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Persistent Reversal of Enhanced Amphetamine Intake by Transient CaMKII Inhibition

Jessica A. Loweth,1,2 Dongdong Li,2 James J. Cortright,2 Georgia Wilke,2 Okunola Jeffyosis,3 Rachael L. Neve,4 K. Ulrich Bayer,2 and Paul Vezina1,2

1Committee on Neurobiology, and Departments of 2Psychiatry and Behavioral Neuroscience and 3Neurobiology, The University of Chicago, Chicago, Illinois 60637, 4Massachusetts Institute of Technology, Cambridge, Massachusetts 02139, and 5Department of Pharmacology, University of Colorado Denver, Aurora, Colorado 80045

Amphetamine exposure transiently increases Ca2+/calmodulin-dependent protein kinase II (CaMKII) α expression in the nucleus accumbens (NAcc) shell and this persistently increases local GluA1 S831 phosphorylation and enhances behavioral responding to the drug. Here we assessed whether transiently interfering with CaMKII signaling using a dominant-negative CaMKII mutant delivered to the NAcc shell with herpes simplex viral vectors could reverse these long-lasting biochemical and behavioral effects observed following exposure to amphetamine. As expected, transient expression of CaMKIIα K42M in the NAcc shell produced a corresponding transient increase in CaMKIIα and decrease in pCaMKIIα (T286) protein levels in this site. Remarkably, this transient inhibition of CaMKII activity produced a long-lasting reversal of the increased GluA1 S831 phosphorylation levels in NAcc shell and persistently blocked the enhanced locomotor response to and self-administration of amphetamine normally observed in rats previously exposed to the drug. Together, these results indicate that even transient interference with CaMKII signaling may confer long-lasting benefits in drug-sensitized individuals and point to CaMKII and its downstream pathways as attractive therapeutic targets for the treatment of stimulant addiction.

Introduction

Previous exposure to psychostimulants such as amphetamine and cocaine enhances subsequent neurochemical and behavioral responding to the drug. These phenomena are thought to contribute to the transition from casual drug use to addiction (Robinson and Berridge, 1993; Vezina, 2004). A brain area prominently linked to the generation of addictive behaviors is the nucleus accumbens (NAcc) shell (Anderson et al., 2008). We recently showed that amphetamine exposure transiently increases expression of the Ca2+/calmodulin-dependent protein kinase II (CaMKII) α isoform in this site and that transient viral overexpression of CaMKIIα mimics the behavioral effects of amphetamine exposure in rats (Loweth et al., 2010). Conversely, NAcc shell infusion of a CaMKII inhibitor (KN93) prevents cocaine-induced reinstatement of drug seeking (Anderson et al., 2008), as well as the expression of enhanced NAcc dopamine overflow (Pierce and Kalivas, 1997), cocaine-induced locomotion (Pierce et al., 1998), and amphetamine self-administration (Loweth et al., 2008), normally observed in sensitized rats. However, KN93 affects not only CaMKII and other protein kinases, but also L-type Ca2+ channels (Li et al., 1992; Gao et al., 2006), and an inhibitor of these channels produces the same biochemical (Pierce and Kalivas, 1997) and behavioral effects (Pierce et al., 1998; Anderson et al., 2008) as KN93. Furthermore, these studies did not directly assess the possibility that transient NAcc shell CaMKII inhibition could have long-lasting effects on behavioral responding to psychostimulants, a possibility suggested by our earlier work showing that transient increases in CaMKIIα in this region lead to long-lasting neuroadaptations that contribute to the maintenance of sensitization (Loweth et al., 2010). Thus, in the present study, we used a CaMKII- and brain region-specific transient inhibition strategy to determine whether interfering directly with endogenous CaMKII signaling within the NAcc shell could persistently reverse long-lasting behavioral manifestations of amphetamine sensitization.

Materials and Methods

Strategy for CaMKII inhibition by viral-mediated expression of a K42M mutant. Replication-deficient herpes simplex virus (HSV) viral vectors were constructed and packaged as previously described by us (Neve et al., 1997; Loweth et al., 2010). These vectors were used because they produce transient overexpression of the transgene that peaks 3–4 d post-infection and returns to baseline by day 8 post-infection (Neve et al., 1997; Carlson and Neve, 2003; Loweth et al., 2010). In addition, local viral infection permits functional testing of specific groups of neurons in specific brain areas (here, the NAcc shell). We chose to transiently express a CaMKIIα K42M mutant that is “kinase dead” due to impaired ATP binding. Such kinase-dead mutants are commonly used as “dominant-negative” mu-
tants to interfere with endogenous kinase functions, and this has been described for CaMKII (e.g., Kühl et al., 2000; Xiao et al., 2005; Garic et al., 2011). Dominant-negative effects can be exerted by competition with endogenous kinase for upstream activators, downstream substrates, or subcellular targeting modules. In the case of CaMKII, which forms dodecameric homooligomers, dominant-negative effects can also be exerted by the formation of heteromeric homooligomers, which interfere with the intra-homooligomer inter-subunit autophosphorylation at T286 that generates Ca$^{2+}$-independent “autonomous” CaMKII activity (for reviews, see Coultrap and Bayer, 2012; Lisman et al., 2012). In addition, the K42M mutant interferes with the targeting of homooligomers to the NMDA-type glutamate receptor subunit GluN2 (Bayer et al., 2006; O’Leary et al., 2011), which is regulated by nucleotide binding to CaMKII (O’Leary et al., 2011) and also generates autonomous activity (Bayer et al., 2001; Bayer et al., 2006). Thus, the K42M mutant of CaMKII has even greater dominant-negative potential than the kinase-dead mutants of most other kinases. The regulation of CaMKII is rather complex (Coultrap and Bayer, 2012), but it should be noted that some cellular functions that simply require Ca$^{2+}$/calmodulin-stimulated CaMKII activity may or may not be affected by the presence of additional kinase-dead mutant. Thus, the designation as a dominant-negative mutant (as done here) is always linked to a specific cellular function; for other functions, a particular kinase-dead mutant may or may not act in a dominant-negative fashion (for further discussion, see Wayman et al., 2011).

Subjects and surgery. Male Sprague-Dawley (locomotion experiments) and Long–Evans rats (self-administration experiments) weighing 250–275 g on arrival were purchased from Harlan Sprague-Dawley and housed individually with food and water available ad libitum. They were implanted with chronic bilateral guide canulae as described previously (Loweth et al., 2010). For the amphetamine self-administration studies, rats were also implanted with intravenous catheters as described previously (Suto et al., 2004). All surgical procedures were conducted using aseptic techniques according to an approved Institutional Animal Care and Use Committee protocol.

Design and procedure. Rats in different groups were exposed to repeated intermittent injections of amphetamine or saline and 2–3 weeks later, infused intracranially into the NAcc shell with HSV-K42M CaMKIIα or a control infusion (mock) consisting of HSV-LacZ or 10% sucrose vehicle. In rats in one experiment were tested for their locomotor response to a systemic amphetamine injection 4 d and again 8 d following injection. Rats in a second experiment were tested for their self-administration of amphetamine before and after HSV infection. Rats in additional groups were killed 4 or 8 d following HSV infection to determine the expression pattern of the transgene and assess its effect on phosphorylation of the AMPA receptor GluA1 subunit at S831, a CaMKII residue. In these cases, brain sections were harvested and subsequently assessed using immunoblotting.

Exposure injections. Exposure injections (5 × 1.5 mg/kg amphetamine or 1.0 ml/kg saline) were administered every 2–3 d as described previously (Vezina et al., 2002; Suto et al., 2004). S (+)-amphetamine sulfate was obtained from Sigma-Aldrich and dissolved in sterile saline. Doses refer to the weight of the salt.

Viral-mediated gene transfer in the NAcc shell. The following HSV vector constructs were used: CaMKIIα K42M and a control vector, LacZ, which encodes the protein β-galactosidase. LacZ and 10% sucrose vehicle were used interchangeably for mock-infection control infusions as they have consistently been found to be without effect (Loweth et al., 2010; Singer et al., 2010). Rats were transported to a biosafety level 2 facility where, as described previously (Loweth et al., 2010; Singer et al., 2010) and according to an approved Institutional Biosafety Committee protocol, they were administered bilateral intracranial microinjections into the NAcc shell of their respective viral vectors or mock control vehicle. Rats were returned to the colony room 24 h later.

Locomotor testing. To assess the effect of transient expression of CaMKIIα K42M in NAcc shell neurons on sensitized locomotor responding to amphetamine, rats were assigned randomly to different groups based on exposure (systemic amphetamine or saline) and infection (NAcc shell HSV-K42M CaMKIIα or mock). Thus, four groups were tested: amphetamine-mock, amphetamine-K42M, saline-mock, and saline-K42M. Two to 3 weeks after the exposure regimen, rats received their respective NAcc shell microinjections and were tested for their locomotor response to amphetamine (1.0 mg/kg, i.p.) 4 and 8 d later. On each test day, locomotor activity was measured 1 h before and 2 h after the amphetamine challenge injection using a bank of 12 activity boxes as described previously (Vezina et al., 2002).

Self-administration training and testing. To assess the effects of transient expression of CaMKIIα K42M on enhanced amphetamine self-
administration, separate rats were assigned randomly to one of four groups based on exposure and infection as described above for locomotor testing. Two to 3 weeks after the last exposure injection, rats were trained to self-administer amphetamine on fixed ratio (FR) schedules of reinforcement and infection (HSV-CaMKII-K42M) infusions. Locomotor responding to AMphetamine was assessed when protein levels were elevated (day 4) and once they had dissipated (day 8). Data are shown to the left as group mean (±SEM) 2 h total locomotor counts following the AMphetamine injection (arrows). In each case, time courses of the locomotor responses are illustrated to the right as group mean (±SEM) locomotor counts obtained before and after the AMphetamine injection (arrows). Transient expression of CaMKII-K42M blocked enhanced AMphetamine-induced locomotion at both days 4 and 8 post-infection. **p < 0.01, ***p < 0.001, AMF exposed mock-infected versus remaining groups; by significant ANOVA followed by LSD tests. n = 6 – 7/group. c. Location of the injection cannula tips for rats included in the data analyses. Symbols indicate group affiliation. Line drawings are from Paxinos and Watson (2005). Numbers to the right indicate the number of millimeters from bregma.

Results

Transient CaMKII inhibition persistently reverses increased GluA1 S831 phosphorylation in the NAcc shell of amphetamine-exposed rats

HSV vectors were used to transiently overexpress an inactive mutant of CaMKIIα (K42M) in NAcc shell neurons (Fig. 1e). As we have previously described for active CaMKIIα (Loweth et al., 2010). Two to 3 weeks after exposure to amphetamine or saline, rats received
CaMKII protein and T286 phosphorylation levels were assessed on days 4 and 8 post-infection (Fig. 1b,c). Consistent with previous findings (Loweth et al., 2010), no changes in either CaMKIIα protein or T286 phosphorylation were detected 2–3 weeks after exposure to amphetamine as displayed in the mock-infected rats. Subsequent infection with HSV-CaMKIIα K42M transiently increased total CaMKIIα protein levels in both amphetamine- and saline-treated rats; these returned to baseline levels by day 8 post-infection (Fig. 1b). In contrast, T286 phosphorylation was transiently decreased, both in absolute terms (data not shown) and as a ratio of phospho-T286 to total CaMKIIα (Fig. 1c), confirming that expression of CaMKIIα K42M interfered with endogenous CaMKII signaling. This effect was also transient and no longer evident by day 8 post-infection (Fig. 1c). The ANOVA conducted on CaMKIIα levels and the ratio pCaMKIIα/CaMKIIα showed significant effects of infection only on day 4 post-infection ($F_{(1,19)} = 20.12, p < 0.001$ and $F_{(1,19)} = 15.90, p < 0.001$, respectively). No other effects, including on day 8 post-infection, were statistically significant. Thus, transient viral expression of CaMKIIα K42M transiently interfered with "autonomy"-inducing CaMKII T286 phosphorylation, which was back to control levels on day 8 post-infection.

As exposure to a sensitizing amphetamine regimen leads to a long-lasting increase in phosphorylation of GluA1 at S831 in the NAcc shell (Loweth et al., 2010), we then assessed the effect of transiently inhibiting CaMKII signaling in this site on phosphorylation of this GluA1 residue. Again, a sustained increase in GluA1 S831 phosphorylation was detected in the NAcc shell of mock-infected rats 2–3 weeks after exposure to amphetamine. This increase in phospho-S831 was reversed by transient and no longer evident by day 8 post-infection ($F_{(1,19)} = 15.90, p < 0.001$, respectively). No other effects, including on day 8 post-infection, were statistically significant. Thus, transient viral expression of CaMKIIα K42M transiently interfered with "autonomy"-inducing CaMKII T286 phosphorylation, which was back to control levels on day 8 post-infection.

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### Transient CaMKII inhibition in the NAcc shell of amphetamine-exposed rats persistently reverses enhanced locomotor response to and self-administration of the drug

As expected, rats exposed to amphetamine 2–3 weeks earlier showed a greater locomotor response to a systemic amphetamine challenge (1.0 mg/kg, i.p.) compared with saline-exposed controls (Vezina, 2004) (Fig. 2). Remarkably, transient CaMKII inhibition in the NAcc shell persistently reversed this expression of locomotor sensitization by amphetamine. The enhanced locomotor response was completely abolished in the HSV-CaMKIIα K42M-infected animals, not only on day 4 (Fig. 2a) but also on day 8 post-infection (Fig. 2b), again a time point when both CaMKIIα expression and T286 phosphorylation had returned to baseline levels (Fig. 1b,c). Significant effects of exposure (day 4, $F_{(1,21)} = 6.30, p < 0.05$; day 8, $F_{(1,21)} = 4.15, p = 0.054$) and infection (day 4, $F_{(1,21)} = 9.20, p < 0.01$; day 8, $F_{(1,21)} = 4.43, p < 0.05$) were detected on both test days. Thus, we next assessed the effect of transient CaMKII inhibition on the enhanced amphetamine intake observed following amphetamine exposure (Vezina et al., 2002; Vezina, 2004). Rats were again randomly assigned to four groups based on condition of exposure (amphetamine or saline) and infection (bilateral NAcc shell HSV-CaMKIIα K42M or mock infection). As expected (Vezina et al., 2002), rats exposed to amphetamine and given the opportunity to self-administer the drug intravenously worked significantly more and as a result obtained more infusions compared with saline-exposed controls (Fig. 3; day −1 pre-infection, exposure, $F_{(1,28)} = 5.41, p < 0.05$). However, the day following infection through day 4 post-infection when expression of the transgene was maximal, amphetamine-exposed rats infected with HSV-CaMKIIα K42M no longer displayed enhanced amphetamine intake, obtaining the same number of infusions as saline-exposed rats and significantly fewer than mock-infected amphetamine-exposed rats (Fig. 3; day 4 post-infection, exposure, $F_{(1,28)} = 3.58, p = 0.07$; $E \times I$, $F_{(1,28)} = 6.69, p < 0.05$). Importantly, this
reversal of enhanced amphetamine self-administration was maintained for the full 12 d of testing post-infection and thus again outlasted the transient interference with CaMKII signaling achieved by HSV-CaMKIIα K42M (Fig. 3; days 8–12 post-infection, E × I, F(1,25) = 4.09, p = 0.05).

Discussion
Expressing a dominant-negative form of CaMKIIα (K42M) that transiently inhibits CaMKII activity in NAcc shell neurons persistently blocked the expression of two major manifestations of sensitization by amphetamine. As expected, previously exposing rats to amphetamine led to sensitized locomotor responding to a challenge injection of the drug and, in a separate group of rats, enhanced drug intake under a PR schedule of reinforcement (Vezina et al., 2002; Vezina, 2004). Transiently inhibiting CaMKII activity in the NAcc shell of these rats blocked these manifestations of sensitization and continued to do so long after expression of the transgene had dissipated, indicating that continued uninterrupted CaMKII activation is necessary for their maintenance. Previous studies suggested that CaMKII activity in the NAcc shell is required for the expression of stimulant sensitization (Pierce and Kalivas, 1997; Pierce et al., 1998; Loweth et al., 2008) and cocaine-induced reinstatement (Anderson et al., 2008). However, these previous studies tested the pharmacological inhibitor KN-93 that is known to affect not only CaMKII and other protein kinases but also to inhibit L-type Ca2+ channels, rendering an interpretation of results that focuses specifically on CaMKII difficult. The use of HSV-mediated delivery of a CaMKIIα mutant in the present experiments circumvented these shortfalls. Importantly, the present study is the first to unequivocally demonstrate that transiently inhibiting CaMKII activity in the NAcc shell can lead to the persistent and selective reversal of enhanced behavioral response to amphetamine. Thus, the evidence reported here identifies CaMKII and the signaling pathways it initiates as attractive therapeutic targets for the treatment of substance abuse.

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