Acute Cocaine Differentially Induces PKA Phosphorylation Substrates in Male and Female Rats

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Abstract

Background: Sex differences in intracellular dopamine pathways may contribute to known sex differences in psychomotor responses to cocaine and differential development of dependence. This study aimed to determine whether there are sex differences in the activation of the extracellular signal-regulated kinases (ERK1/2, or p44/p42 MAPK) and PKA phosphorylation-dependent substrates in the nucleus accumbens (NAcc) of male and female rats at baseline or after acute cocaine administration.

Methods: 60-day-old male and female Fischer rats were injected with saline or cocaine (30 mg/kg) and sacrificed 5, 15, 30, 45 or 90 minutes later. Total locomotor activity, stereotypic, rearing, and ambulatory behaviors was measured for 90 minutes using a two-frame automated Photobeam Activity

Results: Similar to our previous findings, total locomotor activities were higher in female rats after this single cocaine administration. Females had higher levels of phosphorylated PKA substrates after cocaine administration, and this change lasted longer and had a greater magnitude than in cocaine treated male rats.

Conclusion: Taken together, these findings suggest that sex differences in basal and cocaine-induced alterations in PKA signaling activity in the NAcc may contribute to sex differences in psychomotor responses to cocaine.

Keywords: Cocaine; Sex differences; ERK; PKA; Nucleus accumbens; Females

Introduction

Cocaine use among women has steadily increased, rising to approximately 30% of users in the United States. As researchers pay more attention to hormonal effects on drug abuse, it is becoming apparent that men and women react differently to cocaine. Overall, women are more vulnerable to some aspects of cocaine abuse, such as being more sensitive to the addictive properties of cocaine, experiencing more nervousness after intermittent administration of cocaine, taking longer to feel its subjective effects, reporting less euphoria, and having more severe cravings in response to cocaine-associated cues [1-9]. Women also increase their rate of cocaine consumption more rapidly than do men [11,12].

Similar to humans, female rodents also show exaggerated and more robust psychomotor responses to cocaine than do males [13-16]. Females also more quickly develop cocaine-induced conditioned place preference (CPP) with lower doses and more readily acquire cocaine self-administration [17-20]. Taken together, human and animal studies suggest that sex-specific differences exist at all stages of cocaine abuse including induction, maintenance, and relapse.

Sex differences in the mesocorticolimbic dopamine (DA) system-a regulator of cocaine's psychomotor and rewarding effects have been demonstrated [21-24]. As recently reviewed by Becker and Hu [9], there are sex differences in the levels of DA receptors in the striatum, in the efficacy of DA antagonists and agonists to block DA receptors, and in cocaine-induced accumbal DA release/reuptake. The sexually dimorphic pattern in DA system activation after cocaine treatment is postulated to be correlated with sex differences in cocaine-induced DA-mediated intracellular responses.

Behavioral studies have shown a positive association between protein kinase A (PKA) signaling changes and behavioral responses after cocaine administration [25,26]. Specifically, administration of a PKA activator or a PKA inhibitor enhances or dampens, respectively, acute cocaine-mediated locomotor behavior and modulates the cocaine-induced CPP [27,28]. PKA activation also potentiates behavioral sensitization after chronic cocaine administration [28,29]. Moreover, intra-accumbens infusion of PKA inhibitor decreases cocaine self-administration [30]. It is therefore feasible that sex differences in cocaine's modulation of the DA-PKA signaling pathway in the NAcc may contribute to sex differences in the initiation and development of the rewarding properties of cocaine. Indeed, three recent studies have found sexually dimorphic responses to cocaine in the DA-PKA signaling pathway.

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Nazarian et al. [22] demonstrated that PKA protein levels in the NAcc are higher in females both before and after cocaine administration; Lynch et al. [31] showed that protein levels of DA- and CAMP-regulated phosphoprotein of Mr 32 kDa (DARPP-32) are also higher in females after cocaine administration and Zhou et al. [32] found basal and cocaine-induced sex differences in the PKA-DARPP-32 cascade. Together, these studies suggest that sex differences both at baseline and after cocaine-induced alterations in the DA-PKA signaling pathways in the NAcc may contribute to patterns of sexual dimorphism in the initiation and development of addictive responses to cocaine.

The intracellular cascade of extracellular signal-regulated kinases (ERK1/2, or p44/p42 MAPK) -- one of several mitogen-activated protein kinases (MAPK) -- has been postulated to be involved in psychostimulant addiction. In mice, acute cocaine treatment induces a rapid and transient increase in the ERK protein phosphorylation (p-ERK) in the NAcc, which can be abolished by pretreatment with a DA-D1 receptor antagonist [33-35]. Behaviorally, the cocaine-induced hyper-locomotor activity and the development/expression of sensitization and CPP are attenuated by treatment with ERK inhibitors [36-39]. Recent studies on ERK mutant mice indicated that, owing to the hyper-phosphorylation of ERK protein, mutated mice show stronger psychostimulant (e.g., cocaine) induced CPP and behavioral sensitization than do the wild-type controls [40,41].

However, it is not known if the stronger and longer-lasting cocaine responses in females (including CPP and psychomotor activation) are related to a deregulation of the ERK-mediated intracellular responses. Cyclic AMP response element binding protein (CREB) is a transcription factor that is involved in modulating the psychomotor and addictive properties of cocaine [42]. According to one report, enhanced activation of ERK enables and mediates cocaine-induced CREB phosphorylation (pCREB) in the NAcc [36]. However, others have found that cocaine-induced ERK and CREB phosphorylation are dissociated in many brain regions [43,44]. After cocaine administration, pCREB levels in the NAcc are induced for a longer time in male rats, but the magnitude of change in pCREB levels is higher in female rats [22]. Because pCREB is a substrate of the ERK pathway, we hypothesized that p-ERK activation closely follows the previously reported observation for pCREB and that differences in basal or cocaine-induced alterations in ERK levels have a sexually dimorphic pattern. The aim of this study was to test this hypothesis by measuring basal and cocaine-induced ERK proteins and p-(Ser/Thr)-PKA substrates in male and female rats.

Methods

Animals

60-day-old male and female Fischer rats (Charles River, Raleigh, NC) were individually housed in Plexiglas chambers (20 × 20 × 41 cm) layered with beta chips. Rats were given free access to food and water and maintained on a 12 hour light/dark cycle (lights on at 9:00 AM). All animals were housed in controlled temperature chambers (20 ± 1°C) and maintained on a 12 hour light/dark cycle (lights on at 9:00 AM). All rats were weighed, handled, and intraperitoneally injected daily with saline for 5 consecutive days prior to testing. For behavioral testing, 10 rats per group were used. For protein analysis, 4 rats per group were used.

Repeated vaginal lavage attenuates cocaine-induced activity, abolishes estrus cycle effects, and establishes CPP in female rats, thus possibly increasing DA-mediated responses [45]. Therefore, as noted by Walker et al. [45], the use of lavaged female rats could skew female behavioral responses when making side-by-side comparisons with male rats. For this reason, females were randomly assigned to experimental groups without regard to their estrous cycle. Animal care and use was in accordance with the Guide for the Care and Use of Laboratory Animals (NIH publication 85-23, Bethesda, MD) and approved by the Hunter Institution Animal Care Use Committee.

Materials

Cocaine hydrochloride and p-ERK primary antibody were purchased from Sigma Chemical Co. (St. Louis, MO). Phospho-(Ser/Thr) PKA substrate-specific antibody measures the levels of substrate phosphorylation by PKA, a class of kinases referred to as Arg-directed kinases or AGC-family kinases that share a substrate specificity characterized by Arg at position -3 relative to the phosphorylated Ser or Thr. This antibody detects phosphorylation by PKA to substrates that contain a phospho-serine/threonine residue with arginine at the 3rd position (Cell Signaling Technologies; Beverly, MA). Phospho-(Ser/Thr) PKA antibody substrates include Apaf-1, CAMKKII, Caspase-10, Caspase-2, CREB, and NMDAR1, among others (see Cell Signaling #9621). Alpha--tubulin antibody was purchased from Santa Cruz Technologies (Santa Cruz, CA). All appropriateSecondary antibodies were purchased from Amersham Pharmacia (Piscataway, NJ).

Cocaine administration

Cocaine solutions were prepared on the day of testing by dissolving the drug in physiological saline (0.9%). On the day of testing, rats were injected (i.p.) with saline or cocaine (30 mg/kg) and sacrificed 5, 15, 30, 45 or 90 minutes later. Rats were returned to their home cages after drug treatment.

Behavioral measurements

All behavioral measurements were conducted in the rat's home cage. Behavioral activity was measured only for the male and female groups sacrificed at 90 minutes after drug treatment. Total locomotor activity was measured for 90 minutes using a two-frame automated Photobeam Activity System (San Diego Instruments; San Diego, CA). Total activity was determined by total counts of any photobeam interruptions in the lower or upper frame. Stereotypic, rearing, and ambulatory behaviors for these rats were previously reported [22].

Tissue collection

After decapitation (following a brief 20 second exposure to CO2), their brains were removed, flash frozen in 2-methylbutane (-40°C), and stored at -80°C until used. The NAcc was dissected from coronal sections (1 mm thick) using a matrix (ASI Instruments; Warren, MI). The NAcc (including both shell and core) was dissected from coronal sections ranging from 1.70 to 1 mm anterior to the bregma (1 mm thick) by using a brain matrix. Tissue was homogenized with the use of a Polytron handheld homogenizer (Kinematica; Luzern, Switzerland) in homogenizing buffer [HEPES 7.9 (20 mM), KCl (10 mM), EDTA (1 mM), NP40 (0.2%), glycercol (10%), NaAcCl (200 mM), pepstatin, leupeptin, DTT (1 M), aprotinin, PMSF (100 mM), NaF (50 mM), and Na3VO4 (1 mM)]. Total protein content was determined with use of a Bradford assay. Protein samples (40 µg) were boiled in Lammeli buffer containing 5% β-mercaptoethanol and loaded onto 10% SDS-PAGE. Gels were electrophoresed, transferred to nitrocellulose membranes, and blocked for 60 minutes with 5% nonfat dry milk in tris-bufffer-saline-tween (TBST, pH = 7.4) at room temperature. Membranes were probed overnight at 4°C with phospho- (Ser/Thr) PKA substrate antibody.
(1:500) and p-ERK antibody (1:1000). After three washes with TBST, membranes were then incubated with their appropriate secondary antibody (1:1000) for 60 minutes at room temperature, followed by three more washes with TBST. Antibody binding was detected using an enhanced chemiluminescence kit (Amersham Pharmacia; Piscataway, NJ). Resulting films were scanned and quantified with a computer densitometer and ImageQuant software (Molecular Dynamics; Sunnyvale, CA).

To compare sex differences in the protein levels, samples from saline-treated male and female rats were loaded onto the same gel. Within each sex, to determine the time course of changes in protein levels after treatment, all saline- treated samples (5 min - 90 min), or cocaine-treated (5 min - 90 min) samples with saline control (5 min), were run on the same gel. Four sets of gels were run for each determinant. To normalize band intensity to protein levels, membranes were re-probed with α-tubulin antibody (1:1000).

Statistical analysis

The time-course of locomotor counts was analyzed using three-way repeated measures analysis of variance (ANOVAs) for the variables sex (male vs. female), drug (saline or cocaine), and time (18 five minute time blocks for acute cocaine/saline). In addition, two-way ANOVAs (sex × drug) were used to determine differences of accumulative locomotor counts. For post hoc analysis, LSD tests were conducted when appropriate.

All protein levels are expressed as a ratio to α-tubulin levels. Data are presented as mean ± SEM. Within sex, one-way ANOVAs followed by post-hoc LSD analysis were used to determine differences during the time course. For comparison between sexes in cocaine-treated rats, percentage changes of protein levels between the cocaine-treated group and the average of protein levels in saline controls of the same sex were used. This approach was necessary because male and female samples were run in different gels. Independent t-test was used to evaluate any sex difference at each specific time point after cocaine administration. Statistical significance was considered to be p < 0.05 for all analyses.

Results

Overall, cocaine increased locomotor activity in both male and female rats (Drug main effect: F (1, 26) = 73.04, p < 0.001; Figure 1A). Female rats had higher cocaine-induced total locomotor activity [Sex main effect: F (1, 26) = 7.19, p < 0.02; Figure 1A] and longer-lasting total locomotor activity than did male rats [Drug × Sex × Time interaction: F (17, 468) = 2.32, p < 0.01; Figure 1B]. This behavioral activity was significantly higher during the final 40 minutes (time intervals 11-18) of the testing session in female rats (p < 0.05 for all comparisons; Figure 1B).

Although no sex differences in basal p-(Ser/Thr)-PKA substrate protein levels were found (Figure 2A), a sexually dimorphic temporal pattern in p-(Ser/Thr)-PKA substrate protein levels was observed after cocaine treatment. In male rats (Figure 2B), although acute cocaine induced p-(Ser/Thr)-PKA substrate protein levels in the NAcc, one-way ANOVA failed to reach statistical significance (F (5, 23) = 2.01, p = 0.13), as compared with the saline control (p < 0.05). In contrast, in female rats (Figure 2C), a significant time effect was seen after cocaine administration (F (5, 23) = 2.83, p < 0.05); p-(Ser/Thr)-PKA substrate protein levels were increased from 5 to 30 minutes after cocaine treatment as compared with saline controls (p < 0.05 for all comparisons). Sex differences were observed in overall levels of p-(Ser/Thr)-PKA substrate protein (Table 1); female rats had higher p-(Ser/Thr)-PKA substrate protein levels than males regardless of the time after cocaine injection (t (1, 38) = 2.98, p < 0.01). In addition, 5 and 30 minutes after cocaine administration, p-(Ser/Thr)-PKA substrate protein levels in females were significantly higher than in males (t (1, 6) = 2.77, p < 0.05; and t (1, 6) = 2.65, p < 0.05, respectively; Table 1).

Although no sex differences in basal or cocaine-induced protein levels of p-ERK were seen (data not shown), p-ERK protein levels in the NAcc of both sexes were increased after acute cocaine administration (F (5, 23) = 3.02, p < 0.05, and F (5, 23) = 4.25, p < 0.01, for males and females, respectively; Figure 3). Furthermore, the temporal pattern of p-ERK induction after cocaine administration was similar for males and females, respectively; Figure 3).
females. In both sexes, p-ERK protein levels were significantly increased 5 and 30 minutes after cocaine administration as compared with their respective saline controls (p < 0.05 for all comparisons; Figure 3). No sex differences in the magnitude of change of p-ERK protein levels after cocaine administration were observed (Table 1).

**Discussion**

This study further extends observations by Nazarian et al., Zhou et al., and Lynch et al. [22,31,32] by demonstrating that female rats have a functionally augmented PKA signaling transduction pathway. We report here that the phosphorylation of PKA substrates was induced differentially after cocaine administration. Cocaine may modulate ERK pathway signaling through the activation of DA-D1 as well as NMDA receptors [46]; e.g., inhibition of NMDA reduces cocaine-induced ERK phosphorylation [47]. The fact that neither the ERK pattern of phosphorylation nor the levels of activation differed between the sexes suggests that (1) cocaine's effects on calcium-NMDA dependent mechanisms are not sexually dimorphic and/or (2) not all components of the females' DA-intracellular signaling pathway are heightened.

No sex differences in basal phosphorylation patterns of substrates for PKA were observed. However, after cocaine administration, female rats had overall faster (activation after 5 minutes of cocaine administration), longer (males have activation lasting only up to 15 minutes, whereas females have activation lasting up to 30 minutes), and higher levels of PKA phosphorylation of its substrates (females have a higher percentage change in p-(Ser/Thr)-PKA substrate protein levels than do male rats). Nazarian et al. [22] showed that female rats have higher basal protein levels of PKA in the NAcc. Since the phosphorylation pattern of substrates for PKA before cocaine administration did not differ between the sexes, it suggests that the increases of basal PKA protein levels seen in the NAcc of females may lead to functional effects of the PKA pathway.

After acute cocaine administration, female rats have been shown consistently to have greater behavioral responses to cocaine [15,48]. Indeed, we found in this study that female rats exhibited longer-lasting and more robust psychomotor responses to cocaine. It is feasible that the augmented phosphorylation of PKA substrates in the NAcc of females may contribute to the observed increase in behavioral responses after cocaine administration. In fact, because many PKA substrates are also transcriptional factors known to be involved in modulating the psychostimulating and addictive properties of cocaine [22], a prolonged and more robust phosphorylation of PKA substrates may indeed provide more changes in protein activation and transcription in female rats. These changes, in turn, may further impel the differences in psychomotor responses between males and females.

Consistent with previous reports [34,39,43,46,49-51] after a single cocaine injection in male rats, a rapid and transient increase of p-ERK protein levels was observed. To our knowledge this is the first report that, as with males, p-ERK protein levels were also induced in the NAcc of female rats. Nazarian et al. [22] demonstrated that in female rats accumbal pCREB activation was more robust, but of shorter duration, than in male rats (in male rats pCREB protein levels increased 5 to 30 minutes after cocaine administration, whereas in female rats they increased only at 5 minutes after cocaine treatment). It has been postulated that enhanced activation of ERK enables and mediates cocaine-induced...
CREB phosphorylation in the NAcc (see review in [52]). For example, acute cocaine administration augments CREB phosphorylation in an ERK-dependent manner [53-56]. However, others have found that cocaine-induced ERK and CREB phosphorylation were dissociated in the NACC [44]. Independent of ERK, CREB phosphorylation at the S133 residue is regulated by PKA in addition to several other kinases [57,44]. Thus, as postulated by Edwards et al. [44], in some brain areas PKA may be the primary kinase for CREB. Unlike p-CREB [22], in this study p-ERK demonstrated neither a sexually dimorphic temporal pattern of activation nor sex differences in the magnitude of cocaine-induced changes. On the other hand, the greater basal and cocaine-induced PKA levels in females may contribute to sex differences in quantified pCREB phosphorylation intensity, and PKA protein levels between sexes may also underlie the increases of phosphorylation of CREB after cocaine administration. In addition to pCREB, the phosphorylation of S845 GluA1 subunit (pGluA1), one of PKA-mediated phosphorylation substrates, is important for the cocaine-mediated behavioral response [58]. The enhanced pGluA1 has been demonstrated to increase AMPA receptor-mediated excitatory current through the augmentation of GluA1 membrane insertion [59-61]. It is plausible that the differential pGluA1 induction in the NAcc is responsible for the behavioral sex difference after acute cocaine administration. Further, besides PKA, pGluA1 is highly regulated by other phosphatases including calcineurin. Previously, we have shown basal and cocaine-induced sex differences on calcineurin protein expression in the NACC [32]. Taken together, these results indicate that pGluA1 is the potential read-out for the cellular mechanism underlying cocaine-induced behavioral difference in male and female rats. Nevertheless, the pGluA1 is our ongoing research to further decipher the molecular pathway contributing the sex dimorphism of cocaine addiction.

In summary, our results suggest that females have an elevated D1-PKA-mediated intracellular second messenger transduction as compared with male rats. Thus, data presented here further elucidate the neurobiological basis for sex differences in cocaine's rewarding effects. Our results reinforce the recurrent postulate that many aspects of cocaine addiction and responses are gender specific. However, the extent to which sex differences in the activation of D1-PKA-mediated intracellular second messenger transduction contribute to females' greater vulnerability to some aspects of cocaine remains to be determined. Answers to these questions are needed to further understand drug abuse in both males and females and to improve treatment in a manner that will meet the needs of both sexes.

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References


