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## Deficits in ventromedial prefrontal cortex Group1 metabotropic glutamate receptor function mediate resistance to extinction during protracted withdrawal from an extensive history of cocaine self-administration

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### Abstract

Anomalies in prefrontal cortex (PFC) function are posited to underpin difficulties in learning to suppress drug-seeking behavior during abstinence. As Group1 metabotropic glutamate receptors (mGluRs) regulate drug-related learning, we assayed the consequences of extended access to intravenous cocaine (6 hrs/day; 0.25 mg/infusion for 10 days) upon the PFC expression of Group1 mGluRs and the relevance of observed changes for cocaine-seeking. Following protracted withdrawal, cocaine-experienced animals exhibited a time-dependent intensification of cue-induced cocaine-seeking behavior and an impaired extinction of this behavior. These behavioral phenomena were associated with a time-dependent reduction in mGluR1/5 expression within ventromedial PFC (vmPFC) of cocaine-experienced animals exposed to extinction testing, but not in untested ones. Interestingly, pharmacological manipulations of vmPFC mGluR1/5 produced no immediate effects upon cue-induced cocaine-seeking behavior, but produced residual effects on a subsequent test for cocaine-seeking. At 3 days withdrawal, cocaine-experienced rats infused intra-vmPFC with mGluR1/5 antagonists, either before or after an initial test for cocaine-seeking, persisted in their cocaine-seeking akin to cocaine-experienced rats in protracted withdrawal. Conversely, cocaine-experienced rats infused with an mGluR1/5 agonist before the initial test for cocaine-seeking at 30 days withdrawal exhibited a facilitation of extinction learning. These data indicate that cue-elicited deficits in vmPFC Group1 mGluR function mediate resistance to extinction during protracted withdrawal from a history of extensive cocaine self-administration and pose pharmacological stimulation of these receptors as a potential approach to facilitate learned suppression of drug-seeking behavior which may aid drug abstinence.

### Keywords

Group I metabotropic glutamate receptors; prefrontal cortex; incubation of craving; cocaine addiction; cue-induced relapse

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In cocaine addicts, the presentation of cues that have been associated with cocaine consumption induces drug craving and may cause relapse to drug use (Childress et al., 1999, 2008; Volkow et al., 1999; Garavan et al., 2000; Goldstein et al., 2009b; O'Brien, 2009).

The presentation of such cues to cocaine addicts also results in heightened activation of the medial prefrontal cortex (mPFC; Grant et al., 1996; Childress et al., 1999, 2008; Volkow et al., 1999; Garavan et al., 2000; Goldstein & Volkow, 2002; but see Bonson et al., 2002), an area important for the inhibition of inappropriate behaviors and for behavioral flexibility as it pertains to the adjustment of a behavior according to its consequences (Rolls et al., 1994; Damasio, 1996; Konishi et al., 1998; Clark et al., 2004). Finally, the mPFC exhibits hypo-functionality in cocaine addicts (Volkow et al., 1992, 1993; London et al., 2000; Franklin et al., 2002; Matochik et al., 2003; Goldstein et al., 2009a). These observations are consistent with the hypothesis that excessive cocaine consumption results in neural adaptations within the mPFC that render the addict especially vulnerable to cue-induced relapse.

Preclinical studies provide further evidence for the importance of the mPFC for cue-induced cocaine seeking. Cue-induced reinstatement of cocaine seeking behavior, as well as incubation of craving, are paralleled by mPFC activation (Neisewander et al., 2000; Ciccocioppo et al., 2001; Hamlin et al., 2008; Hearing et al., 2008; Koya et al., 2009; Zavala et al., 2008), and inactivation of this brain area attenuates cue-induced reinstatement (McLaughlin and See, 2003; Fuchs et al., 2005), as well as the incubation of, cocaine craving (Koya et al., 2009). More specifically, glutamatergic projections from the mPFC to the nucleus accumbens are hypothesized to be important for cue-induced reinstatement of cocaine seeking behavior (Kalivas and O'Brien, 2008; Uys and LaLumiere, 2008; Van den Oever et al., 2010), and chronic cocaine self-administration produces alterations in indices of mPFC glutamate function (Robinson et al., 2001; Tang et al., 2004; Ben-Shahar et al., 2009, 2012). One component of the glutamatergic system that was shown to play a critical role in cocaine reinforcement and addiction are the Group1 metabotropic glutamate receptors (subtypes, mGluR1 and mGluR5) (Chiamulera et al., 2001; Kenny & Markou, 2004; Olive, 2010). Especially relevant to the current project are demonstrations that systemic administration of the mGluR5 antagonists MPEP or MTEP significantly attenuate cue-induced reinstatement (Backstrom and Hyytia, 2006; Iso et al., 2006; Kumaresan et al., 2009), and that systemic administration of mGluR5 agonist CDPPB facilitated the extinction of cocaine seeking (Cleva et al., 2011). Together, these data suggest that one type of neural adaptation within the mPFC that might develop after excessive cocaine use and be important for cue-induced relapse involves changes in Group1 mGluR function. The present study sought to test this hypothesis by first monitoring changes in the expression Group1 mGluRs within the mPFC after various withdrawal periods, and then manipulating such changes to test their functional significance for the incubation of cue-mediated cocaine-seeking behavior (see Tran-Nguyen et al., 1998; Grimm et al., 2001, 2002).

## Materials and Methods

### Procedure

The general procedure for all experiments described in the current manuscript was identical and consisted of the following steps in chronological order: Lever responding (food) training; Surgery; Self-administration of saline (1 hr or 6 hrs daily session, depending on the specific experiment) or cocaine (6 hrs daily sessions) across 10 daily sessions; and then, 3 or 30 days of withdrawal (rats kept in their home cages). Upon the completion of saline/ cocaine self-administration, the specific procedures varied depending upon the particular experiment. For immunoblotting studies, brain tissue was harvested from withdrawn rats at the 2 different withdrawal time-point or animals were subjected to a 2-hr cue test, followed by harvesting of brain tissue. For some neuropharmacological studies, animals received intra-ventromedial PFC (vmPFC) microinjections, followed by a 30-min cue test and then brains were harvested for histological verification of microinjector placements. For other neuropharmacological studies addressing extinction learning, animals received intra-vmPFC or dorsomedial PFC (dmPFC) microinjections followed by a 30-min cue test and then were

tested, 24 hrs later, in a 2<sup>nd</sup> 30-min cue test in the absence of any further intracranial manipulation, followed by histological verification of microinjector placement. In the following sub-sections, these various steps are described in detail. For clarity this general procedure is outlined in Figure 1.

## Subjects

The subjects were male Sprague-Dawley rats weighing 275–325g at the beginning of each experiment and obtained from Charles River Laboratories (Hollister, CA, USA). Rats were housed in a colony room controlled for temperature (25 °C) and humidity (71 %), under a 12-hr day/12-hr night cycle (lights on at 20:00 hrs). Animals were given *ad libitum* access to food and water, except during operant training for food reinforcement (see Lever Response Training below). The animals were allowed to acclimate to the colony room for at least 3 days following arrival. All experimental protocols were consistent with the guidelines of the NIH *Guide for Care and Use of Laboratory Animals* (NIH Publication No. 80-23, revised 1996) and were reviewed and approved by the University of California, Santa Barbara, Institutional Animal Care and Use Committee.

## Lever Response Training

To promote lever-pressing behavior for subsequent saline and cocaine reinforcement, rats were first trained to lever press on a FR1 schedule of food reinforcement (45 mg pellets; Noyes, Lancaster, NH, USA) in sound-attenuated operant conditioning chambers (30 × 20 × 24 cm high; Med Associates Inc., St. Albans, VT) during either a 16-hr overnight training session or during daily 1-h training sessions (depending upon availability of our chambers) as described previously by our group (e.g., Kerstetter et al. 2008; Ben-Shahar et al. 2009). For this, rats were food deprived for 24 h before the initiation of training and maintained on a restricted diet for the duration of food training. The operant chambers were equipped with two retractable levers, a stimulus light above each lever, a food pellet dispenser outside the operant box and a food trough between the levers, a house light on the wall opposite to the levers, and a speaker connected to a tone generator (ANL-926, Med Associates). During the session, each lever-press on the active lever resulted in delivery of one food pellet. Lever-presses on the inactive lever had no programmed consequences. Rats that failed to exhibit the criterion of a minimum of 200 responses on the active lever during the overnight session or a minimum of 100 responses on the active lever during 2 consecutive 1-hr sessions received additional lever response training sessions until criterion was met. Following successful acquisition of lever-pressing behavior (1 or 2 over-night sessions; 5–7 days of 1-h daily sessions), food was available *ad libitum* for the remainder of the study.

## Surgery

Following lever response training, animals underwent surgical procedures to implant chronic intravenous (IV) catheters and bilateral guide cannulae into the PFC as described previously by our group (e.g., Ben-Shahar et al. 2008; 2009). Under ketamine/xylazine anesthesia (56.25 and 7.5 mg/kg, respectively; Abbott Laboratories, North Chicago, IL, USA), rats were implanted with a chronic silastic catheter (13 cm long; 0.3 mm inner diameter, 0.64mm outer diameter; Dow Corning Corporation, Midland, MI, USA) into the right jugular vein. Atropine (0.04 mg/kg) was administered intramuscularly to minimize respiratory congestion during anesthesia and banamine (2 mg/kg; a non-opiate analgesic) was injected subcutaneously to treat post-surgical pain. Each catheter ran subcutaneously around the shoulder to back where it was secured to a threaded 22 gauge metal guide cannula (Plastics One, Roanoke, VA, USA), which emerged from the midline of the animal's back perpendicular to the dorsal surface. An obturator covered the open end of the cannula to protect from contamination and the cannula was held in place via a small swatch of Bard

Mesh (C.R. Bard Inc., Cranston, RI, USA) to which it was cemented. The mesh was, in turn, laid flat subcutaneously on the animal's back.

Immediately following the IV catheterization and while still under anesthesia, rats in the behavioral studies were transferred to a stereotaxic apparatus and implanted with 30-gauge guide cannulae (Plastics One Inc.) above the vmPFC (AP +2.5 mm, ML +/- 1.0 mm, DV -2.0 mm from Bregma) or above the dmPFC ( AP +2.5 mm, ML +/- 1.0 mm, DV -1.0 mm from Bregma), according to the atlas of Paxinos and Watson (2008), using procedures similar to those described previously by our group (Ben-Shahar and Ettenberg, 1998; Zayara et al., 2011). Four small screws and cranioplastic cement secured the guide cannulae to the skull. Stylets (Plastics One) were placed into each cannula to prevent occlusion. All animals were allowed a minimum of 5 days for recovery and IV catheter patency was maintained by flushing with 0.1 ml of sterile heparin+timentin/saline (60 IU/ml and 100mg/ml, respectively) solution each day.

### Self-administration training

Following surgery, rats (n=10–15 per group/per cohort at the start of each experiment) were trained to self-administer IV cocaine (0.25 mg/0.1ml/infusion; a generous gift from the National Institute on Drug Abuse, Bethesda, MD, USA) during daily 6-hr sessions on a FR1 schedule of reinforcement. At the start of each session, the rat's catheter was connected to a motorized pump (located outside of the sound attenuated chamber) via a liquid swivel as previously described (e.g., Ben-Shahar et al. 2008; Kerstetter et al 2008). Active lever presses resulted in a 5-sec activation of the infusion pump and a 20-sec presentation of a stimulus complex, consisting of activation of the white stimulus light above the active lever and the tone generator (78 dB, 2 kHz), during which responses on the active lever had no consequences. Responses on the inactive lever were recorded, but had no programmed consequences. Rats were trained to self-administer cocaine for 10 sessions. To prevent overdose, the number of cocaine infusions permitted during the first 2 days of training was capped at 100 (day1) and 120 (day2) and rats failing to meet self-administration criterion (minimum of 50 infusions/6-h session for the last 3 days of training) were excluded from the study.

For immunoblotting studies, additional groups of rats were trained to self-administer IV saline (0.1 ml/infusion) during daily 1-h or 6-h sessions (n=12/control group/cohort at the start of the experiment). Two saline controls groups were included in the immunoblotting study as, for practical reasons, we wished to replicate the results of prior studies indicating that extended (6-h) access to IV saline does not produce behavioral or protein changes distinct from those observed in animals with short (1-h) access to IV saline (Ben-Shahar et al., 2004). Our behavioral and immunoblotting results confirmed that there were no differences between the two saline controls (see Results) and thus, only 1h saline controls were included in the behavioral neuropharmacological studies. All self-administration training and testing occurred during the dark phase of the circadian cycle. Upon completion of the 10 days of self-administration training, animals remained in their home cages in the colony room for either 3 or 30 days withdrawal. At these time-points, PFC subregions were harvested from a subset of animals, while another subset of animals were tested for cue-elicited cocaine-seeking over a 2-hr period, followed by harvesting of PFC subregions.

### Tests for Cue-elicited Cocaine-seeking

With the exception of the one subset of withdrawn rats employed for immunoblotting, saline and cocaine self-administering rats were subjected to a test for cue-elicited cocaine-seeking under extinction conditions. For the immunoblotting experiments, the cue tests were 2 h in duration in order to capture potentially relevant changes in receptor protein expression. To

statistically evaluate the presence/absence of response incubation during withdrawal and provide a behavioral correlate for our immunoblotting data, the number of lever-responses on the 2-hr cue tests were analyzed using an IV Treatment (Sal1h, Sal6h, Coc6h) X Withdrawal (3 vs. 30 days) X Lever (active vs. inactive) ANOVA. For the behavioral neuropharmacological experiments, the cue tests were 30 min in duration, as conducted by others (e.g., Koya et al., 2009). The cue test conditions were the same regardless of the duration of testing; testing involved tethering the animals and the lever-press response-contingent presentation of the tone-light cue previously paired with saline/cocaine infusions but no saline/cocaine infusions were delivered.

### Immunoblotting Studies

Our first immunoblotting study was aimed to investigate the influence of a history of cocaine self-administration experience upon the protein expression of mGluR1/5 within PFC subregions at early versus protracted withdrawal. For this study, within each cohort of rats, separate groups of Sal1h, Sal6h and Coc6h rats underwent a 2-h cue test session under extinction conditions at either 3 or 30 days withdrawal (n=12 per group at the start of the experiment) to assay for cue-reinforced lever-pressing behavior. Immediately upon completion of this cue test, animals were killed by rapid decapitation. The dmPFC (anterior cingulate and dorsal prefrontal cortices) and vmPFC (ventral prefrontal and infralimbic cortices) were dissected out over ice in a manner identical to that described by others (Ghasemzadeh et al., 2009). The tissue derived from animals in this study was immunoblotted in 2 separate ways. First, comparisons were made across the tissue from the 3 different self-administration groups separately for each time-point (i.e., Sal1h, Sal6h & Coc6h rats sacrificed at 3 days withdrawal were compared in one assay, and a parallel assay compared Sal1h, Sal6h & Coc6h rats sacrificed at 30 days withdrawal in a separate assay). This “single time-point” analysis was done to enable a comparison of protein expression between the SAL1h and SAL6h rats at each time-point, as well as to determine whether or not a history of cocaine self-administration influenced protein expression relative to a history of saline self-administration at that particular withdrawal time-point. For these assays, the immunoreactivity of the mGluR1 and mGluR5 bands from the Sal6h and Coc6h animals was normalized to the average of that for the Sal1h animals run on the corresponding gel (n=4–5/gel) and the data expressed as a percentage of that averaged Sal1h signal for each gel. The data for the 3-day and for the 30-day assays were analyzed separately using an univariate analysis of variance (ANOVA) across the 3 different self-administration groups to determine the presence/absence of a cocaine effect.

The next analysis examined for proteomic correlates of the increased lever-pressing exhibited by both saline and cocaine self-administering animals at 30 days of withdrawal, by comparing tissue from the Sal6h and Coc6h animals sacrificed at both withdrawal time-points on the same gel. For this “time-course” assay, the immunoreactivity of the mGluR1 and mGluR5 bands were normalized to the average of the Sal6h-3 day withdrawal group run on the corresponding gel (n=3–4/gel) and the data expressed as a percentage of that averaged signal for each gel. The data were analyzed using orthogonal comparisons, within each self-administration group, to determine the presence/absence of time-dependent changes in protein expression. A second, follow-up, study assayed vmPFC and dmPFC tissue from Sal1h and Coc6h rats, sacrificed at 3 versus 30 days withdrawal, in the absence of any cue testing. This study was conducted to determine the extent to which our cocaine-induced changes in mGluR1/5 protein expression reflected alterations in the basal expression pattern of these receptors or some interaction between cocaine experience and the cue-reinforced testing conditions of our animals. The data analysis for this study was identical to that employed for the “time-course” study above.

The immunoblotting procedures for detection of total mGluR1 and mGluR5 levels in brain tissue homogenate were identical to those described recently by our group (e.g., Cozzoli et al., 2009, 2012; Goulding et al., 2011). In brief, tissue was homogenized in a solution consisting of 0.32 M sucrose, 2 mM EDTA, 1% w/v sodium dodecyl sulfate, 50  $\mu$ M phenyl methyl sulfonyl fluoride and 1  $\mu$ g/ml leupeptin (pH=7.2) and 1 mM sodium fluoride, 50 mM sodium pyrophosphate, 20 mM 2-glycerol phosphate, 1 mM p-nitrophenyl phosphate, 1 mM orthovanadate, and 2  $\mu$ M microcystin LR were included to inhibit phosphatases. After centrifugation at 10,000 g for 20 min, the supernatant of the homogenates were quantified for protein content using the Bio-Rad DC protein assay (Bio-Rad, Hercules, CA) and stored at  $-80^{\circ}\text{C}$ . Protein samples (20  $\mu$ g/lane) were subjected to SDS-polyacrylamide gel electrophoresis on Tris-Acetate gradient gels (3–8%) (Invitrogen, Carlsbad, CA). Wet polyvinylidene difluoride (PVDF) (Bio-Rad) membrane transfer was employed, and membranes were pre-blocked with either phosphate-buffered saline containing 0.1% (v/v) Tween 20 and 5% (w/v) nonfat dried milk powder for a minimum of 2 hrs before overnight incubation with primary antibody. An anti-mGluR5 rabbit polyclonal antibody (Millipore, Billerica, MA; 1:1000 dilution) and an anti-mGluR1a mouse polyclonal antibody (Millipore; 1:500 dilution) were used for receptor detection. A rabbit primary anti-calnexin antibody (Stressgen, Ann Arbor, MI; 1:1000 dilution) was used as a control to ensure even protein loading and transfer. Membranes were washed, incubated with horseradish peroxidase-conjugated goat anti-rabbit secondary antibody (Millipore; 1:5,000–1:10,000 dilution) or anti-mouse secondary antibody (Jackson Immuno Research Laboratories, West Grove, PA; 1:10,000) for 90 min, washed again, and immunoreactive bands were detected by enhanced chemiluminescence using either ECL Plus (Amersham Biosciences) or Pierce SuperSignal West Femto (Thermo Fisher Scientific, Rockford, IL). Image J (NIH, Bethesda, MD) was used to quantify immunoreactivity levels.

### Behavioral Neuropharmacological Studies

**Intra-vmPFC delivery of Group1 mGluR compounds: acute effects upon drug-seeking**—A series of behavioral neuropharmacological experiments were conducted to determine the functional relevance of reduced vmPFC Group1 mGluR expression (Figures 2 & 3) for the time-dependent increase in cue-seeking behavior during withdrawal (Figure 2A). For these experiments, Sal1h and Coc6h animals were trained to self-administer for 10 days and then bilaterally infused intra-vmPFC with compounds targeting Group1 mGluRs immediately prior to cue testing using microinjection procedures similar to those described previously (e.g., Ben-Shahar and Ettenberg, 1998; Ben-Shahar et al., 2012; Zayara et al., 2008). It was hypothesized that if the reduction in vmPFC Group1 mGluR expression observed at 30 days withdrawal mediated the time-dependent increase in lever-pressing, then mimicking this reduction earlier in withdrawal should potentiate cocaine-seeking behavior (i.e., render a 3-day withdrawn animal more similar to a 30-day withdrawn animal). Conversely, hyper-stimulating these receptors later in withdrawal should attenuate the heightened cocaine-seeking behavior (i.e., render a 30-day withdrawn animals more similar to a 3-day withdrawn animal). To test this hypothesis with respect to mGluR5, groups of 3 day-withdrawn saline and cocaine self-administering rats ( $n = 5-8$  for saline animals and  $n > 10$  for cocaine animals at the outset of experiment/replicate) were infused at 0.5  $\mu$ l/min with the selective mGluR5 antagonists 2-Methyl-6-(phenylethynyl)pyridine hydrochloride (MPEP; 3  $\mu$ g/side; Sigma-Aldrich, St. Louis, MO) or 3-((2-Methyl-1,3-thiazol-4-yl)ethynyl)pyridine hydrochloride (MTEP; 3  $\mu$ g/side; Tocris BioScience, Ellisville, MO). A parallel study assayed the effects of the selective mGluR1 antagonist (3,4-Dihydro-2H-pyrano[2,3-b]quinolin-7-yl)-(cis-4-methoxycyclohexyl)-methanone (JNJ 16259685; 30 ng/side; Tocris). To test the converse hypothesis, groups of 30 day-withdrawn saline or cocaine self-administering rats ( $n > 8$ /group/replicate) were infused with the mGluR1/5 agonist (*S*)-3,5-Dihydroxyphenylglycine (DHPG; 27.5 ng/side; Tocris

BioScience). The antagonist doses were selected based on published reports indicating an attenuation of drug-taking or drug-seeking behaviors when microinjected into the brain in doses at, or lower than, the doses employed in the present studies (e.g., Backstrom and Hyytia, 2007; Cozzoli et al., 2009, 2012; Gass and Olive, 2009a; Kumaresan et al., 2009; Xie et al., 2010, 2011). The dose of DHPG was selected as it elicits a marked rise in locomotor activity and glutamate release when infused into brain (Swanson and Kalivas, 2000; Swanson et al., 2001; Melendez et al., 2005; Szumlinski et al., 2004). Injectors (projecting 2 mm below the tip of the guide cannulae) were left in place for 1 min following infusion and then animals were tested for cue-maintained lever-pressing behavior in a 30-min test immediately following the intracranial injection. Control animals received intra-vmPFC infusions of 0.1% DMSO vehicle (for JNJ 16259585) or water (for MPEP, MTEP and DHPG). The data for this reiteration of the antagonist study were analyzed using an IV (saline vs. cocaine) X IC (VEH, MPEP, MTEP, JNJ) ANOVA. As only cocaine-experienced animals were assayed in this reiteration of the DHPG study, the data were analyzed using t-tests across the IC factor.  $\alpha = 0.05$  for all analyses.

**Intra-vmPFC infusion of Group1 mGluR compounds: residual effects upon drug-seeking**—We next tested the possibility that the reduction in vmPFC Group1 mGluR expression observed at 30 days withdrawal from cocaine self-administration (Figures 2D, 3A) might render an animal more resistant to extinction if tested repeatedly. To test this hypothesis: (1) animals in the acute MTEP experiment were subjected to a second drug-seeking test (Day 4 of withdrawal) in the absence of any further intracranial manipulations; (2) the JNJ 16259585 study was replicated in a separate group of animals, which were tested again for effects of vmPFC mGluR1 blockade on Day 3 of withdrawal, followed by a second cue test in the absence of any further pretreatment on Day 4 of withdrawal; and (3) the DHPG was replicated in a separate group of animals, which were tested again for effects of vmPFC group 1 mGluR stimulation on Day 30 of withdrawal, followed by a second cue test in the absence of any further pretreatment on Day 31 of withdrawal. To determine whether or not the effects of vmPFC Group1 mGluR antagonism upon behavior on the second cue test reflected an impairment in operant learning during the first cue-reinforced test session or an impairment in memory consolidation post-session, additional cocaine-experienced rats were infused intra-vmPFC with 30 ng/side JNJ 16259585 immediately following the first test session on Day 3 of withdrawal and then tested subsequently for cue-induced lever-pressing behavior the next day in the absence of any further intracranial manipulation. Finally, to confirm that the behavioral effects that we observed resulted from manipulating Group1 mGluR signaling specifically within the ventral region of the medial PFC, additional groups of Coc6h rats were infused intra-dmPFC (injectors projecting 1 mm below the tip of the guide cannulae; see Figure 6) with 30 ng/side of JNJ 16259585 or vehicle immediately prior a test for cue-seeking at 3 days withdrawal, followed by a second cue in the absence of any further pretreatment on Day 4 of withdrawal. The data from these neuropharmacological studies were analyzed initially using an IV (saline vs. cocaine) X IC (VEH, MTEP, JNJ or VEH, DHPG) X Test (Test 1 vs. Test 2) ANOVA (results not reported). In all cases, we observed significant 3-way interactions ( $p < 0.05$ ) and thus, the data were deconstructed first along the IV factor and analyzed using an IC X Test ANOVA to determine whether or not intra-PFC manipulations depended upon the drug experience of the animal. To statistically confirm the presence/absence of extinction, Significant IC X Test interactions were then deconstructed along the IC factor and analyzed using a repeated measures ANOVA across the 2 cue test sessions ( $\alpha = 0.05$  for all analyses). A significant reduction in lever-responding from Test 1 to Test 2 indicated the presence of extinction, while a lack of statistical differences were interpreted to reflect an extinction impairment.

## Histology

Upon the completion of all behavioral pharmacological experiments, animals were killed for histological verification of microinjector placement within the vmPFC or dmPFC. Only animals exhibiting correct placement within ventral prelimbic or within the infralimbic cortices were included in the statistical analyses of the vmPFC neuropharmacological data. Only animals exhibiting correct placement within the dorsal prelimbic or within the anterior cingulate were included in the statistical analyses of the dmPFC neuropharmacological data. Examples of microinjector tip placements from several of the experiments in this study are included in Figure 5. For the most part, 2–3 animals/group were excluded from the statistical analyses of the behavioral data due to misplaced microinjection sites or occlusion of the guide cannulae and the final sample sizes employed in the statistical analyses of the data are indicated in the appropriate figure or table legend.

## Results

### Time-dependent intensification of cue-induced drug-seeking occurs following protracted withdrawal only in cocaine self-administering animals

An analysis of the behavior exhibited by our 3 IV treatment groups during a 2-hr test for cue-reinforced behavior conducted under extinction conditions at 3 versus 30 days withdrawal revealed a significant IV Treatment (Sal1h, Sal6h, Coc6h) X Withdrawal (3 vs. 30 days) X Lever (active vs. inactive) interaction [ $F(2,58)=12.09$ ,  $p<0.0001$ ]. Animals with a 10-day history of extended access IV cocaine self-administration (Coc6h) exhibited a time-dependent increase (~50%) in the number of active lever-presses emitted for the contingent presentation of the tone-light stimulus complex previously paired with cocaine delivery when tested in the absence of further cocaine delivery (Figure 2A) and there was no concurrent change in the number of inactive lever-presses (Figure 2B) [Withdrawal X Lever ANOVA:  $F(1,16)=12.55$ ,  $p=0.003$ ; for active lever,  $F(1,16)=5.29$ ,  $p=0.04$ ; for inactive lever,  $p=0.72$ ]. In addition, as no differences were observed in self-administration behavior during training, between the Coc6h groups tested (and killed) at the different withdrawal time-points, the time-dependent increase in cocaine-seeking behavior could not be explained by group differences in cocaine intake or behavioral responding for drug (Table 1). As expected, animals with a history of either brief or extended access to IV saline (respectively, Sal1h and Sal6h) exhibited significantly lower levels of active lever-pressing than cocaine self-administering animals both during self-administration training (Table 1) and during cue testing (Figure 2). Importantly, in contrast to the cocaine groups, saline groups exhibited a time-dependent increase in responding on both active and inactive levers [for Sal1h: Withdrawal effect:  $F(1,21)=16.93$ ,  $p<0.0001$ ; Withdrawal X Lever:  $p=0.4$ ; for Sal6h: Withdrawal X Lever:  $F(1,21)=8.21$ ,  $p=0.009$ ; one-way ANOVA for active lever:  $F(1,21)=19.40$ ,  $p<0.0001$ ; for inactive lever:  $F(1,21)=4.11$ ,  $p=0.05$ ].

### Cue-induced cocaine-seeking produces subregionally distinct and time-dependent changes in PFC Group1 mGluR expression during withdrawal

We next employed conventional immunoblotting strategies to relate the time-dependent increase in cue-reinforced responding to the levels of mGluR1 and mGluR5 within PFC subregions. A comparison of vmPFC receptor levels between Sal1h, Sal6h and Coc6h animals failed to indicate group differences for any protein when animals were sacrificed immediately following the 2-h test session at 3 days withdrawal (Figure 2C; one-way ANOVAs:  $p>0.05$ ); however, vmPFC levels of both receptors were reduced in Coc6h rats, relative to both saline groups, when animals were sacrificed following testing at 30 days withdrawal (Figure 2D) [mGluR1:  $F(2,33)=13.78$ ,  $p<0.0001$ ; mGluR5:  $F(2,33)=5.74$ ,  $p=0.008$ ; LSD post-hoc tests:  $p<0.05$ ]. As these data suggested a time-dependent decrease in vmPFC mGluR1/5 levels selectively in animals with a history of cocaine self-



administration, as well as no differences between the saline groups ( $p > 0.05$ ), we conducted a second round of immunoblotting to directly compare for changes in protein expression following cue-testing at 3 versus 30 days in Sal6h and Coc6h animals (Figure 3). Orthogonal comparisons between the 2 withdrawal time-points indicated no change in vmPFC mGluR1/5 expression in Sal6h rats (t-tests,  $p > 0.05$ ), while the levels of both receptors were significantly lower within the vmPFC at the 30-day versus 3-day time-point in Coc6h animals (Figure 2A) [mGluR1:  $t(16) = 2.61$ ,  $p = 0.02$ ; mGluR5:  $t(16) = 2.20$ ,  $p = 0.05$ ]. Interestingly, the influence of cocaine experience and withdrawal upon vmPFC mGluR1/5 expression appeared to depend upon the test history of the subjects as no group differences in vmPFC receptor expression were observed when protein was analyzed in test-naïve rats (Figure 3B; t-tests, all  $p$ 's  $> 0.05$ ). Thus, the opportunity to engage in cue-reinforced behavior following protracted cocaine withdrawal, and not cocaine withdrawal *per se*, reduced vmPFC Group1 mGluR expression.

In contrast to the vmPFC, dmPFC levels of mGluR1 and mGluR5 were elevated in Coc6h animals, relative to both saline self-administration groups at both 3 days (Figure 2E) [mGluR1:  $F(2,29) = 3.64$ ,  $p = 0.04$ ; mGluR5:  $F(2,29) = 4.23$ ,  $p = 0.03$ ; LSD post-hoc tests] and 30 days withdrawal (Figure 2F) [mGluR1:  $F(2,32) = 27.98$ ,  $p < 0.0001$ ; mGluR5:  $F(2,32) = 5.83$ ,  $p = 0.007$ ; LSD post-hoc tests]. A direct comparison of dmPFC tissue from Sal6h rats across both withdrawal time-points (Figure 3C) revealed no time-dependent changes in mGluR5, but a significant time-dependent reduction in mGluR1 [ $t(21) = 2.19$ ,  $p = 0.04$ ]. No time-dependent changes in dmPFC receptor expression were apparent in cocaine-experienced animals (t-test:  $p < 0.05$ ), despite their obvious increase in mGluR1/5 expression relative to saline-experienced animals. Although there was a trend for test-naïve, cocaine-experienced animals to exhibit elevated dmPFC mGluR1 expression at both withdrawal time-points, group differences in basal mGluR1 or mGluR5 expression were not statistically reliable (Figure 3D;  $p$ 's  $> 0.05$ ).

### **Intra-vmPFC pharmacological manipulations of Group1 mGluR function do not acutely affect cue-induced seeking behavior during withdrawal**

The average number of infusions earned across the last 3 self-administration sessions were similar in the different intra-vmPFC treatment groups (Table 2;  $p$ 's  $> 0.05$ ). While an intra-vmPFC infusion of the mGluR1a antagonist JNJ16259685 appeared to increase the number of active lever-presses emitted by cocaine rats, the results of the statistical analysis failed to indicate any significant acute effects of our various intra-vmPFC pretreatments upon the behavior of either Sal1h (Table 3) or Coc6h (Figure 4A) animals when tested for cue-reinforced responding at 3 days withdrawal [IV effect:  $F(1,114) = 128.02$ ,  $p < 0.0001$ ; IC effect and IC X IV interaction,  $p$ 's  $> 0.15$ ]. In addition, we failed to observe any influence of our intra-vmPFC pretreatments upon the number of inactive lever-presses emitted during this test session (total inactive lever-presses in 30 min, VEH:  $5.3 \pm 1.7$ ; MPEP:  $4.8 \pm 1.2$ ; MTEP:  $8.4 \pm 2.8$ ; JNJ:  $3.6 \pm 1.0$ ; one-way ANOVA,  $p = 0.36$ ).

Similarly, the data from a study employing only cocaine-experienced animals also failed to indicate any significant effect of an intra-vmPFC infusion of the Group1 mGluR agonist DHPG upon lever-pressing on a test conducted at 30 days withdrawal (Figure 4B) (t-test,  $p > 0.20$ ).

### **vmPFC Group1 mGluR blockade impairs extinction learning during short-term withdrawal from cocaine**

To determine whether the reduction in vmPFC Group1 mGluR expression observed at 30 days withdrawal in Coc6h animals tested for cue-reinforced behavior relates to impaired extinction learning in these animals, we assayed for residual effects of vmPFC Group1

mGluR blockade either before or after the initial test of cue-reinforced behavior (without further pharmacological manipulation), 24 h later. As was the case for the initial studies, there were no significant differences regarding the saline/cocaine intake of the animals during the last 3 days of self-administration training between the different intracranial treatment groups prior to testing (Table 2; all  $p$ 's>0.05).

As was observed previously, manipulations of vmPFC Group1 mGluRs failed to influence the behavior of saline-experienced animals on either test, although all saline groups exhibited greater lever-pressing behavior on the second test, in the absence of intracranial microinjection (Table 3) [Test effect:  $F(1,27)=19.8$ ,  $p<0.0001$ ; all other  $p$ 's>0.50]. A comparison of the behavior of cocaine-experienced rats revealed that the change in behavior across the 2 cue tests depended upon the intracranial treatment of the animals (Figure 5A) [IC Group X Test:  $F(3,37)=3.98$ ,  $p=0.02$ ]. Deconstruction of the significant interaction along the IC Group factor, revealed a test-dependent reduction in responding in rats infused intra-vmPFC with vehicle prior to initial testing for cue-reinforced behavior [ $F(1,10)=9.17$ ,  $p=0.01$ ], indicating extinction learning. Interestingly, cocaine-experienced rats infused intra-vmPFC with either MTEP or JNJ16259685 prior to initial testing failed to show this test-dependent reduction in responding (respectively,  $p=0.27$  and  $p=0.61$ ), indicating impaired extinction learning or memory. In support of the latter possibility, cocaine-experienced animals infused intra-vmPFC with JNJ16259685 immediately following the initial test also failed to exhibit a test-dependent reduction in lever-pressing behavior, compared to controls ( $p=0.45$ ).

We also assayed the acute and residual effects of intra-dmPFC pretreatment with JNJ16259685 upon cue-reinforced responding at 3 days withdrawal. As illustrated in Figure 5B, acute pretreatment with the mGluR1 antagonist into the dmPFC failed to influence cue-reinforced behavior, nor did it alter the test-dependent reduction in behavior observed at 3 days withdrawal [Test effect:  $F(1,14)=5.46$ ,  $p=0.04$ ; IC effect and interaction,  $p>0.90$ ].

### **vmPFC Group1 mGluR stimulation facilitates extinction learning following protracted withdrawal from cocaine**

Here again, no differences were observed for the average number of lever-presses emitted for infusions of either saline or cocaine during self-administration training between the groups slated to receive intra-vmPFC vehicle versus DHPG (Table 2) [IV effect:  $F(1,45)=66.15$ ,  $p<0.0001$ ; no IC effect or interaction,  $p$ 's>0.20]. Similar to the results for the antagonists, intra-vmPFC DHPG pretreatment failed to alter the behavior of saline-experienced animals and all rats exhibited higher responding on the second test, in the absence of any further intracranial manipulation (Table 3) [Test effect:  $F(1,26)=9.07$ ,  $p=0.006$ ; IC effect and interaction,  $p>0.17$ ]. Replicating the data from our initial study (Figure 4B), we failed to detect a significant acute effect of intra-vmPFC DHPG in cocaine-experienced animals upon cue-reinforced behavior at 30 days withdrawal. In fact, an analysis of the time-course (in 5-min intervals) of lever-pressing behavior during the first cue test indicated that within-session extinction did not differ between VEH- and DHPG-pretreated cocaine-experienced rats [Time effect:  $F(5,70)=11.67$ ,  $p<0.0001$ ; Time X IC interaction:  $p>0.05$ ]. However, we observed differential effects of intra-vmPFC pretreatment upon the change in behavior upon subsequent testing (Figure 5C) [Test effect:  $F(1,21)=5.55$ ,  $p=0.03$ ; IC effect:  $p=0.62$ , Test X IC:  $F(1,21)=6.15$ ,  $p=0.02$ ]. As illustrated in Figure 5C, vehicle-infused cocaine rats exhibited no change in behavior across the two cue tests ( $p=0.90$ ), indicating resistance to extinction during protracted withdrawal. In contrast, DHPG-infused cocaine rats exhibited an approximately 30% reduction in lever-pressing during subsequent testing and this was statistically significant [ $F(1,6)=8.63$ ,  $p=0.03$ ].

## Discussion

The present report provides validation of an important role for vmPFC Group1 mGluRs in learning to suppress cocaine-seeking behavior during protracted withdrawal from heavy cocaine intake. Specifically, “incubated” drug-seeking and across-session extinction deficits exhibited by cocaine-experienced rats during protracted withdrawal were associated with reduced vmPFC mGluR1/5 levels, which reflected an interaction between cocaine withdrawal and the opportunity to engage in cue-reinforced behavior under extinction conditions. Interestingly, intra-vmPFC manipulations of mGluR1/5 function had no immediate effects upon cue-reinforced behavior under extinction conditions but produced residual effects upon subsequent behavioral testing, 24 hrs later. Intra-vmPFC mGluR1/5 blockade, either before or after testing for cue-reinforced behavior, impaired, while receptor hyper-stimulation prior to testing facilitated, the manifestation of extinction learning in cocaine-experienced rats during short- and long-term withdrawal, respectively. In contrast, intra-dmPFC mGluR1 blockade failed to influence cue-reinforced behavior. Taken together, these results support the hypothesis that reduced vmPFC Group1 mGluR function is a time-dependent adaptation that manifests when a heavily cocaine-experienced animal is given opportunity to engage in drug-seeking during withdrawal. This reduction in vmPFC Group1 mGluR function impairs the ability to consolidate memories related to new behavior-outcome contingences in the absence of drug, rendering it more difficult to suppress drug craving and drug-seeking behavior when presented with stimuli predictive of drug availability.

### Cue-reinforced behavior and the regulation of PFC Group1 mGluR expression

Alterations in PFC mGluR1/5 expression were observed only in cocaine-experienced animals allowed a single (2-hr) opportunity to engage in cue-reinforced behavior in the absence of cocaine. As reported previously (Ben-Shahar et al., 2009; Hao et al., 2010; Ghasemzadeh et al., 2011), we failed to detect changes in PFC Group1 mGluR expression in cocaine-experienced animals left undisturbed in the home cage. Using procedures akin to those employed in the present study, Ghasemzadeh et al. (2011) also failed to detect changes in either vmPFC or dmPFC mGluR5 expression in cocaine-experienced animals withdrawn for 10 days in the absence of extinction training. However, they also failed to observe receptor changes within the vmPFC of animals with extensive (10 X 6-h sessions) extinction training, but observed *reduced* mGluR5 levels within the dmPFC in behaviorally extinguished animals (Ghasemzadeh et al., 2011). Such data indicate that, akin to reports for mPFC AMPA glutamate receptors in heroin-experienced animals (Van den Oever et al., 2008), mGluR1/5 expression within both the vmPFC and dmPFC appears to be regulated in a complex manner by interactions between cocaine experience, duration of withdrawal and the opportunity to engage in cue-reinforced behavior in the absence of cocaine. Further, given the discrepancies in biochemical results between the present study (1 X 2-hr extinction session) versus the earlier study by Ghasemzadeh and colleagues (10 X 6-hr extinction sessions), it would appear that cocaine’s effects upon mGluR1/5 expression within both the vmPFC and dmPFC depends also on the extent to which animals have undergone extinction training and/or manifest drug-seeking behavior, with vmPFC mGluR1/5 levels inversely related to drug-seeking and dmPFC receptor levels positively related to drug-seeking.

Group1 mGluRs undergo rapid desensitization involving receptor internalization, then degradation and this develops tolerance upon repeated receptor activation (e.g., Aronica et al., 1993; Desai et al. 1995, 1996; Gereau and Heinemann, 1998). Thus, an incubation of cue-elicited glutamate release within the vmPFC might contribute not only to down-regulated receptor expression observed within this subregion during initial extinction learning (present study), but also the apparent tolerance to this effect upon extensive extinction training (Ghasemzadeh et al. 2011). At the present time, it is difficult to envision

how a similar mechanism might be in place to regulate mGluR1/5 expression within dmPFC as receptor levels are increased in animals during initial extinction learning (present study), but decreased in behaviorally extinguished animals (Ghasemzadeh et al., 2011). It is not known whether or not cue-elicited glutamate release within PFC subregions varies as a function of cocaine experience/withdrawal. As Group1 mGluRs are perisynaptic in localization, future studies should focus on elucidating the relation between subregional differences within PFC glutamate receptor expression and extracellular glutamate and how this relation varies as a function of drug-related learning to begin to understand more precisely the mechanisms through which mGluR1/5 is differentially regulated within PFC subregions by the opportunity to engage in cue-reinforced behavior in the absence of cocaine.

### Neuropharmacological manipulations of vmPFC Group1 mGluRs and cue-seeking

Pharmacological manipulations of Group1 mGluR function produced no obvious *acute* effects upon cue-reinforced responding. Thus, neither vmPFC nor dmPFC Group1 mGluRs play a major role in the cognitive, motivational or motor capacity required to engage in cue-reinforced behavior or in the processes that underpin the expression of incubated cue-reinforced behavior during protracted withdrawal. To our knowledge, this study is the first to address the role for PFC Group1 mGluRs during initial extinction of cocaine-seeking. As such, we employed doses of MPEP, MTEP and JNJ 16259685 that are within the range of those demonstrated to maximally inhibit drug reward and/or reinforcement when infused intracranially in other animal models of drug-seeking or –taking and the dose of DHPG employed was at, or above, maximal for increasing locomotion and extracellular glutamate levels when infused into the brain (see Materials and Methods). The fact that we observed residual behavioral effects of our intra-vmPFC pretreatments argues against inadequate antagonist/agonist dosing as an explanation for our negative behavioral data. To date, all of the studies that have examined a potential role for Group1 mGluRs in cue- or context-reinforced behavior have employed animals with extensive extinction training and typically, antagonist infusion into either the nucleus accumbens or hippocampus reduces cocaine-seeking upon cue/context re-exposure (mGluR1: Dravolina et al., 2007; mGluR5: Backstrom et al., 2004; Bespalov et al., 2005; Schroeder et al., 2008; Kumaresan et al., 2009; Martin-Fardon et al., 2009; Moussawi et al. 2009; Gass and Olive, 2009b; Xie et al., 2010; Martin-Fardon and Weiss, 2011; Xie et al., 2011). Although there is some indication that prelimbic and infralimbic cortical inactivation influences drug-seeking behavior only in animals having undergone extinction training (e.g., McFarland and Kalivas, 2001; Capriles et al., 2003; Fuchs et al., 2006; Peters et al., 2008; but see Koya et al., 2009), the role of Group1 mGluRs within PFC subregions in drug-seeking under extinction conditions remains to be examined.

### Neuropharmacological manipulations of vmPFC Group1 mGluRs and behavioral extinction

More ventral aspects of the PFC are posited to play a critical role in extinction of cocaine-seeking behavior (e.g., Schmidt et al., 2005; Peters et al., 2008; Van den Oever et al., 2008; Hayton et al., 2010) and there is an important glutamatergic component in this regard (e.g., Van den Oever et al., 2008; Gass and Olive, 2009b; Hayton et al., 2010; c.f. also Millan et al., 2011). Consistent with this idea, dmPFC mGluR1 antagonist infusion failed to influence the manifestation of extinction learning during early withdrawal, while pharmacological manipulations of vmPFC mGluR1/5 function exerted extinction-modulating effects on the residual tests of cue-reinforced behavior. Mimicking the cocaine-induced reduction in vmPFC mGluR1/5 expression observed following protracted withdrawal was sufficient to elicit an extinction impairment akin to that exhibited by cocaine-experienced animals during protracted withdrawal. Interestingly, a similar MPEP-induced impairment of extinction of conditioned fear has been recently reported (Fontanez-Nuin et al., 2011) and is consistent

with a general role of mGluR1/5 receptors in consolidation of learning (see for example, Riedel et al., 2003). Conversely, “rescuing” vmPFC Group1 mGluR function by high-dose DHPG infusion reversed the extinction deficit and reduced subsequent drug-seeking behavior exhibited by cocaine-experienced rats during protracted withdrawal. Overall, these data are consistent with recent demonstrations that systemic pretreatment with an mGluR5 positive allosteric modulator facilitates behavioral extinction and reduces drug-seeking behavior in cocaine-experienced animals (Cleva et al., 2011) and point to vmPFC Group1 mGluRs as an important neural substrate in mediating these pro-cognitive and anti-addictive effects.

Electrophysiological and immunocytochemical evidence indicates there is hyper-activation of vmPFC neurons by the behavior-contingent presentation of cocaine-paired cues early during extinction training in animals with a history of cocaine self-administration (Koya et al., 2009; Sun and Rebec, 2006). Particularly noteworthy are the parallel results for reduced vmPFC mGluR1/5 expression in the present study and elevated vmPFC levels of phospho-extracellular regulated kinase (ERK) reported by Koya et al. (2009) for cocaine-withdrawn rats assayed under very similar self-administration and cue-reinforced testing conditions to those employed in the present study. The inverse relation between vmPFC Group1 mGluR and phospho-ERK levels is intriguing in light of recent findings that Group1 mGluR stimulation, particularly mGluR1, augments inhibitory currents to a greater degree than excitatory currents within PFC (Sun and Neugebauer, 2011). Thus, while reduced mGluR1/5 expression may very well reflect a compensatory response to cue-elicited increases in vmPFC neuronal activity, such a reduction is predicted to disinhibit glutamatergic output from the vmPFC – a finding supported by a recent *in vivo* microdialysis study of animals responding for cocaine-paired cues without cocaine reinforcement (Suto et al., 2010). It follows then that the cue-elicited reduction in vmPFC mGluR1/5 expression observed during protracted withdrawal could very well instigate a feed-forward mechanism of neuronal hyper-excitability within this structure that might be responsible for maintaining the saliency of drug-associated cues and the capacity of these cues to elicit drug craving even in protracted abstinence (e.g., Volkow et al., 1992; Childress et al., 1999; Goldstein and Volkow, 2002). Thus, the present data pose a time-dependent reduction in vmPFC mGluR1/5 expression as a molecular cordon to addiction recovery that best be overcome by receptor agonist, rather than antagonist, strategies.

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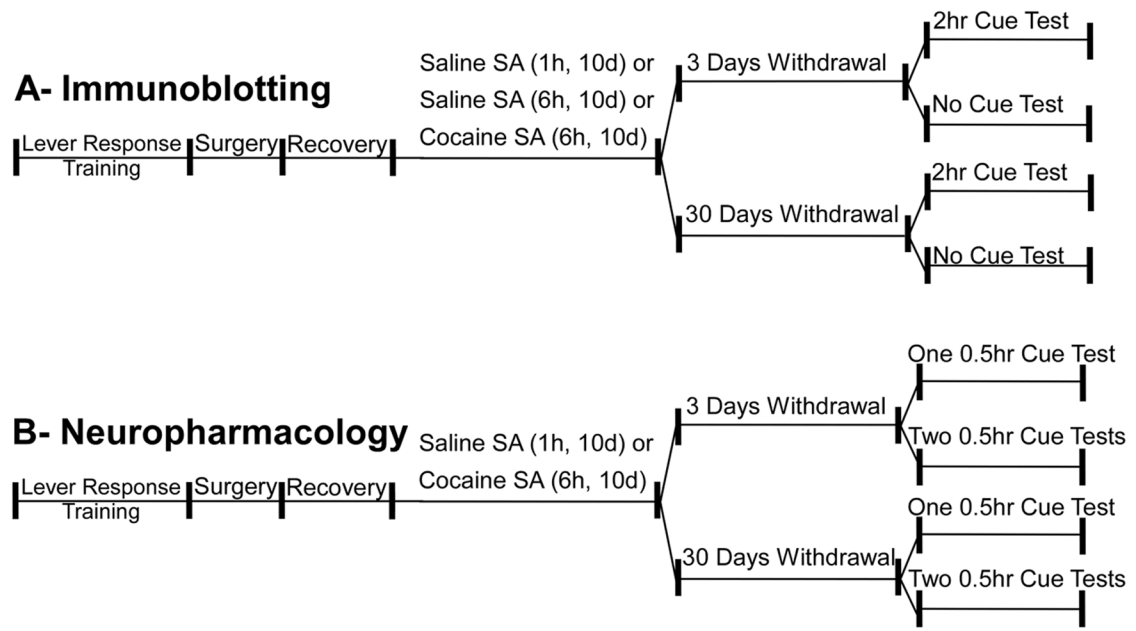
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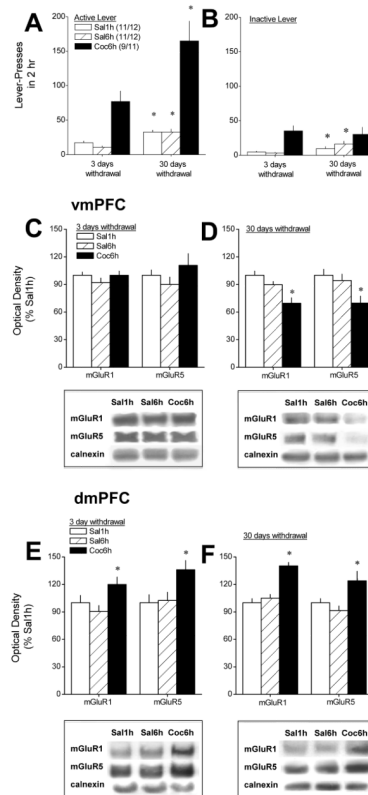
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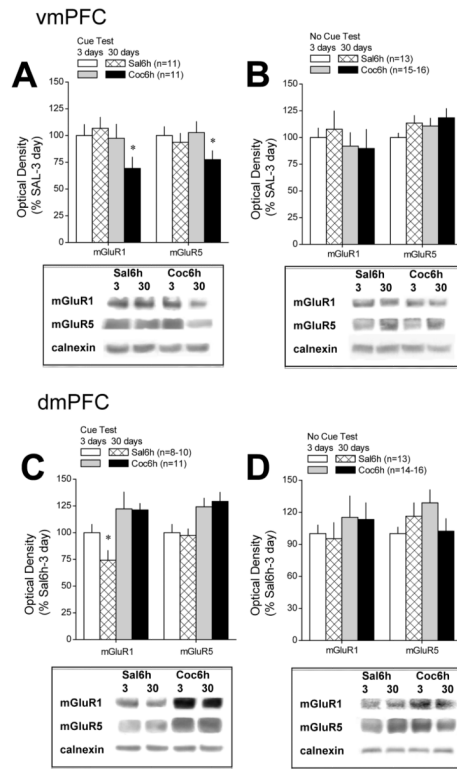


**Figure 1. Schema of general procedures and experimental design**

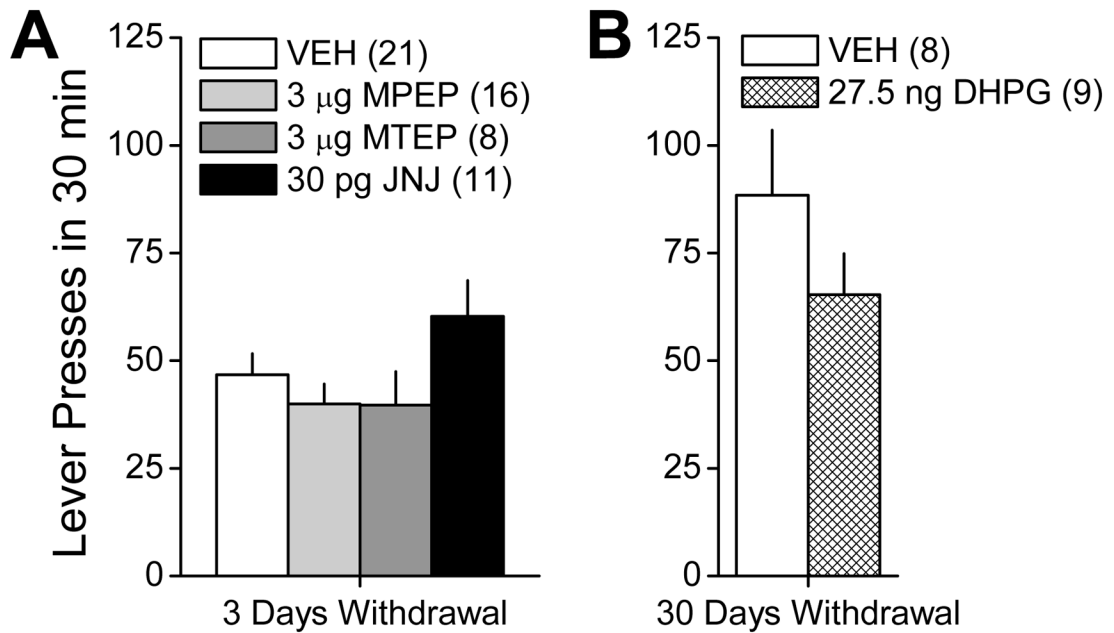
(A) Immunoblotting studies: Following self-administration (10 daily sessions of 1- or 6-hr access to Saline or 6-hr access to Cocaine) animals were divided to two conditions: 3 or 30 days of withdrawal. Following withdrawal, animals were either subjected to a 2-hr cue test, immediately euthanized and the brains harvested for immunoblotting analysis, or were simply euthanized (with no other experimental manipulation) with brains harvested for immunoblotting. (B) Neuropharmacological studies: The procedure was the same as that illustrated in Panel A with one exception: treatment following withdrawal. Here, cohorts of subjects received one or two 30-min cue tests and brains were harvested for histological verification of microinjector placements.



**Figure 2. Protracted withdrawal from self-administered cocaine results in increased reinforced responding concomitant with a reduction in vmPFC Group1 mGluR expression**  
 Summary of the total number of active (A) and inactive (B) lever-presses emitted by animals with a 10-day history of extended access (6h/day) to either cocaine (Coc6h) or saline (Sal6h) and by animals with an equivalent history of limited access (1h/day) to saline (Sal1h) on a 2-h test in which each active lever-press resulted in the presentation of the tone-light stimulus complex previously paired with saline/cocaine infusion, but no saline/cocaine delivery. Tests were conducted in separate groups of animals at either 3 or 30 days withdrawal from saline/cocaine self-administration and sample sizes ranged from 9–12/group/withdrawal time-point. Immediately upon completion of behavioral testing, the dmPFC and vmPFC were dissected out and immunoblotting conducted for mGluR1 and mGluR5. Group differences in vmPFC receptor expression were not observed when assayed at 3 days withdrawal (C), but Coc6h rats exhibited reduced mGluR1/5 when assayed at 30 days withdrawal (D). In contrast, immunoblotting on dmPFC of these same animals revealed elevated receptor levels in Coc6h animals at both 3 and 30 days withdrawal (respectively, E & F). Sample sizes are indicated in Panel A (3 days/30 days withdrawal). For Panel A, \* $p < 0.05$  vs. respective data at 3 days withdrawal. For Panels B–F, \* $p < 0.05$  vs. respective Sal1h group.

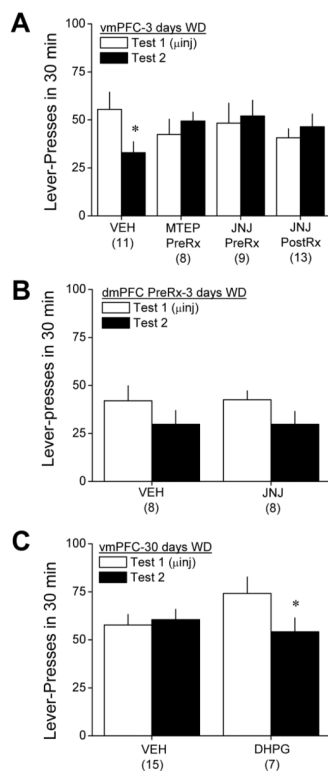


**Figure 3. Differential effects of testing for cue-reinforced behavior upon mGluR1/5 expression within PFC subregions during protracted withdrawal from self-administered cocaine**  
Immunoblotting for the change in total mGluR1 and mGluR5 protein expression within the vmPFC and dmPFC was conducted on tissue derived from saline- and cocaine-experienced animals tested for cue-reinforced behavior over a 2-h period (Cue Test) or left undisturbed in their home cages (No Cue Test) at 3 or 30 days following a 10-day history of extended access (6h/day) to either cocaine (Coc6h) or saline (Sal6h). Sample sizes ranged from 9–12/group/withdrawal time-point. **(A)** A direct comparison of receptor protein expression within the vmPFC of Sal6h and Coc6h animals across the 2 withdrawal time-points confirmed a time-dependent reduction in mGluR1/5 expression in Coc6h animals that were tested for cue-reinforced behavior prior to tissue harvest. **(B)** No statistically reliable change in vmPFC receptor expression was apparent in COC6h rats that were not tested for cue-reinforced behavior prior to tissue collection. **(C)** In contrast to the vmPFC, immunoblotting on dmPFC of these same animals revealed elevated receptor levels in Coc6h animals tested for cue-reinforced behavior at both 3 and 30 days withdrawal. **(D)** However, as also observed for the vmPFC, the changes in dmPFC mGluR1/5 expression was not apparent in animals left undisturbed prior to tissue harvest. Sample sizes are indicated in parentheses. \* $p < 0.05$  vs. respective data at 3 days withdrawal.



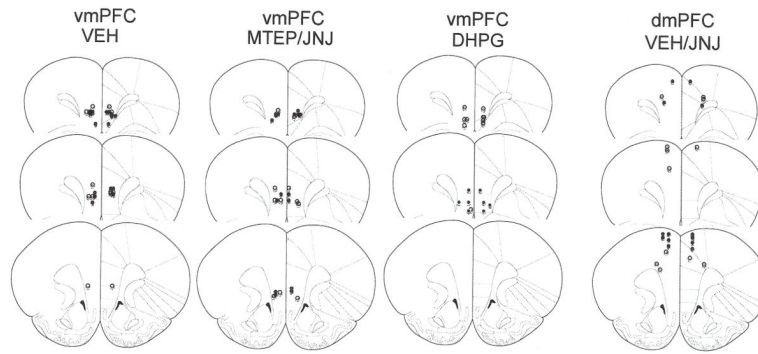
**Figure 4. Intra-vmPFC pretreatment with mGluR-targeting compounds does not acutely influence cue-reinforced responding in cocaine-experienced animals**

(A) When assessed at 3 days withdrawal, an infusion of 3.0  $\mu$ g/side of the mGluR5 antagonists MPEP or MTEP or of 30 pg/side of the mGluR1 antagonist JNJ16259685 (JNJ) failed to significantly influence cue-reinforced lever-pressing behavior exhibited by cocaine-experienced animals during a 30-min test of drug-seeking. (B) Similarly, an infusion of 27.5 ng/side of the mGluR1/5 agonist DHPG also failed to influence lever-pressing behavior when assessed in rats with a history of cocaine self-administration at 30 days withdrawal. Sample sizes are indicated in parentheses.



**Figure 5. Intra-vmPFC pretreatment with mGluR-targeting compounds produces residual effects upon cue-reinforced responding in cocaine-experienced animals**

(A) When tested for cue-reinforced behavior 24 h following an intra-vmPFC infusion of 3.0  $\mu\text{g}/\text{side}$  MTEP prior to initial testing for cue-reinforced behavior (MTEP PreRx) or an intra-vmPFC infusion of 30  $\text{pg}/\text{side}$  of JNJ16259685 either prior to (JNJ PreRx) or immediately following (JNJ Post-Rx) the initial test for cue-reinforced behavior in the absence of any further intracranial manipulation, vehicle (VEH)-pretreated rats exhibited reduced responding (extinction), while no behavioral extinction was apparent in animals pre- or post-treated with mGluR antagonists (B) In contrast to the vmPFC, intra-dmPFC pretreatment with 30  $\text{pg}/\text{side}$  JNJ16259685 failed to influence behavioral extinction at 3 days withdrawal. (C) Interestingly, when assayed at 30 days withdrawal, VEH-pretreated rats exhibited no sign of behavioral extinction when tested for cue-reinforced behavior, while extinction was facilitated in cocaine-experienced animals pretreated with 27  $\text{ng}/\text{side}$  of DHPG. Sample sizes are indicated in parentheses. \* $p < 0.05$  vs. Test 1. +main effect of Test,  $p < 0.05$ .



**Figure 6. Histological verification of microinjector placements within the PFC**

A cartoon depicting the results of our histological examination of microinjector placements conducted upon the completion of testing for the residual effects of an intra-vmPFC infusion of vehicle (VEH), MTEP, JNJ16259685 and DHPG and of an intra-dmPFC infusion of JNJ16259685, as examples of placements for the animals included in the statistical analyses of the data in this study. As exemplified by the saline (open) and cocaine (closed) animals in this figure, only animals exhibiting microinjector placement within the ventral prefrontal cortex, within the infralimbic cortex or at their interface were included in the statistical analyses of the data for vmPFC studies. As exemplified by the VEH (open) and JNJ16259685 (closed) animals in the panel for the dmPFC study, only animals exhibiting microinjector placement within the dorsal prefrontal cortex, the anterior cingulate or their interface were included in the statistical analyses of the data for the dmPFC study.



**Table 1**

Comparison of the number of active and inactive lever-presses, as well as the number of infusions earned, by animals with extended access to cocaine (6h/day; Coc6h) or to saline (Sal6h) and by animals with brief access to saline (1h/day; Sal1h) during the last 3 days of self-administration training prior to being tested for cue-reinforced behavior and tissue harvest for immunoblotting at 3 and 30 days withdrawal. No differences were observed between behavior of animals tested at the 3 versus 30-day time-point within any of the IV treatment groups as revealed by no significant main effect of, or interactions with, the Withdrawal factor as determined by 3-way ANOVA (for active vs. inactive lever-presses) or 2-way ANOVA (for infusions). Numbers in parentheses indicate the final sample sizes employed in the statistical analyses of these data.

IV Treatment	Active Lever-Presses		Inactive Lever-Presses		Infusions	
	3 days	30 days	3 days	30 days	3 days	30 days
Sal1h	5.3 ± 1.0 (11)	16.4 ± 3.4 (12)	1.8 ± 0.7	2.9 ± 0.7	3.2 ± 0.5	8.6 ± 1.4
Sal6h	9.4 ± 2.6 (11)	16.7 ± 2.9 (12)	4.1 ± 1.4	5.2 ± 1.9	7.8 ± 2.2	11.6 ± 2.1
Coc6h	158 ± 12 (7)	139.6 ± 13.4 (10)	3.7 ± 1.4	5.7 ± 2.8	125.3 ± 12.4	121.3 ± 8.9

**Table 2**

Comparison of the average number of infusions earned by saline (1 h/day; Sal1h) and cocaine (6 h/day; Coc6h) self-administering rats during the last 3 days of self-administration prior to studies of the effects of intra-vmPFC manipulations of Group 1 mGluRs at 3 days (antagonist studies) or 30 days (agonist study) withdrawal. No differences in behavior were observed across the different intra-vmPFC pretreatment groups within Coc6h or Sal1h animals prior to testing at either withdrawal time-point. Numbers in parentheses indicate the final sample sizes employed in the statistical analysis of these data, as well as, the cue test data presented in Figures 3 and 4.

	IC Rx	Acute Test		Repeated Testing	
		Sal1h	Coc6h	Sal1h	Coc6h
<i>Withdrawal</i>					
<i>3 days</i>	VEH	9.4 ± 1.7 (18)	191.8 ± 31.2 (21)	5.67 ± 0.73 (13)	127.1 ± 11.3 (13)
	MPEP	11.7 ± 2.6 (5)	160.4 ± 41.7 (16)		
	MTEP			6.25 ± 2.08 (8)	134.79 ± 32.22 (8)
	JNJ16259685 PreRx	5.1 ± 1.5 (10)	246.9 ± 91.9 (12)	7.36 ± 1.57 (9)	103.6 ± 8.09 (9)
	JNJ16259685 PostRx				110.8 ± 6.3 (13)
	dmPFC JNJ16259685			108.1 ± 14.2 (6)	91.7 ± 4.5 (8)
<i>30 days</i>	VEH		112.8 ± 13.2 (8)	6.2 ± 0.9 (14)	150.3 ± 30.6 (11)
	DHPG		109.6 ± 11.4 (9)	4.5 ± 0.6 (12)	115.0 ± 12.8 (9)

**Table 3**

Comparison of the average number of lever-presses emitted by saline-experienced animals during tests for cue-reinforced behavior conducted at 3 days (antagonist studies) or 30 days (agonist study) following the last saline self-administration session. No differences in behavior were observed across the different intra-vmPFC pretreatment groups at either withdrawal time-point. It should be noted that in studies in which saline-experienced animals were tested repeatedly, all groups (incl. vehicle controls; VEH) exhibited lower lever-pressing behavior on the test for the acute effects of vmPFC microinjection (i.e., Test 1), compared to the test for residual drug effects in which the animals were not microinjected prior to behavioral testing (Test 2). Numbers in parentheses indicate the final sample sizes employed in the statistical analysis of these data.

Withdrawal	IC Rx	Acute Test	Repeated Testing	
			Test 1	Test 2
3 days	VEH	6.1 ± 1.3 (18)	2.92 ± 0.9 (13)	8.2 ± 1.6
	MPEP	5.8 ± 1.9 (5)		
	MTEP	1.14 ± 0.7 (7)	2.7 ± 0.7 (7)	7.7 ± 2.1
	JNJ16259685	1.9 ± 0.6 (10)	1.0 ± 0.6 (9)	6.6 ± 2.3
30 days	VEH		9.8 ± 1.9 (13)	12.8 ± 2.0
	DHPG		5.5 ± 1.2 (15)	12.7 ± 2.8