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Methamphetamine preconditioning: Differential protective effects on monoaminergic systems in the rat brain

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Abstract

Pretreatment with methamphetamine (METH) can attenuate toxicity due to acute METH challenges. The majority of previous reports have focused mainly on the effects of the drug on the striatal dopaminergic system. In the present study, we used a regimen that involves gradual increases in METH administration to rats in order to mimic progressively larger doses of the drug used by some human METH addicts. We found that this METH preconditioning was associated with complete protection against dopamine depletion caused by a METH challenge (5 mg/kg x 6 injections given 1 hour apart) in the striatum and cortex. In contrast, there was no preconditioning-mediated protection against METH-induced serotonin depletion in the striatum and hippocampus, with some protection being observed in the cortex. There was also no protection against METH-induced norepinephrine depletion in the hippocampus. These results indicate that, in contrast to the present dogmas, there might be differences in the mechanisms involved in METH toxicity on monoaminergic systems in the rodent brain. Thus, chronic injections of METH might activate programs that protect against dopamine toxicity without influencing drug-induced pathological changes in serotonergic systems. Further studies will need to evaluate the cellular and molecular bases for these differential responses.

Keywords

Methamphetamine; toxicity; preconditioning; hormesis; dopamine; serotonin; striatum

Introduction

Methamphetamine (METH) abuse has reached epidemic proportion in the United States and throughout the world (Rawson *et al.*, 2002). METH addicts suffer from cognitive deficits (Chang *et al.*, 2002; Simon *et al.*, 2002; Sekine *et al.*, 2003; Johanson *et al.*, 2006) and several studies have documented biochemical and structural abnormalities in the brains of chronic abusers (Wilson *et al.*, 1996; Volkow *et al.*, 2001; Jernigan *et al.*, 2005; Chang *et al.*, 2007; reviewed by Cadet and Krasnova, 2007). These neuropathological changes have been replicated in mice, rats, and monkeys in which METH causes depletion of monoamines and loss of dopamine and serotonin transporters (Ricaurte *et al.*, 1980; Ricaurte *et al.*, 1982; Cadet *et al.*, 1994; Itzhak and Ali, 1996; Friedman *et al.*, 1998; Ladenheim *et al.*, 2000; Jayanthi *et al.*, 2005; reviewed by Cadet and Krasnova, 2007; Cadet *et al.*, 2007). The majority of these studies were conducted using model systems where moderate to large doses of METH were injected using short intervals on a single day (see Cadet *et al.*, 2003 for discussion of doses

and scheduling of METH injections). These patterns are therefore more compatible with doses taken by addicts during accidental overdoses (Davidson *et al.*, 2001).

Because METH users increase drug usage progressively (Kramer *et al.*, 1967; Cho and Melega, 2002), several attempts have been made to replicate patterns of drug abuse by METH addicts (Stephans and Yamamoto, 1996; Johnson-Davis *et al.*, 2003; Johnson-Davis *et al.*, 2004; O'Neil *et al.*, 2006; Danaceau *et al.*, 2007). These experiments have suggested that pretreatment with METH can provide some degree of protection against drug-induced damage to monoaminergic systems (Schmidt *et al.*, 1985; Gygi *et al.*, 1996; Stephans and Yamamoto 1996; Segal *et al.*, 2003; Johnson-Davis *et al.*, 2004; Danaceau *et al.*, 2007). Because most of the studies have reported on the effects of pretreatment on the striatal dopamine (DA) system, we have recently begun to investigate the effects of METH pretreatment on monoaminergic systems in various regions of the rodent brain and have reported that there might be some differential effects of METH pretreatment on DA and serotonin (5-HT) systems in the rat brain (Graham *et al.*, 2008). In order to clarify these issues further, we have modified the pretreatment schedule used in the previous study to now include the administration of larger doses of METH during the second week. We have also eliminated the 3-day holiday that was used before the final drug challenges (Graham *et al.*, 2008). Thus, in the present study, the rats were injected with the challenge METH only after a 1-day holiday (see Table I). We reasoned that this approach might allow us to better ascertain the differential protective effects of METH preconditioning on the toxic effects of METH challenge (5 mg/kg x 6 given 1 hour apart in a single day) on the levels of norepinephrine (NE), DA, 5-HT and their metabolites in specific brain regions.

Materials and Methods

Animals

Male Sprague-Dawley rats (Charles Rivers Laboratories, Raleigh, NC) weighing approximately 350–400 g were used in the present study. Animals were housed in pairs in polyethylene cages containing hardwood bedding in a humidity- and temperature-controlled room. Animals were given rat chow and water *ad libitum*. All animal use procedures were performed according to the National Institutes of Health *Guide for the Care and Use of Laboratory Animals* and were approved by the local Animal Care Committee.

Drug Treatment and Tissue Collection

Following habituation, each rat was injected i.p. with either METH-hydrochloride (NIDA, Baltimore, MD) or an equivalent volume of saline as described in Table I. Following the preconditioning period, the saline-pretreated animals received either saline or METH (5 mg/kg x 6 at 1 h intervals) challenges whereas the METH-pretreated rats were then given the METH challenge. The animals were sacrificed 24 h later by decapitation, their brains were quickly removed, brain regions were microdissected over ice, snap frozen on dry ice, and stored at –80°C until used in the HPLC experiments. One hemisphere of each region frontal cortex, striatum, nucleus accumbens, and hippocampus was used to generate the HPLC data.

HPLC

For monoamine analysis, the brain regions were homogenized in 0.01M HClO₄ and centrifuged at 14,000 x g for 15 min. NE, DA, 3,4-dihydroxyphenylacetic acid (DOPAC), homovanillic acid (HVA), 5-HT and 5-hydroxyindoleacetic acid (5-HIAA) levels were analyzed in the brain extracts using HPLC with electrochemical detector as previously described (Ladenheim *et al.*, 2000; Krasnova *et al.*, 2007). Monoamine levels were calculated as ng/mg of tissue weight and shown as percentages of control concentrations for ease of presentation.

Statistical Analysis

Statistical analysis was performed using analysis of variance (ANOVA) followed by Fisher's protected least significant difference (StatView 4.02, SAS Institute, Cary, NC). Values were represented as mean $\pm$ SEM. The null hypothesis was rejected at $p < 0.05$.

Results

Striatum

The effects of METH on DA and its metabolites, DOPAC and HVA, in the striatum are shown in Fig. 1A. ANOVA revealed that the drug caused decreases in DA [$F(2,23) = 3.22$; $p = 0.05$]. Post-hoc analyses showed that this was due to METH-induced decreases (-33.6%) in DA levels in the striata of rats pretreated with saline. The METH-pretreated animals suffered no drug-induced decreases in striatal DA levels (Fig. 1A). The METH injections caused significant changes [$F = 3.32$, $p = 0.05$] in the DA metabolite, DOPAC; post-hoc analyses revealed that these were due to differences between the two METH challenged groups in that the METH/METH group had higher levels of DOPAC than the SAL/METH group (Fig. 1A). HVA levels were not significantly affected after the drug challenge [$F = 2.01$, $p = 0.15$] (Fig. 1A). METH caused no changes in NE levels [$F = 0.769$, $p = 0.47$] in the striata of SAL/METH and METH/METH groups (data not shown).

The effects of METH on the 5-HT system are shown in Fig. 1B. The drug caused significant changes in the levels of 5-HT in the striatum [$F(2,23) = 3.63$, $p = 0.04$]. Post-hoc analyses showed that, in contrast to the observations in the DA system, METH pretreatment did not protect against drug-induced decreases in 5-HT levels in the rat striatum; specifically, the METH challenge caused 15.6% decreases ($p = 0.114$) in the SAL/METH group but 24.2% decreases ($p = 0.013$) in 5-HT levels in the METH/METH group (See Fig. 1B). 5-HIAA levels were not significantly affected by the METH challenge [$F(2,23) = 1.04$, $p = 0.36$] (Fig. 1B).

Nucleus Accumbens

The effects of METH on the DA system in the nucleus accumbens are shown in Fig. 2A. The METH challenge did not cause significant decreases in DA levels in the nucleus accumbens [$F(2,21) = 1.93$; $p = 0.17$]. The METH-induced changes in DOPAC levels almost reached significance by ANOVA [$F = 3.1$; $p = 0.06$]. These changes were due to METH-induced decreases (-29.7%) in the nucleus accumbens of saline-pretreated rats whereas there were no differences in DOPAC levels in the METH/METH group in comparison to SAL/SAL control (Fig. 2A). HVA levels were not affected by the METH challenge [$F = 0.812$, $p = 0.45$]. METH treatments caused no significant changes [$F = 0.492$, $p = 0.61$] in NE levels in the nucleus accumbens (data not shown).

Fig. 2B shows the effects of METH on the 5-HT system in the nucleus accumbens. The METH injections did not cause significant changes by ANOVA [$F(2, 21) = 2.49$, $p = 0.10$]. Further comparisons revealed that 5-HT levels in the SAL/METH group were decreased in comparison to the control group (see Fig. 2B). There were significant METH-induced decreases in 5-HIAA levels in nucleus accumbens [$F = 4.25$, $p = 0.02$]. These decreases, although small, were observed in both the SAL/METH (-17.5%) and in the METH/METH (-14.2%) groups (Fig. 2B).

Cortex

The effects of METH on DA and its metabolites in the frontal cortex are shown in Fig. 3A. The METH challenge caused significant changes in DA levels in the cortex of rats [$F(2, 19) = 3.58$, $p = 0.04$]. Post-hoc analyses revealed that there were significant decreases in the saline-pretreated group but not in the METH/METH group (Fig. 3A). The METH challenge did not

affect DOPAC [$F = 0.25$; $p = 0.77$] or HVA [$F = 0.48$, $p = 0.61$] levels. There were no METH-induced changes [$F = 0.66$, $p = 0.52$] in NE levels in the cortices of rats (data not shown).

The effects of METH on the cortical 5-HT system are shown in Fig 3B. ANOVA did not reveal any significant effects of the drug on 5-HT levels [$F = 2.2$, $p = 0.13$]. Further analyses revealed that 5-HT values were significantly decreased (-24.5%) in the SAL/METH group. METH preconditioned rats did not show significant decreases after the acute drug challenge. 5-HIAA levels were not significantly affected by ANOVA [$F = 2.6$, $p = 0.09$]. However, further analyses revealed significant decreases in the SAL/METH rats that showed 23.9% decreases (Fig. 3B). There were no differences between the SAL/METH and the METH/METH groups.

Hippocampus

The effects of METH on NE, 5-HT and 5-HIAA levels in the hippocampus are shown in Fig 4. The METH challenge caused significant decreases in NE levels in the rat hippocampus [$F(2,23) = 8.4$, $p = 0.001$], with the saline- (-27.2%) and the METH-pretreated (-28.9%) groups showing similar changes (Fig. 4). In addition, the METH challenge caused decreases in hippocampal 5-HT levels [$F = 6.45$; $p = 0.005$]. The SAL/METH (-31.3%) and the METH/METH (-28.4%) groups were similarly affected by the drug challenge. 5-HIAA levels were also significantly affected by the METH challenge [$F = 7.49$, $p = 0.003$], with the SAL/METH and the METH/METH showing 28.9% and 22.6% decreases, respectively (Fig. 4).

Discussion

The main findings of the present paper include METH-induced depletion of striatal DA in saline-pretreated rats as reported by us and others (reviewed by Cadet *et al.*, 2003; Cadet and Krasnova, 2007; Cadet *et al.*, 2007). There were also significant decreases in the levels of DA, 5-HT and 5-HIAA in the frontal cortex and hippocampus and of NE levels in the hippocampus of METH-treated saline-pretreated rats. The decreases in NE in the hippocampus are consistent with those recently reported by Graham *et al.* (2008). The present data are also the first to report on the effects of METH preconditioning on monoamine levels in the nucleus accumbens of rats. METH preconditioning was associated with almost complete protection against drug-induced depletion of DA in the striatum and in the frontal cortex, findings that are consistent with the reports that various pretreatment METH regimens can protect against METH-induced DA depletion (Schmidt *et al.*, 1985; Gygi *et al.*, 1996; Johnson-Davis *et al.*, 2003; Segal *et al.*, 2003; O'Neil *et al.*, 2006; Danaceau *et al.*, 2007; Graham *et al.*, 2008). The METH pretreatment also provided small degrees of protection against 5-HT depletion in the nucleus accumbens and the frontal cortex but not in the striatum or hippocampus. The present results are consistent with our previous report that a somewhat different pattern of METH pretreatment was only able to afford small protective effects against drug-induced toxicity on striatal and hippocampal 5-HT systems (Graham *et al.*, 2008). This study, which uses a pattern of higher doses of METH pretreatment and includes the elimination of the 3-day holiday used in a previous study (Graham *et al.*, 2008), expands on that report and provides strong evidence of a very clear dissociation between the effects of METH preconditioning on dopaminergic and serotonergic systems in the rat brain.

The present data are consistent with reports that repeated sub-toxic stimuli can provide protection against subsequent exposure to more fulminant stressors, a process referred to as preconditioning (Blanco *et al.*, 2006) or hormesis (Calabrese, 2008; Mattson, 2008). Hormesis might be a more general and comprehensive term which covers a more varied picture of sub-toxic or sub-clinical manipulations (Calabrese, 2008). These phenomena include pre-exposure to mild ischemia (Blanco *et al.*, 2006), endotoxins (Rosenzweig *et al.*, 2007), and thermic conditions (Ren *et al.*, 2004). Multiple mechanisms are involved in preconditioning or hormesis (Mattson, 2008). These include activation of cellular defense mechanisms such as antioxidant

enzymes and heat shock proteins as well as reduced inflammatory responses (Hoshida *et al.*, 2002; Glantz *et al.*, 2005; Mattson, 2008).

In the case of METH toxicity, it has become clear that several protoxic events are involved in causing loss of monoaminergic terminals (Cadet *et al.*, 2003; Segura-Aguilar and Kostrzewa, 2004; Cadet and Krasnova, 2007; Cadet *et al.*, 2007). Prominent among these are the production of oxygen-based radicals from DA quinones generated within DA terminals (Lavoie and Hastings, 1999), production of reactive species such as superoxide radicals (Cadet *et al.*, 1994) and nitric oxide (Itzhak and Ali, 2006), and the involvement of mitochondria-generated toxic proteins (reviewed in Cadet *et al.*, 2007). Because mitochondria play such an integral part in METH-induced toxicity, drug preconditioning might affect these organelles in such a way that subsequent exposures to toxic doses of METH might not be able to generate enough oxygen-based radicals to cause degeneration of DA terminals. This reprogramming might be dependent on increased production of chaperones such as Hsp60, Hsp70 and others (Sabil, 2008) that might participate in the maintenance of homeostasis within DA terminals, given the presence of large number of mitochondria in nerve axons (Ly and Verstreken, 2006). Thus, low levels of oxygen-based radicals generated by progressive increases in METH administration might trigger signaling functions that stimulate adaptive responses instead of acute toxic stresses that destroy DA terminals. Protective enzymes such as manganese superoxide dismutase, copper/zinc superoxide dismutase, catalase, and glutathione peroxidase might also be mediators of METH preconditioning because these enzymes would protect against METH-induced ROS generation (see Cadet and Brannock, 1998 for discussion). In addition to regulating the generation of reactive oxygen species and increasing enzyme levels, such an adaptation might also lead to tolerance of glutamate-mediated disturbances of calcium homeostasis within these terminals since excitotoxicity is thought to be involved in METH-induced DA terminal degeneration (see Cadet *et al.*, 2007 for discussion). This suggestion is consistent with the demonstration that sub-toxic levels of activation of the N-methyl-D-aspartate (NMDA) type of glutamate receptors can induce adaptive responses in neurons that become resistant to subsequent severe stressful events (Lin *et al.*, 2008).

It is important to note that the METH-induced inaugural events that led to protection against METH-induced striatal and cortical DA depletion did not have similar protective effects against 5-HT depletion in the striatum and the hippocampus, with only small attenuating effects being observed in the cortex and the nucleus accumbens. These findings suggest that the pattern of METH preconditioning used in the present study was not able to generate similar adaptive changes in 5-HT terminals. These observations are important because it is often assumed that identical mechanisms are involved in METH-induced destruction of DA and 5-HT terminals since various manipulations that protect against damage to DA systems also protect against loss of 5-HT terminals (Cadet *et al.*, 2003; Cadet *et al.*, 2007). The present observations are more in accord with gene expression studies that suggest that diverse stimuli might trigger protection through unique though overlapping processes that generate molecular reprogramming. This reprogramming would, in turn, activate the transcription of protective factors such as chaperones, trophic factors and antioxidants (see Dirnagl *et al.*, 2003 and Stenzel-Poore *et al.*, 2003, for further discussion). Because our observations are different from those of Danaceau *et al.* (2007) who reported METH preconditioning-induced protection against brain 5-HT systems using a different METH pre-treatment paradigm, the present data suggest that a given stimulus might cause differential adaptive responses depending on the patterns of drug administration and the neurotransmitter systems under observation. This suggestion is consistent with the fact that the present study was able to document different patterns of protection (or lack thereof) from those that we previously published (Graham *et al.*, 2008) after increasing the doses of METH used during the pretreatment period and eliminating the long interval before challenging the rats with toxic doses of METH. Thus, this discussion suggests that future studies will need to identify specific patterns of drug

preconditioning that provide protection against specific aspects of METH-induced neurotoxicity and use these paradigms to clarify the cellular and molecular bases of the observed protection.

In summary, chronic injections of METH in a pattern that is reminiscent of that used by METH addicts caused almost complete protection against drug-induced DA depletion in the striatum and cortex. This preconditioning paradigm was not protective against the toxic effects of the drug on 5-HT systems in the striatum and hippocampus. METH-induced NE depletion in the hippocampus was also not prevented. These observations suggest that there might be substantial differences between the mechanisms involved in the METH-induced degeneration of these three monoaminergic systems. More in-depth studies will be needed to clarify similarities and differences between the responses of these monoaminergic systems to METH assaults on their structural integrity. Finally, studies focusing on the dissection of the endogenous mechanisms involved in protecting DA terminals should advance our approaches to the treatment of METH addicted individuals and of Parkinsonian patients who suffer from pathologies in their dopaminergic systems.

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Figure 1.

Effects of METH pretreatment and challenge on markers of striatal DA (A) and 5-HT (B) terminals. Values were normalized to SAL/SAL controls and expressed as percentages of controls $\pm$ SEM. N = 8–10 animals per group. Key to statistics: * $p < 0.05$ in comparison to SAL/SAL group, # $p < 0.05$ in comparison to SAL/METH group.

Figure 2.

Levels of DA, DOPAC and HVA (A) and of 5-HT and 5-HIAA (B) in the nucleus accumbens following METH pretreatment and challenge. Values were normalized to SAL/SAL controls and expressed as percentages of controls $\pm$ SEM. N = 8–10 animals per group. Key to statistics:

* $p < 0.05$ in comparison to SAL/SAL group.

Figure 3.

Effects of METH pretreatment and challenge on markers of cortical DA (A) and 5-HT (B) terminals. Values were normalized to SAL/SAL controls and expressed as percentages of controls $\pm$ SEM. N = 8–10 animals per group. Key to statistics: * $p < 0.05$ in comparison to SAL/SAL group, # $p < 0.05$ in comparison to SAL/METH group.

Figure 4.

Levels of NE, 5-HT, 5-HIAA in the hippocampus following METH pretreatment and challenge. Values were normalized to SAL/SAL controls and expressed as percentages of controls $\pm$ SEM. N = 8–10 animals per group. Key to statistics: ** $p < 0.01$, *** $p < 0.001$ in comparison to SAL/SAL group.

Table 1

Schedule of METH pretreatment and challenges

	Monday	Tuesday	Wednesday	Thursday	Friday	Week 1
9:00	0.5 mg/kg	1 mg/kg	1 mg/kg	1.5 mg/kg		
10:00						
11:00			1 mg/kg	1.5 mg/kg		
12:00						
13:00			1 mg/kg	1.5 mg/kg		
14:00						
15:00		1 mg/kg	1 mg/kg	1.5 mg/kg		
16:00						
	Monday	Tuesday	Wednesday	Thursday	Friday	Week 2
9:00	1 mg/kg	1.5 mg/kg	1.5 mg/kg	2 mg/kg		
10:00			1.5 mg/kg	2 mg/kg		
11:00	1 mg/kg	1.5 mg/kg	1.5 mg/kg	2 mg/kg		
12:00			1.5 mg/kg	2 mg/kg		
13:00	1 mg/kg	1.5 mg/kg	1.5 mg/kg	2 mg/kg		
14:00			1.5 mg/kg	2 mg/kg		
15:00	1 mg/kg	1.5 mg/kg	1.5 mg/kg	2 mg/kg		
16:00			1.5 mg/kg	2 mg/kg		
	Monday	Tuesday	Wednesday	Thursday	Friday	Week 3
9:00	2 mg/kg		5 mg/kg			
10:00	2 mg/kg		5 mg/kg			
11:00	2 mg/kg		5 mg/kg			
12:00	2 mg/kg		5 mg/kg			
13:00	2 mg/kg		5 mg/kg			
14:00	2 mg/kg		5 mg/kg			

Initially the rats were divided into two groups, with one group receiving saline and the other group getting METH pretreatment according to the schedule described during first and second weeks as well as Monday of the third week. The saline pretreatment was followed by with either saline (SAL/SAL) or METH challenges (SAL/METH), the METH pretreatment was followed by METH challenges (METH/METH) on Wednesday of the third week. Rats were killed 24 hours later.