

Reducing Astrocyte Calcium Signaling *In Vivo* Alters Striatal Microcircuits and Causes Repetitive Behavior

Highlights

- Multiple approaches were used to assess striatal astrocyte Ca^{2+} signaling *in vivo*
- Ambient GABA was altered when astrocyte signaling was reduced
- Reduced astrocyte signaling resulted in excessive self-grooming in mice
- Astrocytes contribute to neural circuits *in vivo* and OCD-like phenotypes in mice

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In Brief

The Khakh laboratory evaluated the consequences of genetically attenuating astrocyte calcium signaling in the adult mouse striatum *in vivo*. They discovered excessive self-grooming phenotypes, the mechanisms of which were explored at the molecular, cellular, and *in vivo* levels.



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SUMMARY

Astrocytes tile the central nervous system, but their functions in neural microcircuits *in vivo* and their roles in mammalian behavior remain incompletely defined. We used two-photon laser scanning microscopy, electrophysiology, MINIsopes, RNA-seq, and a genetic approach to explore the effects of reduced striatal astrocyte Ca^{2+} signaling *in vivo*. In wild-type mice, reducing striatal astrocyte Ca^{2+} -dependent signaling increased repetitive self-grooming behaviors by altering medium spiny neuron (MSN) activity. The mechanism involved astrocyte-mediated neuromodulation facilitated by ambient GABA and was corrected by blocking astrocyte GABA transporter 3 (GAT-3). Furthermore, in a mouse model of Huntington's disease, dysregulation of GABA and astrocyte Ca^{2+} signaling accompanied excessive self-grooming, which was relieved by blocking GAT-3. Assessments with RNA-seq revealed astrocyte genes and pathways regulated by Ca^{2+} signaling in a cell-autonomous and non-cell-autonomous manner, including *Rab11a*, a regulator of GAT-3 functional expression. Thus, striatal astrocytes contribute to neuromodulation controlling mouse obsessive-compulsive-like behavior.

INTRODUCTION

Astrocytes tile the entire central nervous system and serve multiple roles, including ion homeostasis, synapse formation,

synapse removal, synaptic regulation, and contributions to neurovascular coupling (Khakh and Sofroniew, 2015). Astrocytes are also proposed to regulate neuronal function via intracellular Ca^{2+} -dependent signaling that has been documented *in vitro* and *in vivo* in diverse species (Bazargani and Attwell, 2016; Shigetomi et al., 2016). Furthermore, astrocyte Ca^{2+} signaling is known to be dysfunctional in mouse models of neurological disease (Khakh and Sofroniew, 2015). However, despite progress, the functional significance of astrocyte Ca^{2+} -dependent signaling for the function(s) of neural microcircuits *in vivo*, for vertebrate behavior, and for brain diseases remains incompletely understood.

A barrier to exploring astrocyte roles *in vivo* has been the inability to reduce Ca^{2+} signaling specifically within astrocytes in the fully developed adult brain in a nucleus-specific manner. Previous studies addressed the functions of astrocyte Ca^{2+} signaling *in vivo* by employing IP3 receptor type 2 (IP3R2) deletion mice, which attenuated Ca^{2+} signals but produced no behavioral consequences (Aguilhon et al., 2010; Petravic et al., 2014). However, the approach was not brain-area specific, the mice lacked IP3R2s throughout the body and throughout development, and some astrocyte Ca^{2+} signaling persisted (Rungta et al., 2016; Srinivasan et al., 2015; Stobart et al., 2018). These are important considerations and the use of IP3R2 deletion mice has led to a confusing picture on the relevance of astrocyte Ca^{2+} signaling (Khakh and McCarthy, 2015; Shigetomi et al., 2016). The suggestion that astrocyte Ca^{2+} signaling has little role in regulating neurons or behavior is at odds with *in vivo* imaging and slice electrophysiology data (Nimmerjahn and Bergles, 2015; Volterra et al., 2014). An additional consideration is the fact that astrocytes change during development, display brain-area specializations, and exhibit diverse types of Ca^{2+} signals (Shigetomi et al., 2016). Furthermore, behavior derives from the output of microcircuits located in



distinct brain nuclei. Therefore, brain-area-specific methods to attenuate astrocyte Ca^{2+} -dependent signaling in adult mice *in vivo* are needed. In these regards, some studies have used overexpression of IP3 binding proteins as a “sponge” to decrease IP3-dependent signaling (Xie et al., 2010). The uncertainty on the functions of astrocyte Ca^{2+} signaling *in vivo* provided the impetus to explore complementary approaches in vertebrates.

We explored the functions of astrocytes within the microcircuits of the striatum, which is the largest nucleus of the basal ganglia, a group of interconnected subcortical nuclei. We define brain microcircuits as comprising neurons and associated cells, such as glia, organized to carry out specific operations within a region of the nervous system (Shepherd and Grillner, 2010).

RESULTS

Reducing Striatal Astrocyte Calcium-Dependent Signaling

To reduce astrocyte Ca^{2+} signaling, we tested isoforms of the plasma membrane Ca^{2+} pump PMCA, which functions to constitutively extrude cytosolic Ca^{2+} (Strehler, 2015). We hypothesized that this would reduce astrocyte Ca^{2+} signaling, because most such signals last seconds. Such an approach would also conceivably be general with regards to the many types of slow Ca^{2+} signals. We focused on PMCA2 because of its high pump rate and because it is unlikely expressed by astrocytes (Srinivasan et al., 2016; Zhang et al., 2014), which we confirmed for adult striatal astrocytes (Figures S1A–S1D). After screening and testing (Figures S1A–S1H), we focused on human PMCA isoform type 2 with the w/b splice variant that lacks cytosolic interaction domains and modified to carry a mCherry tag. We abbreviated the construct as hPMCA2w/b and used “CalEx” for the approach to reflect calcium extrusion. hPMCA2w/b significantly reduced by >95% 50 μM ATP-evoked P2Y1-receptor-mediated Ca^{2+} signals in HEK293 cells (Figures S1E–S1H; $n = 33$, $p < 0.001$, one-way ANOVA).

To deliver hPMCA2w/b to astrocytes *in vivo*, we used local microinjections of adeno-associated viruses (AAV2/5) with an astrocyte-specific GfaABC₁D promoter (Shigetomi et al., 2013). We confirmed that microinjection of AAV2/5-delivered cargo was astrocyte selective and restricted to the striatum, although there was a little expression along the needle tract in the cortex (Figure S2A; $n = 3$ mice): our studies need to be interpreted with this in mind. We microinjected hPMCA2w/b and the genetically encoded calcium indicator GCaMP6f *in vivo*, resulting in expression of both proteins in ~90% of astrocytes in the dorsolateral striatum (Figures 1A–1C; Figures S2B and S2C). Neuronal expression was not observed ($n = 4$ mice; Figure S2D). In controls that expressed tdTomato instead of hPMCA2w/b, we measured frequent, large-amplitude, spontaneously occurring Ca^{2+} elevations that we call Ca^{2+} signals; previously, they have also been called transients, events, fluctuations, and oscillations. Such signals lasted seconds in striatal astrocyte somata, branches, and territories (Figures 1D–1G; $n = 14$ cells, $n = 4$ mice). In hPMCA2w/b-expressing astrocytes, Ca^{2+} signals were reduced by ~70% in terms of amplitude (Figure 1D; $p < 0.001$, unpaired Mann-Whitney tests; Video S1) and were also

significantly shorter in duration in striatal astrocyte somata, branches, and territories (Figure S2E, unpaired Mann-Whitney tests), but the effects on frequency were variable between compartments (Figure S2F). We interpret this to indicate that, by virtue of its Ca^{2+} extrusion ability, hPMCA2w/b significantly reduced the amplitude and shortened the duration of Ca^{2+} signals, but not their occurrence.

Astrocytes respond with global Ca^{2+} elevations to a variety of G-protein-coupled receptor (GPCR) agonists. We next explored whether hPMCA2w/b affected pharmacologically evoked Ca^{2+} elevations in striatal slices. hPMCA2w/b significantly reduced 10 μM phenylephrine (PE)-evoked GPCR-mediated astrocyte Ca^{2+} signals in all astrocyte compartments such that PE failed to evoke significant Ca^{2+} elevations (Figures 1H–1K). hPMCA2w/b also attenuated global Ca^{2+} elevations triggered by intracellular Ca^{2+} store depletion (Figures S2G–S2I). Recent data emphasize the relevance of astrocyte basal Ca^{2+} levels (Rosenegger et al., 2015). We found that hPMCA2w/b significantly reduced astrocyte basal Ca^{2+} levels (Figures S3A–S3C).

IP3R2 deletion reduced striatal astrocyte Ca^{2+} signaling (Jiang et al., 2016). We directly compared the degree of Ca^{2+} signaling attenuation produced by hPMCA2w/b ($n = 14$ cells, 6 mice) and IP3R2 deletion ($n = 14$ cells, 6 mice) mice in relation to controls ($n = 20$ cells, 7 mice) using GCaMP6f (Figure S3D). In regards to spontaneous Ca^{2+} signals, we found that IP3R2 deletion produced a statistically larger reduction of Ca^{2+} signal frequency than hPMCA2w/b across all astrocyte compartments. There was a trend for a larger reduction in Ca^{2+} signal amplitude in IP3R2 deletion mice, but this was only significant for astrocyte branches. However, hPMCA2w/b more strongly attenuated the duration of the Ca^{2+} signals. For PE-evoked responses, both hPMCA2w/b and IP3R2 deletion similarly (>80%) attenuated the responses, but the degree of reduction was statistically greater in IP3R2 deletion mice (Figure S3D). These data compare hPMCA2w/b expressed in the striatum of adult mice with global IP3R2 deletion mice.

Reducing Striatal Astrocyte Calcium-Dependent Signaling Leads to Excessive Self-Grooming

We microinjected hPMCA2w/b (or control AAVs) bilaterally into the dorsolateral striatum to determine how attenuation of Ca^{2+} -dependent signaling affected behavior. There are no past *in vivo* data available to indicate which striatal behaviors astrocyte Ca^{2+} signaling may regulate; we performed several tests and studied the phenotypes that we discovered.

Control and hPMCA2w/b mice were healthy, weighed similarly (26.8 ± 0.5 and 26.3 ± 0.5 g for control and hPMCA2w/b mice, respectively; $n = 8$, $p = 0.543$; Student's *t* test) and performed equally in a rotarod test of motor function (latency to fall was 195.6 ± 19.3 and 162.5 ± 21.6 s for control and hPMCA2w/b mice, respectively; $n = 8$ mice, $p = 0.273$; Student's *t* test). However, in the open field test, hPMCA2w/b mice displayed shorter travel distances and spent less time in the center (Figures 2A–2C; $n = 8$ mice, $p = 0.001$ and $p = 0.003$, two-way ANOVA and Student's *t* test). While avoidance of center regions of the open field arena can be interpreted as reflecting phenotypes related to anxiety, other behavioral measures of anxiety produced no differences. For example, control and hPMCA2w/b

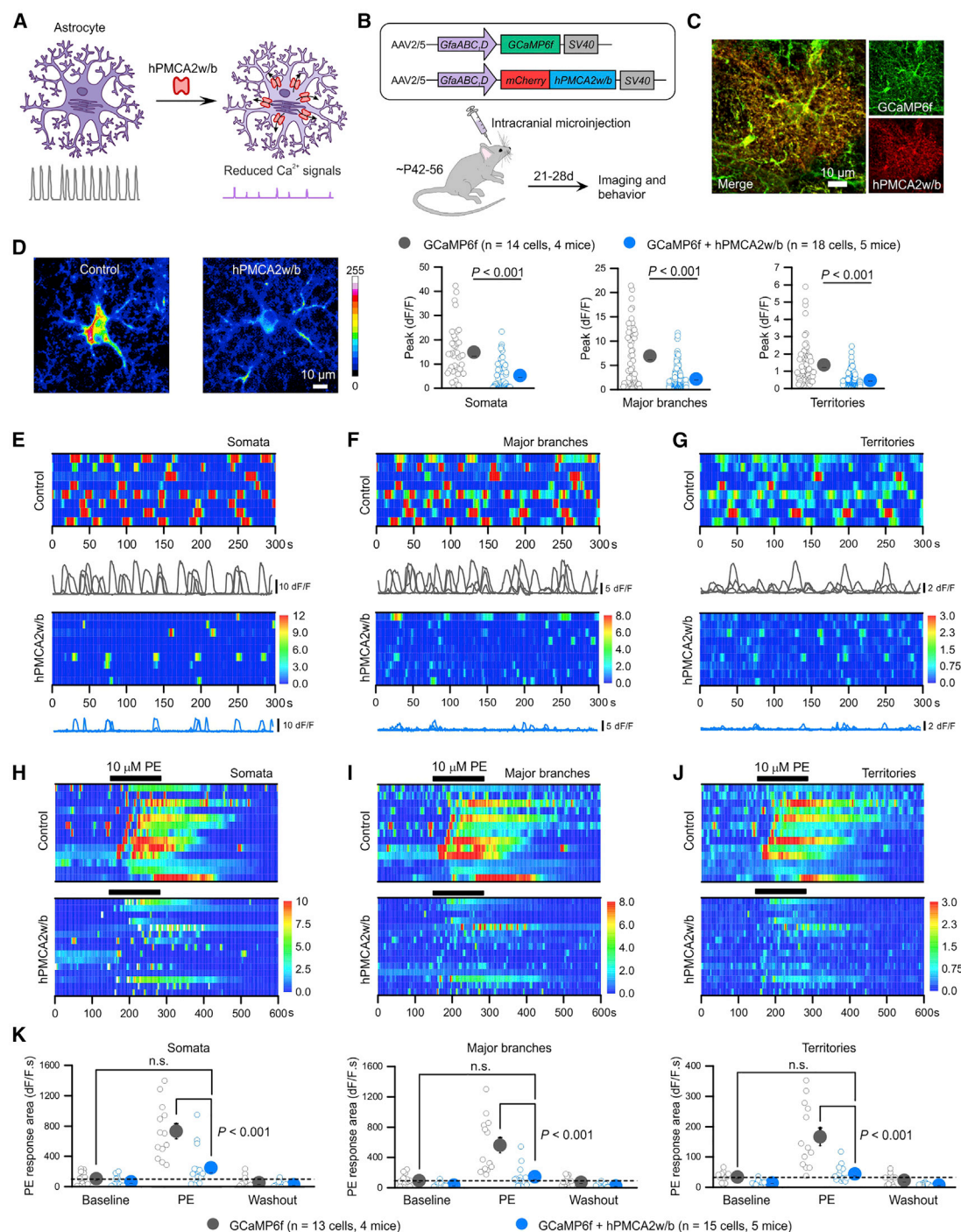


Figure 1. Reducing Ca^{2+} -Dependent Signaling in Striatal Astrocytes with hPMCA2w/b

(A) Schematic of hPMCA2w/b to attenuate Ca^{2+} signals.

(B) AAVs for expressing mCherry-hPMCA2w/b with GCaMP6f in astrocytes.

(C) IHC images showing colocalization of GCaMP6f and hPMCA2w/b in striatal astrocytes (Figure S2).

(D) Z stack images of GCaMP6f-expressing striatal astrocytes for control and hPMCA2w/b mice. Bar graphs: peak amplitude of spontaneous Ca^{2+} signals in somata, major branches, and territories of striatal astrocytes (Mann-Whitney tests).

(E–G) Kymographs and $\Delta\text{F}/\text{F}$ traces of Ca^{2+} signals in somata (E), major branches (F), and territories (G) of control and hPMCA2w/b-expressing astrocytes.

(H–J) Kymographs of cells with Ca^{2+} responses evoked by phenylephrine (PE, 10 μM) in somata (H), major branches (I), and territories (J) of control and hPMCA2w/b-expressing astrocytes.

(K) Summary plots for experiments in (H)–(J). Average data are shown as mean \pm SEM. In some cases, the SEM symbol is smaller than the symbol for the mean.

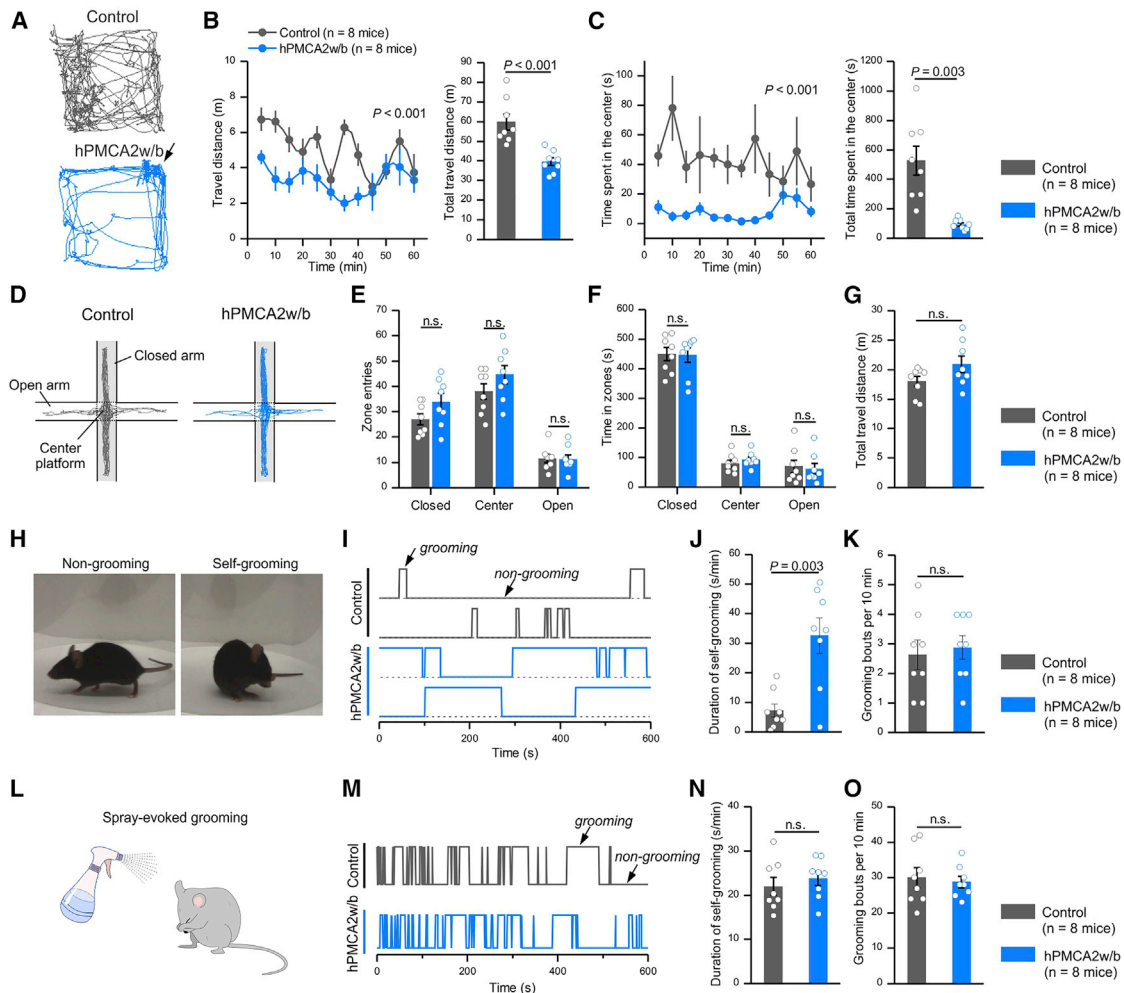


Figure 2. Reducing Ca^{2+} -Dependent Signaling in Striatal Astrocytes Increased Self-Grooming

(A) Representative track traces of locomotor activity in open field tests of control and hPMCA2w/b-expressing mice over 10 min. hPMCA2w/b mice spent extended periods in the corners (arrow).
 (B and C) Travel distance (B) and time spent in the center (C) were decreased in hPMCA2w/b mice. Graphs tracking open field activity over time were assessed with two-way ANOVA tests. The comparisons between bar graphs were with Student's *t* tests or Mann-Whitney tests.
 (D) Representative track traces of locomotor activity in the elevated plus maze for control and hPMCA2w/b mice.
 (E–G) Average data for zone entries (E), time in different zones (F), and the total travel distance (G) between control and hPMCA2w/b mice.
 (H) Images of non-grooming and self-grooming behaviors.
 (I) Behavioral traces of control and hPMCA2w/b-expressing mice for times in self-grooming and non-grooming episodes.
 (J and K) The number of self-grooming bouts (K) and duration in seconds per minute of self-grooming (J) in hPMCA2w/b and control mice (Student's *t* tests).
 (L–O) Grooming evoked by spraying the mice with water for controls and hPMCA2w/b (L): dwell times (M), average duration (N), and number of bouts (O). The average data are shown as mean \pm SEM. In some cases, the SEM bars are smaller than the symbol used to represent the mean.

mice were indiscernible in the elevated plus maze (Figures 2D–2G). Furthermore, hyperlocomotion of hPMCA2w/b and control mice was indistinguishable following cocaine (Figures S4A–S4C), indicating that the dopaminergic pathway was intact and arguing against motor dysfunction. Notably, we found that hPMCA2w/b mice spent long periods in the corners of the open field (Figure 2A), and closer examination showed that hPMCA2w/b mice were excessively self-grooming during such episodes (Figures 2H–2K). The increased self-grooming was observed in the open field and when the mice were placed in a clear chamber (Video S2). We quantified the total duration of

self-grooming by measuring dwell times in grooming and non-grooming states (Figure 2I), confirming that hPMCA2w/b mice self-groomed excessively and significantly by $\sim 450\%$ (Figure 2J; $n = 8$, $p = 0.0034$, Student's *t* test). The number of self-grooming bouts was not significantly different for control and hPMCA2w/b mice (Figure 2K; $p = 0.7008$ Student's *t* test), but the total bout length was significantly longer (Figure 2J). This was also significant for the duration of individual self-grooming bouts (16.6 ± 5.6 s and 60.3 ± 9.9 s for control and hPMCA2w/b, respectively; $n = 7$, $p < 0.001$ Student's *t* test). We assessed stress-related grooming and found equally high levels of grooming in this spray

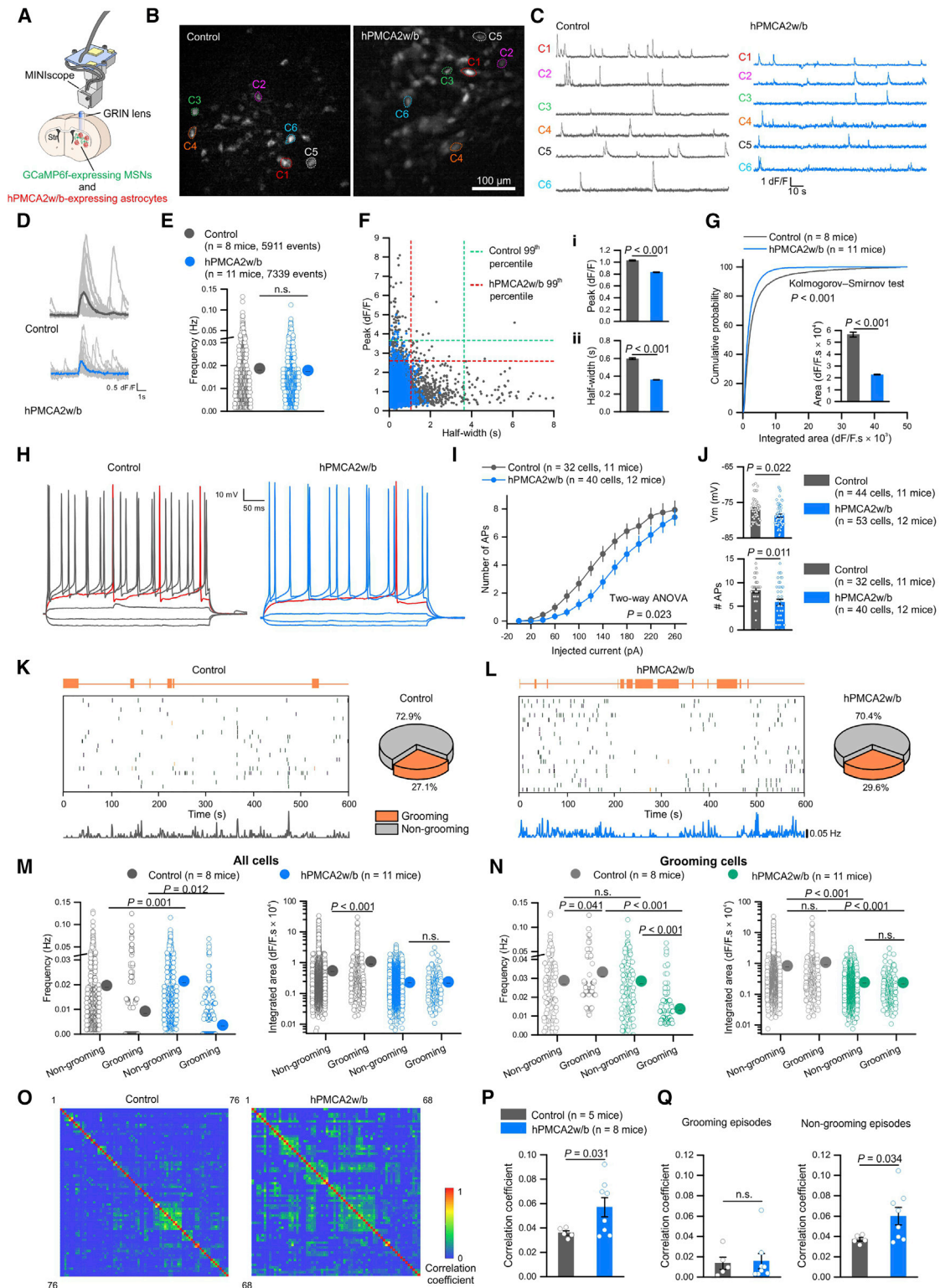


Figure 3. Reducing Striatal Astrocyte Ca^{2+} -Dependent Signaling Altered MSN Activity

(A) The GRIN lens for MINIScope was implanted above the dorsal striatum near GCaMP6f-expressing MSNs and hPMCA2w/b-expressing astrocytes. (B) Z-projected Ca^{2+} images from MINIScope recordings in control and hPMCA2w/b mice. (C) Representative $\Delta F/F$ traces of individual MSNs recorded in (B).

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test (Figures 2L–2O). Self-grooming is an innate behavior for which syntax is controlled by the dorsolateral striatum, and excessive self-grooming is associated with obsessive-compulsive disorder phenotypes (Kalueff et al., 2016).

For the remainder of this study, we focused on self-grooming (Figure 2). For a specific set of experiments, we expressed hPMCA2w/b in hippocampal astrocytes and found that it significantly reduced both spontaneous and PE-evoked Ca^{2+} signals (Figures S5A–S5F; $n = 27$ and 24 cells, $n = 4$ and 6 mice for controls and hPMCA2w/b). However, we did not observe altered self-grooming relative to AAV controls following hippocampal expression (Figures S5G and S5H). These and other controls suggest that the self-grooming phenotype following hPMCA2w/b expression in the striatum is specific.

Reducing Striatal Astrocyte Calcium-Dependent Signaling Alters Microcircuit Function *In Vivo*

The dorsolateral striatum is involved in coordinating motor functions, including self-grooming (Graybiel and Grafton, 2015; Kalueff et al., 2016). To explore how astrocyte hPMCA2w/b affected striatal microcircuits, we expressed GCaMP6f in medium spiny neurons (MSNs) (Figures S6A–S6D) and used Ca^{2+} imaging to measure MSN activity in freely behaving mice with head-mounted MINIsopes (Cai et al., 2016) (Figures 3A and 3B; Video S3; Table S1). We detected activity *in vivo* from ~80 MSNs (Figures 3B–3D; $n = 8$ and 11 mice). MSNs from hPMCA2w/b-expressing mice displayed the same frequency of activity as controls (Figure 3E), but with smaller amplitudes and shorter durations, reflected as smaller integrated areas ($p < 0.001$; Figures 3D, 3F, and 3G). Furthermore, examination of the data showed that hPMCA2w/b caused a preferential loss of large-amplitude MSN activity lasting several seconds (Figure 3F), suggesting that it may reduce MSN excitability associated with bursts of action potentials. To test this, we performed whole-cell patch-clamp recording from MSNs in acute brain slices. Relative to controls, MSNs from mice with astrocyte hPMCA2w/b displayed significantly hyperpolarized resting membrane potentials (~1.6 mV; Figure 3J), higher rheobase and right-shifted input-output curves (Figures 3H–3J), and

decreased numbers of action potential bursts from up-state-like membrane potentials (Figure 3J; $n = 32$ –40 cells from 11–12 mice). Thus, reducing astrocyte Ca^{2+} signaling produced significant effects on MSN activity *in vivo*, which we interpret to reflect reduced MSN excitability (Figures 3A–3J).

We also measured MSN activity in parallel with self-grooming (Figures 3K–3Q; Table S1; $n = 8$ and 11 mice). We found that in control and hPMCA2w/b mice, a similar ~30% fraction of imaged MSNs defined an ensemble active during grooming (Figures 3K and 3L; $p = 0.6574$). In control mice, however, the frequency of MSN activity in these neurons was lower during grooming (Figures 3K and 3M; Table S1), but the event areas were larger (Figure 3M; Table S1). In hPMCA2w/b mice, the frequency of MSN activity was initially higher during non-grooming episodes than in control mice and decreased significantly further during grooming episodes (Figures 3L and 3M; Table S1). Moreover, the event area did not increase during grooming episodes (Figure 3M; Table S1; $p > 0.05$), likely because large-amplitude and long-duration events were markedly reduced (Figure 3F; Table S1). These data establish that a pattern of MSN activity associated specifically with grooming episodes was altered significantly by reducing astrocyte Ca^{2+} signals. In particular, self-grooming episodes were associated with decreased frequency of MSN activity, and this drop in frequency was greater when astrocyte Ca^{2+} signaling was attenuated (Figures 3K–3M; Table S1). The activity of neurons that were active only during grooming is shown in Figure 3N. Next, we evaluated the degree of correlation between active MSNs and found a significantly greater correlation in hPMCA2w/b mice relative to controls irrespective of behavior (Figures 3O and 3P; Table S1; $n = 5$ and 8 mice). Greater correlated activity was also significant for non-grooming behaviors but was equivalently low during grooming (Figure 3Q; Table S1). Thus, a predominant phenotype by reducing astrocyte Ca^{2+} -dependent signaling in the striatum was greater correlated activity across MSNs during non-grooming episodes (Figure 3Q; Table S1). The resting correlation coefficient of ~0.04 observed between ~80 MSNs ($400 \times 400 \mu\text{m}$; Figure 2A) is comparable with silicon probe recordings at ~0.01 (Bakhurin et al., 2016).

(D) Average traces of 30 bouts of MSN activity measured as Ca^{2+} signals from control and hPMCA2w/b mice.

(E) Frequency of MSN activity.

(F) Scatterplot showing the relationship between peak amplitude and half-width of MSN Ca^{2+} signals. Peak amplitude (i) and half-width (ii) were significantly reduced in hPMCA2w/b mice.

(G) Cumulative probability of integrated area of Ca^{2+} events in MSNs from control and hPMCA2w/b mice. Inset: integrated area plot.

(H) Traces of MSN membrane responses to current injection in control and hPMCA2w/b mice.

(I) Relationship of injected current to number of APs.

(J) MSN resting membrane potentials (V_m) and the number of APs at up state V_m values. The p values are from Mann-Whitney tests.

(K) Left: raster plots of Ca^{2+} events with corresponding behaviors from 20 neurons in control mice. Bottom: event frequency of all neurons in single mice with 1 s time bins (76 neurons). Right: pie chart shows the percentage of imaged neurons active during grooming.

(L) Left: raster plots of Ca^{2+} events with corresponding behaviors from 20 neurons in hPMCA2w/b mice. Bottom: event frequency of all recorded neurons in single mice with 1 s time bins (72 neurons). Right: pie chart shows the percentage of imaged neurons active during grooming.

(M) Frequency and integrated area of Ca^{2+} events from all neurons during non-grooming and grooming episodes.

(N) Frequency and integrated area of Ca^{2+} events during non-grooming and grooming episodes for the neurons active during grooming.

(O) Correlation matrices showing the correlation coefficients between Ca^{2+} events of MSN pairs in control and hPMCA2w/b-expressing mice.

(P) Average correlation coefficient of MSN pairs between all Ca^{2+} events was greater in hPMCA2w/b-expressing mice.

(Q) Average correlation coefficients of Ca^{2+} events during non-grooming and grooming episodes in control and hPMCA2w/b-expressing mice. Data were assessed with Mann-Whitney tests or Student's t tests as appropriate. The average data are shown as mean \pm SEM. In some cases, the SEM symbol is smaller than the symbol for the mean.

Astrocyte Signaling Controls Self-Grooming via Tonic GABA-Mediated Neuromodulation

The greater coordination of MSNs covering an anatomically large area ($\sim 160,000 \mu\text{m}^2$) in hPMCA2w/b mice (Figure 3) prompted evaluation of mechanisms. Since MSNs are GABAergic, we examined fast phasic inhibition in control mice and those with hPMCA2w/b. We recorded phasic action-potential-dependent spontaneous inhibitory postsynaptic currents (sIPSCs) (Figure 4A), action-potential-independent miniature inhibitory postsynaptic currents (mIPSCs) (Figure 4B; in 500 nM TTX), evoked IPSCs at different stimuli (Figure 4C), paired-pulse responses (Figures 4D and 4E), and 300 μM GABA puff-evoked responses (Figure 4F). hPMCA2w/b had no significant effect on these metrics (Figures 4A–4F; $p > 0.05$, unpaired Mann-Whitney or Student's t test). We also evaluated fast excitatory synaptic transmission onto MSNs and found that AMPA-receptor-mediated spontaneous excitatory postsynaptic currents (sEPSCs), AMPA-receptor-mediated evoked EPSCs, and NMDA-receptor-mediated evoked EPSCs onto MSNs were not significantly different in hPMCA2w/b versus control (Figures 4G–4J; $n = 12$ cells, 4 mice, unpaired Mann-Whitney or Student's t test). Furthermore, by applying the glycine site agonists and antagonists D-serine (10 μM) and DCKA (10 μM), respectively, we found that there were no differences between controls and hPMCA2w/b in D-serine-mediated regulation of NMDA-receptor-evoked EPSCs (Figure 4K; $n = 12$ cells, 4 mice, unpaired Mann-Whitney or Student's t test). Thus, significantly reducing Ca^{2+} -dependent signaling in striatal astrocytes did not affect fast inhibitory or excitatory synaptic transmission. We do not imply that fast synaptic transmission is not regulated by astrocytes in brain areas that we have not examined.

We found that attenuation of Ca^{2+} -dependent signaling did not alter astrocyte current waveforms in response to 10 mV voltage jumps from -140 to $+40$ mV (Figure 4L). Furthermore, membrane resistance (MOhms), capacitance (pF), macroscopic current-voltage relations, and the resting membrane potential (mV) were all not significantly different between control and hPMCA2w/b groups (Figures 4L–4N; $n = 14$ and 15 cells from 5 control and hPMCA2w/b mice). Insofar as these metrics frequently change during astrogliosis and disease (Tong et al., 2014), these data suggest that the astrocytes were healthy in hPMCA2w/b groups.

We next evaluated tonic inhibition of MSNs from brain slices expressing hPMCA2w/b in astrocytes (Figure 5A). GABA tonic inhibition was measured by applying bicuculline (BIC; 25 μM) and by recording the change in holding current at -60 mV (Figure 5A). Tonic inhibition was significantly smaller in MSNs from slices with reduced astrocyte Ca^{2+} signals (Figures 5A and 5B; $p = 0.002$, one-way ANOVA). This effect was not due to lower expression of GABA_A receptors, because synaptic (Figures 4A–4E) and extrasynaptic responses evoked by GABA puffs were unaffected by hPMCA2w/b (Figure 4F). Astrocytes are known to regulate ambient GABA levels, and astrocytes preferentially express GAT-3 GABA transporters (Chai et al., 2017; Zhang et al., 2014). We explored whether astrocyte hPMCA2w/b led to increased function of GAT-3, as more GABA uptake could result in reduced tonic inhibition (Figures 5A and 5B). We found that MSN tonic inhibition in hPMCA2w/b slices was rescued to control levels by 40 μM SNAP5114

(Figure 5B), a selective GAT-3 blocker. Moreover, immunohistochemistry (IHC) showed that hPMCA2w/b-expressing astrocytes had significantly more GAT-3 (Figure 5C; $n = 14$ –17 cells, $n = 4$ mice, $p = 0.009$, unpaired Mann-Whitney tests) and across whole fields of view (Figures S7A–S7C; $n = 4$ mice, $p = 0.023$), providing evidence for upregulation of GAT-3. In contrast, expression of Kir4.1 and S100 β was unaffected by hPMCA2w/b (Figures 5D and 5E). To further explore GAT-3, we performed western blots from whole-striatum membrane-protein preparations from control and hPMCA2w/b mice (Figure 5F; $n = 7$ mice). We found that attenuation of astrocyte Ca^{2+} signaling with hPMCA2w/b resulted in increased GAT-3 expression (relative to tubulin). However, the expression of GABA transporter GAT-1 was unaffected (Figure 5F).

By recording from D1 and D2 MSNs from *Drd1-cre* and *Adora2a-cre* mouse lines injected with AAV-FLEX-tdTomato, we found that the hPMCA2w/b-mediated loss of tonic inhibition was significant for D1, but not D2, MSNs (Figures 5G and 5H). However, it is important to note that the genetic background of the *Drd1*- and *Adora2a-cre* mice is not identical to wild-type mice. Hence, it is not possible to directly compare absolute amplitudes of responses between mice. Nonetheless, these data illustrate functional selectivity of reducing astrocyte Ca^{2+} -dependent signaling in striatum.

The finding that astrocyte Ca^{2+} signaling regulates tonic inhibition in a GAT-3-dependent manner (Figures 5A–5F) prompted us to explore whether excessive self-grooming in hPMCA2w/b mice (Figure 2) was driven by decreased GABA-mediated tonic inhibition. To test this possibility, we administered SNAP5114 (50 $\mu\text{mol/kg}$ i.p. once per day for 1 week) using published methods (Wu et al., 2014) to hPMCA2w/b mice and found that the excessive self-grooming duration phenotype was rescued when assessed ~ 2 hr after the last dose (Figures 5I and 5J; $n = 8$ mice each, $p > 0.05$, one-way ANOVA). Importantly, the duration of self-grooming was not altered, and there was no effect on the number of grooming bouts in either group (Figures 5I and 5J). Furthermore, the self-grooming phenotype observed in hPMCA2w/b mice was not affected by vehicle (Figures S7D and S7E; $n = 8$ mice).

Taken together, these data (Figures 1, 2, 3, 4, and 5) provide strong evidence that hPMCA2w/b delivered selectively to the dorsolateral striatum leads to excessive self-grooming in a GAT-3-dependent manner, implying that hPMCA2w/b has specific effects within the striatum. Furthermore, in control mice, we found that GAT-3 inhibition did not alter tonic inhibition, which was 7.7 ± 0.8 pA in vehicle and 5.6 ± 0.9 pA in 40 μM SNAP5114 (Figures S7F and S7G; $n = 11$ and 13 cells, 5 mice, $p > 0.05$). However, inhibition of neuronal GAT-1 did, and tonic inhibition was significantly larger at 24.7 ± 3.4 pA ($n = 8$ cells, 4 mice, $p < 0.0001$) in the presence of the GAT-1 blocker NNC711 (10 μM) and was not further altered by SNAP5114 (Figure S7G). These data show that tonic inhibition in the striatal microcircuitry is controlled by GAT-1. However, when astrocyte Ca^{2+} -dependent signaling was reduced, GAT-3 functional expression increased and also contributed (Figure 5). However, astrocyte overexpression of GAT-3 did not cause excessive self-grooming (Figures 5K–5M), indicating that GAT-3 is necessary, but not sufficient, for the phenotype.

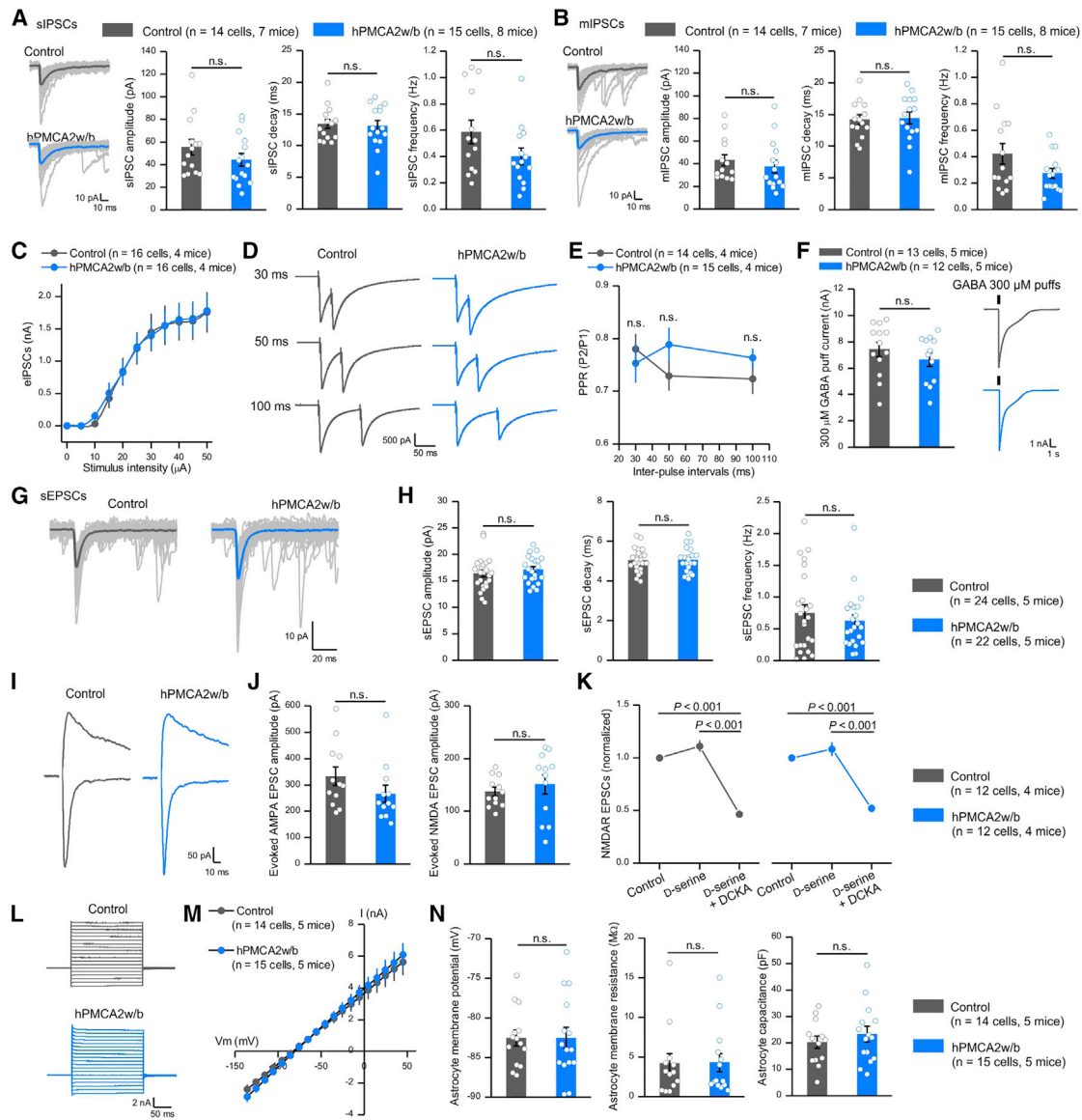


Figure 4. Reducing Striatal Astrocyte Ca^{2+} -Dependent Signaling *In Vivo* Did Not Affect Fast Synaptic Transmission or Astrocyte Properties

(A) Left: individual and average traces of 30 sIPSCs from control and hPMCA2w/b mice. Right: amplitude, decay time, and frequency of sIPSCs recorded from MSNs.

(B) Left: individual and average traces of 30 mIPSCs from control and hPMCA2w/b-expressing mice. Right: mIPSC amplitude, decay time, and frequency.

(C) Evoked IPSCs from control and hPMCA2w/b mice.

(D and E) Representative traces and average data for evoked IPSCs due to paired stimuli in control and hPMCA2w/b mice.

(F) Traces and average data for currents evoked by 300 μM GABA puffs for control and hPMCA2w/b. Data in (A)–(F) were assessed with Student's *t* or Mann-Whitney tests.

(G and H) Traces (G) and average data (H) for sEPSCs in control and hPMCA2w/b mice.

(I and J) Traces and average data for AMPA- and NMDA-receptor-mediated evoked EPSCs. AMPA EPSCs were measured at -70 mV, and NMDA EPSCs were measured at +40 mV, 50 ms after the stimulus.

(K) Normalized NMDA-receptor-mediated evoked EPSCs. Responses to D-serine and DCKA were indiscernible for control and hPMCA2w/b mice.

(L) Current waveforms from astrocytes during 10 mV voltage jumps from -140 mV to 40 mV.

(M) Average current-voltage relations.

(N) Average data for astrocyte membrane potential, membrane resistance, and capacitance for control and hPMCA2w/b groups. Data in (H), (J), and (N) were assessed by Student's *t* or Mann-Whitney tests as appropriate; data in (K) were assessed by two-way ANOVA tests. The average data are shown as mean ± SEM. In some cases, the bars used to represent SEM are smaller than the symbol for the mean.

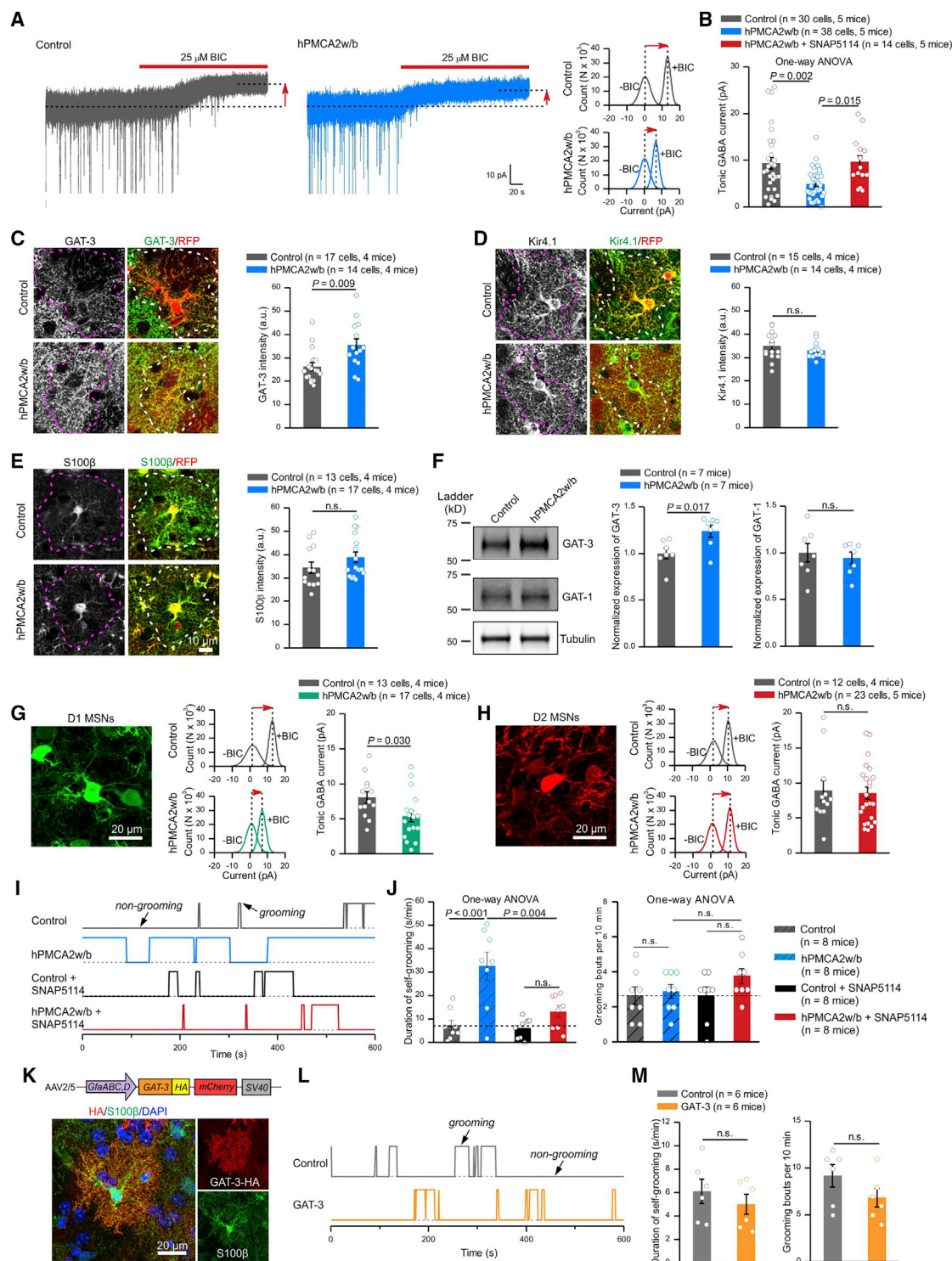


Figure 5. Reducing Striatal Astrocyte Ca^{2+} -Dependent Signaling *In Vivo* Altered MSN Tonic Inhibition, which Contributed to Excessive Self-Grooming

(A) Current recordings in voltage-clamp (−60 mV) from MSNs in control and hPMCA2w/b-expressing mice. Dashed lines and arrows indicate the changes of baseline induced by bicuculline application (BIC; 25 μ M). Histograms show the shift in baseline currents recorded during BIC application.

(B) Tonic GABA currents were reduced in MSNs from hPMCA2w/b-expressing mice, and this effect was rescued by GAT-3 inhibitor SNAP5114 (40 μ M).

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We next explored which aspect(s) of the *in vivo* MSN activity altered in hPMCA2w/b mice was rescued by SNAP5114 by repeating all the *in vivo* experiments in Figure 3 following SNAP5114 treatment (Table S1). Relative to controls, hPMCA2w/b altered eight aspects of MSN activity *in vivo* (Figure 3). They were (1) peak amplitude, (2) half-width and (3) correlation coefficient of MSN events analyzed independently of behavior, (4) frequency, (5) integrated area and (6) correlation coefficient of MSN events during non-grooming episodes, and (7) frequency and (8) integrated area of MSN events during grooming episodes (Figure 3; Table S1). Of these eight changes, five were fully rescued by SNAP5114 (Table S1; 1, 3, 4, 6, and 7, above). We suggest that these five MSN properties drive GAT-3-dependent excessive self-grooming. Of particular interest, the heightened correlated activity across all events (3) and during non-grooming episodes (6) was rescued by SNAP5114, which recalls the loss of tonic inhibition at a cellular level and its rescue by SNAP5114 (Figure 5). Parenthetically, our data show three alterations of MSN activity *in vivo* that were not rescued by SNAP5114 (2, 5, and 8, above), implying specificity of GAT-3 and additional circuit properties unrelated to self-grooming.

Astrocytic Contributions to Excessive Self-Grooming in Huntington's Disease Model Mice

Huntington's disease (HD) strongly affects the striatum and is associated with astrocyte dysfunction (Khakh et al., 2017). At early stages, HD patients display obsessive-compulsive symptoms (Beglinger et al., 2008) that are seen in R6/2 HD-model mice as excessive self-grooming (Steele et al., 2007). In light of our data (Figures 1, 2, 3, 4, 5), we explored whether reduced astrocyte Ca^{2+} signaling in R6/2 mice (Jiang et al., 2016) was accompanied by loss of MSN tonic inhibition and whether excessive self-grooming in R6/2 mice was GAT-3 dependent. We used the R6/2 mouse model, because most past studies on astrocytes have used this, and AAV reagents have only been validated in R6/2 mice, which model early-onset HD.

We found significantly reduced astrocyte Ca^{2+} signals in R6/2 mice relative to non-carrier controls (Figures 6A–6D). We also found that tonic inhibition onto MSNs was significantly reduced in R6/2 mice relative to controls (Figure 6E). Moreover, we observed excessive self-grooming in R6/2 mice, and this was relieved by SNAP5114 to control levels (Figures 6F and 6G). However, the excessive self-grooming phenotype in R6/2 mice

was not affected by vehicle (Figures S7H and S7I; $n = 8$ mice). In contrast to self-grooming, later-stage motor dysfunctions, such as the time taken to clasp limbs when suspended by the tail, forelimb grip strength, and walking stride, that are proposed to be associated with striatal tissue loss were not affected by SNAP5114 (Figures 8H–8L). Our data suggest that some HD phenotypes that occur during early stages, such as repetitive self-grooming, are correlated with reduced Ca^{2+} signaling and are corrected by GAT-3 blockers, recalling our data with wild-type mice and hPMCA2w/b.

Astrocyte-Specific RNA-Seq Following Ca^{2+} Signaling Attenuation *In Vivo*

We performed RNA sequencing (RNA-seq) to explore mechanisms that underlie GAT-3-dependent changes following attenuation of astrocyte Ca^{2+} signaling. First, we crossed astrocyte-specific *Aldh1l1*-cre/ERT2 BAC transgenic mice with RiboTag mice and used offspring that express Rpl22HA in astrocytes to determine how astrocyte gene expression changed following Ca^{2+} attenuation with hPMCA2w/b in relation to controls (Figure 7A). In this approach, we immunoprecipitated astrocyte-specific, actively translated RNA from the whole striatum following dorsolateral hPMCA2w/b microinjections (Figures 7A and 7E). In the second approach (Figures 7C and 7E), we used AAV2/5 GfaABC₁D to deliver a Rpl22-HA cDNA (Sanz et al., 2015) to the same astrocytes that received hPMCA2w/b (or control AAV) in the dorsolateral striatum (Figures S8A–S8D). In this approach, because genes delivered by two AAVs are co-expressed at the ~98% level (Figure S2B), we immunoprecipitated astrocyte-specific RNA mainly from astrocytes with attenuated Ca^{2+} signaling. By comparing RNA-seq data from RiboTag mice (i.e., astrocytes from whole striatum) with that from RiboTag AAV (i.e., astrocytes with attenuated Ca^{2+}), we sought to identify non-cell-autonomous and cell-autonomous changes following attenuation of astrocyte Ca^{2+} signaling. RNA-seq samples were replete with astrocyte markers and depleted of other cell markers for both RiboTag mice and RiboTag AAV data in both control and hPMCA2w/b samples (Figures 7B and 7D; $n = 4$ mice; Table S2).

We evaluated whether hPMCA2w/b caused astrogliosis by assessing the expression of pan-reactive, A1, and A2 markers (Liddel et al., 2017). None of these 37 markers were markedly elevated in hPMCA2w/b samples for either RiboTag mice or

(C) Left: images of GAT-3 IHC in astrocytes expressing control protein or hPMCA2w/b. Right: quantification of GAT-3 fluorescence intensity in hPMCA2w/b-expressing astrocytes relative to those expressing tdTomato (Mann-Whitney test).

(D and E) As in (C), but for Kir4.1 (D) and S100 β (E) immunostaining.

(F) Western blot analysis of striatum membrane protein fractions showing for GAT-3 and GAT-1 levels in hPMCA2w/b groups relative to controls. The gel shown has been cropped to fit in the figure, and the crop lines are shown as a solid border. The data were normalized to tubulin as a loading control.

(G and H) Tonic GABA inhibition was decreased in D1-MSNs (G), but not D2-MSNs (H), in hPMCA2w/b-expressing mice (assessed with Student's *t* or Mann-Whitney tests). The MSNs are colored but were from *Drd1*-cre and *Adora2a*-cre mice with FLEX AAV tdTomato injections.

(I) Behavioral traces of control and hPMCA2w/b-expressing mice under baseline and SNAP5114 treatment showing self-grooming and non-grooming episodes over 10 min.

(J) Increased self-grooming behaviors in hPMCA2w/b-expressing mice were rescued by SNAP5114 (50 $\mu\text{mol/kg}$; data were assessed with one-way ANOVA followed by post hoc Bonferroni tests).

(K) AAV2/5 overexpressing GAT-3 in astrocytes and GAT-3HA detection in S100 β -positive astrocytes.

(L) Traces for self-grooming.

(M) Average data for self-grooming duration and bouts for controls and GAT-3HA-overexpressing mice. Data were assessed with Student's *t* test. The average data are shown as mean \pm SEM.

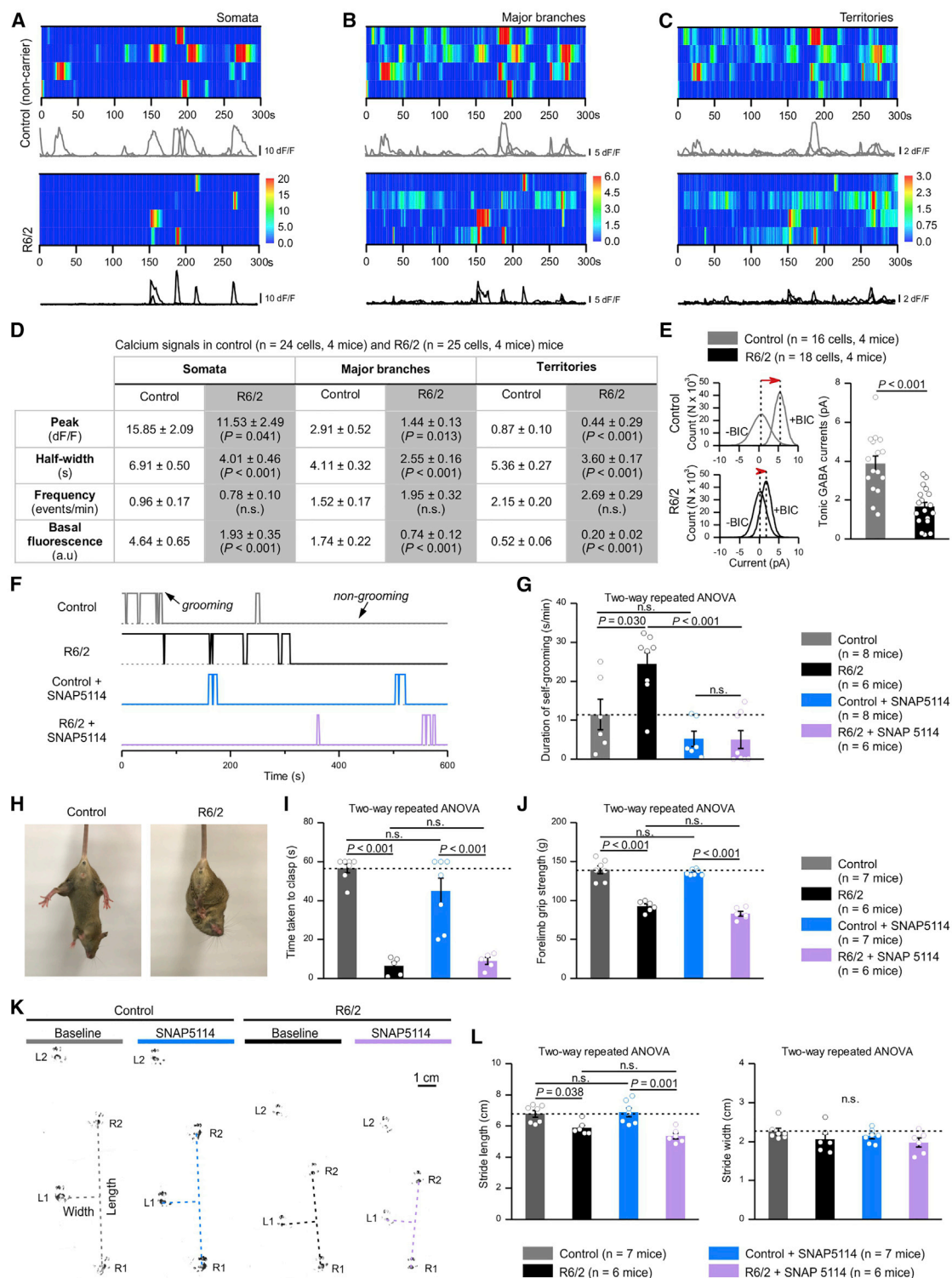


Figure 6. Excessive Self-Grooming in Huntington's Disease Model Mice, 7–8 weeks, Was Accompanied by Reduced Ca^{2+} Signaling and Was GAT-3 Dependent

(A–C) Kymographs and $\Delta F/F$ traces of Ca^{2+} signals in somata (A), major branches (B), and territories (C) of astrocytes in non-carrier control and R6/2 mice.

(D) The inset table summarizes Ca^{2+} imaging data for control and R6/2 mice.

(E) MSNs in R6/2 mice had significantly reduced tonic GABA currents (Student's *t* tests).

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RiboTag AAV, and the increase across the 37 markers was 1 ± 0.2 -fold and 1 ± 0.1 -fold for RiboTag mice and RiboTag AAV (Figure 7F; FDR > 0.05). However, 33/37 of these genes were increased when we used lipopolysaccharide (LPS) as a control to trigger neuroinflammation: the average increase across 37 markers was 16 ± 5 -fold (Figure 7F; some >40-fold). Furthermore, we analyzed the top 100 microglia markers and 23 activated microglia markers from the input RNA-seq (Figures S8E and S8F). For RiboTag mice and RiboTag AAV, 0/123 and 5/123 were altered, respectively, by hPMCA2w/b relative to control. The altered genes changed <2-fold and were *C3ar1*, *Lag3*, *Emr1*, *Lyz2*, and *Ccl3* (Figures S8E and S8F). In contrast, 52/123 genes were altered by LPS (range 1.5- to 16-fold). Our data provide strong evidence that reducing astrocyte Ca^{2+} signaling did not cause astrogliosis or overt microglial activation. The five microglial genes altered by astrocyte Ca^{2+} attenuation suggest microglial functions regulated by astrocytes.

Next, we measured astrocyte-enriched (>2-fold), differentially expressed genes (DEGs; Figure 7G; Figure S8G). We parsed the data by fragments per kilobase per million (FPKM) and significant changes induced by hPMCA2w/b for both RiboTag mice and RiboTag AAV datasets (Figure S8G). By assigning a FPKM of 5 for the level of expression as the cutoff, we detected 36 and 516 significantly DEGs in the RiboTag mice and RiboTag AAV datasets (FDR < 0.05). This fairly low number of DEGs suggests specificity for the functions served by astrocyte Ca^{2+} signaling within striatal astrocytes and supports data showing that attenuated astrocyte Ca^{2+} signaling did not alter astrocyte electrophysiological properties (Figures 4L–4N), marker expression (Figures 5D and 5E), or cause reactivity (Figure 7F).

Figure 7H reports the top 36 up- and downregulated genes for the RiboTag mice and RiboTag AAV datasets. Surprisingly, only five genes were shared. We interpret this to indicate that Ca^{2+} attenuation produced cell-autonomous (RiboTag AAV) and non-cell-autonomous effects on astrocyte gene expression (i.e., the difference between RiboTag AAV and RiboTag mice). This trend is summarized in Figure 7I for all DEGs (FPKM > 5, FDR < 0.05). Remarkably, these analyses show that of the 36 genes altered throughout the striatum, 25 are not altered within the astrocytes that received hPMCA2w/b. We propose that these genes define Ca^{2+} -dependent functions that are regulated in astrocyte-astrocyte networks. This is an unexpected finding from our work. Data in Figure 7 provide a rich resource for exploring Ca^{2+} -dependent astrocyte physiology in the cells that had attenuated Ca^{2+} signals and in those they impact in the striatum.

In order to shed light on the mechanisms that underlie the physiological changes in Figures 1, 2, 3, 4, and 5, we used weighted gene co-expression network analysis (WGCNA) of

RiboTag mice and RiboTag AAV RNA-seq data. WGCNA is a systems biology method for describing the correlation patterns among genes across data-rich samples, which can be used for identifying and interpreting clusters (modules) of highly co-expressed genes. WGCNA revealed three modules in the RiboTag mice data, but none of these differed between control and hPMCA2w/b groups (Figures 8A and 8B). In the RiboTag AAV data, we identified 16 modules (Figures 8C and 8D), some of which (e.g., green, salmon, cyan, and pink) showed differences between control and hPMCA2w/b. Of note, the green module comprised a set of transcripts involved in membrane protein trafficking, Golgi vesicle transport, and protein stabilization (Figure 8D; pink module shown for comparison). Upon closer examination, one downregulated gene belonging to the green module was *Rab11a*. This is notable because *Rab11a* regulates GAT-3 trafficking: a decrease in *Rab11a* increases cell-surface GAT-3 expression by changing the balance between forward trafficking and removal (Zhang et al., 2017). These data suggest that GAT-3 functional upregulation that underlies cellular, neuronal, and behavioral effects (Figures 1, 2, 3, 4, and 5) involves trafficking. Consistent with this, we found no change in GAT-3 (*Slc6a11*) or GAT-1 (*Slc6a1*) FPKM levels (Figure 8E), whereas we detected significantly increased GAT-3 expression by immunostaining (Figure 5C; Figures S7A–S7C), by western blot of membrane protein fractions (Figure 5F), and by measurements of tonic GABA currents (Figures 5A and 5B). Furthermore, the FPKM values for *Rab11a* were lower in hPMCA2w/b groups relative to controls for both RiboTag mice and RiboTag AAV groups (Figure 8F). To assess this rigorously, Figure 8G shows analyses from the DEGs with FDR < 0.1 for *Rab11a*, *Slc6a11*, and two other exemplar kinase genes involved in posttranslational regulation of GABA transporters (*Ddr2* and *Tyk2*). We found no significant change in *Slc6a11* expression, increased *Ddr2* and *Tyk2* expression, and significantly decreased expression of *Rab11a* in hPMCA2w/b relative to controls in both RiboTag mice and RiboTag AAV data (Figure 8G). Table S2 reports expression data for other kinases, phosphatases, and other proteins implicated in GABA transporter regulation. Taken together, our unbiased gene expression survey supports the notion that a posttranslational mechanism is involved in elevated GAT-3 functional expression following attenuation of astrocyte Ca^{2+} signaling and involves altered trafficking caused by the downregulation of *Rab11a*, without a change in *Slc6a11* expression.

DISCUSSION

There are four main findings from this work. First, hPMCA2w/b reduced Ca^{2+} signals in astrocyte somata, branches, and

(F) Behavioral traces of non-carrier control and R6/2 mice under baseline and during SNAP5114 treatment (50 $\mu\text{mol/kg}$) showing self-grooming and non-grooming episodes over 10 min.

(G) R6/2 mice exhibited excessive self-grooming, which were rescued by SNAP5114 treatment (two-way ANOVA followed by a post hoc Bonferroni test).

(H) Forelimb claspings of a R6/2 mouse and a control mouse.

(I) Average time taken to clasp forelimbs was significantly less in R6/2 mice, and SNAP5114 treatment did not improve this.

(J) Forelimb grip strength of R6/2 mice was smaller than controls. SNAP5114 treatment did not change this.

(K) Representative footprint tracks of control and R6/2 mice before and after SNAP5114 treatment. Dotted lines indicate stride lengths and widths.

(L) Average footprint stride length and width of control and R6/2 mice before and after SNAP5114 treatment. The p values were from two-way ANOVA with a post hoc Bonferroni test. The average data are shown as mean \pm SEM.

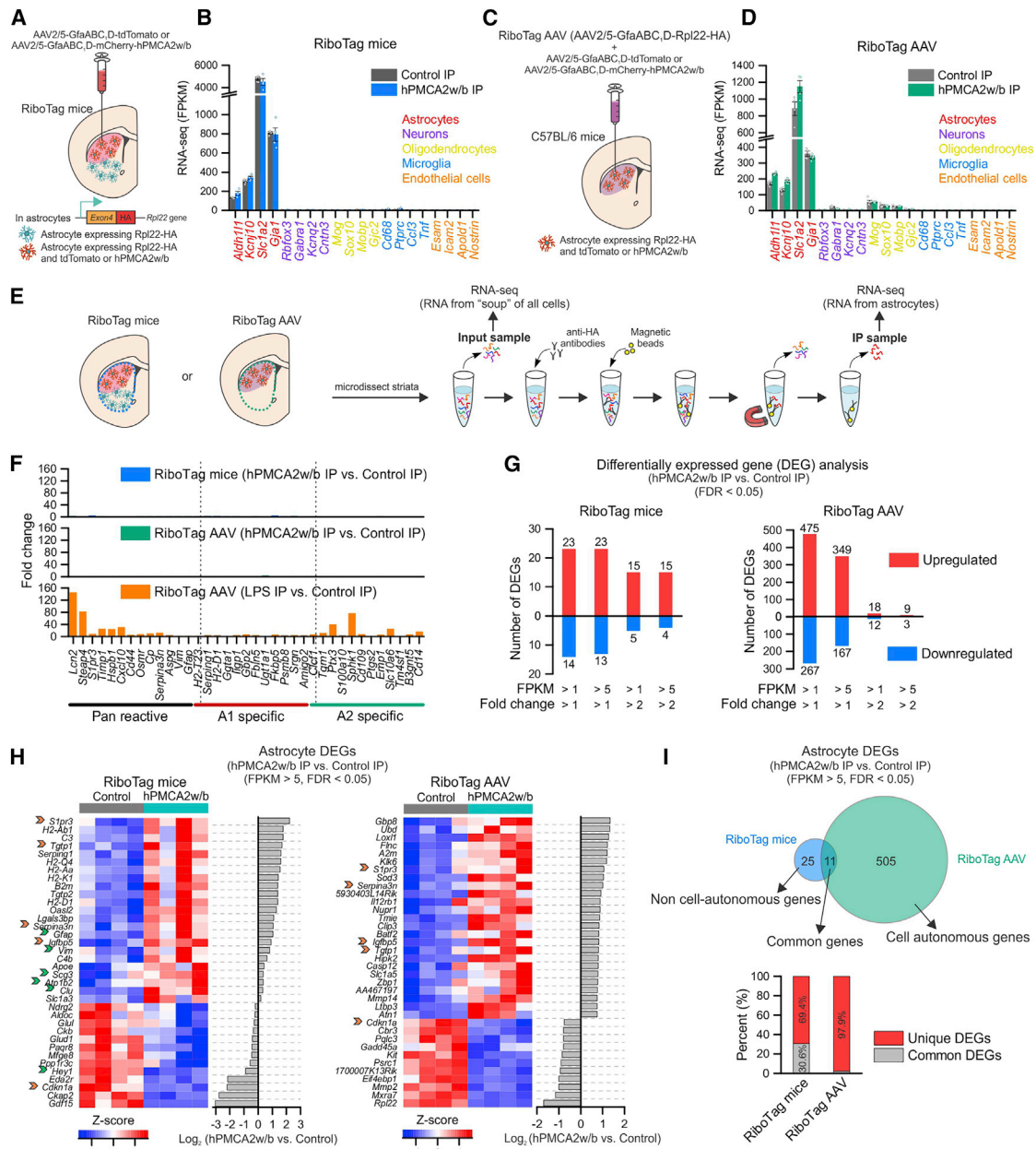


Figure 7. Ca^{2+} Attenuation by hPMCA2w/b Changed Gene Expression in Astrocytes

(A) RiboTag mice were crossed with *Aldh1l1*-cre/ERT2 mice to generate offspring expressing Rpl22HA in astrocytes (green cells in the cartoon). Control or hPMCA2w/b AAVs were injected into the dorsolateral striatum to target astrocytes (red cells). Astrocyte RNA from the whole striatal tissue was extracted; sequencing of IP and input material was performed ("RiboTag mice").

(B) Expression levels (in fragments per kilobase per million, FPKM) of cell-specific markers from RiboTag mice IP samples (n = 4 mice).

(C) C57BL/6 mice received AAVs for Rpl22HA ("RiboTag AAV") together with control or hPMCA2w/b AAVs in the dorsolateral striatum (red cells). RNA from whole striatal tissue was extracted for IP of Rpl22HA; sequencing of IP and input material was performed ("RiboTag AAV").

(D) Expression levels of cell-specific markers in RiboTag AAV IP samples (n = 4 mice).

(E) Schematic of the protocol used.

(F) Fold-change of pan-reactive, A1-, and A2-astrocyte-specific markers between hPMCA2w/b IP and control IP samples in RiboTag mice (top) and RiboTag AAV (middle). As a positive control, i.p. injection of LPS (5 mg/kg) was used.

(G) Differentially expressed gene (DEG) analysis (limma/voom, FDR < 0.05) showing the numbers of up- and downregulated astrocyte genes between hPMCA2w/b IP and control IP in RiboTag mice and RiboTag AAV datasets. These analyses were restricted to genes with >2-fold enrichment in the IP compared with the input, and the plots show different threshold criteria.

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processes, which resulted in GAT-3-dependent excessive self-grooming in mice. Second, reducing striatal astrocyte Ca^{2+} signals reduced tonic inhibition of striatal MSNs in a GAT-3-dependent manner and altered microcircuit activity *in vivo*. Third, at early stages in HD-model mice, striatal astrocyte Ca^{2+} signaling was reduced and was associated with reduced MSN tonic inhibition and excessive self-grooming, which was also GAT-3 dependent. Fourth, two evaluations of astrocyte gene expression revealed molecular changes within astrocytes following attenuation of Ca^{2+} signaling, including of *Rab11a*, a known regulator of GAT-3 functional expression (Zhang et al., 2017). The main findings are schematized in Figures 8H–8J.

The magnitude of striatal astrocyte Ca^{2+} signaling attenuation produced by hPMCA2w/b was smaller than that observed with IP3R2 deletion mice. However, using hPMCA2w/b, we observed clear cellular and behavioral phenotypes. We interpret this to indicate that astrocyte functions within circuits are best explored by performing circuit-specific astrocyte manipulations in the adult brain rather than across the whole brain and throughout development as with IP3R2 deletion mice. This is analogous to the use of optogenetics and pharmacogenetics aimed at exploring neuronal function within anatomically selected areas. By manipulating astrocytes in specific brain areas, it should be possible to determine whether astrocyte function and/or dysfunction contributes to behaviors and disease-related phenotypes encoded by the local neurons. Once such behaviors are discovered, the underlying astrocyte-related molecular, cellular, and circuit mechanisms could be teased apart. By performing such top-down studies for the dorsolateral striatum, our studies emphasize the importance of tonic GABA rather than direct effects of astrocytes on fast neurotransmitter release. Astrocytic regulation of GABA is considered a type of “gatekeeper” within circuits (Beenhakker and Huguenard, 2010; Muthukumar et al., 2014).

Although we focused on the striatum and self-grooming, hPMCA2w/b could be used in other brain areas in order to explore the functions of astrocyte Ca^{2+} signaling. We do not imply that self-grooming is the only or the most important outcome of astrocyte Ca^{2+} signaling in the brain. Hence, systematic use of hPMCA2w/b constructs may reveal additional functions of astrocyte Ca^{2+} signaling *in vivo*. Such functions could include roles at quad-partite synapses (Schafer et al., 2013), release of trophic factors, phagocytosis, homeostasis, neurovascular coupling, and responses to injury, disease, trauma, and infection. hPMCA2w/b could also be improved: the current strategy uses AAV2/5-mediated expression that requires ~14–21 days. It may be possible to deliver the protein more expeditiously using mouse genetics and to engineer it so it can be activated by ligand.

In agreement with past work (Aguilhon et al., 2010; Petravic et al., 2008), we found that attenuation of astrocyte Ca^{2+} signaling did not affect striatal fast synaptic transmission. There are two interpretations of this. First, it is conceivable that the ~70%–80% attenuation of Ca^{2+} signals produced by hPMCA2w/b was insufficient to reveal effects on fast synapses. Second, the result may indicate that astrocyte Ca^{2+} signaling does not in fact regulate fast excitation and inhibition in the striatum. On balance, we favor the latter view, which is consistent with the paucity of astrocyte Ca^{2+} -dependent glutamate release (Sloan and Barres, 2014), including from striatal astrocytes (Chai et al., 2017). Furthermore, the bulk of the evidence to suggest that astrocytes regulate fast synaptic transmission has been gathered with the use of Ca^{2+} chelators such as BAPTA, the validity of which has been questioned (Smith et al., 2018). We do not rule out astrocyte contributions to fast synaptic transmission in brain areas we have not studied.

By using *in vivo* imaging in freely behaving mice, we identified eight alterations of MSN activity that were caused by hPMCA2w/b expression within astrocytes *in vivo*. Of these eight, five were reversed by SNAP5114, which also rescued the self-grooming phenotype and the loss of tonic inhibition triggered by reducing astrocyte Ca^{2+} signals (Figure 8J). These data provide a strong correlation at different biological levels (from molecules to behavior) for a mechanism whereby reduced astrocyte Ca^{2+} signaling resulted in elevated functional activity of GAT-3, reduced ambient GABA, which regulated MSN activity *in vivo* and self-grooming (Figure 8J). Of the five GAT-3-dependent changes, two merit discussion. First, we recorded increased correlated activity between MSNs during non-grooming episodes following hPMCA2w/b expression in astrocytes: this effect was reversed by SNAP5114. We hypothesize that heightened correlated activity during non-grooming acts on downstream projection areas regulating self-grooming and results from reduced tonic inhibition. Second, we recorded a larger drop in the frequency of MSN activity during self-grooming in hPMCA2w/b mice, and this too was reversed by SNAP5114. These data suggest that specific GAT-3-dependent patterns of MSN activity are needed during self-grooming episodes in hPMCA2w/b mice. There is no circuit level model of self-grooming that can be used to model striatal astrocyte data. However, consistent with our data, there is good evidence that the dorsolateral striatum is involved in the execution of sequential patterns related to self-grooming (Kalueff et al., 2016). Furthermore, pharmacologically increasing or decreasing GABA decreases and increases grooming, respectively (Kalueff et al., 2016). Additionally, very precise lesions to the dorsolateral striatum result in the failure to complete self-grooming syntax (Cromwell and Berridge, 1996). This phenotype is separable from the prolonged self-grooming we observed but consistent with our data showing

(H) Heatmaps of the top 36 astrocyte DEGs as ranked by differential expression \log_2 ratio between hPMCA2w/b IP and control IP (FPKM > 5). Orange arrows indicate genes that were common in the top 36 DEGs for both RiboTag mice and RiboTag AAV. Green arrows indicate additional DEGs in RiboTag mice that were also astrocyte DEGs in RiboTag AAV, but not in the top 36 for the latter.

(I) Top: Venn diagram representing DEGs (FPKM > 5) in astrocytes from RiboTag mice and RiboTag AAV revealing putative cell-autonomous genes and non-cell-autonomous genes. Bottom: percent of unique astrocyte DEGs and common astrocyte DEGs between RiboTag mice and RiboTag AAV. Average data are shown as mean \pm SEM. All RNA-seq data were for four separate replicates.

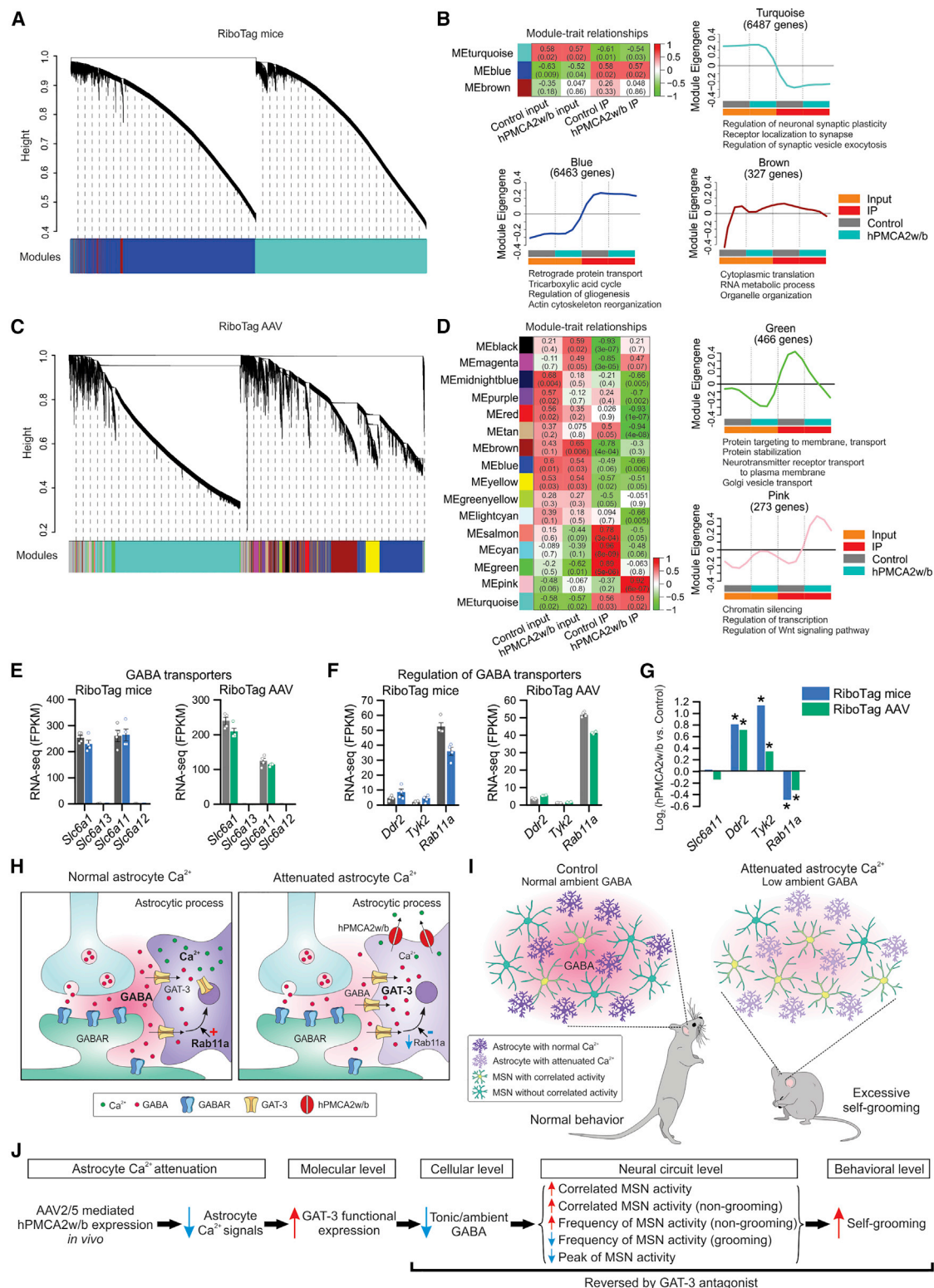


Figure 8. Weighted Gene Co-expression Network Analysis Identified Distinct Modules of Co-expressed Genes Associated with Attenuated Astrocyte Ca²⁺ Signaling

(A) Hierarchical cluster dendrogram with colors underneath denoting three distinct modules of co-expressed transcripts in RiboTag mice RNA-seq data. (B) Heatmap showing module-trait relationships, with each row representing a module. Correlation coefficients between a module eigengene (ME) and trait (coded from -1 to 1) and corresponding p values are shown in each cell. The color indicates the level of correlation (positive correlation in red and negative correlation in blue) (legend continued on next page)

little astrocyte reactivity that would have occurred in the event of an overt injury-related response. A satisfying microcircuit level model for astrocyte regulation of self-grooming is not yet possible, but our study provides an *in vivo* experimental platform and rationale to explore an adult innate vertebrate behavior regulated by astrocytes located within an anatomically important circuitry providing syntax for sequential motor tasks. Our data also show three alterations of MSN activity *in vivo* that were not rescued by SNAP5114, implying specificity of the GAT-3- and astrocyte-related mechanisms as well as the existence of additional circuit properties unrelated to GAT-3.

Excessive self-grooming has also been observed in *Hoxb8* null mutant mice that are proposed to alter neural circuits via microglia (Chen et al., 2010). We did not detect expression of *Hoxb8* in astrocytes or the input samples in control or hPMCA2w/b datasets (Table S2). Hence, *Hoxb8*-related mechanisms are unrelated to our experiments. Excessive self-grooming has been observed in adolescent mice lacking the glutamate transporter GLT1 (Aida et al., 2015). These studies are tangential to our study as we did not delete GLT1, and GLT1 expression did not decrease following Ca^{2+} attenuation (Table S2).

At a molecular level, we found that attenuating astrocyte Ca^{2+} signaling increased GAT-3 functional expression and protein levels. However, RNA-seq showed no change in GAT-3 levels, suggesting that astrocyte Ca^{2+} signaling regulates GAT-3 in a posttranslational way. In accord, WGCNA and DEG analyses identified *Rab11a* as a significantly downregulated gene. Downregulation of *Rab11a* leads to functional upregulation of GAT-3 (Zhang et al., 2017), which supports our observations. In addition, the RNA-seq data provide a rich resource of gene expression changes and WGCNA modules that can be used in future, hypothesis-driven experiments aimed at evaluating astrocyte Ca^{2+} signaling. Pursuing such agnostically derived mechanisms complements strategies exploring other mechanisms such as gliotransmission. It should also be possible to make floxed *Rab11a* allele mice and image *Rab11a* regulation of GAT-3 in reduced preparations. Overall, RNA-seq data provide a valuable resource to design hypothesis-driven experiments to explore astrocyte Ca^{2+} signaling—a significant issue that has been difficult to address.

We have used several approaches to study the adult striatum as a model area to explore how astrocytes contribute to vertebrate microcircuit function (Chai et al., 2017; Jiang et al., 2016; Khakh et al., 2017; Oceau et al., 2018; Srinivasan et al., 2016; Tong et al., 2014). In the present study, we found that striatal

astrocyte Ca^{2+} -dependent signaling affects microcircuit function in brain slices and *in vivo* and that reducing such signaling triggers innate behavioral phenotypes in mice via altered neuromodulatory GABA signaling. Our data show that astrocytes regulate striatal MSNs and a specific behavior via ambient GABA-mediated neuromodulation. Moreover, we show that both experimentally induced and disease-model-associated reductions in striatal astrocyte Ca^{2+} -dependent signaling in adult mice can produce abnormal repetitive behavioral phenotypes that are related to psychiatric disorders (Kalueff et al., 2016).

STAR★METHODS

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- QUANTIFICATION AND STATISTICAL ANALYSIS
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SUPPLEMENTAL INFORMATION

Supplemental Information includes eight figures, two tables, and three videos and can be found with this article online at <https://doi.org/10.1016/j.neuron.2018.08.015>.

correlation in green). Individual plots showing the expression of the module eigengene across sample categories with significantly enriched terms for biological processes revealed by gene ontology enrichment analysis (FDR < 0.05).

(C) Cluster dendrogram identified 16 co-expressed WGCNA modules for RiboTag AAV RNA-seq data.

(D) Module-trait relationships revealed modules correlated with hPMCA2w/b expression in striatal astrocytes. Green module was composed of genes regulating protein transport, which were downregulated in striatal astrocytes expressing hPMCA2w/b. Pink module was enriched in genes regulating chromatin silencing and transcription, which were upregulated in striatal astrocytes expressing hPMCA2w/b.

(E) Expression levels of GABA transporters: *Slc6a1* (GAT-1), *Slc6a13* (GAT-2), *Slc6a11* (GAT-3), and *Slc6a12* (GAT-4) in RiboTag mice and RiboTag AAV RNA-seq datasets.

(F) Expression levels of potential GABA transporter regulators suggested by previous work in RiboTag mice and RiboTag AAV RNA-seq data.

(G) Changes in gene expression of striatal astrocytes expressing hPMCA2w/b, expressed as \log_2 (fold change), in RiboTag mice and RiboTag AAV RNA-seq data. Asterisk (*) indicates differential expression between hPMCA2w/b IP and control IP using edgeR analysis on combined RNA-seq datasets (FDR < 0.1).

(H and I) Cartoon summary of the main findings at synaptic (H) and *in vivo* (I) levels.

(J) Descriptive model based on our work (see Discussion). Average data are shown as mean \pm SEM from four mice in each group.

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AUTHOR CONTRIBUTIONS

X.Y. performed molecular biology, imaging, electrophysiology, and MINIscope experiments. X.Y. and A.M.W.T. performed and analyzed mouse behavior experiments. P.G. and C.J.E. provided access to equipment. J.N. assessed excitatory synaptic transmission. G.C. directed RNA-seq data analyses/bioinformatics. B.S.K. conceived, designed, and directed the project, helped with data analyses, and performed some electrophysiology. B.S.K. and X.Y. made the figures and wrote the paper; others commented.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Rabbit anti-S100 β	Abcam	Cat#ab41548; RRID: AB_956280
Mouse anti-NeuN (clone A60)	Millipore	Cat#MAB377; RRID: AB_2298772
Chicken anti-GFP	Abcam	Cat#ab13970; RRID: AB_300798
Rabbit anti-RFP	Rochland	Cat#600-401-379; RRID: AB_2209751
Chicken anti-RFP	Brecha Lab, University of California at Los Angeles	N/A
Mouse anti-HA 11 antibody (clone 16B12)	Covance	Cat#MMS-101R; RRID: AB_291263
Rabbit anti-GAT-3	Brecha Lab, University of California at Los Angeles	Cat# 374D; RRID: AB_2313749
Rabbit anti-GAT1	Abcam	Cat# ab426; RRID: AB_2189971
Rabbit anti-Kir4.1	Alomone	Cat#APC-035; RRID: AB_2040120
Rabbit anti-DARPP-32	Abcam	Cat#ab40801; RRID: AB_731843
Mouse anti- β -tubulin	Millipore	Cat# 05-559; RRID: AB_309804
Alexa Fluor 488 goat anti-chicken	Molecular Probes	Cat#A11039; RRID: AB_2534096
Alexa Fluor 488 goat anti-rabbit	Molecular Probes	Cat#A11008; RRID: AB_143165
Alexa Fluor 546 goat anti-mouse	Molecular Probes	Cat#A11003; RRID: AB_2534071
Alexa Fluor 546 goat anti-chicken	Molecular Probes	Cat#A11040; RRID: AB_2534097
Alexa Fluor 594 goat anti-rabbit	Molecular Probes	Cat# R37117; RRID: AB_2556545
IRDye 800CW anti-rabbit	LI-COR	Cat#827-08365; RRID: AB_10796098
IRDye 680RD anti-mouse	LI-COR	Cat#926-68170; RRID: AB_10956589
Bacterial and Virus Strains		
pcDNA3.1 mCherry-hPMCA2w/b	This study: Addgene	111570
pcDNA3.1 mCherry-hPMCA2x/b	This study: Addgene	111571
pZac2.1 GfaABC ₁ D mCherry-hPMCA2w/b	This study: Addgene	111568
pZac2.1 GfaABC ₁ D mCherry-hPMCA2x/b	This study: Addgene	111569
pZac2.1 GfaABC ₁ D Rpl22HA	This study: Addgene	111811
AAV5 GfaABC ₁ D cyto-GCaMP6f	Haustein et al., 2014 ; UPenn Vector Core Addgene Vectors	Cat#AV-5-52925 #52925-AAV5
AAV5 GfaABC ₁ D tdTomato	Tong et al., 2014 ; UPenn Vector Core Addgene Vectors	Cat#AV-5-PV3106 44332-AAV5
AAV2/5 GfaABC ₁ D Rpl22HA	This study: UPenn Vector Core	In process
AAV2/5 GfaABC ₁ D mCherry-hPMCA2w/b	This study: UPenn Vector Core	In process
AAV2/5 GfaABC ₁ D mCherry-hPMCA2x/b	This study: UPenn Vector Core	In process
AAV1 Synapsin GCaMP6f	UPenn Vector Core	Cat#AV-1-PV2822
AAV1 CAG-Flex tdTomato	UPenn Vector Core	Cat#AV-1-PV2816
Chemicals, Peptides, and Recombinant Proteins		
Paraformaldehyde, EM grade	EMS	Cat#19202
TTX	Cayman Chemical Company	Cat#14964
Bicuculline	Sigma-Aldrich	Cat#14340
CNQX disodium salt	Abcam	Cat#ab120044
Cyclopiazonic acid (CPA)	Tocris	Cat#1235
Clozapine N-oxide (CNO)	Tocris	Cat#4936
Phenylephrine	Tocris	Cat#2838

(Continued on next page)

Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Papain	Worthington	Cat#LS003126
Ovomucoid trypsin inhibitor	Worthington	Cat#LS003086
EBSS	Sigma-Aldrich	Cat#E7510
Halt protease inhibitor cocktail	Thermo Fisher Scientific	Cat#1861278
Critical Commercial Assays		
Ovation PicoSL WTA System v.2	Nugen	Cat#3312
QIAquick PCR Purification Kit	QIAGEN	Cat#28104
QIAGEN RNeasy Plus Micro Kit	QIAGEN	Cat#74034
Pierce BCA protein assay kit	Thermo Scientific	Cat#23225
Deposited Data		
RNA-seq data	This paper	GEO: GSE114757, GSE114756
Experimental Models: Cell lines		
Human: HEK293 cell line; sex: female	ATCC	Cat# PTA-4488; RRID: CVCL_0045
Experimental Models: Organisms/Strains		
Mouse: <i>Aldh1l1</i> -eGFP	MMRRC	Stock#011015-UCD; RRID: MMRRC_011015-UCD
Mouse: B6N.129-Rp122 ^{tm1.1Psam} /J	Jackson Laboratory	Stock#011029; RRID: IMSR_JAX:011029
Mouse: <i>Aldh1l1</i> -cre/ERT2	Jackson Laboratory	Stock#029655; RRID: IMSR_JAX:029655
Mouse: C57BL/6NTac inbred mice	Taconic	Stock#B6; RRID: IMSR_TAC:b6
Mouse: B6.FVB(Cg)-Tg(Drd1-cre)EY262Gsat/Mmucd	MMRRC	Stock# 030989-UCD; RRID: MMRRC_030989-UCD
Mouse: B6.FVB(Cg)-Tg(Adora2a-cre)KG139Gsat/Mmucd	MMRRC	Stock# 036158-UCD; RRID: MMRRC_036158-UCD
Mouse: B6CBA-Tg(HDexon1)62Gpb/3J	Jackson Laboratory	Stock#006494; RRID: IMSR_JAX:006494
<i>lpr3r2</i> knockout (<i>lpr2</i> ^{tm1.1Chen})	MGI	Cat# 3641042; RRID: MGI:3641042
Oligonucleotides		
qPCR <i>Atp2b1</i> forward primer sequence: 5' TTCCGGAACCAAGCCATC 3'	This paper	N/A
qPCR <i>Atp2b1</i> reverse primer sequence: 5' AGAAACCCCAACAAGGCACA 3'	This paper	N/A
qPCR <i>Atp2b2</i> forward primer sequence: 5' TCGAAGACCCAGTACGACCT 3'	This paper	N/A
qPCR <i>Atp2b2</i> reverse primer sequence: 5' CCCTTCTCGTTGCGAATCCT 3'	This paper	N/A
qPCR <i>Atp2b3</i> forward primer sequence: 5' TTCTGGTTCACGCCATCTCC 3'	This paper	N/A
qPCR <i>Atp2b3</i> reverse primer sequence: 5' TATCTGCTCCCGTACTGGCT 3'	This paper	N/A
qPCR <i>Atp2b4</i> forward primer sequence: 5' TACATCGGGGGCACTCACTA 3'	This paper	N/A
qPCR <i>Atp2b4</i> reverse primer sequence: 5' CAGCCACACTCTGTCTTGT 3'	This paper	N/A
qPCR <i>Arbp</i> forward primer sequence: 5' TCCAGGCTTTGGGCATCA 3'	Jiang et al., 2016	N/A
qPCR <i>Arbp</i> reverse primer sequence: 5' AGTCTTTATCAGCTGCACATCAC 3'	Jiang et al., 2016	N/A
Software and Algorithms		
OriginPro 8	Origin Lab Corporation	RRID: SCR_015636
SciScan 2014	Scientifica	https://www.scientifica.uk.com/products/scientifica-sciscan

(Continued on next page)

Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
pCLAMP10.4	Molecular Devices	RRID: SCR_011323
ClampFit10.4	Molecular Devices	N/A
Fluoview FV3000	Olympus	N/A
ImageJ v.1.51h	NIH	RRID: SCR_003070
MiniAnalysis 6.0.7	Synaptosoft	RRID: SCR_002184
MATLAB	MathWorks	RRID: SCR_001622
R v.3.3.2	R Core Team, 2016	https://cran.r-project.org/bin/windows/base/old/3.2.2/
Ethovision XT	Noldus Information Technology	RRID: SCR_000441
CorelDraw12	Corel Corporation	RRID: SCR_014235
Bioconductor	Bioconductor	http://www.bioconductor.org
Htseq-count	Anders et al., 2015	N/A

CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Baljit S. Khakh (bkhakh@mednet.ucla.edu).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Mouse Models

All animal experiments were conducted in accordance with the National Institute of Health Guide for the Care and Use of Laboratory Animals and were approved by the Chancellor's Animal Research Committee at the University of California, Los Angeles. Male and female mice aged between 6- and 14-week old were used in this study. Mice were housed in the vivarium managed by the Division of Laboratory Animal Medicine at University of California, Los Angeles (UCLA) with a 12 h light/dark cycle and *ad libitum* access to food and water. Wild-type C57BL/6NTac mice were generated from in house breeding colonies or purchased from Taconic Biosciences. Transgenic mice expressing eGFP under an astrocyte specific promoter *Aldh111* (Aldh111-eGFP mice) were maintained as a heterozygous line (Gong et al., 2003). D1-Cre (Strain B6.FVB(Cg)-Tg(Drd1-cre)EY262Gsat/Mmucd) and D2-Cre (Strain B6.FVB(Cg)-Tg(Adora2a-cre)KG139Gsat/Mmucd) BAC transgenic mouse lines were obtained from MMRRC at University of California Davis and maintained on a C57BL/6NTac background. R6/2 and non-carrier wild-type control mice were purchased from Jackson Laboratories (Strain B6CBA-Tg(HDexon1)62Gpb/3J). RiboTag mice were purchased from Jackson Laboratories (Strain B6N.129-Rpl22^{tm1.1P_{Sam}}/J) and crossed with *Aldh111*-Cre/ERT2 mice (Srinivasan et al., 2016) to generate mice carrying both alleles for RNA-seq experiments. *lpr2* knockout (*lpr2*^{tm1.1Chen}) mice were available from previous work (Srinivasan et al., 2015) and were originally obtained from Dr. Ju Chen at University of California–San Diego and maintained as a heterozygous line.

Cell Lines

HEK293 cells (sex: female, RRID: CVCL_0045) were obtained from ATCC and maintained in 25 cm² cell culture flasks in DMEM/F12 media with Glutamax (Invitrogen) supplemented with 10% fetal bovine serum and penicillin/streptomycin. Cells were grown in a humidified cell culture incubation with 95% air / 5% CO₂ at 37°C.

METHOD DETAILS

Molecular Biology and Generation of Adeno-Associated Viruses

Adeno-associated viruses (AAVs) were generated as previously described (Shigetomi et al., 2013). In brief, the hPMCA2w/b and hPMCA2x/b sequences were removed by restriction digestion from Addgene plasmids 47586 and 47585, respectively. Together with a sequence encoding mCherry that was amplified by PCR from pCS2-mCherry, hPMCA2w/b and hPMCA2x/b were incorporated into pZac2.1 vector using In-Fusion Cloning Kits (Takara) to generate plasmid pZac2.1 GfaABC₁D mCherry-hPMCA2w/b (Addgene, 111568) and pZac2.1 GfaABC₁D mCherry-hPMCA2x/b (Addgene, 111569), which carried the minimal GfaABC₁D astrocyte-specific promoter. After verifying the sequences, plasmids were used to make AAVs of the 5 serotype at the Penn Vector Core (AAV2/5 GfaABC₁D mCherry-hPMCA2w/b; 9.2×10^{12} gc/ml and AAV5 GfaABC₁D mCherry-hPMCA2x/b; 1.6×10^{13} gc/ml). For HEK293 cell transfection, mCherry-hPMCA2w/b and mCherry-hPMCA2x/b sequences were cloned into pcDNA3.1 vectors between *Afl*III and *Xba*I sites to generate pcDNA3.1-CMV-mCherry-hPMCA2w/b (Addgene, 111570) and pcDNA3.1-CMV-mCherry-hPMCA2x/b (Addgene, 111571). To generate RiboTag AAV, the cDNA of the ribosomal protein Rpl22 with three HA tags was

amplified from pcDNA3.1-Rpl22-3xHA (Sanz et al., 2009) (a gift from Dr. McKnight) and cloned into pZac2.1 using In-Fusion Cloning Kits (Takara) to generate pZac2.1-GfaABC₁D-Rpl22-HA (Addgene, 111811). The plasmid was then sent to the Penn Vector Core for AAV2/5 production (AAV2/5 GfaABC₁D-Rpl22-HA; 2.1×10^{13} gc/ml). All the new plasmids generated in this study that are listed in [Key Resources Table](#) have been deposited at Addgene in the Khakh laboratory repository (http://www.addgene.org/Baljit_Khakh).

HEK293 Cell Imaging

HEK293 cell imaging methods have been described (Richler et al., 2008). HEK293 cells were prepared for transfection by plating onto six-well plates and transfection was performed when cells reached 80%–90% confluence. For transient expression in HEK293 cells, 0.5–1 μ g plasmid DNA were transfected using Effectene transfection reagent (QIAGEN). Cells were imaged 24–72 hr after transfection in HEK cell recording buffer (150 mM NaCl, 1 mM MgCl₂, 1 mM CaCl₂, 10 mM D-glucose, 10 mM HEPES, with pH adjusted to 7.4). Ca²⁺ signals were recorded under a laser-scanning confocal microscope (Olympus) with a 40 \times water-immersion objective lens (NA 0.8). To activate P2Y₁ receptors, ATP (50 μ M) was bath-applied in the recording solution.

Astrocyte Purification and Quantitative PCR

Fluorescence-activated cell sorting (FACS) was performed to purify striatal astrocytes from Aldh1l1-eGFP mice at P30, as previously described (Jiang et al., 2016). Briefly, the striata from 4 mice were dissected and digested at 36°C for 45 min with papain solution (1 \times EBSS, 0.46% D-glucose, 26 mM NaHCO₃, 50 mM EDTA, 75 U/ml DNase 1, 300 units of papain, and 2 mM L-cysteine). The tissue was then rinsed with ovomucoid solution (1 \times EBSS, 0.46% D-glucose, 26 mM NaHCO₃, 1 mg/ml ovomucoid, 1 mg/ml BSA, and 60 U/ml DNase 1) and mechanically dissociated with borosilicate glass pipettes. A concentrated ovomucoid solution (1 \times EBSS, 0.46% D-glucose, 26 mM NaHCO₃, 5.5 mg/ml ovomucoid, 5.5 mg/ml BSA, and 25 U/ml DNase 1) was layered at the bottom of the cell suspension. Cells were precipitated by centrifuging and the pellet was resuspended and filtered. FACS was performed using a FACSaria II (BD Biosciences) with a 70 μ m nozzle at UCLA Cell Sorting Core. Total RNA was extracted from sorted GFP⁺ cells (RNeasy Plus Micro Kit; QIAGEN) and cDNA was amplified (Ovation PicoSL WTA System V2; Nugen) and purified (QIAquick PCR Purification Kit; QIAGEN). Quantitative PCR was performed in a LightCycler 96 Real-Time PCR System (Roche). Ten nanograms of cDNA were loaded per well and the expression of *Atp2b1*, *Atp2b2*, *Atp2b3*, *Atp2b4* and *Arbp* was analyzed using the primers listed in [Key Resources Table](#). *Arbp* was used as an internal control to normalize RNA content. To calculate the expression of gene of interest, the following formula was used: $2^{-\Delta Ct}$ (Gene of interest-*Arbp*).

Surgical Procedure of In Vivo Microinjection

Surgical procedure of viral microinjection has been described previously (Haustein et al., 2014; Jiang et al., 2014, 2016; Shigetomi et al., 2013). In brief, mice aged 6- to 8-week-old were anesthetized and placed onto a stereotaxic frame (David Kopf Instruments, Tujunga CA). Continuous anesthesia using isoflurane was carefully monitored and adjusted throughout the surgery. Mice were injected with buprenorphine (Buprenex; 0.1 mg/kg) subcutaneously before surgery. Scalp incisions were made and craniotomies (1–2 mm in diameter) above the left parietal cortex were created using a high-speed drill (K.1070; Freedom) for unilateral viral injections while two craniotomies were made above both parietal cortices for bilateral viral injections. Beveled glass pipettes (1B100–4; World Precision Instruments) filled with viruses were placed into the striatum (0.8 mm anterior to the bregma, 2.0 mm lateral to the midline, and 2.4 mm from the pial surface) or the hippocampus CA1 (2 mm posterior to the bregma, 1.5 mm lateral to the midline, and 1.6 mm from the pial surface). For slice Ca²⁺ imaging, 1 μ l AAV2/5 GfaABC₁D GCaMP6f (2.4×10^{13} gc/ml) with either 1 μ l of AAV2/5 GfaABC₁D mCherry-hPMCA2w/b (9.2×10^{12} gc/ml) or 0.5 μ l AAV5 GfaABC₁D tdTomato (5.0×10^{13} gc/ml) were injected at 200 nl/min using a syringe pump (Pump11 PicoPlus Elite; Harvard Apparatus). For MINIscope Ca²⁺ imaging, 0.5 μ l AAV1 Synapsin GCaMP6f (2.0×10^{13} gc/ml) with either 1 μ l of AAV5 GfaABC₁D mCherry-hPMCA2w/b or 0.5 μ l AAV5 GfaABC₁D tdTomato were injected. For D1- and D2-MSN physiological recording, 0.5 μ l AAV1 CAG Flex tdTomato (1.1×10^{13} gc/ml) with either 1 μ l of AAV5 GfaABC₁D mCherry-hPMCA2w/b or 0.5 μ l AAV5 GfaABC₁D tdTomato were injected. For Ca²⁺ imaging in R6/2 and non-carrier control mice, 1.5 μ l AAV5 GfaABC₁D GCaMP6f were injected. For RiboTag mice RNA-seq experiments, 1 μ l of AAV2/5 GfaABC₁D-mCherry-hPMCA2w/b or 0.5 μ l AAV2/5 GfaABC₁D-tdTomato were injected into the striatum of RiboTag mice. For RiboTag AAV RNA-seq experiments, 0.7 μ l AAV2/5 GfaABC₁D-Rpl22-HA with either 1 μ l of AAV2/5 GfaABC₁D-mCherry-hPMCA2w/b or 0.5 μ l AAV2/5 GfaABC₁D-tdTomato were injected into the striatum of C57BL/6NTac mice. Glass pipettes were withdrawn after 10 min and scalps were cleaned and sutured with sterile surgical sutures. Mice were allowed to recover in clean cages with food containing Trimethoprim/Sulfamethoxazole and water for 7 days. Imaging and behavioral experiments were performed at least three weeks after surgeries.

Brain Slice Preparation

For imaging and electrophysiological recordings, mice were transferred from the vivarium to the laboratory during the light cycle between 8 am and 10 am. Brain slices were prepared 30 min to 1 hr afterward and were used for experiments within 8 h of slicing (mostly within 6 hr). Both male and female mice (9–11 weeks old) with AAV injection were anesthetized and decapitated. Coronal slices (300 μ m thick) of the striatum or hippocampus were prepared in ice-cold sucrose cutting solution (30 mM NaCl, 4.5 mM KCl, 1.2 mM NaH₂PO₄, 26 mM NaHCO₃, 10 mM D-glucose, 194 mM sucrose and 1 mM MgCl₂) using a vibratome (DSK Microslicer; Ted Pella, Inc.). Slices were then incubated in artificial cerebrospinal fluid (ACSF) (124 mM NaCl, 4.5 mM KCl, 1 mM MgCl₂,

1.2 mM NaH_2PO_4 , 26 mM NaHCO_3 , 10 mM D-glucose, and 2.0 mM CaCl_2) for 30 min at 32°C and 1 h at room temperature before recording. All the solutions were oxygenated with 95% O_2 /5% CO_2 . Procedures have been described in detail (Jiang et al., 2016; Tong et al., 2014).

Intracellular Ca^{2+} Imaging of Striatal Astrocytes

Ca^{2+} imaging in striatal slices was performed using a Scientifica two-photon laser-scanning microscope (2PLSM) equipped with a MaiTai laser (Spectra-physics). To image GCaMP6f signals, laser was tuned at 920 nm wavelength. The laser power measured at the sample was less than 30 mW with a 40 × water-immersion objective lens (Olympus). Astrocytes located in the dorsolateral striatum and at least 40 μm from the slice surface were selected for imaging. Images were acquired at 1 frame per second using SciScan software (Scientifica). Striatal slices were maintained in ACSF (124 mM NaCl, 4.5 mM KCl, 1 mM MgCl_2 , 1.2 mM NaH_2PO_4 , 26 mM NaHCO_3 , 10 mM D-glucose, and 2.0 mM CaCl_2) through a perfusion system. Phenylephrine (10 μM) was bath-applied to activate GPCR-mediated Ca^{2+} signals. Cyclopiazonic acid (20 μM) was applied to deplete intracellular calcium stores. Slices were pre-incubated with SR101 (1 μM) for 10 min for some imaging experiments. Ca^{2+} signals were processed in ImageJ (NIH) and presented as the relative change in fluorescence ($\Delta F/F$). Peak amplitude, half-width, and frequency of Ca^{2+} signals were analyzed in ClampFit10.4, while integrated areas were calculated in OriginPro 8.

Intracellular Ca^{2+} Imaging of Hippocampal Astrocytes

Ca^{2+} imaging in hippocampal slices was performed using a confocal microscope (Fluoview 1000; Olympus) with a 40X water-immersion objective lens (NA = 0.8). To image GCaMP6f signals, 488 nm line of an Argon laser was used with the intensity adjusted to 10%–14% of the maximum output of 10 mW. The emitted light pathway consisted of an emission high pass filter (505–525 nm) before the photomultiplier tube. Astrocytes located in the hippocampal CA1 and at least 40 μm from the slice surface were selected for imaging. Images were acquired at 1 frame per second. Hippocampal slices were maintained in ACSF (124 mM NaCl, 4.5 mM KCl, 1 mM MgCl_2 , 1.2 mM NaH_2PO_4 , 26 mM NaHCO_3 , 10 mM D-glucose, and 2.0 mM CaCl_2) through a perfusion system. Phenylephrine (10 μM) was bath-applied to activate GPCR-mediated Ca^{2+} signals. Ca^{2+} signals were processed in ImageJ (NIH) and presented as the relative change in fluorescence ($\Delta F/F$). Peak amplitude, half-width, and frequency of Ca^{2+} signals were analyzed in ClampFit10.4, while integrated areas were calculated in OriginPro 8.

MINIscope Implantation and Imaging in the Striatum

Both male and female mice (9 weeks old) were anesthetized and placed onto a stereotaxic frame (David Kopf Instruments, Tujunga, CA) one week after viral injection in the striatum. Dexamethasone (0.2 mg/kg) and carprofen (5 mg/kg) were administered subcutaneously during the surgery. After hair removal and scalp incision, the skull was exposed and the connective tissue underneath was removed. A 2-mm diameter circular craniotomy was made with the center located 0.5 mm from the viral injection site. The cortex underneath the craniotomy was aspirated using a blunt syringe needle connected to a vacuum pump until a layer of corpus callosum was exposed. ACSF was constantly applied to the surgery site to prevent tissue from drying. Sterilized GRIN lens (0.25 pitch, 2 mm in diameter and 4.79 mm in length; Grintech GmbH) attached to a stereotaxic arm was slowly lowered into the aspirated space until the bottom of the lens was 2 mm below the surface of the skull. Implanted GRIN lens was then anchored to the skull as well as a custom-made head bar that was glued to the back of the occipital bone with cyanoacrylate (Krazy glue) and dental cement (Ortho-Jet, Lang Dental). After surgery, mice were given carprofen (5 mg/kg) and amoxicillin (BIOMOX, Virbac Animal Health) for at least 7 days. At least 3 days before imaging, a light-weight aluminum baseplate was glued on top of the dental cement to secure the miniature microscope (MINIscope), a procedure called “baseplating.” During baseplating, the mouse was head-fixed without anesthesia so that the appropriate field of view could be determined based on neuronal activities. Mice were accustomed to head-fixation after GRIN lens implantation and before baseplating. SNAP5114 (50 $\mu\text{mol/kg}$) was intraperitoneally injected for 7 days before imaging. On the day of imaging, MINIscope was placed on top of the baseplates and secured while the mice were awake (see <http://www.miniscope.org/> for further details). Both MINIscope and behavior camera (Logitech) were controlled by a custom data acquisition software written in C++. Images were acquired at 30 frames per second and time stamped by the internal CPU clock for subsequent analysis.

Electrophysiological Recordings in the Striatal Slices

Electrophysiological recordings were performed as described previously by us for striatal slices (Chai et al., 2017; Jiang et al., 2016; Oceau et al., 2018; Srinivasan et al., 2015, 2016; Tong et al., 2014), which uses methods reported earlier in landmark studies of striatal astrocytes (Ademark and Lovinger, 2006). Of note, the brain slice recordings were performed at room temperature (21–23°C). This is because in our past work over a period of 5 years we have found that incubating striatal slices from adult mice (> P60) at ~33°C closer to mouse body temperature caused swelling, appearance of sick cells and the rapid loss of whole-cell seal integrity (particularly for astrocytes – the focus of our work). In order to minimize such effects, we had to balance the requirement for healthy tissue with that of recording at warmer temperatures. By opting to record at room temperature we prioritized the quality of the recordings over temperature. However, the *in vivo* imaging, behavior, immunohistochemistry and RNA-seq work described all reflect measurements at mouse body temperature. Hence, our studies need to be interpreted with the knowledge that the kinetics of the

synaptic responses we report are likely slowed. Our use of room temperature is identical to that used in past striatal astrocyte studies by other labs (Adermark and Lovinger, 2006; D'Ascenzo et al., 2007; Martín et al., 2015).

Whole-cell patch-clamp recordings were made from medium spiny neurons (MSNs) or astrocytes morphologically identified under infrared optics from the dorsolateral striatum. D1- and D2-MSNs were selected based on tdTomato fluorescence. The intracellular solution for MSN IPSCs recordings comprised the following: 138 mM KCl, 10 mM HEPES, 1 mM EGTA, 0.3 mM Na-ATP, 4 mM Mg-ATP, 0.1 mM CaCl_2 , 8 mM Na_2 -phosphocreatine, with pH adjusted to 7.3. The intracellular solution for MSN EPSCs recordings comprised the following: 120 mM CsMeSO_3 , 15 mM CsCl, 8 mM NaCl, 10 mM HEPES, 0.2 mM EGTA, 0.3 mM Na-GTP, 2 mM Mg-ATP, 10 mM TEA-Cl, with pH adjusted to 7.3 with CsOH. The intracellular solution for astrocyte recordings comprised the following: 135 mM potassium gluconate, 3 mM KCl, 10 mM HEPES, 1 mM EGTA, 0.3 mM Na-ATP, 4 mM Mg-ATP, 0.1 mM CaCl_2 , 8 mM Na_2 -phosphocreatine, with pH adjusted to 7.3. To isolate sIPSCs, MSNs were voltage-clamped at -60 mV and pre-incubated with ACSF (124 mM NaCl, 4.5 mM KCl, 1 mM MgCl_2 , 1.2 mM NaH_2PO_4 , 26 mM NaHCO_3 , 10 mM D-glucose, and 2.0 mM CaCl_2) in the presence of $10 \mu\text{M}$ CNQX for 10 min before recording. Note, that all our recordings used whole-cell patch clamp and not perforated patch clamp. Brain slice neuronal perforated patch recordings most successfully employ antibiotics such as nystatin and amphotericin that rapidly form pores in the membrane patch. However, these pores are permeable to monovalent anions and cations, including Cl^- (Lippiat, 2008; Sherman-Gold, 2006). Hence, perforated patch recordings would not allow us to maintain intracellular Cl^- at its endogenous value. The pores in perforated patches do discriminate between monovalent and multivalent cations and larger organic species, thus allowing researchers to preserve intracellular metabolites such as ATP. However, such recordings do not prevent equilibration of Cl^- (Lippiat, 2008; Sherman-Gold, 2006). Gramicidin may more favorably discriminate between monovalent anions and cations, but the procedure is very slow, leads to high access resistances and also creates a Donnan equilibrium potential that in turn causes a voltage offset and change in cell volume. These factors would impair our ability to record tonic currents (which are far smaller than phasic GABA IPSCs and exogenous GABA puff currents). For all these reasons, we chose whole-cell patch clamp, which is the same as that used in past studies on tonic inhibition in MSNs (Kirmse et al., 2009; Wójtowicz et al., 2013). mIPSCs were recorded after incubation with $10 \mu\text{M}$ CNQX and $0.5 \mu\text{M}$ TTX for 10 min. To assess extrasynaptic GABA_A receptor-mediated responses, GABA ($300 \mu\text{M}$) was puffed onto the cell bodies of MSNs from a micropipette connected to Picospritzer III (Parker) for 100 ms to evoke GABA currents. To evoke IPSCs in MSNs, a bipolar stimulating electrode was placed approximately 100 – $150 \mu\text{m}$ from the recorded neuron in the dorsal striatum. Glutamate receptors were blocked by incubation with $10 \mu\text{M}$ CNQX and $25 \mu\text{M}$ APV for 10 min. To assess input-output function, test-stimuli were applied at increasing intensities ranging from 0 to $50 \mu\text{A}$. Stimulation intensities were set to evoke responses at 50% maximal amplitude to induce paired pulse responses. Paired pulses were delivered at three different inter-pulse intervals: 30, 50 and 100 ms apart. To access tonic GABA currents, $25 \mu\text{M}$ bicuculline was bath-applied in the presence of $10 \mu\text{M}$ CNQX, $0.5 \mu\text{M}$ TTX and $25 \mu\text{M}$ APV. To investigate the contributions of GABA transporters to tonic GABA currents, striatal slices were pre-incubated with GAT-1 inhibitor NNC711 ($10 \mu\text{M}$; Tocris Bioscience) or GAT-3 inhibitor SNAP5114 ($40 \mu\text{M}$; Tocris Bioscience) or both for 10 min before recording. To isolate spontaneous EPSCs, MSNs were voltage-clamped at -60 mV and pre-incubated with ACSF in the presence of $25 \mu\text{M}$ bicuculline for 10 min before recording. To assess evoked EPSCs, electrical field stimulation (EFS) were achieved using a bipolar matrix electrode (FHC) that was placed on the dorsolateral corpus callosum to evoke glutamate release from cortico-striatal pathway (Jiang et al., 2016). The MSNs to be assessed were typically located about 250 – $300 \mu\text{m}$ away from the stimulation site. To isolate the AMPAR- and NMDAR-mediated evoked EPSCs, MSNs were voltage-clamped at -70 mV or $+40$ mV in the presence of $10 \mu\text{M}$ bicuculline. The AMPAR-mediated EPSC was measured at the peak amplitude of the EPSC at -70 mV, while the amplitude of the EPSC 50 ms after stimulation at $+40$ mV was used to estimate the NMDAR-mediated component. All recordings were performed at room temperature, using pCLAMP10 (Axon Instruments, Molecular Devices) and a MultiClamp 700B amplifier (Axon Instruments, Molecular Devices). Cells with R_a exceeded $20 \text{ M}\Omega$ were excluded from analysis. Analysis was performed using ClampFit 10.4 software.

Immunohistochemistry (IHC)

Both male and female mice (9–14 weeks old) were transcardially perfused with 0.1 M phosphate buffered saline (PBS) followed by 10% formalin. Brains were dissected out and post-fixed in 10% formalin for several hours and then cryoprotected in 30% sucrose (0.1 M PBS). Serial $40 \mu\text{m}$ sections were collected and incubated with the following primary antibodies overnight at 4°C : rabbit anti-S100 β (1:1,000; Abcam, ab41548), mouse anti-NeuN (1:1,000; Millipore, MAB377), chicken anti-GFP (1:1,000; Abcam, ab13970), rabbit anti-RFP (1:1,000; Rockland, 600-401-379), chicken anti-RFP (1:1,000; a gift from Dr. Brecha's laboratory at UCLA), rabbit anti-GAT-3 (1:500; a gift from Dr. Brecha's laboratory at UCLA), rabbit anti-Kir4.1 (1:1,500; Alomone, APC-035), rabbit anti-DARPP-32 (1:200; Abcam, ab40801) or mouse anti-HA (1:1,000; Covance, MMS-101R). Sections were then incubated with the following secondary antibodies (1:1,000; Molecular Probes): Alexa Fluor 488 goat anti-chicken (A11039), Alexa Fluor 488 goat anti-rabbit (A11008), Alexa Fluor 546 goat anti-mouse (A11003), Alexa Fluor 546 goat anti-chicken (A11040) and Alexa Fluor 594 goat anti-rabbit (R37007) at room temperature for 2 h. Fluorescence imaging was acquired with a $40\times$ oil-immersion objective lens (NA 1.3) under a laser-scanning confocal microscope (Olympus).

Western Blot Analyses

To extract membrane fraction, mice were decapitated and striata were dissected and homogenized in ice-cold buffer (250 mM sucrose, 1 mM EDTA, 10 mM Tris-HCl and protease inhibitor cocktail, adjusted to pH 7.2). The homogenate was centrifuged at

500 × g for 15 min at 4°C. The pellet was discarded and the supernatant was centrifuged twice at 75,000 × g for 30 min at 4°C. The pellet containing the membrane fraction was dissociated in RIPA buffer with protease inhibitors by sonication for 30 min. Protein concentrations were measured using BCA protein assay (Thermo Scientific). Membrane fraction was mixed with 2 × Laemmli buffer containing 8 M urea. The samples were then electrophoretically separated by 12% SDS-PAGE (5 µg protein per lane) and transferred onto nitrocellulose membrane (0.45 µm). The membrane was probed at 4°C overnight with the following primary antibodies: rabbit anti-GAT-3 (1:1,000; a gift from Dr. Brecha's laboratory at UCLA), rabbit anti-GAT-1 (1:500; Abcam) and mouse anti-β-tubulin (1:5,000; Millipore). IRDye 680RD anti-mouse (1:10,000; Li-Cor) and IRDye 800CW anti-rabbit (1:10,000; Li-Cor) secondary antibodies were used and images were acquired on a Li-Cor Odyssey infrared imager. Signal intensities at expected molecular weight were quantified using ImageJ (NIH). GAT-1 and GAT-3 levels were normalized to β-tubulin.

Behavioral Tests

Behavioral tests were performed during the light cycle between 12:00 pm and 6:00 pm. Only male mice (9–13 weeks old) were used in behavioral tests because of sex-dependent differences known for striatal physiology. All the experimental mice were transferred to the behavior testing room at least 30 min before the tests to acclimatize to the environment and to reduce stress.

Open Field Test

The open field chamber consisted of a square arena (28 cm × 28 cm) enclosed by walls made of Plexiglass (19 cm tall). The periphery of the arena was defined as the area within 2.5 cm adjacent to the walls of the chamber and the center of the arena was defined as the area 2.5 cm away from the chamber walls. Each mouse was placed into an open field box to habituate for 20 min. Locomotor activity was then recorded for 1 h using an infrared camera located underneath the open field chamber. Recording camera was connected to a computer operating an automated video tracking software Ethovision XT (Noldus Information Technology). Parameters analyzed included distance traveled and time spent in the center, with 5 min and 1 h time bins. After recordings finished, mice were returned to their home cages.

Acute Locomotor Effects of Cocaine

Each mouse was weighed and received single intraperitoneal injection of saline or cocaine (10 mg/kg) and then placed into the open field chamber. Locomotor activities was recorded for 15 min and distance traveled was assessed using Ethovision XT with 3 min time bins.

Locomotor Sensitization Induced by Cocaine

Each mouse was weighed and received intraperitoneal injection of saline or cocaine (10 mg/kg) for 5 consecutive days. After 3 weeks of withdrawal from repeated cocaine injection, cocaine challenge was performed with mice injected with either saline or cocaine (10 mg/kg) to test the sensitization. Locomotor activity was recorded for 15 min in the open field chamber and distance traveled was assessed using Ethovision XT with 15 min time bins.

Self-Grooming Behavior

The procedure of self-grooming behavior measurement was adapted from previously published study (Kalueff et al., 2007). Mice were placed individually into the open field chambers or plastic cylinders (15 cm in diameter and 35 cm tall), and allowed to habituate for 20 min. Self-grooming behavior was recorded for 10 min. SNAP5114 (50 µmol/kg) was intraperitoneally injected for 7 days and the experiments were performed again. A timer was used to assess the cumulative time spent in self-grooming behavior, which included paw licking, unilateral and bilateral strokes around the nose, mouth, and face, paw movement over the head and behind ears, body fur licking, body scratching with hind paws, tail licking, and genital cleaning. The number of self-grooming bouts was also counted. Separate grooming bouts were considered when the pause was more than 5 s or behaviors other than self-grooming occurred. Self-grooming microstructure was not assessed.

Spray Test

A standard spray bottle was filled with distilled water and the nozzle was adjusted to the “misting” mode. Mice were held by the tails on the bench and sprayed three times from 30 cm away to be adequately covered with mist. Mice were placed individually into the plastic cylinders and grooming behavior was recorded for 10 min following the spray and then analyses as described in the preceding section.

Rotarod Test

Mice were held by the tails and placed on the rod (3 cm diameter) of a single lane rotarod apparatus (ENV-577M, Med Associates Inc.), facing away from the direction of rotation. The rotarod was set with a start speed of 4 rpm. Acceleration started 10 s later and was set to 20 rpm per minute with a maximum speed 40 rpm. Each mouse received two trials 30 min apart and the latency to fall was recorded for each trial. The average latency to fall was used as a measurement for motor coordination (Deacon, 2013).

Elevated Plus Maze

Modified from the protocol described previously (Komada et al., 2008), the elevated plus maze apparatus was a cross-shaped platform that comprised two open arms and two closed arms (29 cm × 9 cm) with a center platform (9 cm × 9 cm). Open arms and closed arms were perpendicular to each other and elevated about 50 cm from the floor. Closed arms had walls (15 cm) to enclose the arms. Each mouse was placed in the center platform with its nose directed toward the same closed arm and allowed to explore the entire apparatus for 10 min. Anxiety-like behavior was evaluated with the number of entries into each compartment and the time spent in each compartment. Total travel distance was also determined. All the parameters were analyzed in Ethovision XT.

Clasping Test

R6/2 mice and non-carrier controls at 10–11 weeks old were used in this test. As previously described (Tong et al., 2014), each mouse was suspended by holding the tail. The experiments were conducted for 1 min. Time taken for the mouse to move both forelimbs close to the body and for both paws to clasp was recorded. The mice that did not clasp in this time period were given a data value of 60 s. SNAP5114 (50 μ mol/kg) was intraperitoneally injected for 7 days and the test was performed again.

Grip Strength Test

R6/2 mice and non-carrier controls at 10–11 weeks old were used in this test. Forelimb strength was measured using a customized grip strength meter (Chatillon Digital force gauge) with an adjustable angel wire mesh grip (San Diego Instruments). Both the grip strength meter and the mesh grip were positioned horizontally. Each mouse was held by the base of the tail and lowered to grasp the mesh with both forelimbs. A steady horizontal force was then applied to extend the mouse away from the meter until it relinquished its grip on the mesh. Maximum pulling force required to separate the mouse from the mesh grip was recorded by the meter. Each mouse was tested in five trials with a 1 min intertrial interval and average maximum pulling force was used to assess the muscle strength. SNAP5114 (50 μ mol/kg) was intraperitoneally injected for 7 days and the test was performed again.

Footprint Test

R6/2 mice and non-carrier controls at 10–11 weeks old were used in this test. A one meter long runway (8 cm wide) was lined with paper. Each mouse with hind paws painted with non-toxic ink was placed at an open end of the runway and allowed to walk to the other end with a darkened box. For the gait analysis, stride length and width were measured and averaged for both left and right hindlimbs over 5 steps. SNAP5114 (50 μ mol/kg) was intraperitoneally injected for 7 days and the test was performed again.

RNA-Seq Analysis of Striatal Astrocyte Transcriptomes

Total RNA was extracted from the striatum of male and female mice (aged 9–10 weeks) as previously described (Chai et al., 2017). For RiboTag mice RNA-seq experiments, tamoxifen (Sigma, 20 mg/ml) was administered intraperitoneally for five consecutive days at 100 mg/kg body weight to *Aldh1l1-Cre/ERT2* \times RiboTag mice. One day after the last tamoxifen injection, 1 μ l of AAV2/5 GfaABC₁D-mCherry-hPMCA2w/b or 0.5 μ l AAV2/5 GfaABC₁D-tdTomato were injected into the striatum of RiboTag mice. For RiboTag AAV RNA-seq experiments, 0.7 μ l AAV2/5 GfaABC₁D-Rpl22-HA with either 1 μ l of AAV2/5 GfaABC₁D-mCherry-hPMCA2w/b or 0.5 μ l AAV2/5 GfaABC₁D-tdTomato were injected into the striatum of C57BL/6NTac mice. Three weeks after the AAV injection, striatal tissues were dissected and homogenized. RNA was extracted from 10%–20% of homogenate after centrifugation as input sample, which contained RNA from all cell types in the striatum (QIAGEN Rneasy Plus Micro #74034). The remaining homogenate was incubated with mouse anti-HA antibody (1:250; Covance, #MMS-101R) for 4 hr followed by the addition of magnetic beads (Invitrogen, Dynabeads #110.04D) for overnight incubation at 4°C. RNA was purified from the immunoprecipitation (IP) sample, which contained astrocyte-enriched RNA (QIAGEN Rneasy Plus Micro #74034). RNA concentration and quality were assessed with Agilent 2100 Bioanalyzer. RNA samples with RNA integrity number (RIN) greater than 7 were used for multiplexed library prep with Nugen Ovation RNA-Seq System V2. For each experiment, all samples were multiplexed into a single pool in order to avoid batch effects (Auer and Doerge, 2010), and sequencing was performed on Illumina HiSeq 4000 for 2 \times 75 yielding between 58 and 83 million reads per sample. Demultiplexing was performed with Illumina Bcl2fastq2 v 2.17 program. Reads were aligned to the mouse mm10 reference genome using the STAR spliced read aligner (Dobin et al., 2013) with default parameters and fragment counts were derived using HTS-seq program. Between 66 and 91% of the reads mapped uniquely to the mouse genome and were used for subsequent analyses. Differential gene expression analysis was performed with genes with CPM > 3 in at least 4 samples, using Bioconductor packages edgeR (Robinson et al., 2010) and limmaVoom (Law et al., 2014) with false discovery rate (FDR) threshold < 0.1 or 0.05 (<http://www.bioconductor.org>). Weighted gene co-expression network analysis (WGCNA) was performed using an R package of WGCNA for finding modules of highly correlated genes. The gene ontology (GO) enrichment analysis was performed using PANTHER overrepresentation test (GO ontology database released 2018-04-04) with FDR < 0.05. RNA-seq data have been deposited in the Gene Expression Omnibus (GEO) repository (<https://www.ncbi.nlm.nih.gov/geo>) with accession numbers GSE114757 and GSE114756.

QUANTIFICATION AND STATISTICAL ANALYSIS

Data from every experiment represent at least four replicates. Sample sizes were not calculated *a priori*. Mice were randomly assigned to different experimental groups. For 2PLSM Ca²⁺ imaging experiments, Ca²⁺ transients recorded in a single optical plane and were measured by plotting the intensity of regions of interest over time. Slow drifts in astrocyte position (~2–5 μ m) were corrected with a custom Plugin in NIH ImageJ. A signal was declared as a Ca²⁺ transient if it exceeded the baseline by greater than twice the baseline noise (SD). Ca²⁺ signals were processed in ImageJ and presented as the relative change in fluorescence ($\Delta F/F$). Peak amplitude, half-width, and frequency of Ca²⁺ signals were analyzed in ClampFit10.4 (Molecular Devices), while integrated areas were calculated in OriginPro 8. To evaluate the resting astrocyte Ca²⁺ level, basal GCaMP6f fluorescence intensity was normalized to SR101 fluorescence intensity. For MINIscope experiments, a custom script written in MATLAB (MathWorks, Natick, MA) was used to correct column-wise ACD variation and small movement artifacts. Active neurons (ROIs) were segmented manually and Ca²⁺ traces were extracted in ImageJ. Peak amplitude, half-width, integrated area and frequency of individual Ca²⁺ events were analyzed in MiniAnalysis Program 6.0.7 (Synaptosoft, Decatur, GA). Ca²⁺ traces were aligned with behavioral analysis using

custom-written scripts in R (version 3.3.2). To assess correlation coefficient between neuronal pairs, Ca^{2+} traces were binned every 1 s and given binary variables without deconvolution. A matrix with columns representing discrete time intervals of 1 s each and rows representing each ROI was created and the Pearson correlation matrix was calculated using custom-written scripts in R. Since the correlation matrix is a symmetric square matrix with 1 s along the main diagonal, the absolute mean correlation of the ROIs was calculated after excluding the values along and below the main diagonal. Results were also confirmed with deconvolved data (Friedrich et al., 2017). For electrophysiological recordings, synaptic currents and tonic GABA currents were analyzed using MiniAnalysis Program 6.0.7 (Synaptosoft, Decatur, GA) or Clampfit10.4 (Molecular Devices).

All statistical tests were performed in OriginPro 8. The graphs were created in OriginPro 8 and assembled in CorelDraw12. Data are presented as mean \pm SEM. Note that in some of the graphs, the SEM bars were smaller than the symbols used to represent the mean. For each set of data to be compared, OriginPro 8 was used to determine whether the data were normally distributed or not. Parametric tests were used for normally distributed datasets while non-parametric tests were applied to data not normally distributed. Paired and unpaired Student's two tailed t tests (as appropriate) and two tailed Mann-Whitney tests were used for most statistical analyses with significance declared at $p < 0.05$, but stated in each case with a precise *P value*. Comparisons of datasets with more than two groups or conditions were calculated with one-way or two-way ANOVA tests followed by Bonferroni post hoc tests. When the *P value* was less than 0.001, it is stated as $p < 0.001$ to save space on the figure panels and tables. When the *P values* was greater than 0.05, it is stated as non-significant (n.s.). Throughout the manuscript, the results of statistical tests (*P values* and n numbers) are reported in the figure panels. N numbers are defined as the numbers of cells/slices or mice throughout.

DATA AND SOFTWARE AVAILABILITY

RNA-seq data have been deposited in the Gene Expression Omnibus (GEO) repository (<https://www.ncbi.nlm.nih.gov/geo>) with accession numbers GEO: GSE114757 and GEO: GSE114756.