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Microglia Disrupt Mesolimbic Reward Circuitry in Chronic Pain

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Chronic pain attenuates midbrain dopamine (DA) transmission, as evidenced by a decrease in opioid-evoked DA release in the ventral striatum, suggesting that the occurrence of chronic pain impairs reward-related behaviors. However, mechanisms by which pain modifies DA transmission remain elusive. Using in vivo microdialysis and microinjection of drugs into the mesolimbic DA system, we demonstrate in mice and rats that microglial activation in the VTA compromises not only opioid-evoked release of DA, but also other DA-stimulating drugs, such as cocaine. Our data show that loss of stimulated extracellular DA is due to impaired chloride homeostasis in midbrain GABAergic interneurons. Treatment with minocycline or interfering with BDNF signaling restored chloride transport within these neurons and recovered DA-dependent reward behavior. Our findings demonstrate that a peripheral nerve injury causes activated microglia within reward circuitry that result in disruption of dopaminergic signaling and reward behavior. These results have broad implications that are not restricted to the problem of pain, but are also relevant to affective disorders associated with disruption of reward circuitry. Because chronic pain causes glial activation in areas of the CNS important for mood and affect, our findings may translate to other disorders, including anxiety and depression, that demonstrate high comorbidity with chronic pain.

Key words: addiction; affective disorder; chronic pain; dopamine; emotion; opioids

Introduction

The pain experience involves activation of multiple brain regions, including those encoding salience and affect (Legrain, 2011; Mouraux et al., 2011). The fact that negative emotional experiences can produce the sensation of pain in the absence of tissue damage and that cognitive processes such as expectation can modulate the effectiveness of analgesia highlights the importance of salience and affect in the experience of pain. Therefore, changes that modulate the activity of motivational, salience, and affective systems will affect how the brain evaluates pain and analgesia.

There is extensive interaction between systems involved in pain processing and other affective and motivational systems, and it has been proposed that aberrant functioning of the affect and salience circuits may contribute to chronic pain (Borsook et al., 2015). It is common for chronic pain to be comorbid with mood disorders (e.g., depression) and it has been posited that dysfunction of mesolimbic circuitry, including dopamine (DA) neurotransmission, precipitates mood disorders and impairs motivated behaviors (Leknes and Tracey, 2008; Cahill et al., 2013). In fact, chronic pain is second only to bipolar disorder as the major cause of suicide among all medical illnesses and the high comorbidity of depression in patients with chronic pain syndromes further highlights the significance of the negative affect associated with chronic pain (Asmundson and Katz, 2009; Elman et al., 2013).

The mesolimbic DA system, comprising dopaminergic neurons in the VTA that project to the NAc, is an integral circuit encoding salience, motivation, and reward (Berridge, 2007). Diseases linked to deficits in the midbrain DA system, including disorders of mood and affect, substance abuse, and Parkinson’s disease, are often comorbid with chronic pain (Jarcho et al., 2012) and can lead to impaired reward behaviors (Flagel et al., 2011; Grob et al., 2012). Imaging studies of chronic pain patients indicate that perturbation of the mesolimbic DA system is linked with chronic pain syndromes (Hagelberg et al., 2003a; Hagelberg et al., 2003b; Wood et al., 2007). Further, total striatal DA levels and opioid-stimulated DA release are...
markedly reduced in animal models of chronic pain (Ozaki et al., 2002; Taylor et al., 2014). Alterations in DA signaling as a result of chronic pain are associated with motivational deficits and reduced rewarding effects of natural and opioid rewards (Martin et al., 2007; Russo and Nestler, 2013; Schwartz et al., 2014).

How chronic pain affects the mesolimbic DA system is relatively unexplored, but there are sufficient reports to validate a role of altered DA signaling, not only in the chronicity of pain, but also in its response to opioid drugs. Here, we describe a mechanism by which chronic pain activates microglia in the VTA to inhibit DA transmission and disrupt reward behaviors. Inhibiting microglial activation restored normal DA signaling and recovered reward behavior. This work identifies microglial activation as being a critical component modulating reward behavior in chronic pain and suggests inhibiting microglial activation as a potential strategy for restoring disrupted DA transmission that is linked to disorders such as depression and addiction.

Materials and Methods

Animals
Male C57BL/6j and GAD65-GFP on C57BL/6j background mice (The Jackson Laboratory), 8–9 weeks old, were housed in groups of four unless otherwise indicated, and kept on a 12 h light/dark cycle with food and water available ad libitum. Male Sprague Dawley rats (Charles River Laboratories), 8–9 weeks old, were housed in groups of two and kept on a 12 h light/dark cycle with food and water available ad libitum. All behavioral tests were performed during the light phase. All procedures were conducted in accordance with the National Institutes of Health’s Guide for the Care of Use of Laboratory Animals and were approved by the University of California–Los Angeles, University of California–Irvine, and Queen’s University Institutional Animal Care and Use Committees.

Peripheral nerve injury
Animals were randomly assigned into two surgery groups, sham or peripheral nerve injury (PNI). Surgery was performed as described previously (Mosconi and Kruger, 1996). All animals received acetaminophen (32 mg/ml, p.o) preoperatively and were anesthetized with gaseous isoflurane (induction at 5% and maintenance at 2.0–2.5% in oxygen). An incision was made in the lateral left thigh followed by blunt dissection to expose the sciatic nerve. For PNI, the nerve was encased with a polyethylene tubing (PE50, 2 mm). The sciatic nerve was isolated and sectioned on a cryostat to confirm cannula placement. The sciatic nerve was used to indicate the bilateral injury. Animals were housed individually and allowed to recover for 7 days after PNI cannula insertion, at which point they received a sham or nerve ligation surgery, as described above (n = 6–9 per group). Six days after nerve ligation surgery, animals were subjected to preconditioning and CPP was performed as above.

During the conditioning sessions, animals received 4 trials with DAMGO (1 or 25 ng/mg/kb, i.p.; n = 5–8) or cocaine hydrochloride (10 mg/kg, i.p.; n = 7) and 4 trials of saline on alternating days, and were confined to the chambers for 15 (cocaine) or 30 (morphine) minutes. In a separate experiment, animals received only 1 trial of morphine (10 mg/kg, i.p.) and 1 trial of saline (n = 7–8 per group). The CPP assay began 6 d after peripheral nerve injury at a time previously shown to be during peak pain behavior (Taylor et al., 2014). On the postconditioning day (day 9 after the conditioning trial assay or day 3 for the 2 conditioning trial assay), no treatment was given to the animals so that they could be tested in a drug-free state. Animals were allowed free access to all chambers of the apparatus and the time spent in the drug-paired chamber measured by motion capture Noldus software was determined over 15–30 min.

Intra-VTA DAMGO. Male Sprague Dawley rats were anesthetized with isoflurane and mounted on a stereotaxic frame to allow implantation of a bilateral intra-VTA cannula (CMA Microdialysis; coordinates from bregma: AP = −5.4, ML ± 0.75, DV = −8.0). Rats were chosen to ensure accurate targeting of drug to the VTA. We have demonstrated comparable systemic opioid CPP responses between rats and mice in control and chronic pain models (Cahill et al., 2013). Animals were housed individually and allowed to recover for 7 d after intra-VTA cannula insertion, at which point they received a sham or nerve ligation surgery, as described above (n = 6–9 per group). Six days after nerve ligation surgery, animals were subjected to preconditioning and CPP was performed as above.

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Drugs
Morphine sulfate (1–10 mg/kg, i.p.; Sigma-Aldrich), cocaine hydrochloride (10 mg/kg, i.p.; Sigma-Aldrich), amphetamine hydrochloride (1 mg/kg, i.p.; Sigma-Aldrich), and n-Ala2, N-MePhe4, Gly-OH-[enkephalin (DAMGO, 1–25 ng; Tocris Biosciences) were dissolved in 0.1M PBS, pH 7.4, and administered at nerve ligation surgery and every 12 h until the end of experimentation. During the conditioned place preference assay (CPP), minocycline was administered at least 6 h before drug conditioning sessions.

CPP
Systemic drugs. CPP is a Pavlovian conditioning test to measure the motivational reward effects of a drug (Cunningham et al., 2006). This test allows for a within-subject analysis of the time spent in the vehicle compared with drug-paired conditioning chamber, which allows for paired analysis to assess the rewarding and/or motivational effects of a drug. CPP was conducted using an unbiased, counterbalanced, two- or three-chamber apparatus. Each square box (28 × 28 × 19 cm for mice; 45 × 45 × 30 cm for rats) was divided into two equal-sized conditioning chambers separated by a guillotine door. The two chambers were distinguished with visual and tactile cues so that the animals could discriminate between chambers. To counterbalance the groups before the conditioning sessions, animals were placed in the CPP apparatus and allowed free access to all chambers. The time spent in each chamber was recorded over 15–30 min using a CCD camera attached to a computer running behavioral tracking software (EthoVision; Noldus). The drug-paired chamber was assigned such that any innate bias for one chamber over the other was balanced between treatment groups. During the conditioning session, animals were confined to only one of the compartments so that they would associate the contextual cues of the compartment with the motivational effects of the stimulus (drug or vehicle, counterbalanced every other day). Conditioning sessions consisted of 8 sessions with 4 trials of morphine sulfate (1 or 10 mg/kg, i.p., n = 5–8) or cocaine hydrochloride (10 mg/kg, i.p., n = 7) and 4 trials of saline on alternating days, and were confined to the chambers for 15 (cocaine) or 30 (morphine) minutes. In a separate experiment, animals received only 1 trial of morphine (10 mg/kg, i.p.) and 1 trial of saline (n = 7–8 per group). The CPP assay began 6 d after peripheral nerve injury at a time previously shown to be during peak pain behavior (Taylor et al., 2014). On the postconditioning day (day 9 after the conditioning trial assay or day 3 for the 2 conditioning trial assay), no treatment was given to the animals so that they could be tested in a drug-free state. Animals were allowed free access to all chambers of the apparatus and the time spent in the drug-paired chamber measured by motion capture Noldus software was determined over 15–30 min.

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Evoked pain tests
Mechanical allodynia. Withdrawal thresholds to a mechanical stimulus applied with calibrated von Frey filaments were measured in all groups (n = 6 per group), as described previously (Chaplan et al., 1994). Mice...
were placed atop a mesh grid floor in a clear Plexiglas enclosure. Von Frey filaments were applied to the plantar surface of the ipsilateral hindpaw in an up–down manner in which filaments with increasing stiffness were applied until a paw retraction was observed. After the first positive reaction, the next less force filament was applied. If no reaction, the next higher force filament was applied; if a reaction was observed, the next less force filament was applied. This was repeated five times per animal and the 50% withdrawal threshold was calculated. Baseline measurements were taken for all animals before surgery and 2 weeks postoperatively.

**Thermal tail withdrawal.** Morphine and cocaine antinociception was measured using the tail withdrawal assay at 2 weeks after nerve injury (n = 6–9 per group). Mice were gently restrained by cupping them in a soft plastic conical sleeve and 2.5 cm of the tail was immersed in 49°C water. The time to tail withdrawal was measured. After 3 basal measurements, mice were injected with morphine (10 mg/kg, i.p.) or cocaine (10 mg/kg, i.p.). Tail withdrawal latencies were measured every 10 or 20 min for 1 h. Mice were returned to their home cages between testing. A cutoff of 15 s was imposed to avoid tissue damage based on baseline latencies for 100. Mice were returned to their home cages between testing. A cutoff of 15 s was imposed to avoid tissue damage based on baseline latencies for 100.

**Immunoblot assay.** Animals were anesthetized with isoflurane and the brains were isolated (n = 4 per group). Brain punches (1 mm × 1 mm × 1 mm) were taken from the VTA and homogenized in radioimmunoprecipitation assay buffer with protease inhibitors. Equal amounts of protein (25 μg) were loaded onto precast 10% polyacrylamide gels (Bio-Rad) and transferred to nitrocellulose membranes. Membranes were incubated with an antibody against BDNF (1:150; Millipore) and visualized with a goat anti-rabbit antibody conjugated to HRP (Jackson Immunoresearch). BDNF levels were normalized to the housekeeping protein β-actin (1:5000; Sigma-Aldrich). Results were analyzed with a computer-assisted densitometry analysis (ImageJ).

**Immunocytochemistry.** Animals (n = 4–7 per group) were transcardially perfused with 4% paraformaldehyde (PFA) in 0.1 M phosphate buffer (PB). The brain was dissected from the skull, postfixed for 30 min in the above-described fixative, and cryoprotected in 30% sucrose before cutting 40 μm sections collected in PBS containing 0.2% Triton X-100. For microglial labeling, sections were incubated with anti-IBA-1 (1:2000; Wako) and a goat anti-rabbit antibody conjugated to HRP (Jackson Immunoresearch). BDNF levels were normalized to the housekeeping protein β-actin (1:5000; Sigma-Aldrich). Results were analyzed with a computer-assisted densitometry analysis (ImageJ).

**Microdialysis.** Mice (n = 4–6 per group) were anesthetized with isoflurane and placed into a stereotaxic frame to allow implantation of a guide cannula (CMA Microdialysis) to an area just dorsal to the right side of the NAC (coordinates from bregma AP +1.5, ML +0.8, DV −3.5). This surgery was performed at the same time as sham and nerve ligation surgery. A separate group received PNI surgery and chronic treatment with minocycline (30 mg/kg, i.p.) every 12 h starting the day of surgery and continuing throughout the experiment. When microdialysis samples were collected, minocycline injection occurred at least 6 h before baseline sampling. Animals were housed individually and allowed to recover for 10–14 d. The day before the experiment, animals were implanted with a concentric microdialysis probe equipped with a 1-mm-long, 300-μm-diameter polyacrylonitril membrane (CMA Microdialysis). Probes were continuously perfused with ACSF containing the following (in mM): 125 NaCl, 2.5 KCl, 0.9 NaHPO4, 5 Na2HPO4, 1.2 CaCl2, and 1 MgCl2, pH 7.4, using syringe pumps at a rate of 1 μl/min. Dialyisates were collected at 4°C in 20 min intervals into plastic microvials preloaded with 2 μl of 12.5 μM perchloric acid/250 μM EDTA. Twelve hours after insertion of microdialysis probes, drug (10 mg/kg morphine, 10 mg/kg cocaine, 1 mg/kg amphetamine, or saline; i.p.) was injected every 1–5 h. Each animal was treated with every drug in one day (saline, morphine, cocaine, and amphetamine). The order of drug treatment was randomized and no effect of drug order was observed. For saline, cocaine, and amphetamine, drugs were given i.h (saline) or 1.5 h (amphetamine and cocaine) apart. Because of the longer effect of morphine on extracellular DA levels, subsequent drug administration was given 5 h after morphine administration. Dialysate samples were analyzed with HPLC analysis (Koizumi et al., 2004). Animals were immediately killed and brains preserved in histological fixative (4% PFA). Brains were coronally sectioned and mounted on glass slides. Sections were counterstained with cresyl violet, dehydrated, and coverslipped to confirm probe placement.
Statistics

All data are expressed as mean ± SEM. For analysis of behavior, Western blotting, and immunohistochemical labeling, data were analyzed using either a one-way or two-way ANOVA followed by post hoc analysis. Fluorescence lifetime Cl− measurements were analyzed by programmed MATLAB software. The average slope (change in Cl− over time) per slice over the first 40 s after high K+ perfusion was averaged across the group. The effects of treatment were analyzed by a one-way ANOVA. Differences were considered statistically different with *p < 0.05.

Results

To test whether chronic pain disrupts DA transmission, we measured evoked extracellular DA release using in vivo microdialysis.

Figure 1. Loss of opioid- and cocaine-evoked DA release in chronic pain. Morphine (10 mg/kg) failed to stimulate extracellular DA in the PNI group, but treatment with minocycline (PNI+MC; 30 mg/kg) restored morphine-evoked extracellular DA levels. Similarly, cocaine-evoked (10 mg/kg) DA levels were lower in the PNI group compared with the sham group and cocaine-evoked extracellular DA could be recovered by pretreatment with minocycline (PNI+MC). Amphetamine-evoked (1 mg/kg) extracellular DA was not significantly different between PNI and sham groups and was not altered by MC treatment. ***p < 0.001. Error bars indicate SEM. Insets depict area under the curve (AUC) for the entire measurement period of DA dialysis for each group. *p < 0.05. Error bars indicate SEM.

Figure 2. Systemic opioid- and cocaine-induced place preference in chronic pain. a, Systemic morphine (10 mg/kg) produced a place preference in control and chronic pain (PNI) groups. When animals received limited conditioning trials (one exposure to 10 mg/kg morphine and one to saline), only the sham group displayed a robust place preference. Systemic low dose of morphine (1 mg/kg) produced a trend toward a place preference, but was not significant in either the sham or the PNI group. *p < 0.05, **p < 0.01. Error bars indicate SEM. b, Systemic cocaine (10 mg/kg) produced a place preference in both sham and PNI groups. **p < 0.01, ***p < 0.001. Error bars indicate SEM. c, Morphine (10 mg/kg) and cocaine (10 mg/kg) antinociception was measured using the tail withdrawal assay. Neither morphine nor cocaine antinociception was significantly different between sham or PNI groups. Further, inhibition of microglia in chronic pain animals (PNI+MC) did not affect morphine antinociception. Error bars indicate SEM. n.s., Not significant; %MPE, percent maximum possible effect.
In the sham surgery group (control), systemic morphine (10 mg/kg) evoked a robust increase in extracellular DA in the NAc (Fig. 1). However, morphine failed to elicit an increase in extracellular DA in the chronic pain group (Fig. 1; F_{pain state(2,138)} = 19.73, p = 0.0001, F_{time(13,138)} = 2.74, p = 0.002, F_{interaction(26,138)} = 1.01, p = 0.45). Cocaine also showed deficits in its ability to evoke DA release in the chronic pain group (F_{pain state(2,52)} = 5.95, p = 0.004, F_{time(6,52)} = 4.28, p = 0.001, F_{interaction(12,52)} = 1.44, p = 0.18) compared with the control group. Interestingly, amphetamine, which causes an increase in extracellular DA by promoting vesicular release and transporter blockade, stimulated extracellular DA levels in both control and chronic pain groups, with no difference observed between groups (F_{pain state(2,38)} = 1.32, p = 0.23, F_{time(6,38)} = 9.1, p = 0.0001, F_{interaction(12,38)} = 1.01, p = 0.46).

To determine whether the loss of evoked release of DA in the mesolimbic system of chronic pain animals was associated with a loss of reward-like behavior, we determined the effects of morphine and cocaine-induced CPP in control and chronic pain animals. Systemic morphine (10 mg/kg) produced a robust place preference in both control and chronic pain groups after conditioning for 8 d (Fig. 2a; F_{drug side(1,15)} = 9.45, p = 0.008; F_{pain state(1,15)} = 0.51, p = 0.49; F_{interaction(1,15)} = 0.0005, p = 0.94). However, when animals were conditioned for only 2 d (once with morphine and once with saline), the chronic pain group failed to show a place preference to morphine, whereas in the control group, morphine place preference was observed (Fig. 2a; F_{drug side(1,26)} = 10.61, p = 0.003, F_{pain state(1,26)} = 0.02, p = 0.90, F_{interaction(1,26)} = 2.80, p = 0.1). To determine whether the chronic pain animals exhibited state-dependent learning to systemic morphine after only two conditioning sessions, mice were injected with morphine (10 mg/kg, i.p.) 72 h after conditioning. A state-dependent effect was evident with a difference score between morphine and saline compartments of 540 ± 198 s (t = 2.732, p < 0.05). A lower dose of morphine (1 mg/kg) produced a trend toward a place preference, but was not significant in either the control or the chronic pain group (F_{drug side(1,8)} = 3.56, p = 0.09, F_{pain state(1,8)} = 2.94, p = 0.12, F_{interaction(1,8)} = 0.002, p = 0.96). Contrary to expectations, systemic administration of cocaine (10 mg/kg) also produced a place preference in both the control and chronic pain groups (Fig. 2b; F_{drug side(1,12)} = 37.78, p = 0.0001, F_{pain state(1,12)} = 0.006, p = 0.94, F_{interaction(1,12)} = 0.45, p = 0.62). To determine whether systemic administration of cocaine or morphine had the potential to produce negative reinforcement in chronic pain animals (via eliciting analgesic effects), we assessed whether cocaine and morphine increased tail withdrawal latencies in both control and chronic pain animals. The analgesic effects of cocaine and morphine on nerve injury-induced pain hypersensitivities of the hindpaw could not be assessed because of drug-induced hyperlocomotion; therefore, the analgesic effects were measured via the tail immersion assay. Systemic morphine (10 mg/kg, i.p.) produced a large antinociceptive response that was not significantly different in chronic pain animals compared...
Figure 4. Chronic pain activates microglia in the VTA. 

**Discussion**

In the present study, we show that chronic pain results in decreased accumbal morphine- and cocaine-stimulated extracellular DA levels that contribute to impaired reward-related...
behaviors. We have identified activated microglia and BDNF as key mediators in this dysfunction of the mesolimbic DA system by disrupting the Cl− gradient in GABAergic interneurons of the VTA.

The loss of morphine- and cocaine-stimulated extracellular DA in the NAc of the chronic pain group was striking (Fig. 1). A prior study conducted 3 d after nerve injury described a partially attenuated morphine-stimulated DA response (Ozaki et al., 2002). This supports our current findings and suggests that dysfunction in mesolimbic DA circuitry becomes increasingly apparent as animals transition from acute to chronic pain. An impaired mesolimbic DA response in chronic pain is also supported by a recent human fMRI study that found dramatically reduced reward-stimulated activity in the VTA of fibromyalgia patients (Loggia et al., 2014).

Despite the lack of morphine-evoked striatal DA release in chronic pain animals, systemic morphine continued to produce a place preference. One hypothesis is that the retained opioid place preference in chronic pain animals despite the absence of DA activation is due to the negative reinforcement (pain relief) and, indeed, morphine antinociception was found to be intact in chronic pain animals (Fig. 2c; Navratilova et al., 2012). It should be recognized that the role of DA signaling in opioid reward is

Figure 5. Chronic pain disrupts Cl− transport in VTA GABAergic through a microglia-BDNF signaling pathway. a, KCC2 labeling in the VTA was primarily localized in processes not labeled for TH. Immunolabeling of KCC2 (green) was identified in non-TH (red)-labeled neurons and the labeling was punctate, which is indicative of terminal arborization. Scale bar, 75 μm. The right micrographs represent a single image of a TH + neuron (middle, arrowhead) surrounded by KCC2-immunoreactive arbors (bottom). The top image is a merge of the bottom two images. b, The chronic pain (PNI) group exhibited decreased KCC2 expression in the VTA that could be recovered by systemic treatment with minocycline (PNI + MC; 30 mg/kg). Left black and white micrographs are representative images of KCC2 labeling in the VTA of sham and PNI groups. The right histogram shows KCC2 quantification. Error bars indicate SEM. *p < 0.05, **p < 0.01. AU, Arbitrary units. c, Top left, KCC2 function was assayed in VTA slices using MQAE FLIM. The VTA was localized with brain atlas coordinates and confirmed with TH labeling (red) after Cl− imaging assay. Bottom left, Representative micrographs showing MQAE loading in GAD65-GFP cells (white arrows) in the VTA (2-photon optical section = 1 μm). Top right, Pseudocolor images showing lifetime maps from control VTA slices in the presence of 2.5 or 15 mM KCl. d, Representative lifetime plots from VTA slices. e, Graph plotted by averaging the slope in the lifetime plots immediately after administration of 15 mM KCl. The PNI group had significantly slower fluorescence lifetime change rate indicative of a slower transport of Cl− into the cell. Normal transport rate was restored in animals pretreated with minocycline (PNI + MC) or in slices treated with a TrkB antagonist (PNI + anti-trkB). (*p < 0.05; n = 7 animals with at least 2 slices per animal, 14–20 slices/condition, and 70–100 cells, on average 5/slice). Error bars indicate SEM.

f, VTA punches from the PNI group showed significantly elevated BDNF protein, which was blocked with pretreatment with MC. *p < 0.05 compared with sham, #p < 0.05 compared with PNI + saline. Error bars indicate SEM.
complex. Several prior studies have indicated that mesolimbic DA activity is not required for opioid place preference (Hnasko et al., 2005; Vargas-Perez et al., 2009) but, rather, that DA release may be involved in the salience or memory processing of rewarding stimuli (Berridge, 2007). Indeed, when only one morphine conditioning trial was run, chronic pain animals failed to display a place preference (Fig. 2a). We suggest that this may be a reflection of a blunted DA release impairing reward-learning behavior in this shortened conditioning trial paradigm. When an opioid agonist (1 ng of DAMGO) was injected directly into the VTA, it failed to produce a place preference in the chronic pain group (Fig. 3a). This further supports the premise that chronic pain leads to a loss of mesolimbic DA-driven reward. Systemic cocaine, like morphine, exhibited a place preference in both control and chronic pain groups. However, cocaine has a well-documented analgesic effect (Fig. 2c; Lin et al., 1989) and it is possible that cocaine place preference could be a result of negative reinforcement as a result of pain relief. The presence of morphine and cocaine place preference after systemic drug administration despite the absence of drug-evoked striatal DA release suggests that circuitry outside of the mesolimbic DA system, possibly related to relief of pain and negative reinforcement, drives place preference in chronic pain states.

One parsimonious hypothesis for the loss of opioid-evoked DA release is a loss of function or expression of the mu opioid (MOP) receptor in the VTA (Nikura et al., 2008). Injection of a higher dose of DAMGO (25 ng) into the VTA produced a full CPP in chronic pain animals, suggesting that MOP receptors are still functional, but higher doses of opioids are needed to evoke increases in extracellular DA in chronic pain animals. This result would be consistent with a desensitized opioid system. To address this potential confound, we have probed the mesolimbic system with nonopioid, DA-stimulating drugs. Loss of DA-mediated reward was not restricted to opioids and a similar phenomenon was observed with cocaine. Injection of cocaine into the NAc produced a place preference in the control group, but not in the chronic pain group. These data, along with microdialysis experiments, serve to reinforce the notion of a hyporeactive mesolimbic DA system to drug reward stimuli in chronic pain.

Our prior studies have determined that chronic pain leads to a major reduction in total tissue levels of striatal DA; however, this was not attributed to a loss of DA cell bodies in the VTA (Taylor et al., 2014). Given our results with cocaine, a drug that blocks the DA reuptake transporter, we challenged the system with amphetamine, a drug that evokes increased extracellular DA via a reverse efflux mechanism. Interestingly, amphetamine-evoked DA levels were equivalent in the control and chronic pain groups. Our data are consistent with increased inhibitory tone on DA neurons in chronic pain animals given the lack of cocaine- or morphine-stimulated extracellular DA levels. However, the unaltered amphetamine-stimulated extracellular DA levels suggest that the free intracellular DA pool is maintained in chronic pain animals.

In this study, we have identified a mechanism by which the opioid- and cocaine-stimulated DA response is blunted in chronic pain animals. We describe a decrease in Cl\(^-\) transport in GABAergic interneurons of the VTA due to a loss of KCC2 expression (Fig. 5). A loss of functional KCC2 has been shown previously to result in a shift in the GABA reversal potential, leading to disinhibition of spinal lamina I neurons (Coull et al., 2003). We identified activated microglia and BDNF as being possible components to this mechanism, in that minocycline treatment or interfering with BDNF signaling in the chronic pain group recovered Cl\(^-\) transport in GABAergic neurons and recovered morphine- and cocaine-stimulated DA release. Activated microglia are implicated in the development of chronic pain (Raghavendra et al., 2002; Ledebor et al., 2006; Loggia et al., 2015) and BDNF released from activated microglia contributes directly to the hyperexcitability of the pain pathway (Coull et al., 2005). Previous reports implicate BDNF-TrkB signaling as a negative modulator of opioid function in the VTA in models of opioid dependence (Koo et al., 2012; Vargas-Perez et al., 2014). Our findings extend these results by implicating KCC2 function on GABAergic neurons as driving this negative modulation in a chronic pain model.

Changes in the DA system likely contribute to the affective disorders associated with chronic pain and may also affect the effectiveness of many analgesics, such as morphine, the therapeutic potential of which is strongly linked to its rewarding capacity. Moreover, this work affirms the importance of microglial activation in modulating reward behavior and suggests microglial inhibitors as an effective therapeutic for disrupted DA transmission. We acknowledge that minocycline may have off-target effects and further studies are warranted to explore the utility of inhibiting microglia in this paradigm using more specific and/or local strategies of inhibition.

In conclusion, it is well established that neuroplastic changes in reward circuitry, including molecular and cellular changes within the mesolimbic DA system, lead to the genesis of negative affect (Koob and Le Moal, 2005; Koob and Volkow, 2010; Koob et al., 2014). Our present data show that chronic pain causes activation of microglia in the VTA that modifies reward by disrupting Cl\(^-\) homeostasis in GABAergic neurons. These findings represent a paradigm shift in the field of chronic pain by identifying a signaling and biophysical mechanism by which microglia alter the reward pathway.

References


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