Cocaine and Pavlovian fear conditioning: Dose–effect analysis

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Abstract

Emerging evidence suggests that cocaine and other drugs of abuse can interfere with many aspects of cognitive functioning. The authors examined the effects of 0.1–15 mg/kg of cocaine on Pavlovian contextual and cued fear conditioning in mice. As expected, pre-training cocaine dose-dependently produced hyperactivity and disrupted freezing. Surprisingly, when the mice were tested off-drug later, the group pre-treated with a moderate dose of cocaine (15 mg/kg) displayed significantly less contextual and cued memory, compared to saline control animals. Conversely, mice pre-treated with a very low dose of cocaine (0.1 mg/kg) showed significantly enhanced fear memory for both context and tone, compared to controls. These results were not due to cocaine’s anesthetic effects, as shock reactivity was unaffected by cocaine. The data suggest that despite cocaine’s reputation as a performance-enhancing and anxiogenic drug, this effect is seen only at very low doses, whereas a moderate dose disrupts hippocampus and amygdala-dependent fear conditioning.

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context and tone fear, and perhaps memory of the shock as well [13,19,22,38,40]. Involvement of both the hippocampus [8,36,53,54] and the amygdala [9,16,21,37] has been implicated in cocaine use and addiction. For example, studies by Thompson et al. [44–46] have shown that cocaine exposure modulates long-term potentiation in the CA1 region of the hippocampus. However, the cognitive and behavioral effects of acute cocaine administration on Pavlovian fear conditioning have yet to be examined.

In the current study, mice were injected with cocaine prior to placement in the conditioning chambers. Following training, subjects were tested for both context and tone memories, off-drug. By one view, acute cocaine administration should potentiate fear conditioning, as cocaine is a nootropic psychostimulant [16] and an anxiogenic [4,42]. Studies have shown that cocaine can inhibit the extinction of a fear-potentiated startle response when paired repeatedly with a nonreinforced CS, suggesting that cocaine can intensify the anxiety-producing qualities of a CS [12]. Increased overall arousal or anxiety could enhance the learning of an anxiety-producing stimulus. Alternatively, cocaine could impair fear conditioning, as it disrupts not only the normal activation of the amygdala and hippocampus, but also areas in the prefrontal cortex related to attention [32]. According to this view, subjects on cocaine may not retain memory of context and tone cues because they would be unable to sufficiently attend to them during training. We were interested in characterizing the effects of cocaine on fear conditioning both with doses similar to those taken by human addicts, as well as lower doses, which we believed more likely to enhance learning. A moderate dose of cocaine produced a remarkable impairment in learning and memory, while a very low dose of cocaine enhanced memory. Several interpretations of these findings are presented.

1. Methods

1.1. Subjects

Sixty-four C57B6J (B6; Jackson Laboratory, West Sacramento, CA) inbred mice were used. Mice were weaned at 3 weeks of age and were group housed (two to five mice per cage) with continuous access to food and water. The vivarium was maintained on a 14:10 light/dark schedule, and all testing was performed during the light phase of the cycle. Mice were at least 8 weeks old before testing. Approximately equal numbers of male and female mice were used. All animal care and testing procedures were approved by the UCSD IACUC and were in accordance with the NIH Principles of Laboratory Animal Care.

1.2. Apparatus

1.2.1. Conditioning context

Three to four mice were tested concurrently, in individual conditioning chambers housed in a windowless room. Each chamber (32 cm wide, 25 cm high, 25 cm deep) was equipped with a speaker in the side wall and a stainless steel grid floor (16 rods, each rod 2-mm diameter, 8-mm center to center; the front wall was clear acrylic, the sidewalls were white acrylic; Med-Associates Inc., St. Albans, VT). A stainless steel drop pan, scented with 7% isopropyl alcohol to provide a background odor, was located beneath each chamber. Between tests, the conditioning contexts were cleaned with 7% isopropyl alcohol solution. Background noise (65-dB) was provided by a HEPA air cleaner and white light was provided by two 100 W bulbs. The mice were continually observed by a wall-mounted color video camera which was connected to a computer and video equipment in an adjacent room. Each chamber was connected to a solid-state scrambler, providing AC constant current shock, and an audio stimulus generator located in an adjacent room, controlled via an interface connected to a Windows computer running Med-PC (Med-Associates Inc., St Albans, VT). Automated assessment of freezing and activity was provided by custom designed software adapted from NIH Image running on an Apple Macintosh G4 [automated algorithm validated elsewhere, 2].

1.2.2. Alternate context

The alternate context for testing tone fear was located in a separate room and differed from the training context along several dimensions. Multiple (three to four) mice were tested concurrently, in individual boxes measuring 30 cm wide, 25 cm high, 24 cm deep, and equipped with a speaker in the side walls. The ceiling, floor, and three interior walls of the chamber were white, while a clear Plexiglas front wall allowed the mice to be continually observed. To create a distinct space, a white plastic, triangular tent was placed inside each box, with each side of the triangle measuring 23 cm. Between tests, the chambers were cleaned and scented with a 5% white vinegar solution. The room was lit with dim red light and an infrared video camera, connected to the Macintosh G4 described above, was used to score freezing.

1.3. Behavior measurement

In order to measure the effect of cocaine on exploratory locomotor activity, baseline activity during training was assessed by counting the number of crossovers each subject performed [e.g. 38]. A single cross-over was defined as the movement of a subject’s entire body from one half of the box to the other. Videotapes of the conditioning sessions were observed using a standard VCR and monitor, and the number of cross-overs was counted during the first 2 min prior to the first tone–shock pairing on the training day.

In addition, to ensure that cocaine did not disrupt shock reactivity, mouse activity burst displayed during the 2 s of shock exposure (Unconditioned Response to shock), as well as activity during the 2 s leading up to the shock were measured as velocity (cm/s) [2]. Full-screen video of this time period was digitized at 10 Hz using NIH Image. X-Y coordinates were obtained for each frame for each mouse using the wand auto-measure tool; these coordinates were imported into Microsoft Excel. Distance traveled (measured in pixels) between successive frames was computed using the distance formula \[ \sqrt{(x_n - x_{n+1})^2 + (y_n - y_{n+1})^2}\]; these values were converted into real distance in centimeters using known landmark distances in the video frame. Distance was then converted into velocity (cm/s) by dividing by time.

1.4. Drugs

Drugs were administered intraperitoneally (i.p.) in a volume of 5–10 ml/kg. Cocaine HCl (Sigma Chemical Co., St. Louis, MO) was dissolved in 0.9% physiological saline. Cocaine injections (salt weight: 15, 5, 1, or 0.1 mg/kg, i.p.) were given 15 min before training. These doses were selected to form a comprehensive dose–effect curve; although 15 mg/kg is usually regarded as a moderate dose, we found an unacceptable lethality rate (about one-third) at the next highest appropriate dose (30 mg/kg), in this strain of mice, whereas lower doses did not produce any lethality.

2. Experimental procedures

2.1. Conditioning

Mice were injected with either saline or cocaine 15 min prior to training. Training consisted of a 2 min baseline period, followed by a 30 s tone (28 kHz, 85 dB, A Scale), with a shock (0.75 mA) administered during the last 2 s of the tone. Two more tone–shock pairings followed, separated by 30 s each. Immediate post-shock freezing was measured for another 5 min. Thus, mice were inside the fear conditioning chambers for a total of 10 min, then returned to their home cages.

2.2. Testing

Mice were returned to the conditioning context without drug 24 hr later and freezing was scored for 5 min. Mice were subsequently placed in the tone test context 48 hr later. Tone testing consisted of a 2 min baseline, followed by a
continuous 3 min tone identical to the training tone. Freezing was scored for the entire 5 min period.

3. Results

3.1. Generalized activity

Cocaine produced an increase in cage cross-overs during the 2 min baseline period on the training day (Fig. 1A). A univariate analysis of variance (ANOVA) confirmed group differences \( [F(4, 59) = 7.1, p = 0.0001] \). Compared to saline control animals, all doses of cocaine produced significant hyperactivity [0.1 mg/kg, \( F(1, 24) = 12.2, p < 0.01 \); 1 mg/kg, \( F(1, 24) = 12.5, p < 0.01 \); 5 mg/kg, \( F(1, 24) = 14.0, p < 0.01 \); 15 mg/kg, \( F(1, 26) = 17.0, p < 0.001 \)], suggesting it is a psychomotor stimulant even at the lowest doses.

3.2. Activity burst velocity

Cocaine produced an increase in baseline velocity in only the animals receiving 15 mg/kg in this very brief sample of velocity, compared to control animals [\( F(1, 26) = 6.97, p < 0.05 \); all others, n.s.; Fig. 1B]. Cocaine had no apparent effect on shock reactivity at any dose, with no difference between doses in activity burst velocity during shock \( [F(4, 59) = 0.8, p > 0.5] \). The shock produced a large increase in velocity compared to baseline in all groups \( [F(4, 59) = 428.0, p < 0.0001] \).

3.3. Training and post-shock freezing

Fig. 2A depicts the acquisition of freezing during training. The first 2 min period is a baseline measurement, and is followed by three tone–shock pairings spread over 3 min. Saline control subjects exhibited a higher level of freezing averaged over the 5 min training session compared with those receiving 1, 5, or 15 mg/kg, but did not differ from those receiving 0.1 mg/kg [0.1 mg/kg, \( F(1, 24) = 2.92, p = 0.10 \); 1 mg/kg, \( F(1, 24) = 5.57, p < 0.05 \); 5 mg/kg, \( F(1, 24) = 25.0, p < 0.0001 \); 15 mg/kg, \( F(1, 24) = 25.0, p < 0.0001 \)]. Subjects receiving 5 or 15 mg/kg cocaine displayed much lower levels of freezing during the three tone–shock pairings. (B) Immediate shock memory: freezing was measured for 5 min immediately following training, on drug. Dose-dependent effects are evident, with animals on 0.1 mg/kg cocaine displaying more freezing than saline control animals. Animals injected with 5 or 15 mg/kg cocaine froze significantly less than controls.

Fig. 1. (A) Average activity: cage cross-overs were counted during the 2 min baseline period before the onset of the tone and shock on training day. Subjects receiving 0.1, 1, 5, or 15 mg/kg cocaine, i.p., 15 min prior to training showed hyperactivity compared to saline control subjects. (B) Shock reactivity: velocity was measured during the 2 s prior to the first shock (baseline) and during the 2 s of the first shock (shock). Only subjects receiving 15 mg/kg showed an increase in velocity during baseline, compared to controls. All groups showed a marked increase in velocity from baseline to shock. The velocity during shock did not differ between groups, showing no dose-dependence in Unconditioned Response.

Fig. 2. (A) Training: freezing was measured continuously throughout the initial, 2 min baseline period, as well as the 3 min period encompassing three tone–shock pairings. Saline control subjects displayed more freezing than most other groups (1.5, and 15 mg/kg cocaine). Subjects receiving 5 or 15 mg/kg cocaine displayed much lower levels of freezing during the three tone–shock pairings. (B) Immediate shock memory: freezing was measured for 5 min immediately following training, on drug. Dose-dependent effects are evident, with animals on 0.1 mg/kg cocaine displaying more freezing than saline control animals. Animals injected with 5 or 15 mg/kg cocaine froze significantly less than controls.
Subjects receiving 5 or 15 mg/kg of cocaine displayed much less freezing than all other groups during the tone–shock pairings ($F$ values $> 25$, $p$ values $< 0.0001$). Fig. 2B depicts the average level of freezing during an immediate memory test that consisted of leaving the mice in the chambers for an additional 5 min, on-drug. Dose-dependent effects on post-shock freezing are evident [ANOVA, $F(4, 59) = 22.1$, $p < 0.0001$], with 0.1 mg/kg producing a nearly significant elevation in freezing, despite producing hyperactivity, compared with controls [$F(1, 24) = 4.1$, $p = 0.05$]. There was no difference between saline and 1.0 mg/kg cocaine groups, while there was a large deficit in freezing in both the 5 mg/kg as well as 15 mg/kg groups, compared with saline controls ($F$ values $> 5$, $p$ values $< 0.05$). Fifteen milligrams per kilogram produced a larger deficit than 5 mg/kg [$F(1, 24) = 8.0$, $p < 0.01$].

This post-shock freezing measurement also, in part, addresses the issue of state-dependency in learning [e.g. 39]. The state-dependency view would predict that the decrement in freezing while on-drug, during training, would be less than that seen off-drug, during testing. However, the freezing decrement seen on-drug is greater than that seen off-drug.

### 3.4. Context fear

One day after training, mice were returned to the context in which they had been trained, off-drug. A dose-dependent effect of cocaine can be seen in the average rate of freezing over the 5 min context test (Fig. 3A) [ANOVA, $F(4, 59) = 7.8$, $p < 0.0001$]. Subjects pre-treated with 0.1 mg/kg during training displayed enhanced freezing, compared to saline controls, during the context test, off-drug [$F(1, 24) = 4.71$, $p < 0.05$]. Mice pre-treated with 15 mg/kg showed a significant decrement in freezing [$F(1, 26) = 8.88$, $p < 0.01$], while those pre-treated with 1 or 5 mg/kg were not significantly different than saline controls ($F$ values $< 1$). In contrast to the immediate memory test, it is unlikely that response competition was a factor, since mice were tested off-drug, suggesting that cocaine given during training somehow interfered with the memory for fear conditioning. Another, although unlikely (see below) explanation is that conditioned hyperactivity, by virtue of drug-environment conditioning, produced an increase in activity in cocaine pre-treated mice that disrupted freezing.

### 3.5. Tone fear

Two days after training, mice were administered the tone test in a novel context, off-drug. A dose-dependent response to the 3 min presentation of the tone is shown in Fig. 3B; there were significant differences [$F(4, 59) = 3.8$, $p < 0.01$]. Mice administered 0.1 mg/kg cocaine, pre-training, displayed an increase in freezing during the tone test compared with saline controls [$F(1, 24) = 4.45$, $p < 0.05$]. Mice pre-treated with either 1 or 5 mg/kg displayed no difference in performance compared with controls ($F$ values $< 0.5$), while mice administered 15 mg/kg showed a decrement in performance [$F(1, 26) = 4.50$, $p < 0.05$]. These findings do not support the view that the cocaine pre-treated mice exhibited deficits in freezing due to conditioned hyperactivity, since cocaine was never paired with this context. In addition, there was no difference in freezing between groups during the 2 min baseline period [$F(4, 59) = 0.9$, $p > 0.45$; data not shown].

Overall, mice exhibited an enhancement of context and tone memory tested off-drug when given 0.1 mg/kg, and a deficit in memory when given 15 mg/kg of cocaine. Five milligrams per kilogram was also able to disrupt immediate memory, while the mice were still on the drug.

Finally, doses of 0.1–15 mg/kg all produced hyperactivity, suggesting these are all still doses at which cocaine is acting as a psychostimulant.

### 4. Discussion

Administration of a very low dose of cocaine before fear conditioning enhanced freezing during both context and tone testing when subjects were off-drug, while a moderate dose disrupted
memory during both tests. These results were surprising, as there are many reasons to expect a moderate dose of cocaine should potentiate fear conditioning. For example, cocaine is thought of as an anxiogenic, amplifying many anxiety and defensive behaviors in rodents [e.g. 4,42,55], and thereby would be predicted to increase the response to fear conditioning. Cocaine has also been shown to increase the fear-potentiated startle response in rats, as well as inhibit extinction to a CS [10,12,52]. In addition, psychostimulants, such as caffeine, are usually thought of as nootropics [e.g. 15,26,49]. This folklore is exemplified by the advertisement campaign of the original Coca-Cola that contained cocaine, declaring it “the brain tonic and intellectual soda-fountain beverage” [51]. However, despite its anxiogenic and nootropic effects, only a very low dose of cocaine increased performance on fear conditioning, while a moderate dose disrupted it.

As fear conditioning is known to require activation of both the hippocampus and amygdala [1,3,19,20,38], one explanation for the results is that cocaine altered the regular functioning of the amygdala, the hippocampus, or both. In another study on fear conditioning, Corodimas et al. [17] discovered the acute administration of caffeine, pre-training, produced a selective deficit in context conditioning only, suggesting a disruption of hippocampal function. Repeated exposure to cocaine can enhance long-term potentiation (LTP) in the CA1 region of the hippocampus [44]. However, this effect may be reversed at high concentrations of cocaine, or after long periods of withdrawal [45,46], and has not been shown with an acute, in vivo exposure to cocaine. It is possible, though, that this enhanced LTP could account for the better performance during testing of the subjects which had received 0.1 mg/kg cocaine before training. Other studies on both acute and frequent cocaine intake have revealed the disruption of amygdala-dependent learning, as well [29,48]. The present effects on both context and tone conditioning suggest that both hippocampus and amygdala-dependent learning can be disrupted even by acute, moderate cocaine administration, while a very low dose can enhance this learning.

One possibility is that cocaine interfered with motivational components of fear conditioning. Cocaine is a powerful local anesthetic, and it is possible that a 15 mg/kg, i.p. injection could have extended some anesthesia to the paws, reducing the painfullness of the footshock. However, measurement of the mouse activity burst UR suggests that footshock reactivity was equivalent in cocaine and saline-treated mice. Alternatively, the cocaine could have induced a pleasurable hedonic state, in which the animals on cocaine may have perceived the shock or fear conditioning experience as less aversive. This possibility would be difficult to explore within the current experiment, but could suggest that all drugs which induce a positive hedonic state may interfere with aversive conditioning. This seems unlikely, as there is evidence that cocaine, despite being thought of as producing a positive hedonic state, amplifies other anxiety and defensive behaviors [4]. Finally, there is also evidence from studies in monkeys that cocaine can disrupt an appetitively motivated working memory task as well [29].

Another possibility is that cocaine directly disrupted conditioning by interfering with learning and memory circuits. Human chronic cocaine users have been shown to have deficits in measures of attention [see 34 for review], while dysfunction in prefrontal cortex activity has been implicated for attentional deficits in several studies of drug abusers [6,7,24,25,30,32,47]. Thus, it is possible that cocaine directly disrupted the circuits required for learning and memory in fear conditioning.

A possible confound in the present study is that we were unable to directly control for state-dependent learning. By this argument, on-drug testing may allow better recall of the memory of the context, tone and shock pairings, after on-drug training. Because the subjects were tested off-drug, one can argue that they were at a disadvantage, compared to controls, because of their different drug state during training and testing sessions. However, subjects which had received 0.1 mg/kg cocaine before training continued to show higher levels of freezing than controls when tested off-drug. State-dependent learning would predict quite the opposite result. Moreover, when the immediate memory test was used to control for state-dependency, mice were actually more sensitive to the disruptive effects of cocaine than when tested off-drug. These findings agree with previous research addressing similar issues. Studies examining the effects of repeated cocaine administration on fear conditioning, using the fear-potentiated startle paradigm, found that state-dependent learning did not satisfactorily account for increased startle amplitudes, after extinction [52]. Overall, there is little evidence that cocaine or other stimulants produce heavily state-dependent learning in humans or in other animals [39].

Although it is likely that hyperactivity directly attributable to the drug is responsible for freezing deficits seen during training, hyperactivity seems an unlikely explanation for deficits in context and tone memory when mice were tested off-drug. Conditioned hyperactivity via drug-environment conditioning could theoretically play a role in the apparent context and tone memory deficit seen in animals given 15 mg/kg cocaine. However, it is unlikely that much conditioned hyperactivity would be produced by a single drug exposure. Moreover, mice also exhibited deficits in tone testing, which was tested in a context never paired with cocaine, suggesting that conditioned hyperactivity played little role in the freezing deficits seen on testing. Finally, subjects which had been administered 0.1 mg/kg cocaine before training exhibited an increased level of activity during the 2 min baseline period, but exhibited increased freezing during both the context and tone tests, off-drug. Taken together this suggests that conditioned hyperactivity is not directly responsible for the memory deficits seen in context and tone conditioning.

An emerging view of addiction states that drugs of abuse hijack existing learning and memory systems normally involved in conditioned reinforcement [28]. If this is true, these neural systems may be unable to properly participate in the forms of memory they normally subserve. Given that cocaine alters the normal functioning of ventral tegmental area (VTA)–nucleus accumbens (NAcc) connectivity, it may disrupt the functional reward system of the brain. Human studies have implicated areas of the amygdala in this disruption, as well, correlating amygdala activation with cocaine craving [e.g. 13,14]. If this is the case, then areas of the brain normally used for encoding the context
and tone information during fear conditioning would be disrupted and unable to encode new information.

In addition, VTA dopamine neurons have been implicated in both the behavioral expression of a fear response as well as memory. For example, lesions to the VTA were shown to inhibit fear-potentiated startle in rats [11]. Also, cocaine administered repeatedly over 5–7 days, but not acutely on a single day, facilitated LTP induction in VTA dopamine neurons [33]. Together, these results suggest that repeated cocaine administration could enhance memory of a fear response via VTA dopamine neurons. These results also predict that an acute, moderate dose of cocaine, as used in the present study, would not be expected to enhance memory encoded via these neurons.

In the present study we chose to focus on a range of cocaine doses, from a low dose which is unlikely self-administered by drug users, to a moderate dose thought to be typical of drug users. Within that range, we found differing results. Future studies will use other psychostimulants [in preliminary studies with modafinil (Provigil®) we have found equivalent results].

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References


