-expressing virus in the CA1 region of the vHPC; infusion site and terminal expression were confirmed by GFP expression (DAPI, blue). Scale bars represent 1 mm. **b**, Biocytin-recovered IL neurons after recording. Scale bar represents 20 μ m. **c**, Intrinsic firing properties of the corresponding interneuron (top) and pyramidal neuron (bottom). **d**, Voltage-clamp (left; red, holding potential of -70 mV; gray, -40 mV) and current-clamp recordings (right) of the recovered interneuron and principal neuron following optical stimulation of vHPC terminals. Note that terminal release in the interneuron caused the neuron to spike (top trace), whereas the pyramidal neuron displayed a small EPSP followed by inhibition. **f**, Synaptic response in a L2/3 principal neuron in the IL showing a monosynaptic EPSC, followed by a feed-forward inhibitory current (black trace). Both the EPSC and IPSC were eliminated in the presence of NBQX and AP5 (purple trace). **g**, EPSC amplitudes (top; L2/3: $n = 27$; L5/6: $n = 14$; one-tailed Mann-Whitney test, $P = 0.046$) and peak conductance ratios of IPSC/EPSC components (bottom; measured at -40 mV; L2/3: $n = 20$; L5/6: $n = 7$; one-tailed Mann-Whitney test, $P = 0.11$) in L2/3 and L5/6 pyramidal neurons. **h**, Comparison of EPSC amplitudes in FS (top left) and non-FS interneurons (top right) revealed significantly larger vHPC input onto FS neurons (bottom; EPSCs for FS neurons: 405 ± 178 pA, $n = 5$; non-FS neurons: 38 ± 13 pA, $n = 8$, from 5 animals; two-sided Mann-Whitney test, $P = 0.0016$). Current injections are shown below the firing traces. Amygdala-projecting neurons in the IL received input from the vHPC. **i**, Schematic illustration of experimental setup. Retrograde markers (red RetroBeads, Lumafluor) were injected into the amygdala and ChR2-expressing virus was injected into the vHPC. **j**, Typical discharge properties of amygdala-projecting pyramidal neurons (amygdala injection with retrograde marker (white bar inset: 1 mm) and recorded amygdala-projecting IL neuron (white bar inset: 20 μ m) shown on the left) to positive and negative current injections (shown below traces). **k**, Typical voltage-clamp (left) and current-clamp (right) responses to optical stimulation of vHPC input to amygdala-projecting IL pyramidal neuron. **l**, Comparisons of the IPSC/EPSC peak conductance ratio (bottom; control neurons: $n = 20$; projection neurons: $n = 9$; two-tailed t test, $t_{27} = 0.42$; $P = 0.68$) and EPSC amplitudes (top) between control neurons (black) and amygdala-projecting neurons (red; control neurons: $n = 16$; projection neurons: $n = 9$; two-tailed t test, $t_{23} = 0.52$; $P = 0.61$). Blue bars illustrate 470-nm light stimulation. Dots represent data from individual neurons. Error bars show mean \pm s.e.m. * $P < 0.05$, ** $P < 0.01$.

combined with optical terminal stimulation of vHPC input revealed targeted, location-specific innervation. We found that vHPC input innervates both the IL and prelimbic (PL) prefrontal cortex, but with rostral and caudal PL neurons receiving significantly smaller input as compared with the IL (Supplementary Fig. 1). In the IL, whose activity is crucial for extinction^{7,8}, both pyramidal neurons and interneurons (Fig. 1b) received excitatory input from the vHPC (Fig. 1c,d) with no failures and low response jitter and latency, consistent with direct monosynaptic connections (Supplementary Fig. 2). Input to interneurons was large (mean excitatory postsynaptic current (EPSC), amplitude: 237 ± 66 pA; $n=9$ of 10) and was able to drive these cells to threshold (Fig. 1d). In contrast, input to pyramidal neurons was subthreshold (Fig. 1d), but was significantly larger to layer 2/3 (L2/3) neurons, with a mean EPSC amplitude of 173 ± 31 pA ($n=27$ of 29), than to L5/6 neurons, which had a mean EPSC amplitude of 81 ± 20 pA ($n=14$ of 14, one-tailed Mann-Whitney test, $P=0.046$; Fig. 1g).

When pyramidal neurons in the IL were depolarized (Fig. 1d), stimulation of vHPC input revealed a delayed disynaptic inhibitory postsynaptic current (IPSC; Fig. 1d) that shunted the excitatory postsynaptic potential (EPSP) and evoked a long-lasting inhibitory synaptic potential (Fig. 1d). This large disynaptic inhibition in pyramidal neurons is consistent with strong vHPC-driven recruitment of local interneurons and feed-forward inhibition (Fig. 2). All of the vHPC-evoked synaptic inputs were blocked by the application of AMPA/kainate- and NMDA-receptor antagonists NBQX and AP5 ($n=3$; Fig. 1f), confirming the glutamatergic nature of this hippocampal projection. The vHPC-driven disynaptic inhibition of principal neurons was large, with a peak IPSC to EPSC conductance ratio of 1.86 ± 0.67 ($n=20$) in L2/3 principal neurons and 2.44 ± 0.68 ($n=7$) in L5/6 principal neurons (Fig. 1g), indicating that vHPC-driven inhibition is nearly twice as large as incoming excitatory input. Interneurons in the mPFC form a diverse population of cells that can be separated by their discharge properties and expression of cellular markers²⁵. Quantification of vHPC input to randomly selected interneurons revealed a distinct innervation pattern, with fast-spiking (FS) interneurons receiving significantly larger input than non-FS interneurons (EPSCs for FS neurons: 405 ± 178 pA, $n=5$; non-FS neurons: 38 ± 13 pA; $n=8$, from 5 animals; two-sided Mann-Whitney test, $P=0.0016$; Fig. 1h).

Feed-forward inhibition in vHPC→IL projections is mediated by parvalbumin-positive interneurons. The vHPC-driven IPSC consisted of two components: a fast GABA_A-receptor mediated component (50–100 ms), which was blocked by the selective GABA_{A/C}-receptor antagonist picrotoxin ($n=3$), and a slower (150–1,000 ms) component, which was blocked by the selective GABA_B-receptor antagonist CGP55845 ($n=4$) (Supplementary Fig. 3). The fast disynaptic IPSC was present in ~88% of L2/3 principal neurons and ~62% of L5/6 neurons, whereas the slow IPSC occurred in ~60% of L2/3 and ~37% of L5/6 principal neurons (Supplementary Fig. 3c), and this inhibitory conductance was capable of suppressing spiking activity of local IL pyramidal neurons (Supplementary Fig. 3d). These results indicate that the effect of vHPC input to pyramidal neurons in the IL is largely inhibitory, and is largest to neurons in L2/3. Given that pyramidal neurons in the IL that project to the basolateral amygdala (BLA) are primarily located in L2/3^{26,27}, we tested whether vHPC input targets amygdala-projecting neurons in the IL by combining optical HPC stimulation with retrograde bead injections into the basolateral and central amygdala (Fig. 1i,j). Amygdala-projecting L2/3 neurons in the IL received excitatory vHPC input (nine of ten) of similar amplitude as that to randomly selected IL pyramidal neurons (Fig. 1l). Moreover, amygdala-projecting neurons received dual component disynaptic IPSCs (eight of nine with fast IPSC components and three of nine with slow IPSC components) and a similar IPSC/EPSC conductance ratio (Fig. 1k,l),

indicating that the vHPC input to the IL modulates amygdala-projecting pyramidal neurons.

Our results indicate that vHPC input to the IL generates strong feed-forward inhibition onto principal neurons that project to the amygdala. To confirm that local IL interneurons are responsible for this dual component feed-forward inhibition, we first used local electrical stimulation in the presence of AMPA/kainate- and NMDA-receptor blockade (NBQX and AP5) to isolate inhibitory synaptic transmission. Stimulation in the IL evoked IPSCs in principal neurons that contained fast and slow inhibitory components (Supplementary Fig. 4), confirming that local IL interneurons evoke dual component inhibition in pyramidal neurons. As with most cortical regions²⁵, the mPFC contains a diverse population of interneurons²⁸ and, in most cortical regions, these interneurons provide fast GABA_A-receptor mediated inhibition. Although GABA_B-receptor mediated inhibition in principal neurons has been known for many years, the predominant subtype of interneuron that mediates this synaptic current are the neurogliaform cells^{29,30}. Given that principal neurons in the IL receive both fast and slow feed-forward inhibition, but the intrinsic firing properties of vHPC-innervated interneurons in the IL are different to those of neurogliaform cells²⁹, we asked which interneuron population provides this inhibition. We therefore selectively expressed ChR2 in parvalbumin (PV)- or somatostatin (SOM)-positive interneurons using PV::Cre and SOM::Cre mice, respectively (Fig. 2a,f). Both cell types had a maintained inward current to prolonged optical stimulation, confirming expression of ChR2 (Fig. 2b). In ex vivo acute brain slices, ChR2-transduced PV⁺ interneurons showed typical FS discharge patterns (Fig. 2c), whereas SOM⁺ interneurons showed a stuttering firing phenotype³¹ (Fig. 2g,h). Whole-cell recordings from IL pyramidal neurons revealed that optical stimulation of PV⁺ interneurons evoked fast GABA_A-receptor-mediated IPSCs in eight of eight neurons with clear, slow GABA_B-receptor-mediated IPSCs in five of eight neurons (Fig. 2d,e). Stimulation of SOM⁺ interneurons evoked GABA_A-receptor-mediated IPSCs in ten of ten neurons, and slow GABA_B-receptor-mediated IPSCs in five of ten neurons (Fig. 2i); data from four PV::Cre and two SOM::Cre animals), with the ratio of the two currents (IPSC_{slow}/IPSC_{fast}) evoked by PV⁺ interneurons being 1.57 ± 0.47 and 2.39 ± 0.74 when evoked by SOM⁺ interneurons (two-sided t test: $t_{(12)}=0.87$, $P=0.40$). Thus, both PV⁺ and SOM⁺ interneurons in the IL can mediate dual component IPSCs (Fig. 2j).

To determine whether vHPC-evoked disynaptic inhibition is mediated by one subtype of interneuron, we tested the effect of selectively silencing these interneurons on vHPC-evoked IPSCs. Using PV::Cre animals, we labeled vHPC input with ChR2 and used an ivermectin-gated chloride channel system (IL injection of AAV-DIO-ivermectin-GFP) to silence PV⁺ interneurons (Fig. 3a). In acute brain slices, application of ivermectin blocked PV⁺ interneuron firing to threshold current injection (Fig. 3b), confirming the expression of the ivermectin-gated chloride channel. Activation of vHPC input to IL principal neurons evoked disynaptic IPSCs in L2/3 pyramidal cells with fast and slow components (Fig. 3c), and application of ivermectin to silence PV⁺ neurons abolished the slow GABA_B-mediated component (three of three), whereas the fast GABA_A-mediated IPSC was blocked in one of three recordings (Fig. 3c), indicating that PV⁺ neurons in the IL can mediate vHPC-driven feed-forward inhibition onto principal neurons. Given that the pharmacological approach is intrinsically slow, to silence interneurons in a more time-locked manner, we applied a dual-optogenetic approach in which vHPC input was recruited using the light-sensitive opsin Chronos, whereas interneurons were silenced using ArchT (Fig. 3d,f). Chronos-driven vHPC input was excited using 420-nm light, whereas ArchT was specifically activated using 640-nm light (Fig. 3e). A prolonged 640-nm light pulse completely blocked Chronos-evoked spiking of

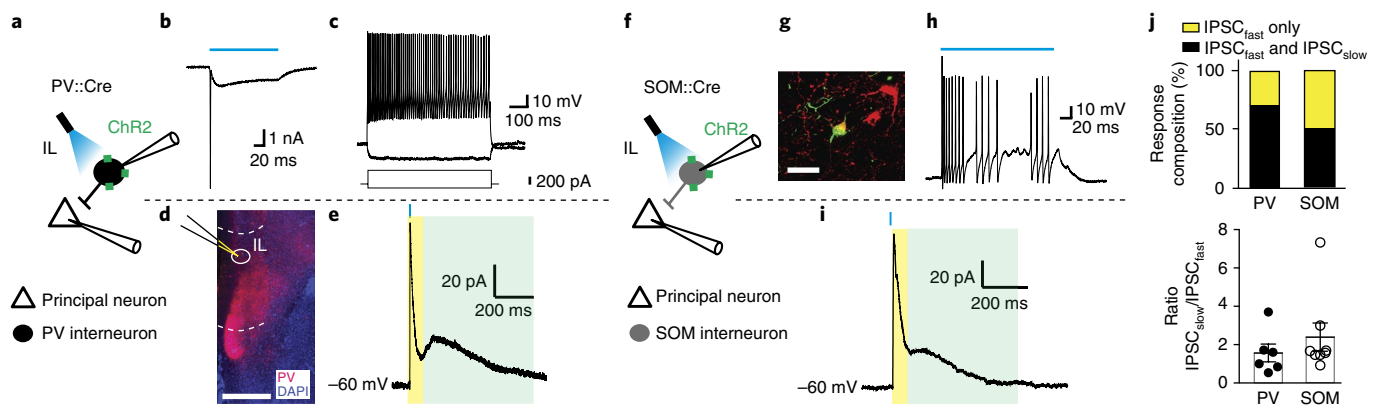


Fig. 2 | Selective optical activation of interneuronal subtypes in the IL evokes slow and fast inhibitory conductances onto pyramidal neurons. **a**, Schematic showing experimental setup of optical stimulation of ChR2-infected PV⁺ interneurons in PV::Cre animals while recording synaptic responses of pyramidal neurons in the IL. **b**, Voltage-clamp trace of an infected PV⁺ interneuron to a prolonged 470-nm light pulse (blue bar; 100 ms) showing a maintained inward current with initial spiking. **c**, Current injection (shown below the trace) revealed a typical regular FS intrinsic discharge pattern. **d**, Coronal section containing the IL showing PV cell labeling (DIO-ChR2-mCherry, red; DAPI, blue) and the recording site (circle). Scale bar represents 1 mm. **e**, PV⁺-interneuron-driven responses in local pyramidal neurons typically evoked an IPSC containing a fast (yellow) and slow component (light green). **f**, Schematic of specific infection of IL SOM⁺ interneurons with ChR2 using SOM::Cre animals to probe SOM⁺-mediated inhibition in the IL. **g**, Example of a biocytin-recovered (green) SOM⁺ interneuron (DIO-mCherry, red). Scale bar represents 20 μ m. **h**, Infected SOM⁺ cells responded to blue light (blue bar; 100 ms) and caused a stuttering discharge pattern. **i**, Optical stimulation of SOM⁺-interneurons evoked IPSCs in local pyramidal neurons with fast (yellow) and slow IPSCs (light green). **j**, The average response composition (top) was similar for both PV⁺ and SOM⁺ interneurons with either only fast IPSCs (yellow) or fast and slow IPSCs (black). Neurons with an IPSC fast (IPSC_{fast}) and slow (IPSC_{slow}) response showed an amplitude ratio (bottom) that was equal for both PV⁺ (filled circle; $n = 6$) and SOM⁺ (empty circles; $n = 8$) driven inhibition.

infected PV⁺ interneurons ($n = 3$; Fig. 3g,h). In IL pyramidal neurons in PV::Cre animals, stimulation with 420-nm light evoked an EPSC-IPSC response (Fig. 3i), and silencing PV⁺ interneurons with 640-nm light during vHPC stimulation eliminated the inhibitory component while preserving the excitatory input ($n = 5$ from 3 animals; Fig. 3i,l). In contrast, silencing SOM⁺ interneurons using ArchT (Fig. 3j) had no effect on vHPC-evoked feed-forward inhibition ($n = 6$ from 2 animals; Fig. 3k,l). Together, these results indicate that vHPC input to the IL innervates both principal neurons and interneurons, but is dominated by the input to PV⁺ interneurons, which drive large disynaptic inhibition of principal neurons in the IL to shape spiking activity.

vHPC→IL projections mediate fear relapse. Considerable work has revealed that the IL is essential for suppressing fear after extinction learning, which is mediated by its excitatory projections to the amygdala^{8, 10, 32}, whereas the vHPC appears to limit fear suppression and promote fear relapse (that is, renewal)¹⁶. In extinction, IL-mediated reduction of fear results from an excitatory projection from IL to the BLA and activation of the intercalated cell clusters, which ultimately reduces output from the central amygdala^{10, 33}. Our recording results suggest that the vHPC may relieve fear suppression and promote fear relapse via feed-forward inhibition of IL principal neurons. To examine this hypothesis, we used designer receptors exclusively activated by designer drugs (DREADDs) to selectively manipulate the activity of vHPC neurons projecting to the IL during presentation of an extinguished CS outside of the extinction context, a situation that leads to fear renewal.

We first confirmed that systemic administration of the DREADD agonist clozapine-*N*-oxide (CNO) decreases spontaneous firing in neurons expressing an inhibitory DREADD in vivo (Supplementary Fig. 5) and that silencing vHPC neurons using the same DREADD receptors attenuates the renewal of extinguished fear (Fig. 4). Specifically, CNO-treated rats expressing hM4D(G_i) receptors in the vHPC exhibited significantly less fear renewal than rats that received control virus without modulating baseline contextual

freezing (Fig. 4a–c). These data confirm the important role of the vHPC in fear renewal¹⁶.

We then specifically manipulated vHPC neurons projecting to IL using an intersectional viral approach. We injected canine adenovirus expressing Cre recombinase into the IL and a Cre-dependent DREADD virus or reporter control virus into the vHPC (Fig. 4d,e). Rats underwent fear conditioning, context exposure and extinction 4 weeks after surgery before receiving a 2-d within-subject renewal test in the conditioning context (that is, an ABA design in which each rat was tested after CNO or vehicle (VEH) administration in a counterbalanced order; Fig. 4a). Silencing vHPC→IL projections with CNO significantly reduced freezing to the extinguished CS relative to VEH treatment without affecting baseline freezing (Fig. 4g). This indicates that inhibiting vHPC neurons projecting to the IL attenuates fear renewal. Notably, CNO administration did not affect fear renewal in rats that expressed a control virus (Fig. 4f), indicating that the effects of CNO (whether directly or through its conversion to clozapine³⁴) were the result of a specific interaction with virally expressed DREADDs.

We next examined whether pharmacogenetically activating IL projectors in the vHPC would induce fear relapse in the extinction context. To this end, we expressed a Cre-dependent excitatory DREADD in vHPC neurons projecting to the IL (Fig. 4d,e). After fear conditioning and extinction, rats received a within-subject retrieval test in the extinction context (that is, an ABB design in which each rat received counterbalanced CNO or VEH administration; Fig. 4a). Activation of vHPC→IL projections increased freezing elicited by the CS in the extinction context (Fig. 4h) without affecting baseline freezing. These results indicate that vHPC→IL projections inhibit the expression of extinction, leading to a relapse of extinguished fear in the extinction context.

Fear relapse requires GABA receptors in the IL. Given that neuronal activity in the IL has been linked to fear suppression⁸, our results suggest that vHPC-mediated GABAergic feed-forward inhibition of IL projection neurons mediates fear relapse. To test this hypothesis,

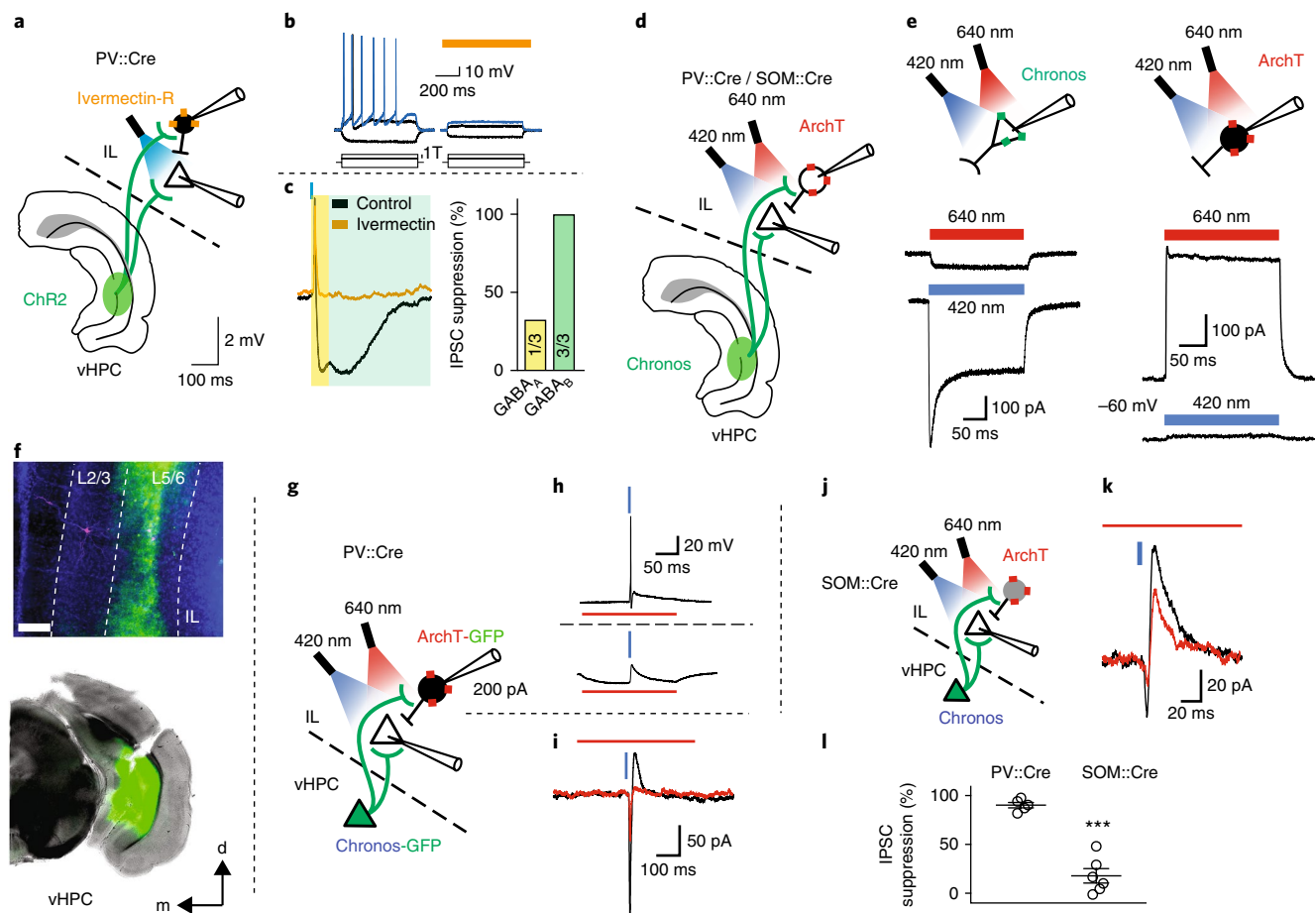


Fig. 3 | vHPC-driven feed-forward inhibition onto IL pyramidal neurons are specifically mediated by PV⁺ interneurons. **a**, Schematic illustrating expression of ivermectin-sensitive chloride channels (red) in IL PV⁺ cells to suppress local feed-forward inhibition mediated by the vHPC (using optical stimulation). **b**, Suppression of neural activity of PV⁺ neurons was confirmed by investigating intrinsic firing (left) that was inhibited in the presence of ivermectin (right; orange bar). **c**, Optical activation of vHPC inputs evoked a disynaptic inhibitory synaptic potential in a pyramidal cell (black trace), and application of ivermectin (30 nM) to silence PV interneurons blocked this inhibition (orange trace). The bar graph shows the suppression of the fast and slow inhibitory conductances (number of times the fast (yellow) and slow (green) IPSC were blocked/total number of investigated neurons). **d**, To achieve better temporal control, we used a dual-optical approach to recruit vHPC inputs using Chronos (a more light sensitive opsin) at 420 nm (blue bars) and to silence local IL PV⁺ neurons using ArchT at 640 nm (red bars). **e**, Left, Chronos-infected cells showed minimal activation using 640-nm light (200-ms pulses), but were strongly activated by 420-nm light. Right, in contrast, ArchT-infected PV⁺ neurons (bottom trace) showed a strong response to 640-nm light, whereas 420-nm light (bottom trace) did not activate the proton pump (ArchT) much. **f**, Examples of viral injection sites into the IL (ArchT) and vHPC (Chronos); white bar inset: 200 μ m. **g**, Illustration showing the dual-optical approach to recruit hippocampal inputs while silencing PV⁺ interneurons using PV::Cre animals. **h**, Chronos-driven spiking activity in ArchT-infected PV⁺ neurons (top trace) was suppressed in the presence of a 640-nm light pulse (200 ms long, $n = 3$). **i**, The vHPC-mediated feed-forward inhibition in pyramidal neurons (black traces; vHPC stimulation: blue bar) was blocked by silencing PV⁺ neurons using 640-nm light during the hippocampal stimulation (red bar; $n = 5$). **j**, Illustration showing the dual-optical approach using SOM::Cre animals (instead of PV::Cre). **k**, vHPC-mediated inhibition in IL pyramidal neurons (black trace) was not blocked in the presence of 640-nm light ($n = 6$). **l**, The bar graph summarizes how often (as a percentage of trials) the disynaptic inhibition in principal cells was blocked during the 640-nm light presentation in PV::Cre (left; $n = 5$) and SOM::Cre animals (right; $n = 6$; two-tailed t test: $t_9 = 8.47$; $P < 0.0001$).

we examined the role of GABAergic transmission in the IL in the expression of fear renewal. Intracranial micro-infusion of muscimol (a GABA_A receptor agonist) into the IL induced relapse of fear in the extinction context (Fig. 5a–c). Notably, IL inactivation did not affect freezing in the earliest trials of the retrieval test (Supplementary Fig. 6), which suggests spontaneous recovery of fear might mask the effects of IL inactivation when brief retrieval tests are used³⁵. Furthermore, intracranial micro-infusions of CGP55845 (a GABA_B receptor antagonist), picrotoxin (a GABA_{A/C} receptor antagonist) or both into the IL significantly reduced fear renewal as compared with vehicle treatment (Fig. 5d–f). These results indicate that GABAergic transmission in the IL regulates fear relapse.

We found that renewal of extinguished fear resulted from the activation of IL projecting vHPC neurons. Activation of these inputs *ex vivo* led to activation of PV⁺ interneurons and inhibition of amygdala-projecting IL pyramidal neurons. These data suggest that activation of PV⁺ interneurons, and the resultant feed-forward GABAergic inhibition of IL pyramidal cells, mediates inhibition in the vHPC→IL circuit in behaving rats. To test this hypothesis *in vivo*, we again expressed excitatory hM3D(G_q) virus in vHPC neurons projecting to the IL (Fig. 5g) and examined Fos expression in vHPC projectors and infralimbic cortical PV⁺ interneurons after CNO injection. Significantly higher percentages of hM3D(G_q)⁺ neurons in the vHPC expressed Fos after CNO injection compared

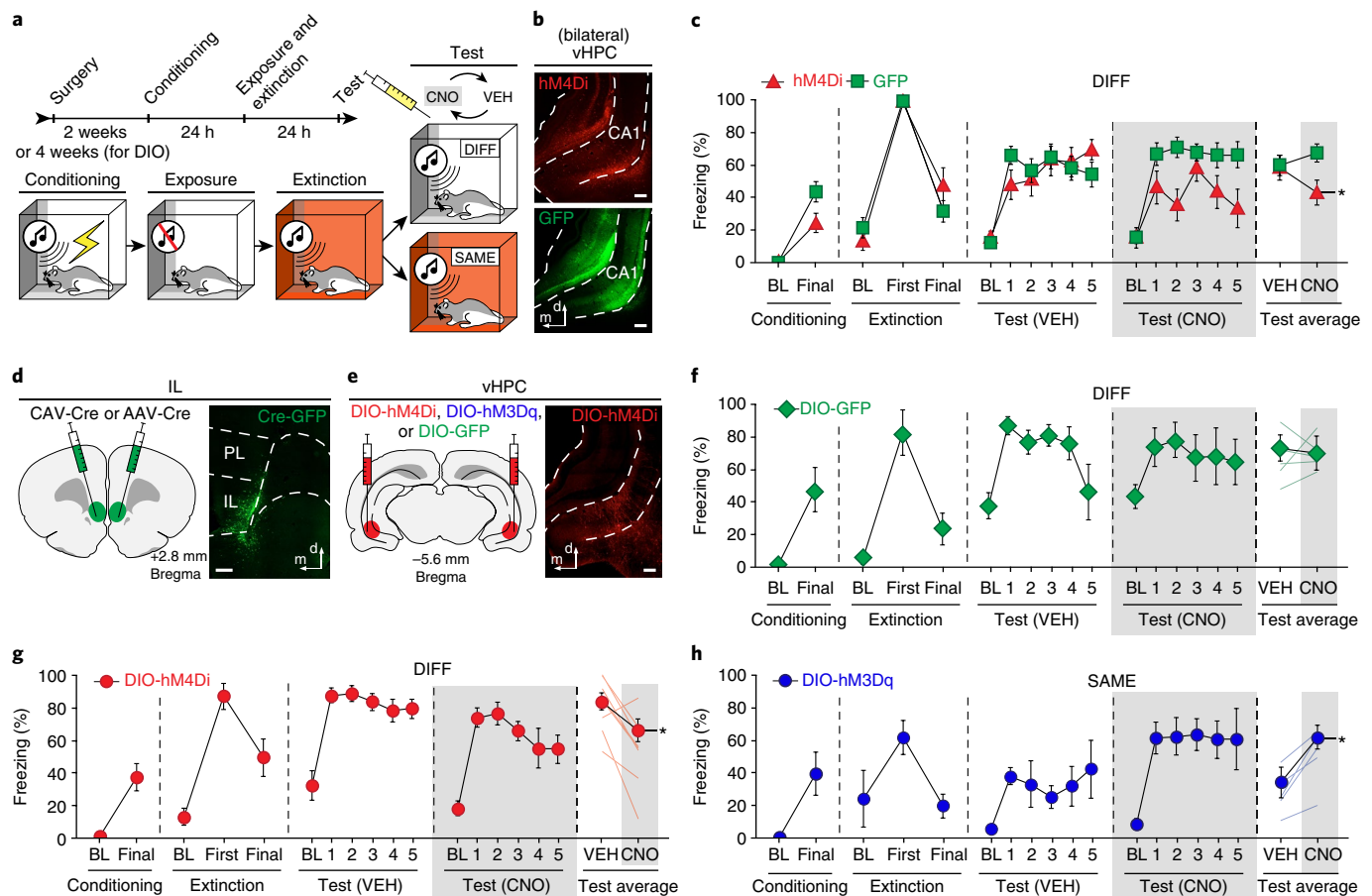


Fig. 4 | vHPC-IL projections bidirectionally modulate fear relapse. **a**, Behavioral design: within-subject ABA renewal (DIFF) or ABB retrieval (SAME) procedure. **b**, Non-Cre-dependent hM4Di(G_s)-expressing virus (top) or control virus (bottom) expression in the vHPC. **c**, Conditioning data revealed the mean percentage of freezing during the pre-conditioning baseline (BL) period and after the final conditioning trial (1 min post-shock). Extinction data revealed mean freezing during the pre-CS baseline period (3 min), as well as during the first and last five-trial blocks (intertrial intervals, ITIs). Test data showed the mean percentage of freezing during the 10-min baseline period and during the 30-s ITIs for each of the five test trials; each animal was tested after VEH or CNO administration in separate counterbalanced tests. The mean freezing across all ITIs is shown in the rightmost panel. Conditioning and extinction were normal (statistical data not shown). CNO- and virus-dependent silencing of the vHPC decreased fear renewal at test (red triangles), $n = 8$; control animals exhibited robust renewal (green squares), $n = 17$ (two-way repeated measures ANOVA, drug \times virus interaction: $F_{1,23} = 5.028$, $P = 0.0349$; two-tailed paired t tests for average freezing (split by virus): hM4Di: $t_7 = 2.783$, $P = 0.0272$; GFP: $t_{16} = -1.178$, $P = 0.2562$). **d**, CAV2-Cre or AAV5-Cre infused into the IL (left); infusion sites confirmed by co-infusion of AAV8-hSyn-GFP (right). **e**, Cre-dependent (DIO) viruses expressing hM4Di(G_s)-mCherry, hM3D(G_s)-mCherry or GFP infused into the vHPC (left); mCherry expression in vHPC→IL neurons in CA1 region (right). **f–h**, Effects of CNO on renewal or retrieval of extinction in rats expressing GFP (**f**), hM4Di (**g**) or hM3Dq (**h**); conditioning and extinction were typical in all cases (statistical data not shown). CNO had no effect on fear renewal in GFP control rats, $n = 5$ (two-tailed paired t test for average freezing, $t_4 = 0.982$, $P = 0.3818$; light lines depict individual animals; **f**). CNO silencing of vHPC→IL decreased fear renewal, $n = 8$ (test average: $t_7 = 2.634$, $P = 0.0337$); baseline freezing was not affected by CNO ($t_7 = 1.692$, $P = 0.1345$; **g**). CNO activation of vHPC→IL led to fear relapse, $n = 5$, (test average: $t_4 = -3.535$, $P = 0.0241$) without affecting baseline freezing ($t_4 = -1.377$, $P = 0.2405$; **h**). Error bars indicate means \pm s.e.m. * $P < 0.05$. Scale bars represent 250 μ m (**b,d,e**).

with the control (Fig. 5b). Notably, this was accompanied by a significant increase in the number of Fos-positive PV⁺ interneurons in the IL, but not PL (Fig. 5i). This suggests that pharmacogenetic activation of VH projectors to the IL recruits PV⁺ interneurons in intact animals, a population of neurons positioned to inhibit IL output and promote fear relapse.

Discussion

Collectively, our data reveal that vHPC projections generate a significant feed-forward inhibition of IL pyramidal neurons by recruiting PV⁺ interneurons. vHPC-mediated inhibition was most pronounced in amygdala-projecting pyramidal neurons, located in layers 2 and 3 of IL. This vHPC-mediated inhibition of IL principal cells had an important role in the relapse of extinguished fear.

Pharmacogenetic activation of IL projectors in the vHPC recruited PV⁺ interneurons in the IL and caused a relapse of extinguished fear; this effect was also observed with direct pharmacological activation of GABA receptors in IL. In contrast, either pharmacogenetic silencing of vHPC→IL projectors or antagonizing GABAergic receptors in the IL diminished fear relapse. Together, these findings reveal that vHPC projections to the mPFC drive feed-forward inhibition of IL principal neurons by PV⁺ interneurons to mediate fear relapse after extinction.

Our findings are consistent with early electrophysiological analyses of vHPC-mPFC projections that have shown a strong inhibitory effect of electrical stimulation of the vHPC on mPFC neuronal activity^{36,37}. Indeed, an inhibitory influence of the vHPC on mPFC function has been reported to influence the expression of learned

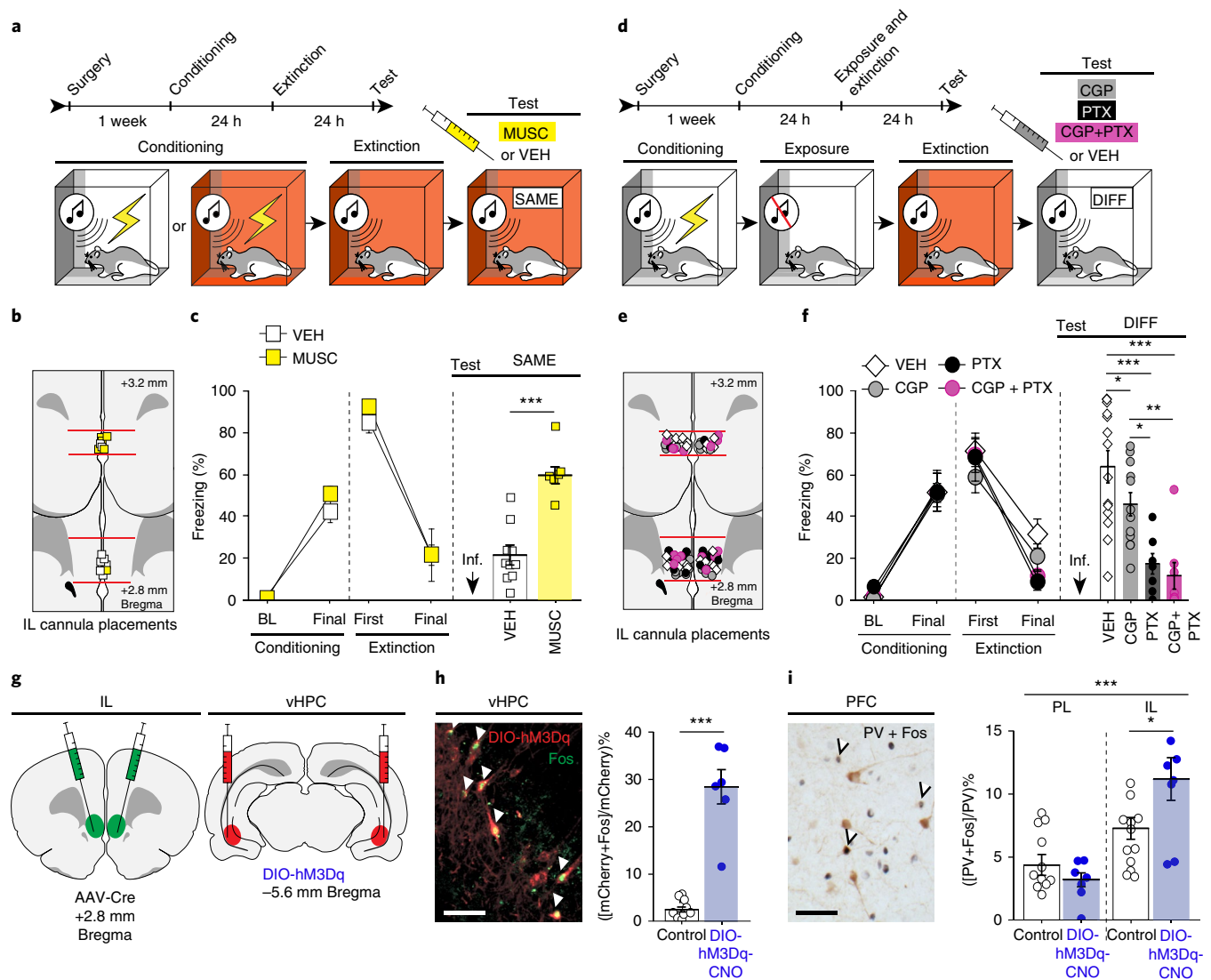


Fig. 5 | Local GABA-mediated signaling in the IL gates fear renewal. **a**, Behavioral design: ABB or BBB retrieval procedure. **b**, Single cannula placements aimed at IL (11°). **c**, Percentage of freezing during the 3-min BL and after the final trial (final, 1-min ITI) during conditioning, during the first and final five-trial blocks (average of five post-CS ITIs) during extinction, and average freezing across 20 CS-alone trials (post-CS ITIs) on the retention test. Conditioning and extinction were typical, with no differences between groups (statistical data not shown). Intra-IL infusions of muscimol (MUSC, $n = 6$) induced relapse of fear in the extinction context compared with vehicle-treated rats (VEH, $n = 10$; $t_{14} = 5.9$, $P < 0.0001$). **d**, ABA renewal procedure. **e**, Bilateral IL cannula placements (30°). **f**, Percentage of freezing during the 3-min BL and after the final trial (final, 1-min ITI) during conditioning, during the first and final five-trial blocks (average of five post-CS ITIs) during extinction, and average freezing across five CS-alone trials (post-CS ITIs) on the retention test. Conditioning and extinction were typical with no differences between groups (statistical data not shown). Intra-IL infusions of CGP55845 (CGP, $n = 12$), picrotoxin (PTX, $n = 7$) or both (CGP+PTX, $n = 8$; VEH, $n = 14$) decreased fear renewal (one-way factorial ANOVA, main effect of drug, $F_{3,37} = 12.470$, $P < 0.0001$; post hoc Fisher's PLSD, $P = 0.0439$, $P < 0.0001$, $P < 0.0001$ for CGP, PTX, and CGP+PTX versus VEH comparisons, respectively; $P = 0.0105$ and $P = 0.0018$ for PTX and CGP+PTX versus CGP comparisons, respectively). **g**, Infusions of AAV5-Cre in the IL (left) and AAV8-hSyn-DIO-hM3D(G_q)-mCherry (or non-DREADD-expressing control virus) in the vHPC (right). **h**, Left, hM3D(G_q)⁺ neurons (mCherry: red) and Fos⁺ nuclei (GFP: green) in vHPC (arrows = double-labeled cells). Scale bar represents 50 μm. Right, percentages of Fos⁺ and mCherry⁺ cells among total mCherry⁺ neurons, comparing controls ($n = 11$: DREADD-expressing rats treated with vehicle ($n = 6$), non-DREADD-expressing rats treated with CNO ($n = 5$) and DIO-hM3Dq-CNO animals ($n = 6$) (two-tailed unpaired t test, $t_{15} = 9.043$, $P < 0.0001$). **i**, Left, PV⁺ neurons (brown in soma) and Fos⁺ neurons (purple/black in nuclei) in PFC (arrows = double-labeled cells). Scale bar represents 50 μm. Right, percentages of PV⁺ and Fos⁺ neurons of total PV⁺ neurons, comparing CNO-treated hM3D(G_q)⁺-expressing animals ($n = 7$) and controls ($n = 12$: hM3D(G_q)⁺-expressing animals rats treated with vehicle ($n = 6$), mCherry-only-expressing rats treated with CNO ($n = 6$); main effect of brain region: $F_{1,11} = 30.5$, $P < 0.0001$; brain region \times percentage interaction: $F_{1,17} = 8.076$, $P = 0.0113$; post hoc Fisher's PLSD, IL: $P = 0.0288$ for hM3D(G_q)⁺-CNO versus controls). Error bars indicate means \pm s.e.m. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.0001$.

fear³⁸. However, in contrast with this work, a recent report suggested that the dominant effect of the vHPC in the mPFC is excitatory³⁹. Specifically, it was reported that optogenetic inhibition of vHPC terminals in the mPFC reduced short-latency (10–20 ms) vHPC-

evoked increases in mPFC multiunit activity in anesthetized mice. We also observed short-latency EPSPs in IL pyramidal neurons, but this excitation was overshadowed by a dominant feed-forward inhibition that lasted hundreds of milliseconds beyond the excitatory

valley (Fig. 1k). Our ex vivo experiments revealed that this vHPC-mediated inhibition was mediated by both GABA_A- and GABA_B-receptor-mediated currents, and inhibition of either receptor type had an effect on fear renewal (Fig. 5). Thus, although short-latency excitatory responses in IL principal cells can be driven by vHPC input, the net effect of feed-forward inhibition on amygdala-projecting neurons is likely to be inhibition over a much longer timescale^{36,37}.

Of course, it is noteworthy that the magnitude of the renewal impairments obtained with pharmacogenetically silencing either the vHPC (Fig. 4c) or vHPC projectors to IL (Fig. 4g) was relatively modest compared with that associated with GABA antagonists infused directly into the IL (Fig. 5f); DREADD-mediated manipulations reduced, rather than eliminated renewal. This disparity might be explained by differences in the inhibitory circuits being influenced by the two manipulations. For example, intracranial drug infusions affect both local inhibitory circuits and extrinsic feed-forward inhibition in the IL that includes, but is not limited to, that mediated by vHPC projections. Moreover, intersectional DREADD expression only targets a subset of vHPC neurons projecting to the IL; inhibition of this discrete population of neurons would not necessarily be expected to recapitulate the behavioral impairments associated with a broad manipulation of IL activity. Conversely, pharmacogenetic activation of vHPC projectors to IL produced significant increases in freezing (Fig. 4h) that were absent during the corresponding block of test trials (that is, trials 1–5) in rats receiving intra-IL muscimol infusions (Supplementary Fig. 6). In this case, the circuit-specific DREADD manipulation appeared to be more effective than the nonselective pharmacological manipulation in revealing a behavioral effect. In both cases, however, pharmacogenetic manipulations of vHPC projectors to IL and GABAergic manipulations of IL produced qualitatively similar outcomes, suggesting that they operate on similar neural processes. Nonetheless, further experiments that directly manipulate vHPC projections onto discrete IL interneuron populations are required to determine the specific contribution of those projections to the regulation of IL function and fear expression.

Our data complement a large body of evidence in both rats and humans implicating the mPFC in the regulation of emotional responses^{2,40,41}. The IL, in particular, is thought to dampen learned fear responses by inhibiting the amygdala^{42–44}. Indeed, prefrontal-amygdala circuits have been posited to be important for both the encoding and retrieval of extinction memories¹⁰. However, it has recently been reported that optogenetically silencing the IL³⁵ or its terminals in the amygdala⁴⁵ impairs the acquisition of extinction, but does not affect the retrieval of extinction memories³⁵. This contrasts with our results, which reveal robust effects of IL manipulations on extinction retrieval and renewal. One possibility is that these recent optogenetic studies only targeted IL principal cells (or their projections)^{35,45}. Consistent with this, another study found that optogenetic inhibition of both principal cells and inhibitory interneurons in the IL produced extinction retrieval deficits; inhibition of principal cells alone did not produce a deficit⁴⁶. This suggests that both inhibitory interneurons and principal cells are recruited in the regulation of extinction retrieval, which is consistent with our observations; future studies should nevertheless attempt to directly modulate the activity of interneurons of the IL during extinction retrieval and relapse.

Taken together, our results reveal a previously unknown hippocampal-prefrontal circuit for the context-dependent regulation of memory retrieval. Specifically, we found that the renewal of extinguished fear resulted from vHPC-mediated inhibition of mPFC circuits involved in suppressing fear. The engagement of this vHPC-IL circuit occurred when animals experienced an extinguished CS in a context in which the meaning of the CS was ambiguous (such as the original conditioning context or a novel context). We and others

have shown that activity in the hippocampus is increased by unexpected events, including experiencing an extinguished CS outside the extinction context^{13,18,22,47,48}. In these instances, it is likely that the hippocampus retrieves information about the context in which stimuli have previously occurred to influence prefrontal cortical processes involved in emotional regulation¹⁴. Of course, it has also been appreciated that prefrontal cortical circuits are involved in directing memory retrieval by the hippocampus based on contextual information⁴⁹. Ultimately, oscillatory networks that regulate bidirectional information flow through hippocampal-prefrontal circuits may generate context-appropriate behavioral responses⁵⁰.

In conclusion, we uncovered a previously unknown circuit for fear relapse that is centered on vHPC projections to the infralimbic cortex. We found that afferent activity in IL-projecting cells of the vHPC was associated with activation of GABAergic PV⁺ interneurons and strong inhibition of IL projection neurons. We suggest that the function of this circuit is to promote the return of fear under conditions in which it is warranted, including in contexts in which an extinguished CS is potentially dangerous (that is, the conditioning context or a novel context). The renewal of fear in these situations is adaptive in many cases, but can interfere with the efficacy of cognitive-behavioral therapies designed to limit fear after trauma. We conclude that vHPC synapses on IL interneurons are a unique target for therapeutic interventions aimed at broadening the generalization of extinction memories, thereby reducing the possibility that pathological fear relapses after therapy.

Methods

Methods, including statements of data availability and any associated accession codes and references, are available at <https://doi.org/10.1038/s41593-018-0073-9>.

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Author contributions

S.M. and P.S. supervised all of the experiments. S.M., P.S. and R.M. designed the experiments. R.M., J.J., T.D.G., T.F.G., Q.W., G.M.A. and P.J.F. collected the data. R.M., J.J., T.D.G., T.F.G., Q.W., G.M.A., P.J.F., S.M. and P.S. analyzed the data. R.H. and J.E.P. generated and provided AAVDJ/8 viral vectors. T.L. and J.W.L. generated and provided the ivermectin construct. R.M., J.J., T.D.G., S.M. and P.S. wrote the manuscript. All of the authors read and edited the manuscript.

Competing interests

The authors declare no competing financial interests.

Additional information

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Methods

Subjects. Adult male Wistar rats (170–240 g), PV::Cre knock-in (C57BL/6 strain) mice (for the local optical PV stimulation, the ivermectin-receptor silencing experiments, and the dual-optical approach; 20–26 g), and SOM::Cre knock-in (C57BL/6 strain) mice (19–25 g; all animals were bred in house) were used in the ex vivo electrophysiological experiments. The animals were housed in a 12/12-h light/dark cycle with access to food and water ad libitum. Adult male Long-Evans rats (200–224 g; Blue-Spruce, Harlan) were used for the behavioral experiments. The Long-Evans rats were individually housed in a 14/10-h light/dark cycle and had access to food and water ad libitum. Rats for behavioral testing were handled for 5 d after arrival. All experimental procedures were performed in accordance with the protocols approved by the University of Queensland Animal Ethics Committee and the Animal Care and Use Committee at Texas A&M University.

Viruses and drugs. AAV5-CamKII α -hM4D(G_i)-mCherry (3.4×10^{-12} vg/ml), AAV5-CamKII α -mCherry (6.4×10^{-12} vg/ml), AAV5-hSyn-DIO-hM4D(G_i)-mCherry (5.7×10^{-12} vg/ml), AAV8-hSyn-DIO-GFP ($\sim 6 \times 10^{-12}$ vg/ml), AAV5-hSyn-DIO-hM3D(G_q)-mCherry (6.7×10^{-12} vg/ml), and AAV8-CamKII α -GFP (5.6×10^{-12} vg/ml) vectors were obtained from the University of North Carolina Vector Core and Addgene. CAV2-Cre (8.7×10^{-12} vg/ml) vectors were obtained from the Institute of Molecular Genetics of Montpellier. AAV5-CMV-HI-eGFP-Cre-WPRE-SV40 (1.62×10^{-13} vg/ml), AAV2/5-CAG-ChR2-Venus-WSV40 (1.1×10^{-13} vg/ml), AAV2/5-hSyn-hChR2(H134R)-eYFP-WPRE-hGH (1.42×10^{-13} vg/ml), AAV2/1-pSyn-Chronos-EGFP (2.41×10^{-12} vg/ml), and AAV5-EF1a-dflox-hChR2(H134R)-mCherry-WPRE (1.31×10^{-13} vg/ml) vectors were obtained from University of Pennsylvania Vector Core. EF1a-D20-ArchT-EGFP (1.3×10^{-11} vg/ml) was produced in-house at the University of Queensland. R.H. and J.E.P. packaged and provided AAVDJ/8-CamKII α -hM4D(G_i)-mCherry (1.23×10^{-13} vg/ml) vectors. For use in the experiments involving designer receptors exclusively activated by designer drugs (DREADDs)³⁴, CNO was obtained from the National Institute of Mental Health (NIMH; Chemical synthesis and drug supply program). CNO was first dissolved in DMSO and then mixed with sterile saline (VEH, 2.5% DMSO). The action of CNO has been suggested to be mediated by its metabolite, clozapine³⁴. To control for this, CNO was also tested in a non-DREADD-expressing GFP control (Fig. 4). Drugs used in the electrophysiological experiments were obtained from Tocris Bioscience. For the behavioral experiments, muscimol (MUS) and CGP55845 (CGP) were obtained from Sigma-Aldrich, and picrotoxin (PTX) was obtained from Tocris Bioscience.

Surgical procedures. For the ex vivo electrophysiological experiments, mice (P25–45) were anesthetized with intraperitoneal injections of ketamine (10%) and xylazine (5%, 10 ml/kg; rats received an additional injection of 0.4 ml/kg of Zoletil). Animals were then placed in a stereotaxic frame. An incision at the midline was made using a single-edged blade. For the investigation of hippocampal projections to the mPFC, viruses (0.3–0.5 μ l of AAV2/5-CAG-ChR2-Venus-WSV40 or AAV2/5-hSyn-hChR2(H134R)-eYFP-WPRE-hGH) were injected in the vHPC of rats. The coordinates (all in mm) used for vHPC stereotaxic injections in the rats are (anterior/posterior, medial/lateral, dorsal/ventral): -6.0 , ± 5.3 , -4.0 . For the investigation of amygdala-projecting IL neurons in rats, retrograde beads (0.3–0.5 μ l; Lumafluor) were injected in the BLA (A/P, M/L [angle], D/V): -2.3 , ± 1.2 (22°), -9.3 . For the study of local PV-projections, 1–2 μ l of AAV5-EF1a-dflox-hChR2(H134R)-mCherry-WPRE were injected into the IL of PV::Cre mice (for the silencing approach using ivermectin-receptors^{32,33}). Coordinates for the IL injections are (A/P, M/L [angle], D/V): 2.5 , ± 2.9 (34°), -4.5 . For the dual-optogenetic approach using Chronos⁵⁴ and ArchT³⁵, AAV2/1-pSyn-Chronos-EGFP (vHPC) and EF1a-D20-ArchT-EGFP (IL) were injected using the following coordinates in either PV::Cre or SOM::Cre mice: (A/P, M/L, D/V) vHPC: -3.3 , ± 3.1 , -3.5 ; IL: 1.8 , ± 0.4 , 2.0 (the injection was targeted slightly more dorsal than usual due to issues with strong expression at the core infection site). The incision was then disinfected and closed using vetbond tissue adhesive and stitched. Baytril (0.1 μ l/g) and Metacam (0.4 μ l/g) were each diluted into 0.5 ml of saline and then injected subcutaneously.

For all behavioral experiments, all surgery and group assignments were randomized for cage position in the vivarium. In the experiment examining the role of DREADD-mediated inactivation of the vHPC during fear renewal (Fig. 4a–c), rats were deeply anesthetized with isoflurane (5% for induction; $\sim 2\%$ during surgery), and were placed into a stereotaxic apparatus for viral injection. The scalp was shaved, treated with povidone-iodine, and a small incision was made to expose the skull. Small holes were drilled in the skull and rats were bilaterally injected with either AAVDJ/8-CamKII α -hM4D(G_i)-mCherry ($n = 14$) or AAV8-CamKII α -GFP ($n = 18$) into the vHPC. Four bilateral injections (0.4 μ l/injection) were made into the vHPC at two different A/P levels (coordinates [in mm from bregma] are shown in A/P, M/L, D/V): -5.2 , ± 6.0 , -6.5 ; -5.2 , ± 6.0 , -5.3 ; -6.1 , ± 6.0 , -6.1 ; -6.1 , ± 6.0 , -5.0 (all D/V coordinates are measured from dura). Viruses were injected (0.15 μ l/min) using an injector connected to polyethylene tube and a Hamilton syringe (10 μ l) mounted on an infusion pump. Following the infusions, virus was allowed 5–10 min for diffusion before removing the injectors. Once completed, the incision was closed and rats were returned to their homecages to recover for 2 weeks following surgery. Rats with unilateral, off-target, or no

viral expression were excluded from the final analyses. In addition, two rats were excluded because of a technical issue during tissue collection resulting in a loss of vHPC tissue. Based on these criteria, twenty-five rats are included in the final analyses (Fig. 4a–c; hM4Di, $n = 8$; GFP, $n = 17$).

For the intersectional DREADD experiments (Fig. 4d–h), rats underwent surgery as described above. To pharmacogenetically inhibit vHPC neurons projecting to IL, rats ($n = 14$) were bilaterally injected with CAV2-Cre in the IL and AAV5-hSyn-DIO-hM4D(G_i)-mCherry in the vHPC. Pharmacogenetic excitation of this projection was achieved by bilaterally injecting rats ($n = 8$) with AAV5-CMV-HI-eGFP-Cre-WPRE-SV40 in the IL and AAV5-hSyn-DIO-hM3D(G_q)-mCherry in the vHPC. Control rats ($n = 5$) received CAV2-Cre in the IL and AAV8-hSyn-DIO-GFP in the vHPC. All the viruses for this experiment were injected (0.1 μ l/min) as described above. Four injections (0.5 μ l/injection) were made into the vHPC at the coordinates described above. One injection (~ 1.5 – 1.8 μ l) was made in the IL in each hemisphere at the following coordinates (mm from bregma): $+2.8$ (A/P), ± 3.0 (M/L), -4.9 (D/V), with a 30° angle. The viruses were allowed to diffuse for 5–10 min per injection before removing the injectors. Once completed, rats were placed back in their home cages to allow for viral expression for at least 4 weeks. Rats with unilateral, off-target, or no viral expression were excluded from the analyses. Specifically, six rats infected with AAV5-hSyn-DIO-hM4D(G_i)-mCherry and three rats infected with AAV5-hSyn-DIO-hM3D(G_q)-mCherry were excluded based on the aforementioned criteria, resulting in the final group sizes (Fig. 4d–h): DIO-GFP ($n = 5$), DIO-hM4Di ($n = 8$), DIO-hM3Dq ($n = 5$).

For the behavioral experiment involving pharmacological inactivation of the IL with a GABA receptor agonist (Fig. 5a–c and Supplementary Fig. 6), rats ($n = 32$; 16 rats per group: vehicle [VEH] and muscimol [MUS]) were implanted with a single guide cannula (26 gauge, 8 mm; Small Parts) in the IL (A/P: $+2.65$, M/L: ± 1.0 , D/V: -4.1) at an 11° angle. Cannulas were secured to the skull with jeweler's screws and dental cement. Stainless steel obturators (30 gauge, 9 mm; Small Parts) were placed in each guide cannula and were changed twice before behavioral tests. Rats recovered from surgery for one week before the onset of behavioral training and testing. Rats with cannulas terminating outside the borders of the IL were excluded from the final analyses. Based on these criteria, sixteen rats are included in the final analyses (Fig. 5a–c and Supplementary Fig. 6; VEH, $n = 10$, MUS; $n = 6$).

For the behavioral experiment consisting of pharmacological activation of the IL with GABA receptor antagonists (Fig. 5d–f), rats ($n = 56$) were implanted with bilateral guide cannulas (26 gauge, 8 mm) in the IL ([mm from bregma] A/P: $+2.7$, M/L: 3.0 , D/V: -4.9 at 30° angle). The guide cannulas were secured as above with jeweler's screws and dental cement. Stainless steel obturators (30 gauge, 9 mm) were inserted into the cannulas and removed and replaced with clean obturators twice before the behavioral tests. Rats recovered for one week following surgery in their homecages before the behavioral procedures. Rats with off-target cannula placements (including unilateral misses) were excluded from all of the analyses (ten vehicle [VEH] rats, four CGP55845 [CGP] rats, and one picrotoxin [PTX] rat were excluded). Accordingly, forty-one rats are included in the final analyses (Fig. 5d–f; VEH, $n = 14$; CGP, $n = 12$; PTX, $n = 7$; CGP+PTX, $n = 8$).

For those animals used in the Fos analyses (Fig. 5g–i), rats were bilaterally injected with AAV5-CMV-HI-eGFP-Cre-WPRE-SV40 in IL and AAV5-hSyn-DIO-hM3D(G_q)-mCherry in vHPC ($n = 16$; eight rats per group: VEH, CNO). As an additional control, another group of rats was bilaterally injected with AAV5-CMV-HI-eGFP-Cre-WPRE-SV40 in the IL and AAV5-hSyn-DIO-mCherry in the vHPC ($n = 8$, CNO). Injection procedures and locations were identical to those in the DIO-DREADD experiment. As with the abovementioned experiments, rats with off-target viral infection were excluded from the final analyses. In addition, five rats were excluded because of a technical issue in collecting tissue at the level of the PFC resulting in poorly stained tissue. A total of 19 rats were included in the final analyses for Fos analyses of the PFC (Fig. 5i; DIO-hM3Dq-CNO, $n = 7$; DIO-hM3Dq-VEH, $n = 6$; DIO-mCherry-CNO, $n = 6$; the latter two groups were collapsed for a total of 12 control animals). For the Fos analyses of the vHPC (Fig. 5h), another two rats were excluded due to a technical issue in collecting vHPC tissue. Accordingly, a total of 17 rats were included in the vHPC Fos analyses (Fig. 5h; DIO-hM3Dq-CNO, $n = 6$; controls, $n = 11$: DIO-hM3Dq-VEH, $n = 5$; DIO-mCherry-CNO, $n = 6$).

Finally, for the in vivo electrophysiological recordings anesthetized rats (three total) received bilateral stereotaxic infusions (2.0 μ l/per infusion; 0.3 μ l/min with 5 min of diffusion time) of AAV5-CamKII α -hM4D(G_i)-mCherry ($n = 1$) or AAV5-CamKII α -mCherry ($n = 2$) in IL (A/P: $+2.7$, M/L: ± 3.0 , D/V: -5.1 at a 30° angle). Following viral infusions, rats were implanted with a 16-channel microelectrode array (Innovative Neurophysiology, Durham, NC) within the right hemisphere targeting the mPFC (8 channels in IL; 8 channels in PL). The 2×8 array was comprised of two rows of 50- μ m diameter tungsten wires terminating at different lengths (6.9 mm for targeting PL, and 8.0 for targeting IL). From center-to-center, the wires were spaced 200 μ m apart. The array was implanted such that its long axis was parallel to the A/P plane of the brain, and that its centermost wires targeted IL as follows: A/P: $+2.7$, M/L: $+0.35$, D/V: -5.1 (from skull surface). Dental cement was used to secure the array to the skull. Animals recovered from surgeries for 2 weeks before the onset of recordings.

Slice electrophysiology. Similar to prior reports⁵⁶, electrophysiological experiments commenced at least 28 d after the virus injections to allow for terminal expression. Single-cell recordings of neurons and application of brief light pulses (470 nm; 5 ms) using an LED source (CoolLED) were performed to study the location and pharmacological properties of synaptic connections onto mPFC neurons. Only injected animals with at least one successful synaptic response at the projection site were included in the analyses. Light stimulation was kept maximal, except for cases where a decrease in light stimulation eliminated the polysynaptic component, but didn't diminish the initial component.

Animals were anesthetized with isoflurane (1 ml; applied in enclosed container) and then decapitated. Brains were rapidly removed and placed in ice cold cutting solution containing the following chemicals (mM): NaCl 118, KCl 2.5, NaHCO₃ 25, glucose 10, MgCl₂ 4, CaCl₂ 0.5 and NaH₂PO₄ 1.2. Coronal brain slices (300 µm) were prepared using a vibratome (Leica VT 1000 S). Slices were allowed to recover in oxygenated (95% O₂ and 5% CO₂) artificial cerebrospinal fluid (aCSF) containing the following chemicals (mM): NaCl 118, KCl 2.5, NaHCO₃ 25, glucose 10, MgCl₂ 1.3, CaCl₂ 2.5 and NaH₂PO₄ 1.2 at 35 °C for at least 30 min, then kept at room temperature (22–23 °C) for at least another 30 min before experiments. Slices were transferred to the recording chamber as required and were continuously perfused with oxygenated aCSF through a gravity fed system and maintained at 30–32 °C. In cases where either a virus or a tracer was injected, slices containing the injection sites were also kept to determine localized injection sites. Slices were visualized using an upright microscope (BX50WI, Olympus Optical, Tokyo, Japan) with a 5× NA 0.1 or 40× NA 0.8 objective and infrared and differential interference contrast optics. Fluorescent neurons were visualized by using an LED system (pE-2, CoolLED) and YFP/RFP filter sets (Olympus). Electrodes (3–7 MΩ) were filled with pipette solution containing chemicals (mM): KMeSO₄ 135, NaCl 7, HEPES 10, Mg₂ATP 2, Na₂GTP 0.3, EGTA 0.3, biocytin 8 (pH 7.3 with KOH; osmolality ~290 mOsm/kg). Signals were recorded using a patch clamp amplifier (Multiclamp 700B, Axon instruments). Sampling rate was 20 kHz and signals were digitized at 2 kHz (Instrutech, ITC-18). All data were acquired, stored and analyzed using Axograph X (Axograph, V 1.2.1). For all the voltage-clamp recordings, neurons were held at –60 to –70 mV by injecting current if needed, and neurons with resting membrane potential above –55 mV were excluded from the analysis. For current-clamp recordings, recordings were corrected for bridge-balance and pipette resistance. All investigations were performed on the ipsilateral site of injection. Access resistance was 5–25 MΩ and was monitored throughout the experiment (neurons with access resistance changes >25% were excluded). Drugs were bath applied by using a gravity-fed system that allowed continuous change of solutions. For optogenetic stimulation, 470-nm light pulses were applied with a CoolLED system (pE-2) attached to the upright microscope (Olympus BX51WI). 5-ms pulses were applied to evoke postsynaptic responses and 200-ms pulses were used to study infected neurons. Maximal light output at 470 nm was measured at 15 mW/mm² (ThorLabs, optical power meter). Response latency was calculated by measuring the delay between the onset of the light pulse and the onset of the EPSC (5%). The response jitter was calculated by measuring the s.d. of 8–15 individual responses to optical stimulation. Only responses >50 pA were used for the analysis. For the dual-optical approach, a fully reflective mirror (Chroma) was selected for the stimulation, and band-pass filters (for 400-nm light, the light was selected from the 400/470-nm LED array module [LAM]; for 640 nm, a bandpass filter of 641/75 nm was used in between the light source [GYR CoolLED LAM] and the microscope). 640-nm light was presented every second time during the 400 nm light pulse presentation. Electrophysiological responses were analyzed using AxoGraph. For synaptic responses, an average of 5–10 traces were taken for analysis. For optogenetic stimulation, only samples with at least one synaptic response to maximal intensity light pulses were used for the statistics. Electrical stimulation (at 0.1 ms using a stimulator-box) was done by using a patch-clamp pipette filled with 3 M saline and the ground loop wire was placed in the bath. Intrinsic firing properties were analyzed at current injections of twofold threshold firing. Drug application of Picrotoxin (100 µM), CGP55845 (1 mM), NBQX (20 mM) and APV (100 mM) (all from Tocris) were done through a gravity fed system. For the comparison between PL and IL and pyramidal neurons and interneurons, respectively, EPSCs were compared in animals with recordings from both regions and subtypes, respectively. Analytical tests were performed with SPSS (IBM) or Prism (GraphPad). Outliers (if applicable) were determined by Grubb's test (<http://graphpad.com/quickcalcs/Grubbs1.cfm>).

In vivo electrophysiology. 2 weeks after surgery, freely moving rats underwent three 70-min single-unit recording sessions across three consecutive days (one session/day) in a standard testing chamber. Following a 10-min baseline period, rats were injected (i.p.) with CNO (1 or 3 mg/kg) or vehicle (counterbalanced) and remained in the chamber for 60 min. Extracellular single-unit activity was recorded automatically using a multichannel neurophysiological recording system (OmniPlex, Plexon) as previously described³⁷. One of the recording wires (located in PL) served as a reference for the wideband signal recorded on each channel. Signals were amplified (8,000×) and digitized (40-kHz sampling rate) and saved on a computer for offline sorting and analysis. After high-pass filtering the signal at 600 Hz, waveforms were manually sorted using two-dimensional principal component analysis (Offline Sorter, Plexon). Isolated waveforms and their

respective timestamps were then imported to NeuroExplorer (Nex Technologies) for further analysis. The analyses of neural activity focused on changes in spontaneous single-unit firing rate during each recording session. For each session, each neuron was z-score normalized to its 10-min baseline firing rate. Data are plotted in 20-s bins across the duration of the session.

Behavioral apparatus. Behavioral testing was conducted in two distinct rooms in the laboratory. Eight identical conditioning chambers (30 × 24 × 21 cm; MED Associates) in each room were used in all behavioral experiments. The chambers consisted of aluminum sidewalls and Plexiglas ceilings, rear walls, and hinged front doors. The chamber floors consisted of 19 stainless steel rods that were wired to a shock source and a solid-state grid scrambler (MED Associates) for delivery of footshock (US). A speaker mounted on one wall of the chamber was used for delivery of the acoustic CS, and ventilation fans and house lights were installed in each chamber to allow for the manipulation of contexts. Each chamber rests on a load-cell platform that is used to record chamber displacement in response to each rat's motor activity and is acquired online via Threshold Activity software (MED Associates). Absolute values of the load-cell voltages are computed and multiplied by 10 to yield a scale that ranges from 0 to 100. For each chamber, load-cell voltages are digitized at 5 Hz, yielding one observation every 200 ms. Freezing is quantified by computing the number of observations for each rat that has a value less than the freezing threshold (load-cell activity = 10). Freezing is only scored if the rat is immobile for at least 1 s. Because all behavioral measurements of freezing were performed using this automated system, no blinding to group assignments was necessary.

Sensory stimuli were adjusted within these chambers to generate distinct contexts (A and B). For context A, a 15-W house light mounted on the sidewall was turned on, and the white room light remained on. Ventilation fans (65 dB) were turned on, cabinet doors were left open, and the chambers were cleaned with 1% ammonium hydroxide. Rats were transported to context A in white plastic boxes. For context B, house light and white room light were all turned off, and fluorescent red room light was turned on. Ventilation fans were turned off, the cabinet doors were closed and the chambers were cleaned with 1% acetic acid. Rats were transported to context B in black plastic boxes. In each context, stainless steel pans were filled with a thin layer of the respective odors of the contexts and inserted below the grid floor. Context B in the muscimol experiment consisted of 70% ethanol odor in place of acetic acid, cupboard doors remained open, black plastic floors were placed on the grid floor, and rats were transported in 5-gallon buckets with bedding (all other features matched context B as described above).

Behavioral procedures. We used standard procedures for Pavlovian fear conditioning, extinction, and retrieval testing^{6,38}. Rats were randomly assigned to each experimental group, and each training or testing squad was counterbalanced to equally represent each group in the squad. In addition, rats were randomly assigned to the testing chambers so as to counterbalance the placement of group assignments in each context and squad. Freezing behavior (as a percentage of a block of time or trial[s]) served as the dependent measure of fear in all training and test sessions. For the experiment involving pharmacogenetic inactivation of the vHPC during the renewal of fear (Fig. 4a–c), animals first underwent fear conditioning consisting of five tone (CS; 10 s, 80 dB, 2 kHz) footshock (US; 2 s, 1.0 mA) pairings with 60-s ITIs after a 3-min baseline period in the context (context A). Rats remained in the chamber for 60 s following the final CS-US pairing, and were then returned to their homecages. On days 2–5, rats underwent extinction training in which they received context exposure to context A for 35.5 min in the morning, and 45 CS-alone trials (30-s ITIs; a total of 35.5 min) in context B in the afternoon (3-min baseline). 24 h after the final extinction session, rats received a within-subject renewal test in context A over two consecutive days; VEH or CNO (3 mg/kg) was administered 30 min before each test in a counterbalanced order. Each test session consisted of a 10-min baseline followed by five CS-alone (30-s ITIs) presentations; the long baseline period was included to assess any nonspecific effects of drug treatment on conditional freezing.

For the intersectional DREADD experiments, rats received fear conditioning followed by extinction as described above (only 2 d of extinction were administered in this experiment). Rats in the DIO-hM4Di and DIO-GFP groups received within-subject renewal tests as described above in context A; whereas DIO-hM3Dq rats received within-subject extinction retrieval tests in context B; thirty minutes before each test, rats received systemic injections of CNO (1–3 mg/kg) or VEH in a counterbalanced order. Each test session consisted of a 10-min baseline followed by five CS-alone (30-s ITIs) presentations.

For the behavioral experiment involving pharmacological blockade of the IL during the retrieval of extinction (Fig. 5a–c and Supplementary Fig. 6), animals underwent fear conditioning (identical to that described above) in either context A or context B (rats conditioned in the two contexts did not differ and were ultimately collapsed). 24 h after conditioning, rats were placed in context B and underwent a single session of extinction training (identical to that described above). 24 h after extinction, the rats underwent a retrieval test consisting of a second extinction session in context B immediately after an intra-IL infusion (0.2 µl; 0.1 µl/min for 2 min) of either SAL or MUS (0.8 mM). The procedure was identical for the experiment examining the effects of GABA receptor antagonism in

the IL during fear renewal (Fig. 5d–f), except that rats received intra-IL infusions (0.3 µl/hemisphere; 0.3 µl/min) of CGP (10 mM), PTX (0.33 mM), CGP+PTX cocktail (from 10 mM and 0.33 mM stocks, respectively), or SAL in context A. For the intracranial drug infusions, injection cannulas were connected via water-filled polyethylene tubing to gas-tight Hamilton syringes (10 µl) that were secured to an infusion pump (KD Scientific). Drug or vehicle was loaded into the injection cannulas and pressure-injected following the parameters described above and based on prior work (injection cannulas remained in the guides for 1 min following infusion to allow for adequate diffusion)³⁹.

Histology. At the conclusion of behavioral testing or in vivo recordings, rats were overdosed with sodium pentobarbital (Fatal Plus; 100 mg/ml, 0.5 ml) and were transcardially perfused with saline and 10% formalin. Brains were extracted and post-fixed in formalin solution for 24 h at 4 °C and transferred to 20% sucrose (4 °C). Brains were flash frozen with dry ice and sectioned (40 µm) on a cryostat maintained at –20 °C. For non-Cre-dependent viral experiments, sections were wet-mounted to subbed slides and coverslipped with Fluoromount (Sigma-Aldrich) and imaged on a Zeiss microscope (via Axio Imager 2). Cre-dependent viral tissue was collected into wells for immunohistochemistry. For cannula-implanted rats, thionin-stained coronal sections containing IL were imaged on a Leica microscope (MZ FLIII).

Immunohistochemistry. For biocytin recovery and immunohistochemistry in the slice electrophysiology experiments, brain slices were fixed with 4% paraformaldehyde in 0.1 M PBS for 40–60 min at room temperature. Slices were washed three times with 0.1 M PBS, then blocked with a blocking solution containing PBS (0.1 M), bovine serum albumin (3%), saponin (0.1%) and sodium azide (0.05%) for 1 h at room temperature. Slices were washed with PBS and incubated in primary antibodies including anti-TBR1 and streptavidin (Alexa fluor 488/555/647 at 1:1,000, Invitrogen) for 1–3 d at room temperature or at 4 °C. Sections were washed three times for 15 min each time in PBS and Alexa Fluor-conjugated species-specific secondary antibodies (1:1,000, Jackson Biosciences or Invitrogen) for at least 5 h at room temperature (22–23 °C). After a triple wash with PBS, the brain slices were mounted in glycerol and PBS solution (50% and 50%) or PBS only for tissues containing retrograde tracers. Slices were imaged using an upright microscope (5× or 20×, Zeiss, Axio Imager) or a confocal system (20×, Zeiss LSM510). Images were produced by flattening z-stacks (1-µm intervals) to a maximum projection image using Zen 2011 software (Zeiss).

For DIO-DREADD-expressing animals, immunohistochemistry was performed on free-floating brain sections containing the vHPC. The tissue was washed three times in 1× Tris-buffered saline (TBS, pH 7.4). Brain sections were then incubated in 10% normal donkey serum (NDS) in 1× Tris-buffered saline with Tween (TBST) for 1 h at room temperature followed by two washes in TBST for 5 min. Tissue was then incubated in primary antibody solution in TBST with 2% NDS (rabbit anti-mCherry antibody at 1:2,000; Abcam) for 48 h at 4 °C. Brain sections were then washed three times in TBST for 10 min and were incubated in secondary antibody solution in TBST with 2% NDS (AF594-conjugated donkey anti-rabbit antibody at 1:200; Abcam) for 2 h at room temperature (22–23 °C). Tissue was washed three times in TBS for 10 min and then was mounted on subbed slides in 0.9% saline and coverslipped with Fluoromount. Brain sections containing the IL were wet mounted to microscope slides and coverslipped with Fluoromount for imaging. For Fos immunohistochemistry, the tissue was processed exactly as for mCherry immunohistochemistry except rabbit anti-c-Fos antibody (1:5,000; Millipore) and AF488-conjugated donkey anti-rabbit antibody (1:500, Life Technologies) were used for primary and secondary antibodies, respectively.

For PV/Fos dual-staining, rats were sacrificed 120 min after CNO or vehicle administration. 18 h post-fix, brains were moved into 30% sucrose solution for 3 d. 40-µm-thick brain sections were collected for staining. Brain sections were washed three times in TBST and then were incubated in 0.3% H₂O₂ for 15 min. The tissue was washed in TBS three times and was incubated in rabbit anti-c-Fos primary antibody (1:1,000; Millipore) overnight. Brain tissue was washed three times in TBS followed by 1 h incubation in a biotinylated goat anti-rabbit secondary antibody (1:1,000; Jackson ImmunoResearch), amplification with the avidin biotin complex at 1:1,000 (ABC; Vector labs), and visualization with 3,3'-diaminobenzidine (DAB) + nickel ammonium sulfate to yield a purple/black nuclear reaction product. The tissue was subsequently incubated in a mouse

anti-PV primary antibody (1:5,000; Sigma) overnight and biotinylated goat anti-mouse secondary antibody (1:500; Jackson ImmunoResearch) for 1 h, amplified with the avidin biotin complex at 1:1,000 (ABC; Vector labs), and visualized with 3,3'-diaminobenzidine (DAB) to yield a light brown somatic stain. Slices were mounted and coverslipped with mounting medium (Permount; Sigma), and stored at room temperature (22–23 °C) until photographed using a Zeiss microscope. Images for cell counting were generated and counted (using ImageJ) following standard procedures (including, when possible, the blinding of the experimenter to group assignments of the subjects) and based on prior work^{18,22}. Three bilateral images of the vHPC were selected (–5.52 mm, –5.76 mm and –6.00 mm from Bregma) for mCherry-hm3D(G_q) and Fos double-labeling. For mPFC analyses, three bilateral images of the PL and IL were selected (PL: +4.2 mm, +3.7 mm and +3.2 mm from bregma; IL: +3.7 mm, +3.2 mm and +3.00 mm from bregma) for PV and Fos double-labeling.

Statistics. Paired *t* tests were used to analyze inputs into mPFC subregions, interneuronal subtypes and IL sublayers. Unpaired *t* tests were used to analyze PV⁺- and SOM⁺-mediated IPSC amplitudes. Paired *t* tests were used to analyze data in DIO-hm4D(G_i), DIO-GFP, and DIO-hm3D(G_q) experiments separately. Unpaired *t* tests were used to analyze the PV/Fos dual-staining experiment. IL microinfusion and the non-Cre-dependent and Cre-dependent virus behavioral data were analyzed with repeated measures analysis of variance (ANOVA). Post hoc comparisons in the form of Fisher's protected least significant difference (PLSD) post hoc tests, which were performed after a significant overall *F* ratio. No statistical methods were used to pre-determine sample sizes, but our sample sizes are similar to those in prior reports and are typical for the field. The data distribution was assumed to be normal, but this was not formally tested. All data are represented as means ± s.e.m.

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

Data and code availability. The data that support the findings of this study are available from the corresponding author upon reasonable request. No custom code was used in the current work.

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► Experimental design

1. Sample size

Describe how sample size was determined.

No statistical methods were used to predetermine sample size, but the numbers for groups in each sample were based on those in previously published studies.

2. Data exclusions

Describe any data exclusions.

Outliers (where noted) were determined using a Grubbs' test.

3. Replication

Describe the measures taken to verify the reproducibility of the experimental findings.

All attempts at replication of included data were successful.

4. Randomization

Describe how samples/organisms/participants were allocated into experimental groups.

Animals for behavioral experiments were randomly selected to initially receive either the control or drug injection.

5. Blinding

Describe whether the investigators were blinded to group allocation during data collection and/or analysis.

All behavioral measurements of freezing were performed using an automated system, hence no blinding was necessary.

Note: all in vivo studies must report how sample size was determined and whether blinding and randomization were used.

6. Statistical parameters

For all figures and tables that use statistical methods, confirm that the following items are present in relevant figure legends (or in the Methods section if additional space is needed).

- | | |
|--------------------------|---|
| n/a | Confirmed |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> The <u>exact sample size</u> (<i>n</i>) for each experimental group/condition, given as a discrete number and unit of measurement (animals, litters, cultures, etc.) |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> A description of how samples were collected, noting whether measurements were taken from distinct samples or whether the same sample was measured repeatedly |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> A statement indicating how many times each experiment was replicated |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> The statistical test(s) used and whether they are one- or two-sided
<i>Only common tests should be described solely by name; describe more complex techniques in the Methods section.</i> |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> A description of any assumptions or corrections, such as an adjustment for multiple comparisons |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> Test values indicating whether an effect is present
<i>Provide confidence intervals or give results of significance tests (e.g. <i>P</i> values) as exact values whenever appropriate and with effect sizes noted.</i> |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> A clear description of statistics including <u>central tendency</u> (e.g. median, mean) and <u>variation</u> (e.g. standard deviation, interquartile range) |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> Clearly defined error bars in <u>all</u> relevant figure captions (with explicit mention of central tendency and variation) |

See the web collection on [statistics for biologists](#) for further resources and guidance.

► Software

Policy information about [availability of computer code](#)

7. Software

Describe the software used to analyze the data in this study.

Data were analyzed and plotted using AxoGraph X (V 1.2.1), Zen (Zeiss; 2011), Neuroexplorer, ImageJ, StatView, Graphpad, Prism, and SPSS.

For manuscripts utilizing custom algorithms or software that are central to the paper but not yet described in the published literature, software must be made available to editors and reviewers upon request. We strongly encourage code deposition in a community repository (e.g. GitHub). *Nature Methods* [guidance for providing algorithms and software for publication](#) provides further information on this topic.

► Materials and reagents

Policy information about [availability of materials](#)

8. Materials availability

Indicate whether there are restrictions on availability of unique materials or if these materials are only available for distribution by a third party.

No unique materials were used.

9. Antibodies

Describe the antibodies used and how they were validated for use in the system under study (i.e. assay and species).

For mCherry immunohistochemistry:
Rabbit anti-mCherry primary antibody (Abcam, ab167453)
Donkey anti-rabbit secondary antibody (Abcam, ab150076)

For Fos immunohistochemistry:
Rabbit anti-c-Fos primary antibody (Millipore, ABE457)
Goat anti-rabbit secondary antibody (Jackson ImmunoResearch, 111-065-003)

For PV/Fos dual immunohistochemistry:
Rabbit anti-c-Fos primary antibody (Millipore, ABE457)
Goat anti-rabbit secondary antibody (Jackson ImmunoResearch, 111-065-003)
Mouse anti-parvalbumin primary antibody (Millipore, MAB1572)
Goat anti-mouse secondary antibody (Abcam, ab150113)
ABC kit (Vector, PK6100)

All the antibodies were tested using immunohistochemistry before use. Different dilutions of the primary antibody (including control without primary) were used to stain sample rat brain tissue to find the optimal primary antibody concentration. Tissue stained without primary antibody helped rule out the possibility of non-specific binding of secondary antibody to the tissue. Finally, a negative tissue sample (tissue without the target protein) was stained using optimal primary and secondary antibody to rule out non-specific binding of the antibodies.

10. Eukaryotic cell lines

- State the source of each eukaryotic cell line used.
- Describe the method of cell line authentication used.
- Report whether the cell lines were tested for mycoplasma contamination.
- If any of the cell lines used are listed in the database of commonly misidentified cell lines maintained by [ICLAC](#), provide a scientific rationale for their use.

No eukaryotic cell lines were used.

No eukaryotic cell lines were used.

No eukaryotic cell lines were used.

No eukaryotic cell lines were used.

► Animals and human research participants

Policy information about [studies involving animals](#); when reporting animal research, follow the [ARRIVE guidelines](#)

11. Description of research animals

Provide all relevant details on animals and/or animal-derived materials used in the study.

Adult male Wistar rats (170-240g), PV-Cre knock-in (C57BL/6 strain) mice (for the local optical PV stimulation, the ivermectin-receptor silencing experiments, and the dual-optical approach; 20-26g), and SOM-Cre knock-in (C57BL/6 strain) mice (19-25g) were used in the ex-vivo electrophysiological experiments. Adult male Long-Evans rats (200-224 g; Blue-Spruce, Harlan, Indianapolis, IN) were used for the behavioral experiments.

12. Description of human research participants

Describe the covariate-relevant population characteristics of the human research participants.

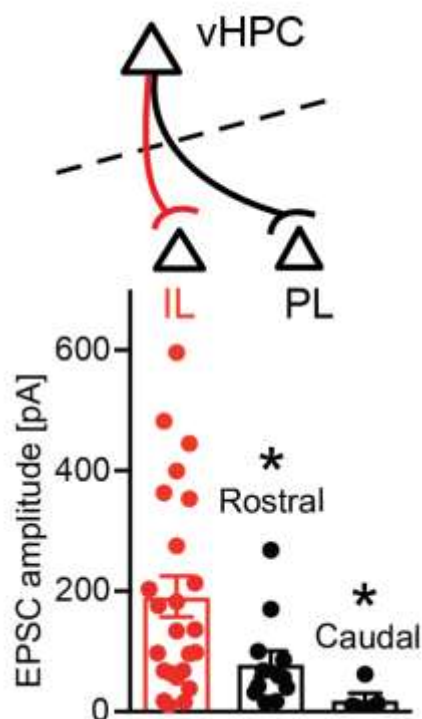
This study did not involve human research participants.

In the format provided by the authors and unedited.

Hippocampus-driven feed-forward inhibition of the prefrontal cortex mediates relapse of extinguished fear

Roger Marek^{1,6}, Jingji Jin^{2,6}, Travis D. Goode^{2,6}, Thomas F. Giustino², Qian Wang³, Gillian M. Acca², Roopashri Holehonnur⁴, Jonathan E. Ploski⁴, Paul J. Fitzgerald², Timothy Lynagh⁵, Joseph W. Lynch¹, Stephen Maren^{2*} and Pankaj Sah^{1*}

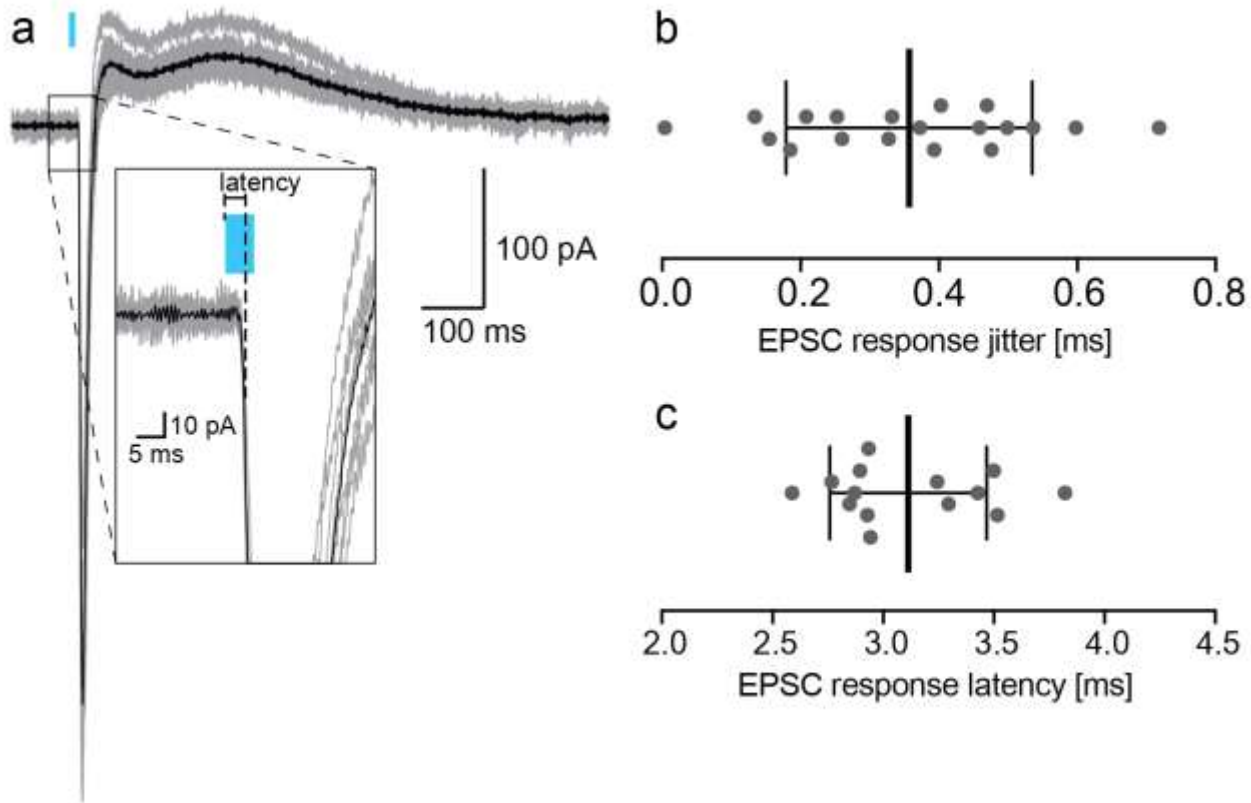
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Supplementary Figure 1

HPC projections predominantly target IL neurons in the mPFC.

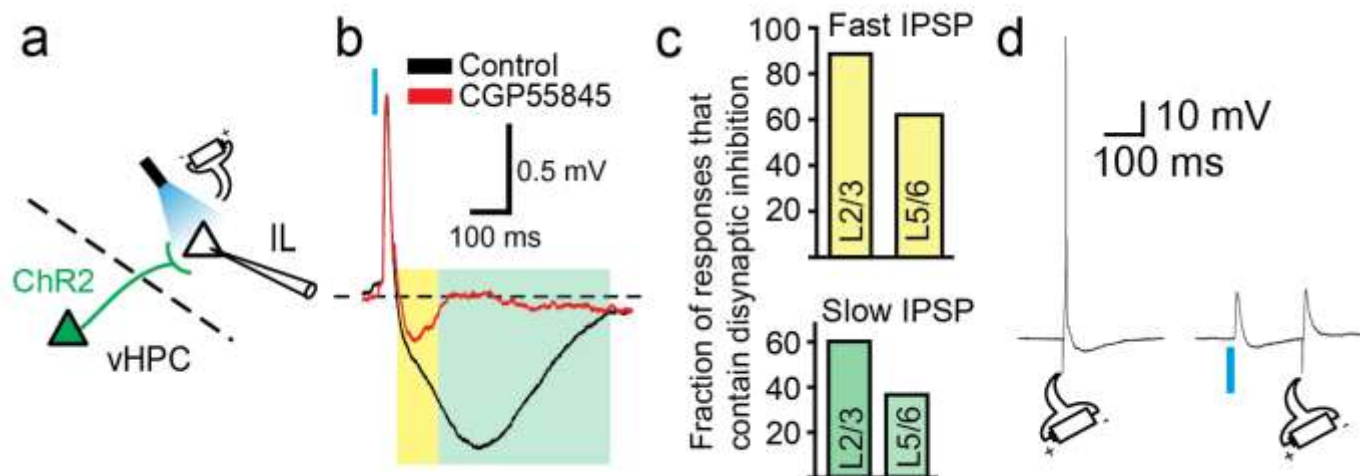
Comparison of hippocampal inputs onto principal neurons in the IL and PL reveal significantly larger EPSCs in IL neurons compared to caudal or rostral PL responses (rostral PL: $n = 12$; caudal PL: $n = 5$; IL: $n = 21$ from 7 animals; one-way ANOVA with Dunnett's multiple comparison test, $F_{2,35} = 6.62$, $*P = 0.0037$). Dots represent data from individual cells. Error bars indicate means \pm s.e.m.



Supplementary Figure 2

Hippocampus-driven IL responses are monosynaptic, time-locked responses.

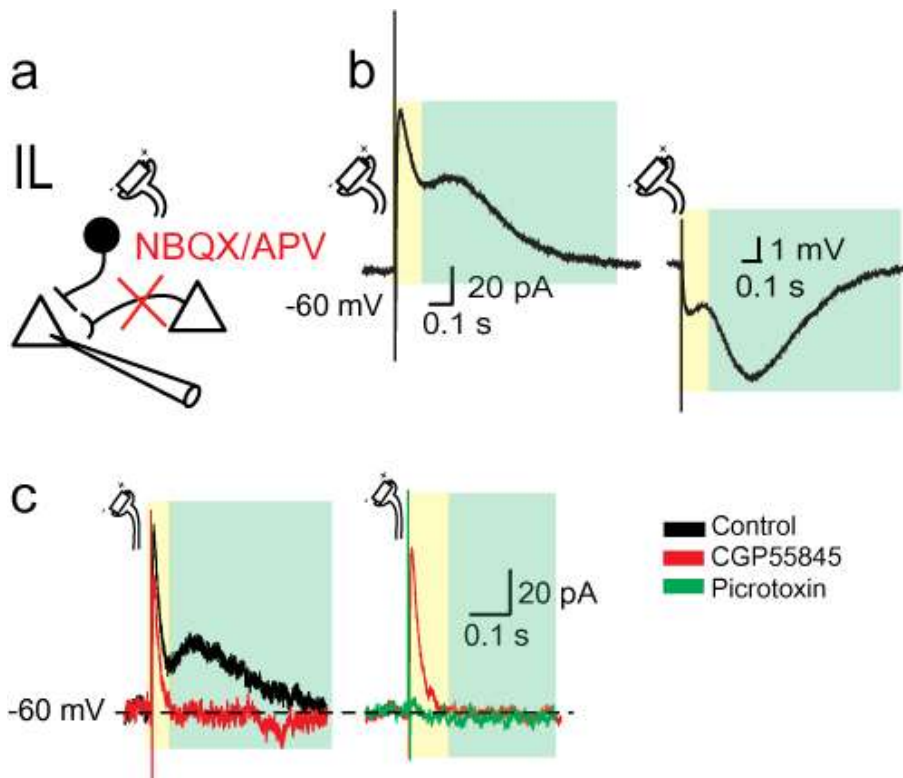
a, Example of a HPC-driven response in a pyramidal neuron, containing the initial excitatory response, followed by an inhibitory component. The 5-fold enlarged inset shows the time-locked onset of individual responses (grey) to the light stimulation (blue) without any response-failures. Averaged response shown in black. Analysis of EPSCs show very low response jitter (**b**) (individual jitter of neurons shown as grey dots; $n = 19$) and response latency (**c**) ($n = 14$), both typical parameters for monosynaptic responses. Error bars indicate means \pm s.e.m.



Supplementary Figure 3

Quantification of vHPC-evoked inhibitory conductances in IL principal cells.

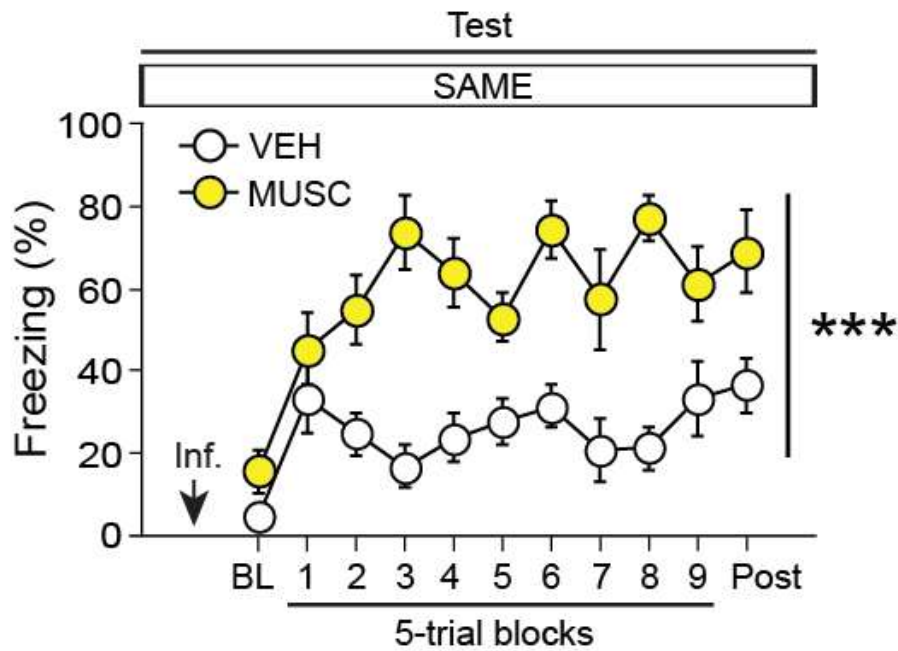
a, Schematic showing experimental setup. Optical terminal stimulation of ventral hippocampal inputs and whole-cell recording of pyramidal neurons in the IL. For the spiking suppression experiments (**d**), local electrical stimulation (battery symbol) was used. **b**, Current-clamp responses to hippocampal terminal release before (black trace) and after (red trace) the application of the GABA_B-receptor antagonist CGP55845 ($n = 3$), revealing the fast (yellow) and slow IPSCs (green). **c**, Bar graphs with response patterns of IL principal neurons to optical hippocampal stimulation containing fast (top, yellow) and slow (bottom, green) inhibitory conductances (percentage of the total). **d**, Investigation of spiking suppression of IL principal neurons by the hippocampus. Electrical suprathreshold stimulation was used to evoke spiking (left) while presenting optical hippocampal stimulation 150 ms before the spiking event ($n = 8$). Blue bars represent optical stimulation.



Supplementary Figure 4

Feed-forward inhibition onto pyramidal neurons is mediated by local IL interneurons.

a. Schematic for electrical stimulation (battery symbol) of IL tissue *ex vivo* in the presence of AMPA- and NMDA-receptor antagonists NBQX and APV, respectively, to isolate GABAergic transmission. **b.** Voltage clamp (left) and current-clamp recordings (right) revealed inhibitory responses that contain both fast (yellow) and slow (green) inhibitory components. **c.** Application of the GABA_B-receptor antagonist CGP55845 (red trace) isolated the fast, inhibitory conductance, which was blocked by the GABA_{A/C}-receptor antagonist picrotoxin (green trace). Holding voltage: -60 mV.



Supplementary Figure 6

Pharmacological inactivation of the IL impedes retrieval of extinguished fear.

Test data show mean baseline freezing (3 min), mean freezing during nine 5-ITI blocks (30-sec ITIs) and a during a post-trial period (150 sec) following infusions of muscimol or vehicle into the IL (MUSC, $n = 6$; VEH, $n = 10$; repeated measures ANOVA, main effect of drug: $F_{1,14} = 35.78$, $***P < 0.0001$). Corresponding conditioning and extinction data are shown in Fig. 4. Error bars indicate means \pm s.e.m.