



HHS Public Access

Author manuscript

Neuropharmacology. Author manuscript; available in PMC 2016 August 01.

Published in final edited form as:

Neuropharmacology. 2015 August ; 95: 468–476. doi:10.1016/j.neuropharm.2015.03.036.

Dopaminergic and Cholinergic Modulation of Striatal Tyrosine Hydroxylase Interneurons

Oswaldo Ibáñez-Sandoval^a, Harry S. Xenias^b, James M. Tepper, and Tibor Koós

Abstract

The recent electrophysiological characterization of TH-expressing GABAergic interneurons (THINs) in the neostriatum revealed an unexpected degree of diversity of interneurons in this brain area (Ibáñez-Sandoval et al., 2010, Unal et al., 2011, 2013). Despite being relatively few in number, THINs may play a significant role in transmitting and distributing extra- and intrastriatal neuromodulatory signals in the striatal circuitry. Here we investigated the dopaminergic and cholinergic regulation of THINs *in vitro*. We found that the dominant effect of dopamine was a dramatic enhancement of the ability of THINs to generate long-lasting depolarizing plateau potentials (PPs). Interestingly, the same effect could also be elicited by amphetamine-induced release of endogenous dopamine suggesting that THINs may exhibit similar responses to changes in extracellular dopamine concentration *in vivo*. The enhancement of PPs in THINs is perhaps the most pronounced effect of dopamine on the intrinsic excitability of neostriatal neurons described to date. Further, we demonstrate that all subtypes of THINs tested also express nicotinic cholinergic receptors. All THINs responded, albeit differentially, with depolarization, PPs and spiking to brief application of nicotinic agonists. Powerful modulation of the nonlinear integrative properties of THINs by dopamine and the direct depolarization of these neurons by acetylcholine may play important roles in mediating the effects of these neuromodulators in the neostriatum with potentially important implications for understanding the mechanisms of neuropsychiatric disorders affecting the basal ganglia.

Keywords

neostriatum; GABAergic; TH; Dopamine; ACh; plateau potential

© 2015 Published by Elsevier Ltd.

Corresponding Authors: James M. Tepper, Ph.D., Center for Molecular and Behavioral Neuroscience, Rutgers The State University of New Jersey, 197 University Avenue, Newark NJ 07102, jtepper@andromeda.rutgers.edu, 973-353-3618 (phone), 973-353-1588 (fax), Tibor Koós, Ph.D., Center for Molecular and Behavioral Neuroscience, Rutgers The State University of New Jersey, 197 University Avenue, Newark NJ 07102, tibkoos@yahoo.com, 973-353-1080x3638 (phone), 973-353-1588 (fax).

^aDepartamento de Fisiología, Universidad Autónoma de San Luis Potosí, Av., Venustiano Carranza, San Luis Potosí, México.

^bDepartment of Physiology, Feinberg School of Medicine, Northwestern University Chicago, IL

Publisher's Disclaimer: This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final citable form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

1. Introduction

Until recently, the neostriatum has been thought to contain only a few types of GABAergic interneurons in comparison to the large diversity of GABAergic cell types in the neocortex or the hippocampus (Freund and Buzsaki, 1996; Tepper et al., 2010; DeFilipe et al., 2013). This picture changed significantly with the introduction of transgenic reporter mouse lines that revealed the existence of 5 new electrophysiologically distinct cell types, more than doubling the number of interneuron classes recognized in this brain area (Ibáñez-Sandoval et al., 2010; Ibanez-Sandoval et al., 2011). In addition to a neuropeptide Y (NPY) expressing neuron described in a NPY-GFP line (Ibanez-Sandoval et al., 2011), the newly discovered interneurons include 4 additional types of GABAergic neurons that were termed THinterneurons (THINs) reflecting their initial identification in a TH-EGFP strain (Ibáñez-Sandoval et al., 2010). The function of THINs remains unclear. Their small population size and connectivity place some important constraints on the possible function of these neurons. One particularly interesting possibility is that these neurons may distribute intra- and extrastriatal neuromodulatory signals to projection neurons.

In the neostriatum, dopamine (DA) and acetylcholine (ACh) are 2 major neuromodulators that exert pronounced effects on most functions of the basal ganglia. Here we investigated how these neuromodulators control the intrinsic electrophysiological properties of THINs. Since preliminary experiments indicated that the most salient effect of these modulators was the triggering and enhancement of a semi-stable depolarizing state we characterized in more detail this important dynamic feature of THINs.

2. Materials and Methods

2.1. Subjects

We used transgenic mice Tg (Th-EGFP) DJ76Gsat/Mmmc (GENSAT; Gong et al., 2003), obtained from the Mutant Mouse Regional Resource Center at UCLA and bred in our colony at Rutgers for all experiments. Hemizygous progeny were mated to wild type FVB or Swiss Webster mice each generation thereafter. All offspring were genotyped from tail samples and only those expressing the EGFP transgene were used in these experiments. Henceforth these mice are referred to as EGFP-TH mice.

All procedures were performed with the approval of the Rutgers University Institutional Animal Care and Use Committee and in accordance with the NIH *Guide to the Care and Use of Laboratory Animals* and all efforts were made to minimize the number of mice used and any possible discomfort.

2.2. Preparation of brain slices

Experiments were performed on brain slices obtained from adult EGFP-TH mice older than one month of age. Mice were deeply anesthetized with 150 mg/kg ketamine and 30 mg/kg xylazine i.p. and transcardially perfused with ice-cold, modified Ringer's solution containing (in mM) 248 sucrose, 2.5 KCl, 7 MgCl₂, 23 NaHCO₃, 1.2 NaH₂PO₄, 7 glucose, 1 ascorbate, 3 pyruvate, and bubbled with 95% O₂ and 5% CO₂ (pH 7.3). The brain was quickly removed into a beaker containing ice-cold oxygenated Ringer's and trimmed to a

block containing the striatum. Coronal or para-horizontal sections (250–300 Nm) were cut in the same medium using a Vibratome 3000 and immediately transferred to normal Ringer's solution containing (in mM) 124 NaCl, 2.5 KCl, 1.2 NaH₂PO₄, 26 NaHCO₃, 1.3 MgCl₂, 2 CaCl₂, 10 glucose, 1 ascorbate, 3 pyruvate, and 0.4 myo-inositol that was heated to 34°C and continuously bubbled with 95% O₂ and 5% CO₂ (pH 7.3) for one hour prior to recording and then maintained at room temperature until use. In some experiments, we substituted in an equimolar manner: choline Cl for NaCl, choline bicarbonate for NaHCO₃ and sucrose for CaCl₂. Slices were transferred to the recording chamber and submerged in continuously flowing oxygenated buffer (2–4 ml/minute) which was heated with an inline heater (Warner Instruments) to approximately 33°C.

2.3. Fluorescence and DIC Imaging and Recording

Slices were initially visualized under epifluorescence illumination with a high sensitivity digital frame transfer camera (Cooke SensiCam) mounted on an Olympus BX50-WI epifluorescence microscope and a 40× long working distance water immersion lens. Once an EGFP-TH neuron was identified, visualization was switched to infrared differential interference contrast (IR-DIC) microscopy for the actual patching of the neuron, usually performed under current clamp.

Micropipettes for whole cell recording were constructed from 1.2 mm o.d. and 0.94 mm i.d. borosilicate pipettes (Harvard apparatus) on a Narishige PP-83 vertical puller. The standard internal solution for whole cell current clamp recording was (in mM): 130 K gluconate, 10 NaCl, 2 MgCl₂, 10 HEPES, 3 Na₂ATP, 0.3 GTP, 1 EGTA plus 0.1–0.3% biocytin (pH 7.3 – 7.4). These pipettes typically exhibited a DC impedance of 4–6 MΩ measured in the recording chamber.

Current clamp recordings were made with a Neurodata IR-283 current clamp amplifier and voltage and/or current clamp recordings were made with a Multiclamp 700B amplifier (Molecular Devices, Sunnyvale, CA) whose output was filtered online with a second order Bessel filter at 1 kHz and digitized at 20–40 KHz with either a CED Micro 1401 Mk II and a PC running Signal v. 4 (Cambridge Electronic Design, Cambridge UK) or an ITC-16 and a Mac running Axograph. Acquired data was stored on a PC or Mac for offline analysis.

At the completion of the experiments slices containing biocytin-injected neurons were fixed by immersion in 4% paraformaldehyde-0.5% glutaraldehyde for 30 minutes at room temperature or microwaved to 60°C for 12 sec and stored overnight at 4°C.

2.4. Pharmacology

Drugs were applied in the perfusion medium or locally via a micropipette using a Picospritzer (General Valve, Fairfield, NJ), at 20 psi / 30 – 100 ms, at 0.1 Hz. Cobalt chloride, nimodipine, flufenamic acid, SCH-23390 hydrochloride, SKF-38393 hydrochloride, dopamine, choline bicarbonate and amphetamine were purchased from Sigma-Aldrich and tetrodotoxin (TTX), carbamylcholine chloride (carbachol), mecamlamine hydrochloride (MEC), Dihydro-β-erythroidine hydrobromide (DHβE) and methyllycaconitine citrate (MLA) were purchased from Tocris. Cobalt chloride was

prepared as an equimolar substitute for sucrose in modified Ringer's solution. Nimodipine and flufenamic acid were dissolved in dimethyl sulfoxide. All other drugs were dissolved freshly in Ringer's solution.

2.5. Extracellular stimulation

Stimulating electrodes consisted of concentric bipolar electrodes of 25 Nm at the tip, and 1 k Ω DC resistance were used (FHC, Bowdoinham, ME). Electrodes were placed onto the surface of the slice within 200 to 500 Nm of the recorded cells. Stimuli consisted of single square wave pulses (typically 0.01 – 1 mA, 200 Ns duration at 0.1 – 0.5 Hz) and was generated by a Winston A-65 timer and SC-100 constant current stimulus isolation unit (eg., Lee and Tepper 2007; Ibáñez-Sandoval et al., 2010).

2.6. Biocytin Histochemistry

Slices containing biocytin-filled neurons were transferred into 4% paraformaldehyde overnight. In some cases, the thick sections were resectioned on a Vibratome at 100 Nm. Sections were washed for 3 \times 10 minutes in 0.1 M phosphate buffer (PB) followed by 10% methanol and 3% H₂O₂ for 15 minutes, and incubated with avidin-biotin-peroxidase complex (Vector Laboratories; 1:200) and 0.1% Triton X-100 overnight at 4° C. After washing 6 \times 10 minutes in 0.1 M PB the sections were reacted with 3,-3'-diaminobenzidine (DAB; 0.025%) and H₂O₂ (0.0008%) in PB. In some cases nickel intensification (Adams, 1981) was used (2.5 mM nickel ammonium sulfate and 7 mM ammonium chloride in the DAB and H₂O₂ incubation). The sections were then postfixed in osmium tetroxide (0.1% in PB) for 30 minutes, dehydrated through a graded series of ethanol, followed by propylene oxide, and infiltrated overnight with a mixture of propylene oxide and epoxy resin (Durcupan; Fluka Chemie, Buchs, Switzerland). The sections were then transferred to fresh resin mixture for several hours and flat embedded between glass slides and coverslips and polymerized at 60°C for 24 hours.

2.7. Statistical Analysis

Most numerical values are reported as the mean \pm SEM. Data were analyzed by using ANOVA with Prism (GraphPad Software), followed with Tukey's post hoc tests to multiple comparisons means. Differences were considered to be significant at $p < 0.05$.

3. Results

3.1. Electrophysiological and anatomical properties of striatal TH interneurons

Ex vivo whole-cell recordings were obtained in current clamp mode from fluorescent neurons in the striatum of adult EGFP-TH mice. In agreement with previously reported data (Ibáñez-Sandoval et al., 2010), several distinct subtypes of THINs could be characterized on the basis of their intrinsic electrophysiological properties (input resistance, R_{in} , resting membrane potential (RMP), action potential duration at half-amplitude ($AP_{50\%}$), their ability or inability to maintain firing throughout modest depolarizing current injections and the presence of a low-threshold spike (LTS). These were : Type I ($R_{in} = 554 \pm 48 M\Omega$, RMP = -63 ± 2 mV and $AP_{50\%} = 1 \pm 0.08$ ms; $n = 45/68$ [66%]), Type II ($R_{in} = 379 \pm 79 M\Omega$, RMP = -75 ± 2 mV and $AP_{50\%} = 0.45 + 0.03$ ms; $n = 8/68$ [12%]), and Type IV ($R_{in} = 486$

$\pm 67 \text{ M}\Omega$, $\text{RMP} = -75 \pm 1 \text{ mV}$, $\text{AP}_{50\%} = 0.61 + 0.06 \text{ ms}$ and LTS; $n = 15/68$ [22%]). Type III striatal THINs that made up only about 5 % of the total population in our original study (Ibáñez-Sandoval et al., 2010) were not encountered in the present study, while the remaining subtypes were present in roughly the same proportions as reported previously. Representative examples of responses to current injection and the resulting current-voltage curves for Type I, II and IV THINs are shown in Figure 1.

Striatal THINs frequently (46%, 21/45 of Type I, and 25%, 2/8 of Type II) exhibited an intrinsic PP in response to strong depolarizing current pulses delivered at rest as previously described (Ibáñez-Sandoval et al., 2010). PPs ranged in duration from 15 ms to over 5 sec. A typical example of a PP from a Type I striatal THIN is shown in Figure 1.

During whole cell recordings, all neurons were filled with biocytin for subsequent anatomical study. Eleven neurons were chosen for further anatomical analysis. These neurons exhibited morphologies consistent with our previous reports (Ibáñez-Sandoval et al., 2010, Tepper et al., 2010). Striatal THIN somata were medium sized ($14.5 \pm 1 \mu\text{m} \times 10 \pm 0.5 \mu\text{m}$) and emitted from 2 to 5 thick primary dendrites that branched modestly, forming a simple dendritic arborization rarely exceeding 300 microns in diameter. The dendrites of many Type I neurons were sparsely invested with spine-like spines processes. The most characteristic feature of the THINs was their axon arborization. The axon originated from the soma or proximally from a primary dendrite, and branched extensively creating a dense arborization overlapping and extending beyond the dendritic field. Axons exhibited very prominent, evenly spaced varicosities, presumably boutons en passant, throughout the axonal field. A typical example is shown in Figure 1. The other three subtypes of THINs were aspiny, exhibited moderately varicose dendrites, and were indistinguishable anatomically from one another.

3.2. Ionic characterization of plateau potential of striatal TH interneurons

The ionic basis of the intrinsic PP was analyzed in current clamp recordings from a subset of striatal THINs exhibiting robust PPs. In this subset, the mean PP duration was $1258 \pm 344 \text{ ms}$ (range, 495 ms to 2626 ms, $n = 6$). PPs were characterized by a strong and sustained depolarization of the membrane potential, upon which truncated spikes sometimes rode. Substitution of Na^+ by choline (Fig. 2A1, red traces; Na^+ -free), abolished the spikes and significantly reduced the PP duration to $405 \pm 122 \text{ ms}$ (range: 59 to 869 ms, control vs. Na^+ -free, $p < 0.05$, $n = 6$). Subsequent substitution of Ca^{2+} by sucrose (Fig. 2A1, gray traces; Ca^{2+} -free), eliminated the PP entirely, reducing the duration to $5 \pm 3 \text{ ms}$ (range, 0 to 17 ms, control vs. Ca^{2+} -free, $p < 0.001$, $n = 6$). These results demonstrate that a sodium conductance was responsible for the generation of the spikes seen riding on the PP, which also contributes to the PP itself, and that Ca^{2+} entry was essential for the activation of the PP. Washing in normal buffer caused a partial recovery of both spikes and the PP at a duration of $795 \pm 199 \text{ ms}$ (range, 323 to 1323 ms, control vs. recovered $p > 0.05$, $n = 6$), as shown in Figure 2A1. Addition of either nimodipine (10 μM) or flufenamic acid (100 μM) completely abolished the PP (Fig. 2A1, blue traces) (range, 0 to 28 ms, control vs. NIM/FFA $p < 0.001$, $n = 6$). Taken together, these results confirm our previous results that the intrinsic PPs are due to a Na^+ and L-type Ca^{2+} dependent calcium-activated nonselective cation

current, I_{CAN} , as suggested previously (Ibáñez-Sandoval et al., 2010). I_{CAN} is likely related to a melastatin-related transient receptor potential cation channel (TRPM), most likely TRPM2 (Fleig and Penner, 2004; Hill et al., 2004; Lee and Tepper, 2007). Further, the strong inhibitory effect of nimodipine applied alone demonstrates that under these conditions L-type channels supply the intracellular Ca^{2+} required for I_{CAN} activation (Fleig and Penner, 2004; Hill et al., 2004; Lee and Tepper, 2007).

In striatal THINs that did not exhibit intrinsic PPs, TEA (20 mM) was capable of inducing their appearance, as illustrated in Figure 2B. In striatal THINs that exhibited short duration intrinsic PPs, 66 ± 34 ms (range, 10 to 299 ms, $n = 8$), TEA significantly prolonged the PP to 429 ± 65 ms (range, 227 to 800 ms, control vs. TEA, NIM and TTX $p < 0.001$, $n = 8$), even in the presence of nimodipine (10 μ M) and TTX (1 μ M). These results suggest that most or all striatal THINs are capable of generating a PP, but that under most conditions, it is partially or completely masked by a constitutively active TEA sensitive potassium conductance, similar to results reported previously for GABAergic neurons from substantia nigra pars reticulata (Lee and Tepper, 2007).

TEA promotes calcium influx into the cell by a different mechanism than that which is involved in control conditions. TEA-induced PPs are sustained in the presence of L-type Ca^{2+} channel blockade and TTX, but are abolished by flufenamic acid 10 ± 4 ms, (range, 0 to 32 ms, $n = 8$) as shown in Figure 2. Thus, although the PP is normally activated by L-type Ca^{2+} channels under normal conditions, the calcium conductance does not underlie the TEA induced PP itself.

These results indicate that both intrinsic and TEA-induced PPs are generated by the same current, I_{CAN} , similar to plateau potentials exhibited by other neurons in the basal ganglia (Baufreton et al, 2003; Hill et al, 2004; Lee and Tepper, 2007).

3.3. Dopamine modulation of plateau potentials in striatal THINs

Because of the importance of dopaminergic modulation in the basal ganglia and because 6-OHDA lesions reduce or abolish PPs in striatal THINs (Unal et al., 2013), we investigated dopaminergic modulation of the PP in striatal THINs.

In several striatal neurons, both those that exhibited intrinsic PPs and as well as those that did not (mean PP duration 43 ± 12 ms, range, 2 to 163 ms, $n = 12$), we explored the effects of amphetamine, which increases the extracellular DA concentration by causing DA release by reverse transport as well as by blocking reuptake (Khoshbouei et al., 2003; Khalig et al, 2005). In the presence of 50 μ M amphetamine (Fig. 3, compare black and blue traces), PPs could be induced or were increased in duration, 354 ± 30 ms (range, 335 to 577 ms, control vs. amphetamine $p < 0.001$, $n = 12$). PPs induced or prolonged by amphetamine were, like intrinsic PPs, completely blocked by nimodipine (5 μ M) or flufenamic acid (100 μ M) (Fig. 3, red traces), 36 ± 6 ms (range, 10 to 82 ms, amphetamine vs. NIM/FFA $p < 0.001$, $n = 12$). Thus, PPs can be induced or prolonged by endogenously released DA.

Similarly to amphetamine, 30 μ M DA induced or increased the duration of the PP 219 ± 34 ms (range, 69 to 386 ms; Fig. 3, compare black and green traces) to 710 ± 134 ms (range,

376 to 1932 ms, control vs. DA $p < 0.001$ $n = 11$). The effects of exogenous application of DA were also eliminated by nimodipine (5 μM) or flufenamic acid (100 μM), (Fig. 3, red traces), 22 ± 3 ms (range, 9 to 38 ms, DA vs. NIM/FFA $p < 0.001$, $n = 11$).

3.4. Dopamine effects on the plateau potential are mediated through D1/D5-like DA receptors

One way that DA could activate or facilitate PPs in striatal THINs is by enhancement of L-type Ca^{2+} currents that promote I_{CAN} activation. Previous reports showed that D1/D5-like receptor activation enhanced the L-type Ca^{2+} current and discharge of action potentials in striatal SPNs (Hernández-López et al., 1997; Surmeier et al., 1995). Thus, we explored whether DA effects on PPs in striatal THINs were mediated by DA D1/D5 receptors.

DA (30 μM) was bath applied and enhanced PPs as described above from a duration of 275 ± 61 ms (range, 34 to 475 ms; Fig. 4A1, compare black and red traces) to 2129 ± 862 ms (range, 641 to 6141 ms, $p < 0.01$, $n = 6$). Subsequent addition of 5 μM SCH-323390, a D1/D5 DA receptor antagonist, reversed the effect of DA as indicated by a significant reduction of the duration of the PP to 532 ± 92 ms (range, 357 to 965 ms; Fig. 4A2 see green traces, $p > 0.05$, $n = 6$). In some cases the PP was completely eliminated by SCH-323390 (Fig. 4A2). The effect of the antagonist (in the continued presence of DA) recovered partially after washout of SCH-23390 1811 ± 936 ms (range, 593 to 6429 ms; Fig. 4A2, compare green and blue traces, $p < 0.01$, $n = 6$). In THINs exhibiting DA-evoked PPs, the addition of 10 μM NIM or 100 μM FFA completely blocked the PPs; 51 ± 7 ms (range; 36 to 80 ms; Fig. 4A2, compare blue and magenta traces, ns, $n = 6$). In another group of striatal THINs that did not exhibit PPs in control, long lasting PPs could be induced by the D1/D5-receptor agonist SKF-38393 (5 μM) and these responses could be blocked by co-application of SCH-23390 (5 μM), a selective antagonist for D1/D5-like DA receptors (Fig. 4C1 and Fig. 4C2).

3.7. Striatal TH interneurons express nicotinic cholinergic receptors

Although ACh exerts important effects in the neostriatum through muscarinic receptors, in this study we were primarily interested in the possibility of fast especially nicotinic receptor mediated regulation of THINs. Therefore, we tested the effects of local pressure application of 100 μM carbachol on 3 of the 4 electrophysiologically defined subtypes of THINs including Type I ($n = 7$), II ($n = 5$) and IV ($n = 8$) neurons. Interestingly, the responses to carbachol were subtype-dependent. Type I THINs, the most common subtype comprising $> 65\%$ of THINs, responded to brief carbachol application with a strong depolarization that elicited an initial, approximately 100 ms long high frequency burst of rapidly adapting and accommodating spikes, similar to the bursts elicited with strong intracellular depolarizing current injections. The spikes were followed by a long-lasting, slowly decaying depolarization lasting for several hundred ms. We also examine the receptor subtype involved. At least 3 distinct brain nicotinic acetylcholine receptors can be distinguished using pharmacological means. This include the MLA and bungarotoxin sensitive Type 1 receptors that are primarily composed of $\alpha 7$ subunits, the $\beta 2$ subunit containing Type 2 receptors which can be identified by low micromolar block with DH β E and finally a less well characterized MLA and DH β E insensitive class of receptors (Type 3 receptors) that are

blocked by low micromolar MEC (Alkondon and Albuquerque, 1993). The depolarization was resistant to the Type I nicotinic receptor antagonist, MLA (500 nM), and the Type 2 nicotinic receptor antagonist, DH β E (1 μ M). However, all responses to carbachol were completely abolished by the weakly subtype-selective nicotinic receptor antagonist mecamylamine (MEC) applied at a concentration (5 μ M) at which Type 1 and 2 receptors are not affected Figure 5A1. In contrast, Type II THINs, responded to similar carbachol application with small-amplitude, brief depolarizations that rarely elicited action potentials and never triggered PPs (Figure 5A2). Finally, Type IV THINs, the second most abundant THIN subtype comprising approximately 25% of striatal THINs, also responded to carbachol with a robust, long lasting depolarization that produced sustained firing, similar to the responses of Type I THINs. The pharmacological profile of Type IV THINs was the same as that of the Type I neurons.

Carbachol elicited similar, large amplitude, long-lasting membrane depolarization in the presence of TTX (1 μ M) in Type I (n = 4) and Type IV (n = 3), THINs (Figure 7B) that was completely blocked by MEC (5 μ M). These data show that at least 2 types of THINs express nicotinic receptors. Interestingly, however, and similar to previous responses observed in striatal FSIs that also exhibit strong excitatory nicotinic responses to exogenous agonist application (Koós and Tepper, 2002), local electrical stimulation of the striatum failed to evoke any nicotinic synaptic response in THINs of any subtypes tested (data not shown).

4. Discussion

The present experiments examined the effects of DA and ACh on striatal THINs. The first principal finding was that intrinsic PPs elicited by depolarizing pulses in Type I and Type II THINs were facilitated by DA acting through a D1/D5-like DA receptor, and that in Type I and Type II THINs that did not exhibit intrinsic PPs, as well as in Type IV THINs that were never observed to exhibit intrinsic PPs, these plateaus could be elicited by D1/D5 receptor stimulation. In all cases, the PPs were primarily mediated by the non-selective cation conductance, I_{CAN} .

Finally, all subtypes of striatal THINs tested responded to brief pressure application of the non-selective cholinergic agonist, carbachol, with depolarization and excitation, whose strength greatly varied by subtype. In all cases, the response was shown to be due to direct activation of a nicotinic receptor distinct from the DH β E sensitive and MLA sensitive receptors.

4.1. Plateau potentials in striatal THINs are mediated by I_{CAN}

The present results elaborate on our earlier findings regarding the ionic mechanism of PPs and identify some of the specific contributions of I_{CAN} and L-type channels to the initiation and maintenance of PPs. Deletion of sodium from the extracellular solution greatly reduced the duration of the PPs but did not completely blocked them while subsequent removal of calcium and/or blockade of L-type Ca²⁺ channels by nimodipine completely eliminated the remaining PP. The PP was insensitive to TTX. In addition, the PP was completely blocked by flufenamic acid an antagonist of a melastatin-related transient receptor potential cation channel isoform (TRPM2) that underlies a calcium activated nonselective cationic

conductance, I_{CAN} (Fleig and Penner, 2004; Hill et al., 2004; Lee and Tepper, 2007; Lee et al., 2013). Moreover, as shown earlier (Ibanez_sandoval et al., 2011) blockade of L-Type channels prevents PPs in THINs in response to response to depolarization. Therefore, the most likely ionic mechanism of the initiation of the PP is Ca^{2+} entry through L-type channels in response to depolarization leading to activation of I_{CAN} . I_{CAN} activation may also involve Ca^{2+} induced Ca^{2+} release (CICR) from intracellular stores initiated by Ca^{2+} entry through L-type channels and possibly aided by a non-conventional $G_{q/11}$ -coupled IP_3 signaling pathway of D_1 DA receptors (Rashid et al., 2007).

Interestingly, although many Type I and Type II THINS expressed intrinsic THINS, in those cells that did not, PPs could be elicited in essentially all striatal THINS (including Type IV THINS, c.f. Figure 2B) by blocking a TEA-sensitive potassium conductance, as reported for other I_{CAN} expressing neurons of the basal ganglia (Lee and Tepper, 2007).

It is less clear what mechanism accounts for the termination of the PPs. Slow inactivation of L-type channels is one possibility. Alternatively if activation of I_{CAN} relies on calcium induced calcium release from intracellular stores store depletion or desensitization of IP_3 receptors may be involved.

4.2. Plateau potentials in striatal THINS are modulated by dopamine

The duration of intrinsic PPs was greatly increased by bath application of DA or the selective D1/D5 DA receptor agonist, SKF38393. Further PPs could be elicited in striatal THINS that did not exhibit intrinsic PPs by application of exogenous DA or amphetamine-induced release of endogenous DA. These effects were abolished by concomitant or subsequent application of the selective D1/D5 DA receptor antagonist, SCH-23390. These effects are consistent with a recent study that showed that intrinsic PPs in striatal THINS were greatly reduced in duration or completely eliminated in striata of mice that had received complete denervation of the nigrostriatal dopaminergic pathway one week earlier (Unal et al., 2013).

Like intrinsic PPs, dopamine-induced or facilitated PPs were completely abolished by nimodipine suggesting that dopamine promotes Ca^{2+} influx through L-type calcium channels or by flufenamic acid, that also blocks a non-selective cation conductance activated by calcium influx through L-type Ca^{2+} channels (Fleig and Penner, 2004; Hill et al., 2004; Lee and Tepper, 2007; Lee et al., 2013).

We did not attempt to identify the exact ionic mechanism of the dopaminergic enhancement of PPs. DA is well known to enhance L-type currents through D1/D5 receptors in a number of different neurons including SPNs (Surmeier et al., 1995; Hernández-López et al., 1997). Since the DA induced or modulated PPs in THINS were L-type channel dependent a similar mechanism may underly the effect of DA. Previous studies utilizing D1 knockout mice showed that the excitatory effects of dopamine and D1/D5 DA agonists on both striatal FSIs (Bracci et al., 2002) and PLTS (Centonze et al., 2002) interneurons persist in the D1 knockout mice suggesting that they are in fact mediated by the D5 receptor (Centonze et al., 2003). Both receptor subtypes have been shown to be expressed by various striatal neurons, with the D1 receptor being present in much greater abundance, especially by SPNs.

Interneurons, on the other hand, including PV+ FSIs, somatostatin- expressing PLTS neurons as well as cholinergic interneurons express relatively high levels of the D5 receptor while CR interneurons express low to moderate levels (Rivera et al., 2002). Perhaps D5 receptor expression is a general property shared by all striatal interneurons and that these receptors are also responsible for the dopaminergic facilitation of PPs in striatal THINs.

It is important to note that the effect of dopamine on PPs is the most dramatic direct effect on the excitability of any striatal neurons described to date. Of particular interest, this effect was readily elicited by endogenously released dopamine and therefore its magnitude is probably well within what can be expected to occur under physiological elevations of extracellular dopamine concentration *in vivo*.

4.4. Striatal TH interneurons express Type 3 nicotinic ACh receptors

Striatal ACh interacts with both metabotropic muscarinic and ionotropic nicotinic receptors at a variety of cellular and subcellular locations. Muscarinic receptors are widely expressed postsynaptically on striatal SPNs and interneurons where they modulate excitability in a complex state- and cell-type specific but generally facilitatory way (Figuroa et al., 2002; Perez-Rosello et al., 2005; Shen et al., 2005, 2007) Muscarinic receptors are also present on GABAergic (Koós and Tepper, 2002) and glutamatergic terminals (Pakhotin and Bracci, 2007) where they generally exhibit powerful presynaptic inhibitory effects (Zhou et al., 2003). Nicotinic receptors, in contrast, are largely restricted to striatal interneurons (Goldberg et al., 2012) where they exert excitatory effects on GABAergic interneuron firing (English et al., 2011; Koós and Tepper, 2002), and are also present on DA terminals (Marshall et al., 2000) where it has been proposed that they may elicit DA release independent of DA neuronal activity (Cachope et al., 2012; Threlfell et al., 2012).

Striatal THINs exhibit a depolarization in response to locally applied carbachol that was unaffected by MLA or DH β E, but abolished by mecamylamine, a selective Type 3 nicotinic receptor agonist. This nicotinic receptor profile differs from that striatal FSIs and NPY-NGF GABAergic interneurons (Koós and Tepper, 2002, English et al., 2012) but is consistent with a recent report by Luo and colleagues (2013) that found the same antagonist sensitivities for striatal Type I THINs as we report here, and additionally demonstrated that striatal Type I THINs respond to cytisine, a selective α 3 β 4 agonist nicotinic agonist. Here we show additionally that although all THINs tested exhibited a nicotinic excitation with a similar or identical pharmacology, there appeared to be subtype-specific differences in the strength of the excitation. The reasons for these differences remain unclear.

4.5. Physiological relevance

The nonspecific cationic conductance, I_{CAN} , is ubiquitously expressed in neurons from a number of different basal ganglia nuclei, including nigrostriatal dopaminergic neurons (Ping and Shepard, 1999; Yamashita and Isa, 2003, 2004), GABAergic substantia nigra pars reticulata projection (Lee and Tepper, 2007; Lee et al., 2013), glutamatergic subthalamic nucleus neurons (Baufreton et al., 2003), as well as striatal SPNs (Bao et al., 2005; Vergara et al., 2003) and THINs, and likely causes a generalized increase in the excitability of these neurons, amplifying excitatory afferents and affecting their firing pattern (Yamashita and

Isa, 2003; Lee and Tepper, 2007). DA activates THINS directly through a D1/D5 receptor, which in turn leads to powerful feed-forward inhibition of SPNs (Ibáñez-Sandoval et al., 2010). THINs are also subject to fast nicotinic excitation by ACh released from striatal cholinergic interneurons, as are NPY-NGF interneurons, which also provide feed-forward inhibition of SPNs (Ibáñez-Sandoval et al., 2011; English et al, 2012).

During the learning process in primates a cue associated with reward or other salient event evokes a pause in the tonic activity of cholinergic interneurons (CIN) in the striatum sometimes preceded by an increase in firing and often followed by a rebound excitation (Aosaki et al., 1994, 1995, 2010), while the same stimulus triggers an increase in the firing rate of midbrain dopaminergic neurons and striatal DA release concomitant with the pause part of the CIN sequence. While the pause may not affect the ongoing firing rate of striatal SPNs (English et al., 2012; but the situation may be different in the ventral striatum; see Witten et al., 2010), the rebound causes a powerful feed-forward disinaptic inhibition of SPNs. The SPN inhibition is caused by Type 2 nicotinic receptor mediated cholinergic activation of NPY-NGF GABAergic interneurons that produce a GABA_A IPSC in the SPN that exhibits unusually slow kinetics (Ibáñez-Sandoval et al., 2011). The rebound also activates a recurrent IPSC in the CIN mediated through a different, as yet unidentified, GABAergic interneuron that is also activated by Type 2 nicotinic receptors but that displays typical fast GABA_A kinetics (Sullivan et al., 2008; English et al., 2012).

Thus, the arrival of a salient, reward predicting stimulus will first trigger a dopaminergic response that will increase the excitability of striatal THINs, followed a few hundred ms later by a slower onset of prolonged activation of both THINs and NPY-NGF interneurons as the CINs rebound from the pause. The CIN rebound may also elicit a prolonged secondary release of DA by acting directly at nigrostriatal terminals (Cachope et al., 2012; Threlfell et al., 2012).

Consequently, the activation of feed-forward inhibition of SPNs following dopaminergic and cholinergic activation involves at least two different populations of GABAergic interneurons that are activated sequentially and whose inhibition of SPNs exhibits different time courses and kinetics. The precise roles that these two forms of feed-forward inhibition play in the signaling of salience and reward in striatum are not yet known, but for striatal THINs alone, the initial facilitation of their excitability by the nigrostriatal dopaminergic reward prediction error signal followed by the subsequent activation by the CIN rebound suggests that they play a role distinct from that of FSIs and NPY-NGF interneurons during the acquisition of associative learning in striatum.

Acknowledgements

This research was supported, in part, by NIH Grants 5R01NS034865 (J.M.T.), 1R01NS072950 (T.K. and J.M. T.) and Rutgers University. We thank Fulva Shah and Parth Gandhi for excellent technical assistance.

References

Adams JC. Heavy metal intensification of DAB-based HRP reaction product. *J. Histochem. Cytochem.* 1981; 29(6):775. [PubMed: 7252134]

- Alkondon M, Albuquerque EX. Diversity of nicotinic acetylcholine receptors in rat hippocampal neurons. I. Pharmacological and functional evidence for distinct structural subtypes. *J. Pharmacol. Exp. Therapeutics*. 1993; 265:1455–1473.
- Aosaki T, Graybiel AM, Kimura M. Effect of the nigrostriatal dopamine system on acquired neural responses in the striatum of behaving monkeys. *Science*. 1994; 265:412–415. [PubMed: 8023166]
- Aosaki T, Kimura M, Graybiel AM. Temporal and spatial characteristics of tonically active neurons of the primate's striatum. *J. Neurophysiol*. 1995; 73:1234–1252. [PubMed: 7608768]
- Aosaki T, Miura M, Suzuki T, Nishimura K, Masuda M. Acetylcholine–dopamine balance hypothesis in the striatum: An update. *Geriatr. Gerontol. Int*. 2010; 10(Suppl. 1):S148–S157. [PubMed: 20590830]
- Bao L, Avshalumov MV, Rice ME. Partial mitochondrial inhibition causes striatal dopamine release suppression and medium spiny neuron depolarization via H₂O₂ elevation, not ATP depletion. *J. Neurosci*. 2005; 25:10029–10040. [PubMed: 16251452]
- Baufreton J, Garret M, Rivera A, de la Calle A, Gonon F, Dufy B, Bioulac B, Taupignon A. D5 (not D1) dopamine receptors potentiate burst-firing in neurons of the subthalamic nucleus by modulating an L-type calcium conductance. *J. Neurosci*. 2003; 23:816–825. [PubMed: 12574410]
- Berke JD, Okatan M, Skurski J, Eichenbaum HB. Oscillatory entrainment of striatal neurons in freely moving rats. *Neuron*. 2004; 43:883–896. [PubMed: 15363398]
- Berke JD. Fast oscillations in cortical-striatal networks switch frequency following rewarding events and stimulant drugs. *Eur J Neurosci*. 2009; 30:848–859. [PubMed: 19659455]
- Berke JD. Functional properties of striatal fast-spiking interneurons. *Front Syst Neurosci*. 2011; 5:45. [PubMed: 21743805]
- Bibbig A, Faulkner HJ, Whittington MA, Traub RD. Self-organized synaptic plasticity contributes to the shaping of gamma and beta oscillations in vitro. *J Neurosci*. 2001; 21:9053–9067. [PubMed: 11698616]
- Bolam JP, Hanley JJ, Booth PA, Bevan MD. Synaptic organization of the basal ganglia. *J. Anat*. 2000; 196(Pt4):527–542. [PubMed: 10923985]
- Bracci E, Centonze D, Bernardi G, Calabresi P. Dopamine excites fast-spiking interneurons in the striatum. *J. Neurophysiol*. 2002; 87:2190–2194. [PubMed: 11929936]
- Buzsaki G. Neural syntax: cell assemblies, synapse ensembles, and readers. *Neuron*. 2010; 68:362–385. [PubMed: 21040841]
- Cachope R, Mateo Y, Mathur BN, Irving J, Wang HL, Morales M, Lovinger DM, Cheer JF. Selective activation of cholinergic interneurons enhances accumbal phasic dopamine release: setting the tone for reward processing. *Cell Rep*. 2012; 2:33–41. [PubMed: 22840394]
- Centonze D, Bracci E, Pisani A, Gubellini P, Bernardi G, Calabresi P. Activation of dopamine D₁-like receptors excites LTS interneurons of the striatum. *Eur. J. Neurosci*. 2002; 15:2049–2052. [PubMed: 12099911]
- Centonze D, Grande C, Usiello A, Gubellini P, Erbs E, Martín AB, Pisani A, Tognazzi N, Bernardi G, Moratalla R, Borrelli E, Calabresi P. Receptor subtypes involved in the presynaptic and postsynaptic actions of dopamine in striatal interneurons. *J. Neurosci*. 2003; 23:6245–6254. [PubMed: 12867509]
- DeCoteau WE, Thorn C, Gibson DJ, Courtemanche R, Mitra P, Kubota Y, Graybiel AM. Learning-related coordination of striatal and hippocampal theta rhythms during acquisition of a procedural maze task. *Proc Natl Acad Sci U S A*. 2007; 104:5644–5649. [PubMed: 17372196]
- DeFelipe J, López-Cruz PL, Benavides-Piccione R, Bielza C, Larrañaga P, et al. New insights into the classification and nomenclature of cortical GABAergic interneurons. *Nat. Rev. Neurosci*. 2013; 14:202–216. [PubMed: 23385869]
- Deister CA, Chan CS, Surmeier DJ, Wilson CJ. Calcium-activated SK channels influence voltage-gated ion channels to determine the precision of firing in globus pallidus neurons. *J Neurosci*. 2009; 29:8452–8461. [PubMed: 19571136]
- Dubois A, Savasta M, Curet O, Scatton B. Autoradiographic distribution of the D1 agonist [3H]SKF 38393, in the rat brain and spinal cord. Comparison with the distribution of D2 dopamine receptors. *Neuroscience*. 1986; 19:125–137. [PubMed: 2946980]

- Egorov AV, Draguhn A. Development of coherent neuronal activity patterns in mammalian cortical networks: common principles and local heterogeneity. *Mech Dev.* 2013; 130:412–423. [PubMed: 23032193]
- English DF, Ibáñez-Sandoval O, Stark E, Tecuapetla F, Buzsáki G, Deisseroth K, Tepper JM, Koós T. GABAergic circuits mediate the reinforcement-related signals of striatal cholinergic interneurons. *Nat. Neurosci.* 2011; 15:123–130. [PubMed: 22158514]
- Figuroa A, Galarraga E, Bargas J. Muscarinic receptor involved in the subthreshold cholinergic actions of neostriatal spiny neurons. *Synapse.* 2002; 46:215–223. [PubMed: 12373736]
- Fleig A, Penner R. The TRPM ion channel subfamily: molecular, biophysical and functional features. *Trends Pharmacol. Sci.* 2004; 25:633–639. [PubMed: 15530641]
- Galarraga E, Hernández-López S, Reyes A, Miranda I, Bermudez-Rattoni F, Vilchis C, Bargas J. Cholinergic modulation of neostriatal output: A functional antagonism between different types of muscarinic receptors. *J. Neurosci.* 1999; 19:3629–3638. [PubMed: 10212321]
- Gerfen CR, Surmeier DJ. Modulation of striatal projection systems by dopamine. *Annu Rev. Neurosci.* 2011; 34:441–466. [PubMed: 21469956]
- Goldberg JA, Ding JB, Surmeier DJ. Muscarinic modulation of striatal function and circuitry. *Handb Exp. Pharmacol.* 2012; 208:223–241. [PubMed: 22222701]
- Gong S, Zheng C, Doughty ML, Losos K, Didkovsky N, Schambra UB, Nowak NJ, Joyner A, Leblanc G, Hatten ME, Heintz N. A gene expression atlas of the central nervous system based on bacterial artificial chromosomes. *Nature.* 2003; 425:917–925. [PubMed: 14586460]
- Graybiel AM. Habits, rituals, and the evaluative brain. *Annu. Rev. Neurosci.* 2008; 31:359–387. [PubMed: 18558860]
- Guzman JN, Hernandez A, Galarraga E, Tapia D, Laville A, Vergara R, Aceves J, Bargas J. Dopaminergic modulation of axon collaterals interconnecting spiny neurons of the rat striatum. *J. Neurosci.* 2003; 23:8931–8940. [PubMed: 14523095]
- Hammond C, Bergman H, Brown P. Pathological synchronization in Parkinson's disease: networks, models and treatments. *Trends Neurosci.* 2007; 30:357–364. [PubMed: 17532060]
- Hawes SL, Gillani F, Evans RC, Benkert EA, Blackwell KT. Sensitivity to theta-burst timing permits LTP in dorsal striatal adult brain slice. *J Neurophysiol.* 2013
- Hernández-López S, Bargas J, Surmeier DJ, Reyes A, Galarraga E. D₁ receptor activation enhances evoked discharge in neostriatal medium spiny neurons by modulating an L-type Ca²⁺ conductance. *J. Neurosci.* 1997; 17:3334–3342. [PubMed: 9096166]
- Hill K, Benham CD, McNulty S, Randall AD. Flufenamic acid is a pH-dependent antagonist of TRPM2 channels. *Neuropharmacology.* 2004; 47:450–460. [PubMed: 15275834]
- Ibáñez-Sandoval O, Tecuapetla F, Unal B, Shah F, Koós T, Tepper JM. Electrophysiological and morphological characteristics and synaptic connectivity of tyrosine hydroxylase-expressing neurons in adult mouse striatum. *J. Neurosci.* 2010; 30:6999–7016. [PubMed: 20484642]
- Ibáñez-Sandoval O, Tecuapetla F, Unal B, Shah F, Koós T, Tepper JM. A novel functionally distinct subtype of striatal neuropeptide Y interneuron. *J. Neurosci.* 2011; 31:16757–16769. [PubMed: 22090502]
- Kahlig KM, Binda F, Khoshbouei H, Blakely RD, McMahon DG, Javitch JA, Galli A. Amphetamine induces dopamine efflux through a dopamine transporter channel. *Proc. Natl. Acad. Sci. U. S. A.* 2005; 102:3495–3500. [PubMed: 15728379]
- Khoshbouei H, Wang H, Lechleiter JD, Javitch JA, Galli A. Amphetamine-induced dopamine efflux. A voltage-sensitive and intracellular Na⁺-dependent mechanism. *J. Biol. Chem.* 2003; 278:12070–12077. [PubMed: 12556446]
- Koós T, Tepper JM. Dual cholinergic control of fast-spiking interneurons in the neostriatum. *J. Neurosci.* 2002; 22:529–535. [PubMed: 11784799]
- Lee CR, Tepper JM. A calcium-activated nonselective cation conductance underlies the plateau potential in rat substantia nigra GABAergic neurons. *J Neurosci.* 2007; 27:6531–6541. [PubMed: 17567814]
- Lee CR, Machold RP, Witkovsky P, Rice ME. TRPM2 channels are required for NMDA-induced burst firing and contribute to H₂O₂-dependent modulation in substantia nigra pars reticulata GABAergic neurons. *J Neurosci.* 2013; 33:1157–1168. [PubMed: 23325252]

- Leventhal DK, Gage GJ, Schmidt R, Pettibone JR, Case AC, Berke JD. Basal ganglia beta oscillations accompany cue utilization. *Neuron*. 2012; 73:523–536. [PubMed: 22325204]
- Lomb NR. Least-squares frequency analysis of unequally spaced data. *Astrophys Space Sci*. 1976; 39:447–462.
- Luo R, Janssen MJ, Partridge JG, Vicini S. Direct and GABA-mediate indirect effects of nicotinic ACh receptor agonists on striatal neurones. *J Physiol*. 2013; 591:203–217. [PubMed: 23045343]
- Marshall DL, Redfern PH, Wonnacott S. Presynaptic nicotinic modulation of dopamine release in the three ascending pathways studied by in vivo microdialysis: Comparison of naive and chronic nicotine-treated rats. *J Neurochemistry*. 2002; 68:1511–1519.
- Matsuda W, Furuta T, Nakamura KC, Hioki H, Fujiyama F, Arai R, Kaneko T. Single nigrostriatal dopaminergic neurons form widely spread and highly dense axonal arborizations in the neostriatum. *J Neurosci*. 2009; 29:444–453. [PubMed: 19144844]
- Ng GY, Mouillac B, George SR, Caron M, Dennis M, Bouvier M, O'Dowd BF. Desensitization, phosphorylation and palmitoylation of the human dopamine D1 receptor. *Eur J Pharmacol*. 1994; 267:7–19. [PubMed: 7515822]
- Pakhotin P, Bracci E. Cholinergic interneurons control the excitatory input to the striatum. *J Neurosci*. 2007; 27:391–400. [PubMed: 17215400]
- Perez-Rosello T, Figueroa A, Salgado H, Vilchis C, Tecuapetla F, Guzman JN, Galarraga E, Bargas J. Cholinergic control of firing pattern and neurotransmission in rat neostriatal projection neurons: Role of $Ca_v2.1$ and $Ca_v2.2$ Ca^{2+} channels. *J Neurophysiol*. 2005; 93:2507–2519. [PubMed: 15615835]
- Ping HX, Shepard PD. Blockade of SK-type Ca^{2+} -activated K channels uncovers a Ca^{2+} -dependent slow afterdepolarization in nigral dopamine neurons. *J Neurophysiol*. 1999; 81:977–984. [PubMed: 10085326]
- Popescu AT, Popa D, Pare D. Coherent gamma oscillations couple the amygdala and striatum during learning. *Nat Neurosci*. 2009; 12:801–807. [PubMed: 19430471]
- Power JM, Sah P. Competition between calcium-activated K^+ channels determines cholinergic action on firing properties of basolateral amygdala projection neurons. *J Neurosci*. 2008; 28:3209–3220. [PubMed: 18354024]
- Rashid AJ, So CH, Kong MMC, Furtak T, El-Ghundi M, Cheng R, O'Dowd BF, George SR. D1–D2 dopamine receptor heterooligomers with unique pharmacology are coupled to rapid activation of $G_q/11$ in the striatum. *PNAS* 2007. 2007; 104(2):654–659.
- Rivera A, Alberti L, Martín AB, Narváez JA, de la Calle A, Moratalla R. Molecular phenotype of rat striatal neurons expressing the dopamine D5 receptor subtype. *Eur. J. Neurosci*. 2002; 16:2049–2058. [PubMed: 12473072]
- Ruf T. The Lomb-Scargle Periodogram in Biological Rhythm Research: Analysis of Incomplete and Unequally Spaced Time-Series. *Biol Rhythm Res*. 1999; 30:178–201.
- Seeman P. Dopamine receptors and the dopamine hypothesis of schizophrenia. *Synapse*. 1987; 1:133–152. [PubMed: 2905529]
- Scargle JD. Studies in astronomical time series analysis. II - Statistical aspects of spectral analysis of unevenly spaced data. *ApJ*. 1982; 263:835–853.
- Schultz W. Predictive reward signal of dopamine neurons. *J Neurophysiol*. 1998; 80:1–27. [PubMed: 9658025]
- Sharott A, Moll CK, Engler G, Denker M, Grun S, Engel AK. Different subtypes of striatal neurons are selectively modulated by cortical oscillations. *J Neurosci*. 2009; 29:4571–4585. [PubMed: 19357282]
- Shen W, Hamilton SE, Nathanson NM, Surmeier DJ. Cholinergic suppression of KCNQ channel currents enhances excitability of striatal medium spiny neurons. *J. Neurosci*. 2005; 25:7449–7458. [PubMed: 16093396]
- Shen W, Tian X, Day M, Ulrich S, Tkatch T, Nathanson NM, Surmeier DJ. Cholinergic modulation of Kir2 channels selectively elevates dendritic excitability in striatopallidal neurons. *Nat. Neurosci*. 2007; 10:1458–1466. [PubMed: 17906621]
- Sciamanna G, Wilson CJ. The ionic mechanism of gamma resonance in rat striatal fast-spiking neurons. *J Neurophysiol*. 2011; 106:2936–2949. [PubMed: 21880937]

- Sullivan MA, Chen H, Morikawa H. Recurrent inhibitory network among striatal cholinergic interneurons. *J. Neurosci.* 2008; 28:8682–8690. [PubMed: 18753369]
- Surmeier DJ, Bargas J, Hemmings H C-Jr, Nairn AC, Greengard P. Modulation of calcium currents by a D1 dopaminergic protein kinase/phosphatase cascade in rat neostriatal neurons. *Neuron.* 1995; 14:385–397. [PubMed: 7531987]
- Surmeier DJ, Carrillo-Reid L, Bargas J. Dopaminergic modulation of striatal neurons, circuits, and assemblies. *Neuroscience.* 2011; 198:3–18. [PubMed: 21906660]
- Tepper JM, Tecuapetla F, Koós T, Ibáñez-Sandoval O. Heterogeneity and diversity of striatal GABAergic interneurons. *Front. Neuroanat.* 2010; 4:150. [PubMed: 21228905]
- Threlfell S, Lalic T, Platt NJ, Jennings KA, Deisseroth K, Cragg SJ. Striatal dopamine release is triggered by synchronized activity in cholinergic interneurons. *Neuron.* 2012; 75:58–64. [PubMed: 22794260]
- Unal B, Ibáñez-Sandoval O, Shah F, Abercrombie ED, Tepper JM. Distribution of tyrosine hydroxylase expressing interneurons with respect to anatomical organization of the neostriatum. *Front. Syst. Neurosci.* 2011; 5:41. [PubMed: 21713112]
- Unal B, Shah F, Kothari J, Tepper JM. Anatomical and electrophysiological changes in striatal TH interneurons after loss of the nigrostriatal dopaminergic pathway. *Brain Struct Funct.* 2013 in press.
- Vergara R, Rick C, Hernández-López S, Laville JA, Guzman JN, Galarraga E, Surmeier DJ, Bargas J. Spontaneous voltage oscillations in striatal projection neurons in a rat corticostriatal slice. *J. Physiol. (Lond).* 2003; 553:169–182. [PubMed: 12963790]
- Witten IB, Lin SC, Brodsky M, Prakash R, Diester I, Anikeeva P, Gradinaru V, Ramakrishnan C, Deisseroth K. Cholinergic interneurons control local circuit activity and cocaine conditioning. *Science.* 2010; 330:1677–1681. [PubMed: 21164015]
- Yang J, Ye M, Tian C, Yang M, Wang Y, Shu Y. Dopaminergic modulation of axonal potassium channels and action potential waveform in pyramidal neurons of prefrontal cortex. *J Physiol.* 2013; 591:3233–3251. [PubMed: 23568892]
- Yamashita T, Isa T. Fulfenamic acid sensitive, Ca²⁺-dependent in-ward current induced by nicotinic acetylcholine receptors in dopamine neurons. *Neurosci. Res.* 2003; 46:463–473. [PubMed: 12871768]
- Yamashita T, Isa T. Enhancement of excitatory postsynaptic potentials by preceding application of acetylcholine in mesencephalic dopamine neurons. *Neurosci. Res.* 2004; 49:91–100. [PubMed: 15099707]
- Zhou FM, Wilson CJ, Dani JA. Cholinergic interneuron characteristics and nicotinic properties in the striatum. *J. Neurobiol.* 2002; 53:590–605. [PubMed: 12436423]
- Zhou FM, Wilson CJ, Dani JA. Muscarinic and nicotinic cholinergic mechanisms in the mesostriatal dopamine systems. *Neuroscientist.* 2003; 9:23–36. [PubMed: 12580337]

Highlights

- Striatal TH interneurons express calcium and sodium dependent plateau potentials
- The plateau potentials are enhanced by dopamine or agonists of D1/D5 DA receptors
- Plateau potentials are mediated by I_{CAN}
- Striatal TH interneurons are also excited by ACh acting through Type 3 nicotinic receptors

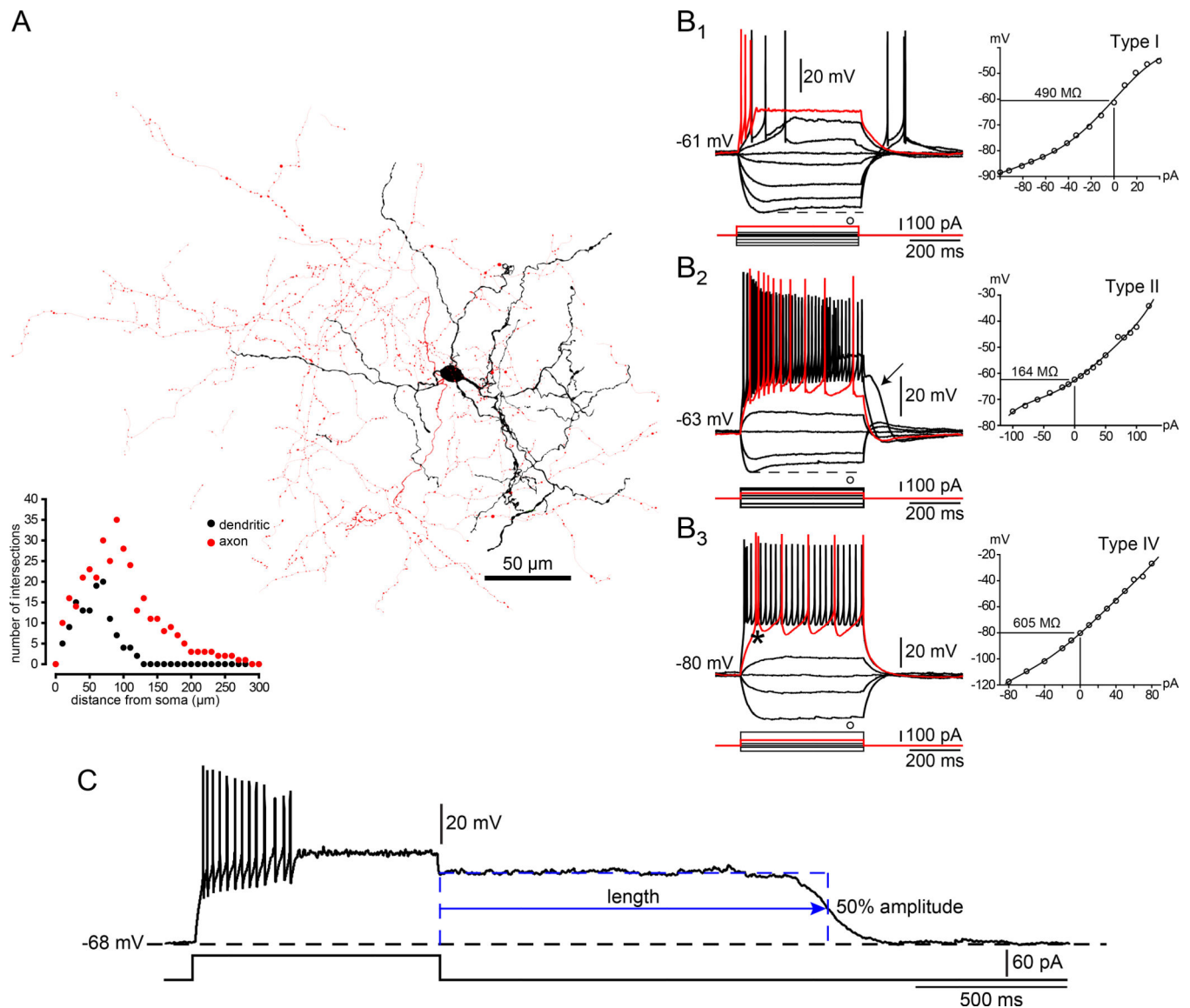


Figure 1. Anatomical and electrophysiological characteristics of striatal THINs recorded in adult EGFP-TH mice. **A.** Drawing tube reconstruction of a representative striatal Type I THIN filled with biocytin after whole-cell recording. Dendrites are drawn in black and the axon in red. Inset: Sholl plot data. **B.** Examples of voltage responses to hyperpolarizing and depolarizing current injection pulses illustrate the typical passive electrophysiological properties that distinguish Type I (B₁), Type II (B₂) and Type IV (B₃) striatal THINs. **C.** Type I THINs frequently exhibit (~41%) an intrinsic plateau potential (PP) following depolarizing from rest that long outlasts the stimulus, illustrated here to show methods used to quantify PPs.

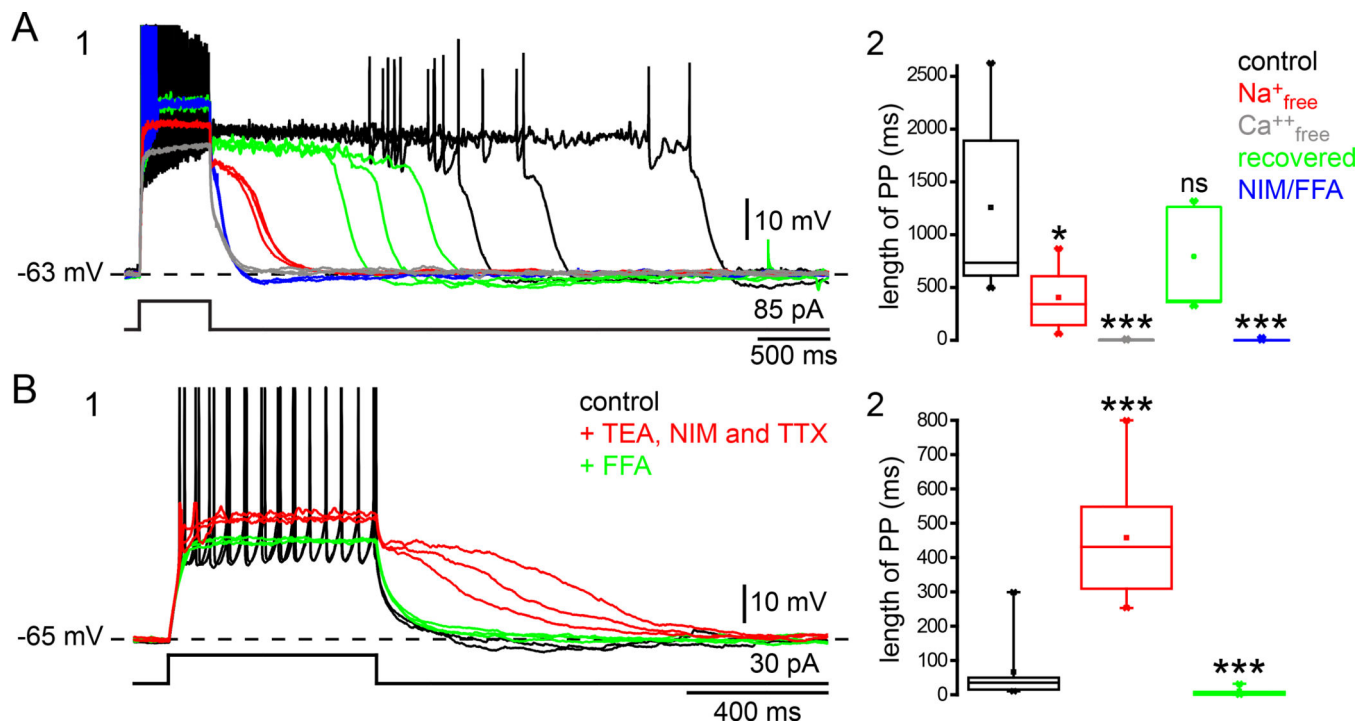


Figure 2.

Ionic mechanisms underlying the PP. A1. PPs evoked under control conditions (black traces) are reduced by over 75% when Na^+ was replaced by choline (red traces), illustrated here in a Type I THIN. Replacement of Ca^{2+} by sucrose completely eliminated the PP (gray traces). Partial recovery of the PP when buffer returned to control conditions (green traces). Either nimodipine (NIM; 10 μM), an L-type calcium channels blocker or flufenamic acid (FLU; 100 μM), a calcium-activated nonselective cation conductance blocker, completely blocked the PP (blue traces, data for NIM and FFA combined). A2. Summary box plots of six experiments. B1. Example of a striatal Type IV THIN that did not exhibit intrinsic PPs (black traces). In these cases, addition of TEA (20 mM) consistently unmasked PP (red traces), even in presence of NIM (10 μM) and TTX (1 μM). TEA evoked PPs were also completely eliminated by FFA (100 μM) (green traces). B2. Summary box plots of seven experiments. Box borders represent 25th and 75th percentiles. Whiskers represent data minima and maxima. Line represents median. Point represents mean. * indicates $p < 0.05$ and *** indicates $p < 0.001$ with respect to control. ns indicates $p > 0.05$ with respect to control. Action potentials are truncated to illustrate subthreshold events.

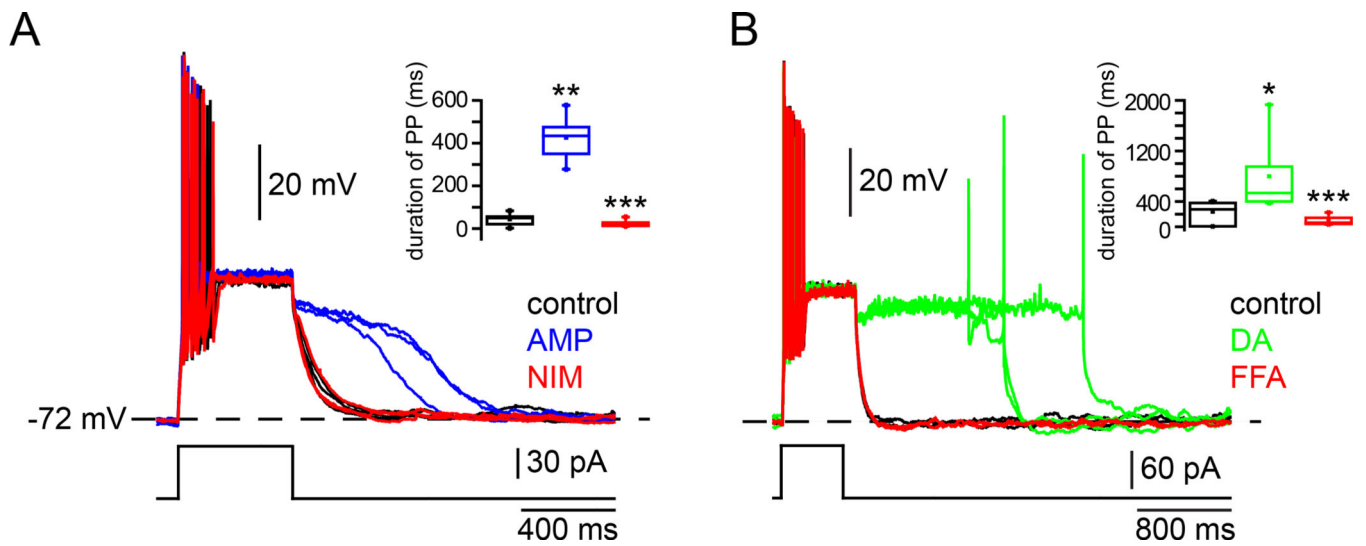


Figure 3. Dopaminergic modulation of PPs in striatal TH interneurons. A. Type I THIN that does not exhibit an intrinsic PP (black traces). D-Amphetamine (AMP; 50 μM) a PP (blue traces). The amphetamine-induced PP was blocked by nimodipine (NIM; 10 μM) or flufenamic acid (FFA; not shown). B. Another Type I THIN did not exhibited a plateau potential (black traces). Dopamine (DA; 30 μM) induces a long duration PP. Similar to the neuron in A either FFA or NIM (not shown) completely blocked the plateau potential.

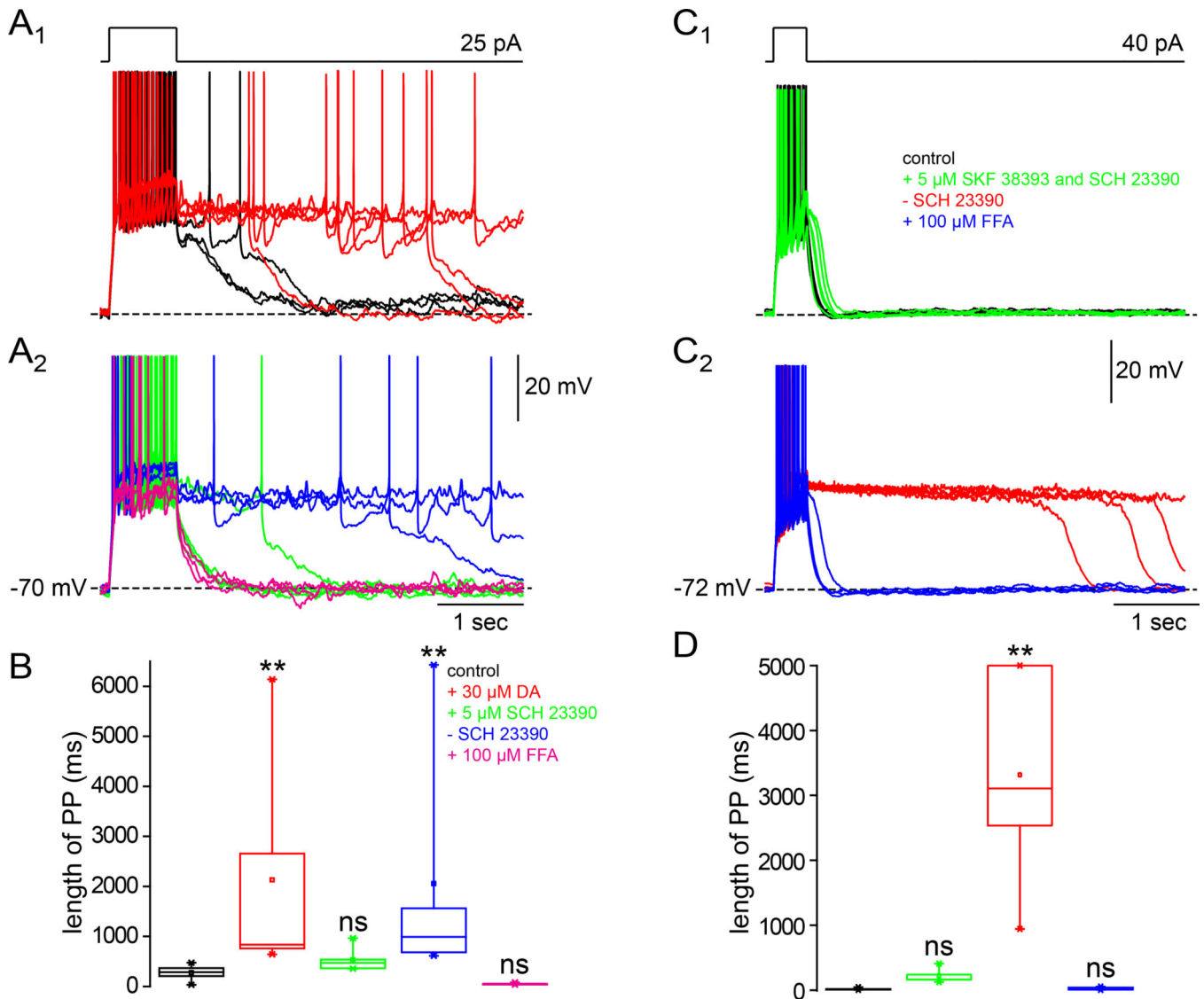


Figure 4.

Intrinsic PPs are facilitated by DA acting through D1/D5 DA receptors in striatal THINs. A1. Intrinsic PP in a Type I THIN (black traces) is greatly prolonged by addition of 30 μM DA (red traces). A2. Subsequent addition of SCH-23390 (5 μM), a selective D1/D5 DA receptor antagonist, first reduced and then completely blocked PPs (green traces). This effect washed out completely when the SCH-23390 was removed and the PP returned (blue traces). Either FFA (100 μM) or NIM (10 μM; not shown), completely blocked the PP. B. Summary box plots of six experiments on effects of DA and SCH 23390. See Figure 2 legend for explanation. C1. Striatal Type IV THIN does not exhibit PP (black traces) under in control condition. Simultaneous addition of both SCH-23390 (5 μM) and 5 μM SKF-38393 does not affect the PP (green traces). C2. Subsequent removal of SCH23390 reinstates the PP. The PP is completely blocked by addition of 100 μM FFA. Summary box plots of five experiments. Action potentials are truncated to illustrate subthreshold events.

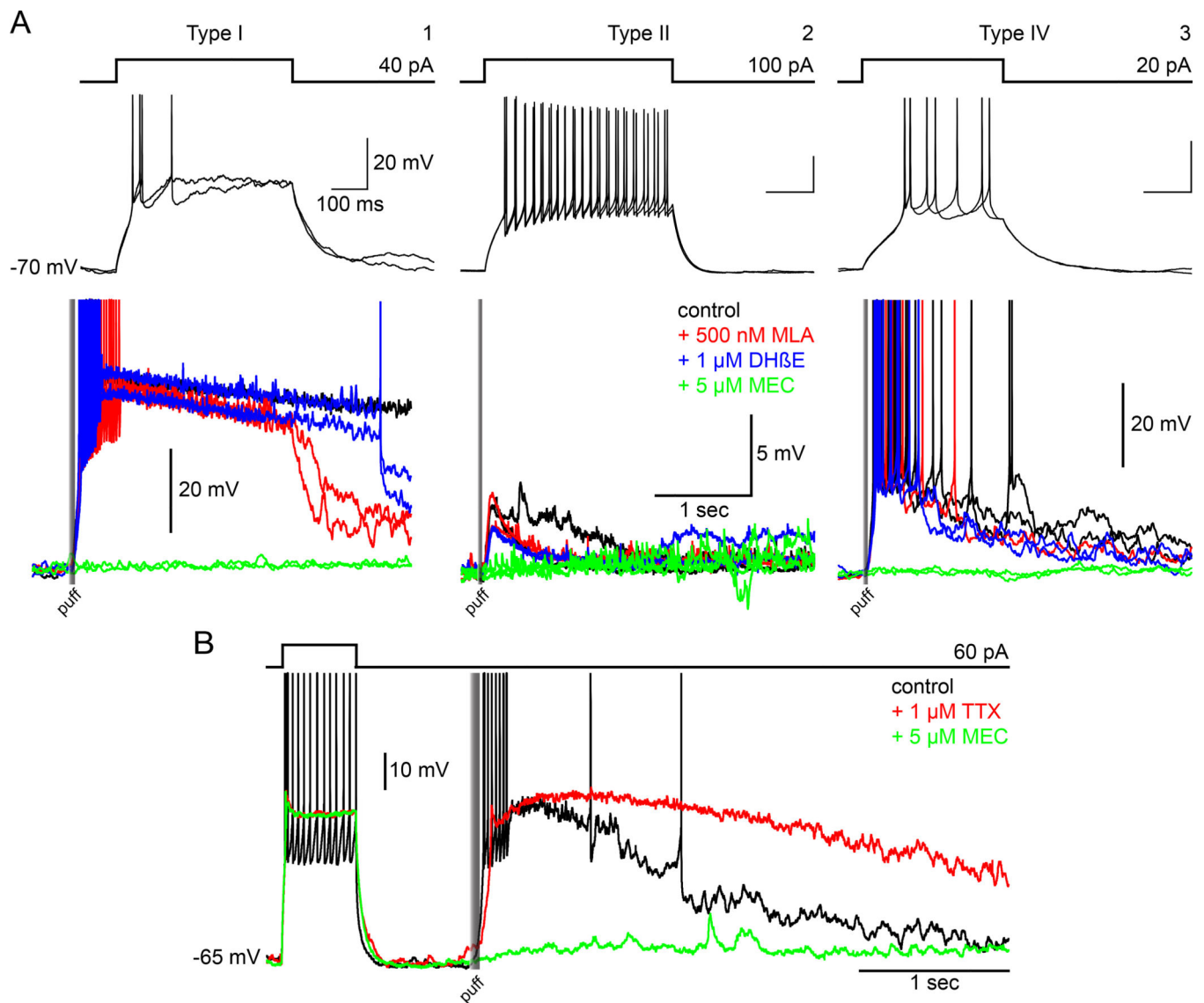


Figure 5. Striatal THINs are excited by Type 3 nicotinic cholinergic receptors. A1–3. Type I, II and IV striatal THINs are depolarized by local pressure application of the non-selective cholinergic agonist, carbachol. The effects of carbachol vary in intensity with THIN subtype. Type I shows a strong depolarization accompanied the onset of action potentials, followed by a long duration depolarization with spikes completely blocked, consistent with the response of Type I THINs to current injection (1). In contrast, Type II THINs showed a small but consistent depolarization that was accompanied on some occasions by action potentials (not shown) (2). Type IV THINs also exhibited a strong depolarization with spikes throughout, consistent with the response of this Type IV THINs to current injection (3). For all cell types, carbachol-induced depolarization was unaffected by MLA (500 nM), an antagonist of Type 1 nicotinic receptors (red traces) or by DHβE (1 μM), a Type 2 nicotinic receptor antagonist (blue traces) but was totally blocked by MEC (5 μM), a Type 3 nicotinic receptor antagonist (green traces). Note different voltage

calibrations for 1 and 3 vs. 2. B. The effect of carbachol persists even in presence of TTX (red trace), showing that it is a direct postsynaptic effect. In the present of TTX, MEC (5 μ M) still completely blocked the effects of carbachol (green trace). Action potentials are truncated to illustrate subthreshold events.

Author Manuscript

Author Manuscript

Author Manuscript

Author Manuscript