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Effects of drugs of abuse on putative rostromedial tegmental neurons, inhibitory afferents to midbrain dopamine cells.

Running title: Modulation of RMTg neurons by drugs of abuse.

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Abstract

Recent findings have underlined the rostromedial tegmental nucleus (RMTg), a structure located caudally to the ventral tegmental area, as an important site involved in the mechanisms of aversion. RMTg contains γ -aminobutyric acid (GABA) neurons responding to noxious stimuli, densely innervated by the lateral habenula and providing a major inhibitory projection to reward-encoding midbrain dopamine (DA) neurons.

One of the key features of drug addiction is the perseverance of drug seeking in spite of negative and unpleasant consequences, likely mediated by response suppression within neural pathways mediating aversion. To investigate whether the RMTg plays a role in the mechanisms of addicting drugs, we studied acute effects of morphine, cocaine, the cannabinoid agonist WIN55212-2 (WIN) and nicotine on putative RMTg neurons. We utilized single unit extracellular recordings in anesthetized rats and whole-cell patch clamp recordings in brain slices to identify and characterize putative RMTg neurons and their responses to drugs of abuse.

Morphine and WIN inhibited both firing rate in vivo and excitatory postsynaptic currents (EPSCs) evoked by stimulation of rostral afferents in vitro, whereas cocaine inhibited discharge activity without affecting EPSC amplitude. Conversely, nicotine robustly excited putative RMTg neurons and enhanced EPSCs, an effect mediated by $\alpha 7$ -containing nicotinic acetylcholine receptors.

Our results suggest that activity of RMTg neurons is profoundly influenced by drugs of abuse and, as important inhibitory afferents to midbrain DA neurons, they might take place in the complex interplay between the neural circuits mediating aversion and reward.

Keywords

Lateral habenula, aversion, addiction, patch clamp, electrophysiology, Fos, rat.

Introduction

Recent studies have provided information on how brain regions encoding aversion and reward are integrated. Mesolimbic dopamine (DA) neurons encode rewarding and appetitive stimuli with phasic excitation, whereas reward omission induces phasic inhibition (Schultz, 2007a, b). Aversive stimuli have been reported to induce both excitation (Brischoux *et al*, 2009) and inhibition (Ungless *et al*, 2004) of DA neurons, a heterogeneous response correlated to specific subgroups of DA neurons (ventral and dorsal, respectively) within the ventral tegmental area (VTA). On the other hand, glutamatergic neurons in the lateral habenula (LHb), an epithalamic region involved in the mechanisms of fear, anxiety and stress, respond in a reverse fashion, being inhibited by rewards and excited by aversive stimuli (Hikosaka *et al*, 2008; Matsumoto and Hikosaka, 2007). Noteworthy, activity of DA and LHb neurons appears to be causally correlated, since electrical stimulation of the LHb inhibits DA neurons (Christoph *et al*, 1986; Ji and Shepard, 2007; Matsumoto and Hikosaka, 2007). However, the sparse innervation of DA neurons by excitatory LHb afferents (Brinshawitz *et al*, 2010; Omelchenko *et al*, 2009) unlikely explains this inhibition, and the presence of an area intermediate between the LHb and the VTA was originally postulated. Elegant studies by Jhou and colleagues (Jhou *et al*, 2009a; Jhou *et al*, 2009b) revealed that γ -aminobutyric acid (GABA) neurons in the rostromedial tegmental nucleus (RMTg), a region also denominated as "tail" of the VTA (Perrotti *et al*, 2005), integrate input from the LHb and send output projection to midbrain DA neurons. As described, the missing link between LHb and midbrain DA neurons is provided. Notably, both RMTg and LHb neurons are phasically activated by aversive stimuli and inhibited by appetitive stimuli (Jhou *et al*, 2009a). In turn, RMTg neurons form inhibitory synapses with DA neurons in the VTA and the substantia

nigra (Balcita-Pedicino *et al*, 2009), and are now accounted among the major inhibitory afferents to mesencephalic DA neurons (Jhou *et al*, 2009a). The electrical activity of DA neurons, and the resulting DA output in terminal areas, crucially depends on the balance between excitatory and inhibitory inputs (Marinelli *et al*, 2006). Thus, the characterization of this novel inhibitory pathway to DA neurons is of interest to understand how DA neurons respond to behavioral and pharmacological stimuli.

Understanding the relationships between brain areas encoding reward and aversion, such as the VTA and the RMTg, is also relevant when it concerns the effects of addicting drugs. Since drug abuse potential can be envisaged as a balance between rewarding and aversive effects (Hutchison and Riley, 2008; Simpson and Riley, 2005; Wise *et al*, 1976), it can be hypothesized that the components of the appetitive properties of addicting drugs might result from both direct activation of neural reward pathways and suppression of responses in neural circuits mediating aversion. Indeed, one of the hallmarks of drug addiction is the perseverance of drug taking despite of negative and unpleasant consequences of drug use. Recent studies indicate that drugs of abuse affect GABA neurons in the RMTg: these neurons, and notably those projecting to the VTA, express Δ FosB (Kaufling *et al*, 2009; Perrotti *et al*, 2005) and c-Fos following repeated cocaine exposure (Geisler *et al*, 2008). Additionally, RMTg neurons are immunoreactive against μ -opioid receptors (Jhou *et al*, 2009b) and supposedly respond to opioid administration (Jhou *et al*, 2009c).

To determine further whether RMTg neurons are a primary target of drugs of abuse, we took advantage of extracellular single unit recordings in anesthetized rats together with whole-cell patch clamp recordings in brain slices. We characterized the electrophysiological features of putative neurons in the RMTg, identified by their excitatory responses to both a noxious stimulus

(i.e. paw pinch) and LHb stimulation. We also studied the effects of the prototypical psychostimulant and opioid (cocaine and morphine, respectively), the type 1-cannabinoid (CB1) receptor agonist WIN55212,2 and nicotine on firing rate in vivo and on excitatory postsynaptic currents (EPSCs) in vitro.

Materials and Methods

In vivo electrophysiology.

Male Sprague Dawley albino rats (Harlan Nossan, San Pietro al Natisone, Italy) weighing 250 to 350g were used in all in vivo experiments. Experiments were performed in strict accordance with the Guidelines for the Care and Use of Mammals in Neuroscience and Behavioral Research (National Research Council 2004) and EEC Council Directive of 24 November 1986 (86/609).

All efforts were made to minimize pain and suffering and to reduce the number of animals used.

Animals were housed in groups of three to six in standard conditions of temperature and humidity under a 12 h light/dark cycle (with lights on at 7:00 A.M.) with food and water available *ad libitum*. We anesthetized rats with urethane (1300 mg/kg, i.p.), cannulated their femoral vein for intravenous administration of pharmacological agents, and placed them in the stereotaxic apparatus (Kopf, Tujunga, CA, USA) with their body temperature maintained at $37 \pm 1^\circ\text{C}$ by a heating pad.

Experiments in the RMTg. The skull surface was exposed and a burr hole was drilled over the RMTg (7.0-7.4 mm posterior to bregma, 0.8-1.0 mm lateral to the midline, 6.5-7.5 mm ventral to the cerebellar cortex) for the insertion of a recording electrode. For orthodromic stimulation experiments, a Formvar-coated stainless steel bipolar electrode (250 μm tip diameter) was inserted in the ipsilateral LHb (1.9 mm posterior to bregma, 0.7 mm lateral to the midline, 4.7 mm ventral to the cerebral cortex) with an inclination of 20° anteroposterior on the coronal plane.

Structures were localized according to the stereotaxic atlas of Paxinos and Watson (2007). Single unit activity of RMTg cells was recorded extracellularly by glass micropipettes filled with 2% Pontamine sky blue dye dissolved in 0.5 M sodium acetate (impedance 5-10 M Ω). Putative GABA-containing neurons in the RMTg were first isolated and identified according to previously described electrophysiological characteristics (Jhou *et al*, 2009a), including a relatively high spontaneous firing rate (>10Hz) and a biphasic and short (<1.5 ms, see Results for more details) action potential. Once a cell was selected, stimuli (~0.5 mA) were delivered to the LHb at 1 Hz. Responses to electrical stimulation of the LHb were evaluated and a peri-stimulus time histogram (PSTH) was generated on-line (Spike 2 software, CED, Cambridge, UK) for each neuron. Only RMTg neurons with a robust excitatory response (latency range 2-8 ms) to LHb stimulation were selected in the present study. We did not include cells whose onset latencies were longer than 8 ms following LHb stimulation because they could exhibit a polysynaptic response component.

The extracellular neuronal signal was filtered (bandpass 1-3 KHz) and amplified (Neurolog System, Digitimer, Hertfordshire, UK), displayed on a digital storage oscilloscope (TDS 3012, Tektronix, Marlow, UK) and digitally recorded. Experiments were sampled on- and off-line by a computer connected to CED Power 1401 laboratory interface (Cambridge Electronic Design, Cambridge, UK). The spontaneous firing rate was recorded for 2-3 min to establish a baseline measure of firing rate. Interspike intervals and coefficient of variation (CV = standard deviation of interspike intervals divided by the mean interspike interval; a measure of firing regularity) were also determined.

Drugs were administered *in bolus* i.v. (1 ml/kg of body weight). Changes in firing rate and regularity were calculated by averaging the effects of the drugs for the 2 min period following

drug administration and comparing them to the mean of the pre-drug baseline. When drugs were administered, only one cell was recorded per rat.

At the end of recording sessions, DC current (15 mA for 5 min) was passed through the recording micropipette in order to eject Pontamine sky blue for marking the recording site. Brains were then rapidly removed and fixed in 4% paraformaldehyde solution. The position of the electrodes was microscopically identified on serial sections (60 μ m) stained with Neutral Red.

Experiments in the VTA. The scalp was retracted and one burr hole was drilled above the VTA (6.0 mm posterior from bregma, 0.3–0.6 mm lateral from midline) for the placement of a recording electrode. To evaluate the inhibitory input arising from the RMTg to the VTA, a Formvar-coated stimulating stainless steel bipolar electrode (250 μ m tip diameter) was inserted in the ipsilateral RMTg (9.6 mm posterior from bregma, 0.8 mm lateral from the midline, 7.0 mm ventral from the cortical surface) with an inclination of 20° anteroposterior on the coronal plane. Single unit activity of neurons located in the VTA (V 7.0–8.0 mm from the cortical surface) was recorded extracellularly with the same instruments previously described for the RMTg experiments. Single units were isolated and identified according to already published criteria (Grace and Bunney, 1983, 1984; Ungless *et al*, 2004). We recorded VTA DA neurons when all criteria for identification were fulfilled: firing rate ≤ 10 Hz, duration of action potential ≥ 2.5 ms, inhibitory responses to hindpaw pinching. Once a cell was selected, stimuli (~ 0.5 mA) were delivered to the RMTg at 1 s intervals and stimulation currents were progressively adjusted to induce an inhibitory response after each stimulus. The experimental protocol was essentially that published by Ji and Shepard (2007). Responses to electrical stimulation of RMTg were

evaluated and a peri-stimulus time histogram (PSTH) was generated on-line (Spike 2) for each neuron. Histological verification was carried out as described above.

In vitro electrophysiology.

Experiments in the RMTg. Whole cell patch clamp recordings from Sprague Dawley rat cells were as described previously (Melis *et al*, 2008). Briefly, male rats (14-31 d) were anesthetized with halothane and killed. A block of tissue containing the midbrain was rapidly dissected and sliced in the horizontal plane (300 μm) with a vibratome (VT1000S, Leica) in ice-cold low- Ca^{2+} solution containing (in mM): 126 NaCl, 1.6 KCl, 1.2 NaH_2PO_4 , 1.2 MgCl_2 , 0.625 CaCl_2 , 18 NaHCO_3 , and 11 glucose. Slices were transferred to a holding chamber with artificial cerebrospinal fluid (ACSF, 37° C) saturated with 95% O_2 and 5% CO_2 containing (in mM): 126 NaCl, 1.6 KCl, 1.2 NaH_2PO_4 , 1.2 MgCl_2 , 2.4 CaCl_2 , 18 NaHCO_3 , and 11 glucose. Slices (two per animal) were allowed to recover for at least 1 hr before being placed (as hemislices) in the recording chamber and superfused with the ACSF (37° C) saturated with 95% O_2 and 5% CO_2 . Putative RMTg cells were visualized with an upright microscope with infrared illumination (Axioskop FS 2 plus, Zeiss), and whole-cell current- and voltage-clamp recordings (one per hemislice) were made by using an Axopatch 200B amplifier (Molecular Devices, CA). Current-clamp experiments were made with electrodes filled with a solution containing the following (in mM): 144 KCl, 10 HEPES, 3.45 BAPTA, 1 CaCl_2 , 2.5 Mg_2ATP , and 0.25 Mg_2GTP (pH 7.2-7.4, 275-285 mOsm). All voltage-clamp recordings were made with electrodes filled with a solution containing the following (in mM): 117 Cs methansulfonic acid, 20 HEPES, 0.4 EGTA, 2.8 NaCl, 5 TEA-Cl, 2.5 Mg_2ATP , and 0.25 Mg_2GTP (pH 7.2-7.4, 275-285 mOsm). Experiments were

begun only after series resistance had stabilized (typically 15-40 M Ω). Series and input resistance were monitored continuously on-line with a 5 mV depolarizing step (25 ms). Estimated resting potential was calculated by averaging V_m values measured 1 ms before the initiation of each spontaneous action potential. Data were filtered at 2 kHz, digitized at 10 kHz, and collected on-line with acquisition software (pClamp 8.2, Molecular Devices, CA). A bipolar stainless steel stimulating electrode (FHC, USA) was placed 150 μ m rostral to the recording electrode to stimulate (duration 50 μ s) the afferent fibers at a frequency of 0.1 Hz. Paired stimuli were given with an interstimulus interval of 50 ms, and the ratio between the second and the first EPSC was calculated and averaged for a 5 min baseline. Each slice received only a single drug exposure.

Stimulation of RMTg FOS expression and immunofluorescent staining.

In a separate set of experiments, animals were first prepared for in vivo recordings as described above. Intraperitoneal injection of (+) methamphetamine hydrochloride (10 mg/kg in 0.9% saline; 1 ml/kg) given two hours prior sacrifice was used to produce Fos expression in the RMTg, as previously described by Jhou et al. (2009b). See “Supplementary Materials and Methods” for detailed information.

Drugs.

Morphine and cocaine hydrochloride were purchased from S.a.l.a.r.s (Como, Italy) and Akzo Pharmadivision Diosynth (Oss, Netherlands), respectively. Nicotine ((-)-nicotine hydrogen

tartrate), mecamylamine and (+) methamphetamine hydrochloride were purchased from Sigma (Milano, Italy). WIN55212-2, methyllycaconitine and AM281 were purchased from Tocris (Bristol, UK). Naloxone was purchased from RBI (Natick, MA, USA). SR141716A was a generous gift of Sanofi-Aventis Recherche (Montpellier, France). For experiments in vivo WIN55212-2 and SR141716A were emulsified in 1% Tween 80, then diluted in saline and sonicated. All other drugs were diluted in saline. Nicotine solution was adjusted to pH=7.0 with NaOH. For experiments in vitro, all drugs were applied in known concentrations to the superfusion medium and, with the exception of WIN 55,212-2 and AM281, were dissolved in saline. WIN 55,212-2 and AM281 were dissolved in DMSO as stock concentrations and the final concentration of DMSO was < 0.01 %.

Statistical Analysis.

Drug-induced changes in rate and regularity of action potential activity were calculated by averaging the effects after drug administration (2 min) and normalizing to the pre-drug baseline. All the numerical data are given as mean \pm SEM. Data were compared and analyzed by using two-way ANOVA for repeated measures (treatment x time), or one-way ANOVA or Student's *t* test for repeated measures, when appropriate. *Post hoc* multiple comparisons were made using the Dunnett's or Bonferroni's tests. Statistical analysis was performed by means of the Graphpad Prism software (La Jolla, CA). The significance level was established at $p < 0.05$.

Results

Electrophysiological characterization of RMTg neurons

We recorded the spontaneous activity of single putative RMTg neurons ($n=41$, from 9 rats) in urethane-anesthetized rats. Relatively high spontaneous firing rate (>10 Hz) and short (<1.5 ms), biphasic action potential of RMTg cells recorded extracellularly were consistent with those recently described in behaving rats (Jhou *et al*, 2009a). The action potential waveform of RMTg neurons ($n=41$), filtered using standard bandpass settings of 1-3 KHz, were characterized by an initial negative phase followed by a positive one (see Fig. 1a). The mean duration from the start to the positive trough was 1.26 ± 0.04 ms (range 0.90-1.49 ms, $n=41$). The width of the neuronal action potentials was longer in our study when compared with that of Jhou and collaborators (2009a) because of the difference in the experimental conditions (anesthetized versus behaving rats, filtering parameters). In general, RMTg neurons fired regularly at an average basal rate of 15.5 ± 1.0 Hz ($n=41$) (Fig. 1a). The regularity of firing was quantified by the CV of interspike intervals (mean CV= $36.8\pm 3.3\%$, $n=41$) (see Material and methods for CV detail). A typical interspike interval histogram showing the relatively constant interspike intervals, and their unimodal distribution peaking at around 50 ms, is shown in Figure 1a.

All in vitro data presented were obtained from cells ($n= 57$) located within the RMTg, identified in the horizontal slice as the region posterior (~ 1 mm) to the VTA (Fig. 1b), which is in turn placed medially to the medial terminal nucleus of the accessory optic tract (MT) and laterally to the interpeduncular fossa (IPF) (Paxinos and Watson) (2007). Under current-clamp mode in a whole-cell configuration, all of the neurons recorded ($n= 22$) were spontaneously active with a

mean firing rate of 4.7 ± 0.3 Hz within a range of 3-9 Hz (Fig. 1c). The estimated resting membrane potential was -44.5 ± 9.9 mV, and putative RMTg cells fired short-duration spikes (mean width at threshold: 1.77 ± 0.01 ms), whose amplitude and input resistance were 65.0 ± 0.8 mV and 304 ± 37 G Ω , respectively. These electrophysiological features are similar to those described for GABAergic neurons of the neighbor pars reticulata of the substantia nigra recorded under the same conditions (Atherton and Bevan, 2005). EPSCs were evoked by repetitively stimulating the afferents (duration 50 μ s, frequency 0.1 Hz, intensity 0.01-0.30 mA) with a stimulating electrode that was placed rostro-laterally (~ 150 μ m) from the recording electrode (Fig. 1b), and pharmacologically isolated by continuous perfusion of the GABA_A antagonist picrotoxin (100 μ M). Bath application of 10 μ M 6-cyano-7-nitroquinoxaline-2,3-dione (CNQX) completely suppressed these currents (Fig. 1c) indicating that they are AMPA-mediated EPSCs. EPSCs had a half rise time of 1.36 ± 0.20 ms ($n=35$) and a decay time constant of 4.71 ± 0.06 ms ($n = 35$). Although we cannot identify definitively the sources of excitatory afferents in this slice preparation, we can assume that most of the rostral inputs electrically stimulated in our preparation are presumably originating from the LHb, given that it is one of the major excitatory sources to the RMTg passing through the VTA (Jhou *et al*, 2009a).

Jhou and co-workers (2009a) recently showed that aversive stimuli induce Fos expression in VTA-projecting RMTg neurons, and phasically stimulate their electrical activity in behaving rats. Notably, a substantial number of VTA-projecting RMTg cells receives a major afferent from the LHb (Jhou *et al*, 2009b), thus corroborating the idea that RMTg might signal aversive events and inhibit midbrain DA neurons. Hence, *in vivo* we first tested the responses of putative RMTg neurons to a standard aversive somatosensory stimulus (hindpaw mechanical

stimulation), which is known to exceed the threshold of pain in freely moving animals (Cahusac *et al.*, 1990).

Following 2-3 minutes of stable baseline firing rate, a brief (approximately 2-4 s) pinch was applied to the hindpaw ipsilateral to the recording site and the response analyzed during the 5 s after stimulus onset (Fig. 2a). In agreement with recent data (Jhou *et al.*, 2009a), we found that a majority (73.2 %, 30 out of 41) of putative RMTg neurons were significantly stimulated by paw pinch. The overall response of RMTg neuronal population to the pinch stimulus is illustrated in Fig. 2a: phasic excitation peaked at around 2 s after the onset of the paw pinch ($237.3 \pm 40.9\%$ of baseline firing rate; $n=12$; $F_{13,43}=6.7$; $p<0.0001$; one way ANOVA and Dunnett's test).

We next examined the response of putative RMTg neurons to electrical stimulation of the LHb. Single-pulse stimulation (0.5 mA, 0.5ms, 1 Hz) of the LHb enhanced spiking activity in 46.3% (19/41) of RMTg neurons sampled with a mean onset latency of 5.5 ± 0.3 ms after current ejection (Fig. 2b). The vast majority (15 out of 19) of RMTg neurons responding to LHb stimulation were also "pinch-excited". The remaining (22 out of 41) cells tested failed to respond to LHb stimulation.

Notably, Ji and Shepard (2007) reported that LHb stimulation transiently suppressed the activity of 97% of midbrain DA neurons through a GABA_A receptor-mediated mechanism. This inhibition unlikely is monosynaptic, being relayed by GABA neurons in the RMTg, which receive a major LHb input, and heavily project to midbrain DA neurons (see Jhou *et al.*, 2009b). To investigate this possibility, in a separate series of experiments, the effect of RMTg stimulation on the spontaneous firing rate of DA neurons in the VTA was examined. As expected, single-pulse electrical stimulation (0.5 mA, 0.5 ms, 1 Hz) of the RMTg resulted in a short-term, though complete, suppression of VTA DA neuron activity (Fig. 2c). Peristimulus

histograms and CUMSUM plots demonstrating the typical responses of these cells to a 0.5 mA current pulse are illustrated in Figure 2c. The duration of inhibition averaged 124.8 ± 13.2 ms ($n=10$) (see Fig. 2c). The stimulating and recording sites were verified to be within the RMTg and the VTA, respectively (Paxinos and Watson, 2007).

Methamphetamine-induced Fos expression within RMTg recording sites

The RMTg is a newly characterized brain region and poorly defined anatomically. Although we selected for our experiments only cells fulfilling precise electrophysiological criteria, the correct localization of recording sites is crucial. Figure 3 shows that these sites were localized within the RMTg region as recently defined (Jhou *et al*, 2009a; Jhou *et al*, 2009b; Kaufling *et al*, 2009) (Fig. 3a and b). Additionally, the RMTg is reliably identified by accumulation of Fos-immunoreactive (Fos-IR) neurons following pharmacological stimuli, such as an acute methamphetamine administration (Jhou *et al*, 2009b). To verify that the recording sites (marked with Pontamine sky blue) were located in the RMTg, we carried out electrophysiological experiments as described above and thereafter induced Fos expression by injecting methamphetamine (10 mg/kg, i.p., 2 hours before sacrifice). We used a specific antibody for c-Fos raised against a synthetic peptide corresponding to amino acids 4-17 of human c-Fos, which reacts with rat Fos proteins but not with Jun protein. As previously described (Jhou *et al*, 2009b; Kaufling *et al*, 2009), two hours after methamphetamine administration Fos-IR cells were found in the RMTg, which resides within and ventral to the superior cerebellar decussation (xscp) (Fig. 3). As shown in Figure 3c, clusters of Fos-IR neurons are observed in the RMTg, symmetrically

to the electrophysiological recording site marked with Pontamine sky blue (which appears red at wavelength 594-617).

Effects of drugs of abuse on RMTg neurons

The firing rate of GABA neurons within the RMTg is inhibited by natural rewards like food or reward predictive stimuli, and, conversely, stimulated by aversive stimuli (Jhou *et al*, 2009a). However, it is not known whether exposure to drugs of abuse influence RMTg neuronal activity. Ergo we investigated the electrophysiological effects of four major addicting drugs (i.e. morphine, cocaine, the cannabinoid receptor agonist WIN55212-2, and nicotine) on putative RMTg neurons both in vivo and in vitro. In vivo all neurons were identified by their excitatory responses to both electrical stimulation of the LHb and paw pinch. Only cells whose correct localization was confirmed histologically were included in the study.

Morphine

GABA neurons projecting from the RMTg to the VTA are immunoreactive against μ -opioid receptors (Jhou *et al*, 2009b). Accordingly, rats self-administer the μ -opioid receptor agonist endomorphin-1 into the RMTg, but not into the regions dorsal, ventral, or lateral to it (Jhou *et al*, 2009c). Therefore, we first assessed if the μ -opioid receptor agonist morphine modulates the neuronal activity of RMTg neurons. We selected 7 neurons that responded to both LHb stimulation and paw pinch. Systemic administration of morphine (4 mg/kg, i.v.) strongly inhibited the spontaneous firing rate of putative RMTg neurons ($49.7 \pm 8.6\%$ of baseline; $n=7$;

$F_{5,30}=13.02$, $p<0.0001$; one way ANOVA and Dunnett's test) (Fig. 4a and c). This effect required the activation of μ -opioid receptors, as pretreatment with the μ -opioid receptor antagonist naloxone (0.1 mg/kg, i.v., 4 min before morphine) fully prevented morphine-induced suppression of RMTg neuron activity ($103.5\pm 7.2\%$ of baseline level; $n=4$; $F_{1,45}=14.55$; $p<0.01$ vs. morphine; two way ANOVA and Bonferroni's test) (Fig. 4b and c). Morphine did not significantly affect firing regularity (basal CV: $45.1\pm 9.8\%$; morphine CV: $46.3\pm 7.2\%$; $p>0.05$, Student's t -test).

To investigate whether or not μ -opioid receptor activation might affect excitatory synaptic transmission onto putative RMTg cells in vitro, we measured EPSCs recorded from these neurons while holding them in a voltage-clamp mode at -70 mV. Bath application of morphine at a concentration of 1 μ M (3 min) significantly reduced EPSCs (by $40.0 \pm 15.0\%$, $n=5$; $F_{18,72}=3.417$; $p=0.0001$; one-way ANOVA for repeated measures) (Fig. 4d). Morphine did not significantly affect either the mean value of the decay time constant (5.06 ± 0.10 ms and 5.14 ± 0.05 ms in the absence and presence of morphine, respectively) or the holding current (5.56 ± 3.01 pA). However, a decreased input resistance (from 0.42 ± 0.05 to 0.32 ± 0.06 G Ω ; paired t -test, $p<0.05$) was detected, which might contribute to the reduction in EPSC amplitude. The effect of morphine was reversible on wash out, and it was fully abolished in the presence of naloxone (0.1 μ M; $n= 5$; $F_{1,152}= 23.77$; $p=0.0012$; two-way ANOVA) (Fig. 4e). The reduction of EPSC amplitude observed could be explained by either a presynaptic effect on glutamate release, a postsynaptic change of the AMPA receptors, or a combination of these. We, therefore, measured the paired-pulse ratio, whose modifications are considered to reflect changes in transmitter release (Melis *et al*, 2002; Nie *et al*, 2004). Morphine did not modify the paired-pulse

ratio (from EPSC2/EPSC1= 0.88 ± 0.10 to EPSC2/EPSC1= 0.91 ± 0.10 ; $n=5$, paired t -test $t=0.20$; $p=0.849$) (Fig. 4f), suggesting that it might reduce EPSC amplitude at a postsynaptic site.

Cocaine

GABA neurons in the RMTg have been demonstrated to display strong Fos, FosB and Δ FosB activation after both acute and chronic exposure to psychostimulants (Colussi-Mas *et al*, 2007; Geisler *et al*, 2008; Jhou *et al*, 2009b; Kaufling *et al*, 2009; Perrotti *et al*, 2005), thus suggesting a potential role for this area in psychostimulant reward-related effects. Therefore, we investigated the possibility that cocaine might influence RMTg neuron discharge activity. We recorded 7 cells identified by both their excitatory response to LHb stimulation and paw pinch. Acute injection of cocaine (1 mg/kg, i.v.) caused a transient depression of firing rate of putative RMTg neurons ($82.1 \pm 6.0\%$ of baseline level; $n=7$; $F_{5,30}=2.58$; $p<0.05$; one way ANOVA and Dunnett's test) (Fig. 5a and b). The inhibitory effect of cocaine was less pronounced and shorter than the one induced by morphine. There was also no significant change in CV after the psychostimulant administration (basal CV: $30.9 \pm 4.6\%$; cocaine CV: $34.3 \pm 6.2\%$; $p>0.05$, Student's t -test).

Since cocaine-induced plasticity at excitatory synapses in the VTA plays a crucial role in those adaptations contributing and promoting addictive behaviors (Chen *et al*, 2010), we decided to examine whether it could trigger similar changes in the RMTg. Bath application of cocaine (1 μ M, 5 min) caused no change in all measured parameters: in fact, neither EPSC amplitude ($108.0 \pm 16.7\%$ of baseline; $n=5$; $F_{18,72}= 0.80$; $p=0.68$; one-way ANOVA) (Fig. 5c and d) nor paired-pulse ratio (from EPSC2/EPSC1= 0.97 ± 0.10 to EPSC2/EPSC1= 1.03 ± 0.10 ; $n=5$, $t=0.5583$;

$p=0.6$; paired t -test) were affected. Additionally, no changes were detected in either mean value of the decay time constant (being 5.03 ± 0.08 ms and 4.87 ± 0.05 ms in the absence and presence of cocaine, respectively) or the holding current (being the change of -0.76 ± 4.15 pA) or input resistance (from 0.39 ± 0.05 to 0.48 ± 0.1 G Ω). This lack of effect could be ascribed to the specificity of acute effect of cocaine on DA neurons, which appears to be merely on NMDA-mediated EPSCs (Schilstrom *et al*, 2006).

WIN55212-2

Cannabinoid type 1 (CB1) receptors are widely distributed in the central nervous system, being mostly located in presynaptic terminals (Freund *et al*, 2003) where they modulate both inhibitory and excitatory neurotransmission in neuronal subpopulations relevant to responses to rewarding and aversive stimuli (reviewed in Moreira and Lutz, 2008). Moreover, since DA neurons are excited by cannabinoids (French *et al*, 1997; Gessa *et al*, 1998), we hypothesized that inhibition of RMTg neurons could contribute to this effect. Hence, we thought to examine the effect of acute exposure to the CB1 receptor agonist WIN55212-2 (WIN) on 7 putative RMTg neurons, which were identified by their excitatory responses to both LHb stimulation and paw pinch. Similarly to morphine, WIN (0.5 mg/kg, i.v.) produced a profound and long-lasting decrease in discharge frequency ($46.9 \pm 6.7\%$ of baseline level; $n=7$; $F_{5,30}=19.44$; $p<0.0001$; one way ANOVA and Dunnett's test) (Fig. 6a and c), whereas no considerable changes were observed in the regularity of firing (basal CV: $31.0 \pm 5.0\%$; WIN CV: $38.5 \pm 3.6\%$; $p>0.05$, Student's t -test).

To clarify the involvement of CB1 receptors in this effect, we administered the CB1 receptor antagonist SR141716A (SR, 0.5 mg/kg, i.v.) 4 minutes before WIN (Fig. 6b and c). SR, while

per se ineffective, fully prevented WIN-induced depression of putative RMTg neurons ($102.6 \pm 3.1\%$ of baseline level; $n=4$; $F_{1,45}=16.08$; $p<0.01$; two way ANOVA and Bonferroni's test) (Fig. 6b and c).

Since CB1 receptor activation by endogenous ligands has been shown to modulate synaptic transmission in many brain regions (Kano *et al*, 2009), we examined whether WIN would affect excitatory transmission onto putative RMTg cells. WIN ($1 \mu\text{M}$, 5 min) significantly and irreversibly reduced EPSC amplitude (by $30.0 \pm 4.0\%$, $n= 5$; $F_{18,90}= 3.62$; $p<0.0001$; one-way ANOVA) (Fig. 6d). Probably due to its high lipophilicity, WIN effect did not wash out. However, when it was co-applied with the potent and selective CB1 receptor antagonist AM281 ($0.5 \mu\text{M}$), WIN-induced inhibition of EPSC amplitude was fully prevented ($n=5$; $F_{1,171}= 5.30$; $p<0.05$; two-way ANOVA) (Fig. 6e). WIN did not affect either the mean value of the decay time constant (being 4.66 ± 0.2 ms and 4.86 ± 0.3 ms in the absence and presence of WIN, respectively) or the holding current (being the change of 4.67 ± 2.60 pA) or the input resistance (from 0.49 ± 0.05 to 0.4 ± 0.05 G Ω). Since probability of release is inversely related to paired-pulse ratio, if WIN decreases EPSC amplitude through activation of presynaptic CB1 receptors, one would expect an increased paired-pulse ratio in the presence of this drug. WIN-induced decrease of EPSC amplitude was accompanied by an increased paired-pulse ratio (from $\text{EPSC}_2/\text{EPSC}_1= 0.95 \pm 0.05$ to $\text{EPSC}_2/\text{EPSC}_1= 1.45 \pm 0.20$; $n=5$; $t=3.039$; $p=0.019$; paired *t*-test) (Fig. 6f), thus suggesting that WIN reduced probability of glutamate release via activation of presynaptic CB1 receptors.

Nicotine

Acute injections of nicotine induce c-Fos expression specifically in non-DA neurons of the most caudal tier of the rat VTA (Pang *et al*, 1993). Moreover, agonists at nicotinic acetylcholine receptors (nAChRs) have been reported to be more readily self-administered into the posterior than anterior VTA (Ikemoto *et al*, 2006). Thus, we next tested whether nicotine would affect the spontaneous activity of RMTg cells. We recorded 7 putative RMTg neurons previously identified by both LHb stimulation and paw pinch. After 2-3 minutes of stable baseline activity, we administered nicotine (0.2 mg/kg, i.v.) (Fig. 7a and c). Nicotine caused a robust stimulation of firing rate ($198.8 \pm 29.2\%$ of baseline level; $n=7$; $F_{5,30}=6.63$; $p<0.001$; one way ANOVA and Dunnett's test) (Fig. 7c), without significantly affecting spiking regularity (basal CV: $34.1 \pm 3.8\%$; nicotine CV: $33.0 \pm 3.8\%$; $p>0.05$, Student's *t*-test). We then asked if the nAChR antagonist mecamylamine (2 mg/kg, i.v.) would antagonize nicotine's actions on putative RMTg cells. Administration of mecamylamine 4 minutes before nicotine did not alter spontaneous activity, but completely prevented nicotine-induced excitation ($100.7 \pm 10.7\%$ of baseline level; $n=4$; $F_{1,45}=18.98$; $p<0.01$; two way ANOVA and Bonferroni's test) (Fig. 7b and c), thus supporting the role of nAChRs in the observed effects.

In vitro, nicotine (1 μ M, 2 min) also produced a long-lasting enhancement of EPSC amplitude (by $40.0 \pm 7.0\%$ in the presence of nicotine; $n= 5$; $F_{18,72}= 10.13$; $p<0.0001$ one-way ANOVA) (Fig. 7d). This effect was accompanied by an increased mean value of decay time constant (from 4.5 ± 0.1 ms to 5.8 ± 0.2 ms; $p=0.005$ paired *t*-test), which corresponds well with the increased EPSC amplitude. We also observed that the decay time constant of EPSCs showed a positive correlation with peak amplitude (Fig. 7d). This correlation was presumably not filtered by properties of the cell membrane, since neither the holding current (being 44.2 ± 13.2 pA and 47.9 ± 14.6 pA for baseline and nicotine, respectively) nor the input resistance (from 0.34 ± 0.1 to

0.38±0.1 MΩ) changed during nicotine application. The positive relationship between the decay time constant and the peak amplitude might suggest a dependency of amplitude on the rate of glutamate diffusion from the terminal to the synaptic cleft (Glavinovic and Rabie, 1998; Hirasawa *et al*, 2001). Notably, nicotine-induced potentiation of EPSC amplitude was sensitive to methyllycaconitine (5 nM), an antagonist of $\alpha 7$ -subtype nAChR subunit ($n=5$; $F_{1,152}= 17.87$; $p=0.0029$; two-way ANOVA) (Fig. 7e). Since nicotine-induced enhancement of EPSC amplitude is associated with a decrease in the paired-pulse ratio (from EPSC2/EPSC1= 1.01 ± 0.07 to EPSC2/EPSC1= 0.82 ± 0.05 ; $n=5$; $t=2.45$; $p=0.03$; paired *t*-test) (Fig. 7f), altogether these data suggest that $\alpha 7$ subunits are the major components of nAChRs located on glutamatergic presynaptic terminals onto RMTg neurons, and might mediate the long-lasting excitation of putative RMTg neurons observed *in vivo*.

Discussion

Here we studied putative RMTg neurons that were excited by LHb stimulation and aversive stimuli (e.g. paw pinch). Conversely, we show that VTA DA neurons are inhibited by RMTg stimulation. On these bases, we hypothesized that these anatomical and functional aspects of RMTg neurons would be consistent with sensitivity to addicting drugs, which are known to interfere with reward and aversion mechanisms. Indeed, here we found that these cells, selected for their excitatory response to both LHb stimulation and a noxious stimulus, are inhibited by morphine, cocaine and the cannabinoid agonist WIN. Additionally, patch-clamp experiments demonstrate that EPSCs recorded from putative RMTg neurons and evoked by stimulation of rostral fibres were unchanged by cocaine, whereas inhibited by both WIN and morphine, with presynaptic and postsynaptic mechanisms, respectively. Notably, we also found that nicotine exerts opposite effects to those of morphine and WIN, since it strongly enhances firing rate of putative RMTg neurons and persistently increases the magnitude of EPSCs. Accordingly, converging anatomical, physiological and behavioral studies show that the RMTg encodes aversive stimuli similarly to the LHb and opposite to the VTA. RMTg neurons are considered as an intermediate site between the LHb, which sends an extensive excitatory innervation, and VTA DA neurons, which in response are uniformly inhibited.

The depressant effect of morphine on putative RMTg neurons is consistent with the finding that these cells display immunoreactivity against μ -opioid receptors (Jhou *et al*, 2009b), and supports the hypothesis that opioids excite VTA DA neurons also by inhibiting RMTg neurons. Notably, RMTg cells exert a tonic inhibition on DA cells (Ikemoto, 2010). Consistently, rats will self administer opioids into the RMTg but not in the adjoining regions (Jhou *et al*, 2009c). Morphine

also depressed EPSCs without changing the paired-pulse ratio, although decreasing the input resistance, thus suggesting that it does not affect the probability of glutamate release and that the site of action is at postsynaptic μ -opioid receptors. However, the presence of Cesium in the recording pipette argues against activation of G-protein coupled inward rectifier K^+ (GIRK) channels as contributing factors to the decreased input resistance (Davila *et al*, 2003; Lesage *et al*, 1995).

Differently from μ -opioid receptors, CB1 receptors in the CNS are mostly localized presynaptically on a large variety of axon terminals to suppress neurotransmitter release (Freund *et al*, 2003). It is not currently known whether CB1 receptors are localized within the RMTg. However, the LHb displays a relative abundance of CB1 mRNA, when compared with other thalamic nuclei (Matsuda *et al*, 1993), suggesting that CB1 receptors may be localized presynaptically at excitatory afferents in the RMTg to control glutamate release. Alternatively, CB1 receptors might also be located on axon terminals of output neurons of the periaqueductal gray, one of the major afferents to the RMTg (Jhou *et al*, 2009a). The observations that the CB1 agonist WIN depresses EPSC amplitude on putative RMTg neurons, induces a paired-pulse facilitation, and inhibits firing rate *in vivo* are consistent with a presynaptic locus of CB1 receptors, whose activation leads to a decreased probability of glutamate release. Additionally, one can speculate that inhibition of RMTg neurons by WIN might be one of the mechanisms contributing to cannabinoid-induced excitation of DA neurons *in vivo* (French *et al*, 1997; Gessa *et al*, 1998).

Cocaine was shown to evoke Δ Fos B expression on GABA neurons in the VTA tail (Perrotti *et al*, 2005), and this indicates that psychostimulants might affect DA neurons through the RMTg-VTA projection. Here we confirmed the finding that methamphetamine induces Fos expression

within RMTg. Prior studies show that psychostimulants activate RMTg Fos much more than in adjacent regions, but in our study Fos levels detected in RMTg neurons do not appear to be strikingly higher than surrounding areas. It must be pointed out that, differently from procedures for methamphetamine-induced Fos activation described in previous papers, here rats were further prepared for electrophysiological recordings with anesthesia, surgery, electrode insertion and dye ejection. Those additional procedures might have interfered with Fos expression by increasing aspecific expression within surrounding areas.

Cocaine was found to depress firing rate of putative RMTg neurons in vivo. The mechanism underlying this effect remains to be established. Since cocaine does not affect the magnitude of EPSC, and neither changes the paired-pulse ratio nor the membrane properties of these cells, we can only hypothesize that effects in vivo result from indirect actions of cocaine on RMTg neurons, possibly through a long synaptic loop, which does not directly involve effects on excitatory afferents from the LHb.

Differently from the other drugs studied here, nicotine induces a robust and persistent excitation on putative RMTg neuron firing rate and increases both EPSC amplitude and the mean value of the decay time constant, while decreases the paired-pulse ratio. Taken together, our findings are indicative of the enhancement of glutamate release from excitatory afferents likely induced by $\alpha 7$ -containing nAChRs located on presynaptic terminals. Habenular afferents are likely to express $\alpha 7$ -containing nAChRs, given that intense in situ hybridization for transcripts encoding these receptors was detected in the habenula (Seguela *et al*, 1993). We cannot exclude, however, that nicotine excites RMTg neurons also by acting at somatodendritic nAChRs. Hence, subcellular localization of nAChRs is relevant for the functional impact of their activation: nAChRs are expressed both on presynaptic terminals, where they directly modulate GABA

release independent of action potential firing (Fisher *et al*, 1998; Lu *et al*, 1999; Radcliffe *et al*, 1999), and on cell bodies, where modulation of GABA release depends on discharge activity (Alkondon *et al*, 1997; Alkondon *et al*, 1999; Frazier *et al*, 1998). Extensive evidence indicates that nicotine stimulates discharge rate of VTA DA neurons (Mereu *et al*, 1987; Pidoplichko *et al*, 1997). In apparent contradiction with those findings, our results suggest that DA neurons might be inhibited by nicotine-induced excitation of RMTg neurons. However, a transient inhibition of DA neurons precedes long-lasting excitation in some instances (Erhardt *et al*, 2002) and the net overall excitatory effect of nicotine on VTA DA neurons is considered to result from the combination of fast sensitizing stimulation of GABA release and slowly sensitizing potentiation of glutamate release within the VTA, as elegantly demonstrated by Mansvelder and colleagues (2002). Thus, it is conceivable that stimulation of RMTg neurons by nicotine may contribute to the elevation of GABA levels in the VTA, which is ultimately overcome by a larger and long-lasting potentiation of glutamate release inducing DA neuron excitation (Mansvelder *et al*, 2002). Consistently, in the VTA, nAChR-induced modulation of GABA transmission was completely tetrodotoxin-sensitive, implying that these receptors are expressed far away from terminals (Mansvelder *et al*, 2002).

The inhibition evoked by addicting drugs, with the exception of nicotine, on discharge activity and/or EPSCs on putative RMTg neurons is consistent with the notion that aversive stimuli encoded by the LHb, relayed to the RMTg and finally conveyed to the VTA, might be attenuated in their magnitude by concomitant administration of addicting drugs. Hence, RMTg lesions blunt inhibitory behavioral responses, such as freezing elicited by aversive stimuli and open arm avoidance in the elevated plus maze (Jhou *et al*, 2009a). Consistently, both opioids and cannabinoids reduce the magnitude of aversion-elicited inhibitory behavioral responses (Haney

and Miczek, 1994; Moreira *et al*, 2009; Vivian and Miczek, 1998). On the other hand, opposite to morphine and cannabinoid agonists, nicotine enhances behavioral responses to aversive stimuli, since it enhances contextual fear as evaluated by measuring freezing, a natural fear response in rodent (Gould and Wehner, 1999). Data on the effects of cocaine are conflicting, possibly because it is difficult to separate the locomotor activating effects of the drug: in one study, impaired freezing was associated with locomotor sensitization (Morrow *et al*, 1995), whereas in other studies enhanced conditioned startle, possibly related to decreased freezing (Gordon and Rosen, 1999), or no effect on fear conditioning, were observed (Burke *et al*, 2006). Our results demonstrate that putative RMTg neurons are a site of action of major addicting drugs. Being an important relay inhibitory region to the VTA, the RMTg is involved in the regulation of firing discharge of VTA DA neurons, and, therefore, might play an important role in the mechanisms and functional aspects of drugs of abuse.

Supplementary information

Supplementary information is available at the *Neuropsychopharmacology* website.

Disclosure/Conflict of Interest

The authors declare that, except for income received from their primary employer, no financial support or compensation has been received from any individual or corporate entity over the past 3 years for research or professional service and there are no personal financial holdings that could be perceived as constituting a potential conflict of interest.

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Titles and legends to figures

Figure 1.

Characterization of neurons in the rostromedial tegmental nucleus (RMTg).

(a) Traces illustrating representative extracellular recordings of a putative GABA neuron in the RMTg of an anesthetized rat. The left trace shows typical superimposed spike waveforms characterized by an initial negative deflection followed by a positive phase. The right trace shows the typical fast firing rate and regular pattern of a RMTg neuron. The representative interspike interval histogram demonstrates the regularity of discharge pattern.

(b) Identification of RMTg in the horizontal section in the brain slice preparation. The RMTg was identified as the region posterior to the ventral tegmental area (VTA), which is located medially to the medial terminal nucleus of the accessory optic tract (MT) and laterally to the interpeduncular fossa (IPF). The stimulating electrode is placed rostro-laterally to the recording electrode (at a distance of $\sim 150 \mu\text{m}$), whose position is posterior to the VTA (at a distance of $\sim 1 \text{ mm}$). A recorded putative RMTg neuron is visualized in the inset. SNC, substantia nigra pars compacta, SNR, substantia nigra pars reticulata. The number indicates the ventral coordinate with reference to the horizontal plane passing through the skull surface at bregma, as in Paxinos and Watson (2007). (c) The electrophysiological properties exhibited by the neuron in (b) were similar to all of the neurons recorded in the study. RMTg cell physiology includes short-duration spikes (left panel, superimposed traces), regular spontaneous activity at rest (middle panel). The right panel displays typical whole-cell patch clamp recordings and shows that bath application of $10 \mu\text{M}$ 6-cyano-7-nitroquinoxaline-2,3-dione (CNQX) completely suppressed evoked

postsynaptic currents (EPSCs) when cells are held at -70 mV, indicating that they are AMPA-mediated. Traces show paired EPSCs from a typical experiment, before (basal, black line) and during superfusion of CNQX (grey line).

Figure 2.

Neurons in the rostromedial tegmental nucleus (RMTg) exhibit a fast and phasic increase in firing rate after paw pinch, are excited by lateral habenula and inhibit dopamine neurons.

(a) Typical recording from a putative RMTg neuron showing the rapid and phasic increase in firing activity during paw pinch. The average rate histogram (1 s bin) is shown ($n=12$) ($***p<0.001$, ($**p<0.01$; one-way ANOVA and Dunnett's test). (b) Superimposed traces acquired from a digital storage oscilloscope showing a relatively constant latency of the orthodromic responses of RMTg neuron after LHb stimulation (top). The arrowhead indicates the artifacts produced by LHb stimulation. The coronal plate indicates the position of the stimulating electrode. Representative peristimulus time-histogram displaying the typical response of a recorded putative RMTg neuron during LHb stimulation (bottom). (c) Traces acquired from a digital oscilloscope (top) and peristimulus time histogram of the same cell (middle), comprising 100 consecutive sweeps, showing that RMTg stimulation induces a complete inhibition of discharge activity of a VTA DA neuron (top) for ~100 ms. Stimuli consisted of rectangular current pulses (0.5 mA, 0.5 ms) delivered 1 s after the start of each sweep (middle). Graph showing the average CUMSUM plots of 10 VTA DA neurons under RMTg stimulation (bottom). The graph was generated by adding the contents of consecutive bins

in the corresponding peristimulus time histograms to create a cumulative total average and SEM of the number of spikes recorded across trials.

Figure 3

Methamphetamine-induced Fos immunoreactivity within the rostromedial tegmental nucleus (RMTg).

The rat from which the brain sections were taken received an injection of methamphetamine (10 mg/kg, i.p.) 2 hours prior to sacrifice. **(a)** Schematic redrawing of coronal plates (based on Paxinos and Watson, 2007) including the RMTg (bregma AP-7.2 and - 7.44) showing Pontamine sky-blue labeling sites (black dots). **(b)** Microphotograph of a transmitted light image showing the Pontamine sky blue dye (arrow). **(c)** This section shows Fos immunostaining (green). The arrow indicates the dye, which appears red at wavelength 594-617. **(d)** Higher magnification of the white boxed area in (c) revealing Fos-positive nuclei inside the RMTg (arrowheads). 4N, trochlear nucleus; DR, dorshal raphe nuclei; lfp, longitudinal fasciculus of the pons; ml, medial lemniscus; Pn, pontine nuclei; tth, trigeminothalamic tract; xscp, superior cerebellar peduncle decussation; Scale bars: 1000 μm in (b-c); 100 μm in (d).

Figure 4.

Effects of morphine on rostromedial tegmental nucleus (RMTg) neurons.

(a) Representative firing rate histogram showing the inhibitory effects of intravenous morphine (Mor, 4 mg/kg injected at arrow) on discharge activity of an individual putative RMTg neuron.

(b) Representative firing rate histogram showing that naloxone (Nlx, 0.1 mg/kg, i.v. injected 4 min before morphine) fully prevents morphine-induced inhibition of firing rate of a RMTg neuron. **(c)** Graph illustrating the time course of morphine's effects (4 mg/kg, i.v.) on firing rate of putative RMTg neurons with and without naloxone (0.1 mg/kg, i.v.) pretreatment. Naloxone blocked the inhibition of RMTg neurons induced by morphine (arrow). Results are means with vertical bars representing the SEM of firing rate expressed as a percentage of the baseline (BAS). *** $p < 0.001$ vs. baseline, one-way ANOVA and Dunnett's test.

(d) Morphine induces inhibition of glutamatergic synaptic transmission in rat RMTg cells. A typical whole-cell patch clamp recording showing that bath application of morphine (1 μ M) inhibits EPSC amplitude, when cells are held at -70 mV. The grey line represents mean EPSC amplitude. The inset shows single EPSC from a typical experiment, before (black line) and during (grey line) superfusion of morphine. **(e)** Morphine reduces EPSC amplitude through activation of μ -opioid receptors. All data are normalized to the respective baseline (5 min of baseline). Black bar shows time of superfusion of morphine in the presence (grey circles) and absence (open circles) of the μ -opioid antagonist naloxone (100 nM). SEM bars are smaller than symbols in some cases. The inset shows 12-trace averages of EPSCs in the absence (black line) and presence (grey line) of naloxone. Black bars represent time of morphine application. **(f)** Morphine does not change the paired-pulse ratio of EPSCs. The top graph plots the paired-pulse ratio for each of the experiments in (e) before (open circles) and during (grey circles) the application of morphine, while the bottom graph plots the averaged paired-pulse ratio in a bar graph form. Representative traces are shown in the inset.

Figure 5.

Cocaine's effects on rostromedial tegmental nucleus (RMTg) neurons.

(a) Representative rate histogram illustrating the inhibition of firing rate of a putative RMTg neuron after cocaine administration (1 mg/kg injected i.v. at arrow). (b) Graph showing the time course of cocaine's effects (1 mg/kg, i.v.) on firing rate of putative RMTg neurons. Results are means with vertical bars representing the SEM of firing rate expressed as a percentage of baseline (BAS). * $p < 0.05$ vs. baseline, one-way ANOVA and Dunnett's test.

(c) Cocaine does not affect glutamatergic synaptic transmission in rat putative RMTg cells. A typical whole-cell patch clamp recording showing the effect of bath application of cocaine (1 μ M) on EPSC amplitude, when cells are held at -70 mV. The grey line represents mean EPSC amplitude. The inset shows single EPSC from a typical experiment, before (black line) and during (grey line) application of cocaine. (d) Cocaine effect on EPSC amplitude. All data are normalized to the respective baseline (5 min of baseline). Black bar shows time of superfusion of cocaine. SEM bars are smaller than symbols in some cases. The inset shows 12-trace averages of EPSCs in the absence (black line) and presence (grey line) of cocaine. Black bars represent time of cocaine application. (e) Cocaine does not change the paired-pulse ratio of EPSCs. The top graph plots the paired-pulse ratio for each of the experiments in (e) before (open circles) and during (grey circles) the application of cocaine, while the bottom graph plots the averaged paired-pulse ratio in a bar graph form. Representative traces are shown in the inset. Lines (black) represent EPSC amplitude before and during (grey line) application of cocaine.

Figure 6.

Responses of rostromedial tegmental nucleus (RMTg) neurons to the cannabinoid receptor agonist WIN55212-2 (WIN).

(a) Typical firing rate histogram showing the inhibitory response of a putative RMTg neuron after WIN (0.5 mg/kg, i.v.) administration (arrow). (b) Representative rate histogram showing that the CB1 antagonist SR141716A (SR, 0.5 mg/kg, injected i.v. 4 min before WIN, at arrows) blocks WIN-induced inhibition of firing rate. (c) Graph showing the time course of WIN-induced inhibition of putative RMTg neurons, and that this inhibition by WIN was prevented by the administration (arrow) of SR (0.5 mg/kg, i.v.). Results are means with vertical bars representing the SEM of firing rate expressed as a percentage of baseline (BAS). *** $p < 0.0001$ vs. baseline, one-way ANOVA and Dunnett's test.

(d) WIN inhibits glutamatergic synaptic transmission in rat putative RMTg cells. A typical whole-cell patch clamp recording showing that bath application of WIN (1 μ M) inhibits EPSC amplitude, when cells are held at -70 mV. The grey line represents mean EPSC amplitude. The inset shows single EPSC from a typical experiment, before (black line) and during (grey line) perfusion of WIN. (e) WIN reduces EPSC amplitude through activation of CB1 receptors. All data are normalized to the respective baseline (5 min of baseline). Black bar shows time of superfusion of WIN in the presence (grey circles) and absence (open circles) of the CB1 receptor antagonist AM281 (500 nM). SEM bars are smaller than symbols in some cases. The inset shows 12-trace averages of EPSCs in the absence (black line) and presence (grey line) of AM281. Black bars represent time of WIN application. (f) WIN enhances the paired-pulse ratio of EPSCs, producing pair-pulse facilitation. The top graph plots the paired-pulse ratio for each of the experiments in (e) before (open circles) and during (grey circles) the application of WIN, while the bottom graph plots the averaged paired-pulse ratio in a bar graph form. Representative traces

are shown in the inset, where the EPSCs are represented before (black lines) and during WIN (grey lines) application. * $p < 0.05$, Student's t -test.

Figure 7.

Rostromedial tegmental nucleus (RMTg) neurons are excited by nicotine.

(a) Representative rate histogram showing that nicotine (Nic, 0.2 mg/kg, i.v., arrow) strongly excites firing activity of a putative RMTg neuron. (b) This example shows that the non-selective nicotinic receptor antagonist mecamylamine (Mec, 2 mg/kg, i.v. 4 min before nicotine, arrow) fully prevents the effects of nicotine. (c) Graph showing the time course of the stimulation of firing rate induced by nicotine. Mecamylamine pretreatment fully prevents nicotine-induced stimulation of firing rate. Results are means with vertical bars representing the SEM of firing rate expressed as a percentage of the baseline (BAS). ** $p < 0.01$, *** $p < 0.001$ vs. baseline, one-way ANOVA and Dunnett's test.

(d) Nicotine enhances glutamatergic synaptic transmission in rat putative RMTg cells. A typical whole-cell-patch clamp recording showing that bath application of nicotine (1 μ M; Nic) increases EPSC amplitude, when cells are held at -70 mV. The grey line represents mean EPSC amplitude. The inset shows single EPSC from a typical experiment, before (black line) and during (grey line) perfusion of nicotine. The graph shows that the decay time constant of EPSCs is positively correlates with peak amplitude. (e) Nicotine enhances EPSC amplitude through activation of $\alpha 7$ -containing nicotinic receptors. All data are normalized to the respective baseline (5 min of baseline). Black bar shows time of superfusion of nicotine in the presence (grey circles) and absence (open circles) of the $\alpha 7$ -subunit receptor antagonist methyllycaconitine (5

nM; Mla). SEM bars are smaller than symbols in some cases. The inset shows 12-trace averages of EPSCs in the absence (black line) and presence (grey line) of MLA. Black bars represent time of nicotine application. **(f)** Nicotine reduces the paired-pulse ratio of EPSCs, producing paired-pulse depression. The top graph plots the paired-pulse ratio for each of the experiments in E before (open circles) and during (grey circles) the application of nicotine, while the bottom graph plots the averaged paired-pulse ratio in a bar graph form. Representative traces are shown in the inset, where the EPSCs are represented before (black lines) and during nicotine (grey lines) application. * $p < 0.05$, Student's t -test.













