The transcription factor cAMP response element binding protein (CREB) is implicated in the actions of drugs of abuse in several brain areas, but little information is available about a role for CREB in the ventral tegmental area (VTA), one of the key reward regions of the brain. Here, we demonstrate that chronic exposure to drugs of abuse induces CREB activity throughout the VTA. Using viral-mediated gene transfer, we expressed green fluorescent protein (GFP)-tagged CREB or mCREB (a dominant-negative form of CREB) in the VTA and, using a conditioned place-preference paradigm, found that CREB activation within the rostral versus caudal subregions of the VTA produces opposite effects on drug reward. We identified VTA subregion-specific differences in the proportion of dopaminergic and GABAergic neurons and in the dopaminergic projections to the nucleus accumbens, another brain region implicated in drug reward, and suggest that this may contribute to behavioral differences in this study. We also measured expression levels of tyrosine hydroxylase and the AMPA glutamate receptor subunit GluR1, both of which are known to contribute to drug reward in the VTA, and found that both of these genes are upregulated following the expression of CREB-GFP and downregulated following expression of mCREB-GFP, raising the possibility that CREB may exert its effects on drug reward, in part, via regulation of these genes. These results suggest a novel role for CREB in mediating drug-induced plasticity in the VTA and establish two functionally distinct subregions of the VTA in which CREB differentially regulates drug reward.

Key words: cocaine; reward; ventral tegmental area; CREB; mesocortical; mesolimbic; morphine

Introduction

Repeated exposure to drugs of abuse causes complex molecular adaptations in discrete brain regions, including the reward circuitry of the brain (Wise, 1996; Koob and Le Moal, 2001; Nestler, 2001). An example of this plasticity is activation of the transcription factor cAMP response element (CRE)-binding protein (CREB) in several regions following chronic morphine, cocaine, or amphetamine exposure (Guitart et al., 1992; Konradi et al., 1994; Lane-Ladd et al., 1997; Barrot et al., 2002; Shaw-Lutchman et al., 2002, 2003; Walters et al., 2003). CREB regulates the expression of genes by interacting with CREs in their regulatory regions. Several signal transduction pathways activate CREB, suggesting that this transcription factor may integrate signaling information from multiple sources and thereby mediate complex forms of neural plasticity (Bailey et al., 1996; Daniel et al., 1998; Shaywitz and Greenberg, 1999; Mayr and Montminy, 2001; Nestler, 2001).

Prominent in brain reward circuitry is the ventral tegmental area (VTA). Drug induction of phospho-CREB in the VTA has been reported recently (Walters et al., 2003), but direct evidence of the functional role of CREB in the VTA is still lacking. Chronic drug exposure alters expression of many proteins in this region, including components of the cAMP, Ras, and phospholipase C/ IP3 signaling cascades, all of which can activate CREB (Berhow et al., 1996; Lane-Ladd et al., 1997; Wolf et al., 1999). In addition, some studies have reported increased levels of the glutamate receptor subunits GluR1 and NMDA receptor 1 in the VTA after...
chronic morphine, cocaine, or ethanol administration (Ortiz et al., 1995; Fitzgerald et al., 1996) (but see Carlezon and Nestler, 2002; Wolf, 2003). These effects could lead to glutamatergic-mediated increases in intracellular Ca\(^{2+}\) and CREB activation (Perkinton et al., 1999; Rajadhyaksha et al., 1999; Riccio and Ginty, 2002). These lines of evidence led us to hypothesize that drug-induced changes in CREB function in the VTA might mediate some biochemical and behavioral adaptations to chronic drug exposure.

Well established, but often overlooked, functional differences in rostral versus caudal VTA were first demonstrated by Ikemoto et al. (1997b, 1998) using intracranial self-administration of GABAergic compounds. They found that GABA\(_A\) antagonists were reinforcing in rostral VTA, whereas GABA\(_A\) agonists were reinforcing in caudal VTA, and speculated that these differences could be related to the differential expression of GABA\(_A\) receptors in GABAergic and dopaminergic neurons in the VTA (McBride et al., 1999). Similar rostral versus caudal VTA differences in glutamatergic regulation of drug reward have been reported previously (Carlezon et al., 2000). Consequently, we focused on differences in the effects of CRE in rostral and caudal VTA and considered a possible role for GABAergic neurons in this phenomenon.

We now report that CREB-mediated transcription is upregulated in the VTA in response to chronic drug exposure, and that this upregulation has different behavioral consequences depending on the subregion of the VTA involved. Careful anatomical analysis confirmed the presence of two distinct subregions of VTA (rostral and caudal) that differ in both neuronal population and projection to the nucleus accumbens (NAc), which could explain why these two subregions differentially regulate reward.

Materials and Methods

Animals. The CRE-reporter mice used in this study express a transgene containing CRE-consensus sequences in tandem, upstream of a minimal regulatory sequence. In experiments examining CREB activity and behavior, both Cre-LacZ and CREB-reporter mice were used for all experiments. For each experimental animal, a littermate of the same gender and bearing the reporter transgene was used as a control. Adult male Sprague Dawley rats (Charles River, Kingston, NY) were used for all other experiments and also were maintained under a 12 h light/dark cycle with food and water available ad libitum. Genotyping was performed by PCR, and male and female mice (between 8 and 12 weeks of age) heterozygous for the transgenic sequence were used for all experiments. For each experimental animal, a littermate of the same gender and bearing the reporter transgene was used as a control. Adult male Sprague Dawley rats (Charles River, Kingston, NY) were used for all other experiments and also were maintained under a 12 h light/dark cycle with food and water available ad libitum. Rats were used for viral and tracer injections and for studies of target genes because of the larger size of their brains and the ability to more easily discriminate between VTA subregions. In contrast, the Cre-LacZ mice provide a unique means of studying regulation of CRE-mediated transcription in the brain in vivo, which is not possible in the rat.

Drug administration to transgenic mice. In experiments examining CREB activation after chronic morphine administration, morphine was administered chronically twice daily by intraperitoneal injection of an escalating dose of morphine sulfate over 5 d (10, 20, 40, 80, and 100 mg/kg) or saline injections of an equal volume (Shaw-Lutcheman et al., 2002). To ensure that Cre-LacZ activation was not attributable to the acute effects of morphine, animals were killed with an overdose of pentobarbital 12 h after the final injection and perfused transcardially with 4% paraformaldehyde in 0.1M phosphate buffer. At this time point, levels of CRE activity returned to normal levels in the VTA and other brain regions after a single injection of morphine (100 mg/kg) (Shaw-Lutcheman et al., 2002).

Immunohistochemistry and anatomical analysis. Brains were postfixed for 12 h and then cryoprotected in 20% glycerol in 0.1 M phosphate buffer for 12 h. Brains were then cut (40 μm sections) coronally on a freezing microtome and stored in PBS with 0.05% sodium azide. In experiments using transgenic mice, LacZ expression was detected with a polyclonal antibody to β-galactosidase (β-gal) (1:5000; Biogenesis, Poole, UK), and some sections were double labeled for tyrosine hydroxylase (TH) (1:1000; Chemicon, Temecula, CA) to determine whether the β-gal-positive neurons were dopaminergic. To directly test the influence of herpes simplex virus (HSV) vectors on CRE-mediated transcription, mice positive for the CRE-LacZ transgene were injected with HSV-CRE-GFP-green fluorescent protein (GFP), HSV-mCREB-GFP (a dominant-negative form of CREB with a mutation at the serine 133 phosphorylation site that prevents phosphorylation and transactivation) or HSV-Cre-GFP (a viral vector expressing the bacterial protein Cre recombinase tagged with GFP, which is inactive in the absence of a floxed gene) 3 d before use. In all cases, antibody labeling was visualized using cyanine 2 (Cy2) and Cy3 fluorophore-labeled secondary antibodies (1:200; Jackson ImmunoResearch, West Grove, PA), and sections were mounted onto slides in DPX (a mixture of diystere, tricresyl phosphate, and xylene) mounting media (Fluka BioChemika, Ronkonkoma, NY). Seven sections from injected mice were taken at five anatomical points throughout the VTA: −2.92, −3.08, −3.16, −3.28, and −3.40 mm bregma; the first three points were within the territory of the rostral VTA, whereas the latter two points were in the territory of the caudal VTA. Total CRE-LacZ-positive neurons with expression above basal background levels were counted using a technique described previously (Barrot et al., 2002). In experiments quantifying transgenes and cell type in rats, sections were labeled with either an antibody to TH (see above) or to GAD65 and GAD67 (1:10,000; Sigma) and either an antibody to β-gal (see above) or GFP (1:1000; Upstate Biotechnology, Waltham, MA) depending on the transgene expressed.

Viral vectors. Construction and packaging of viral vectors has been described previously (Neve et al., 1997). Transgenes (LacZ, Cre recombinase-GFP, CREB-GFP, and mCREB-GFP) were inserted into the HSV PrcUC ampiclon under the control of an HSV 4/5 immediate early gene promoter, packaged with helper virus 5dl1.2, purified on a sucrose gradient, pelleted, and resuspended in 10% sucrose. The average titer of viral stocks was 2.0 × 10^10 infectious units/ml. CREB-GFP and mCREB-GFP fusion proteins were created by adding a GFP tag to the N terminus of CREB and mCREB. Previous studies have indicated that tagging the N terminus of CREB does not interfere with its functional activity (Chao et al., 2002). Functional activity of these constructs was confirmed in cell culture, in which overexpression of CREB-GFP increased while overexpression of mCREB decreased CRE-mediated transcription (data not shown). After intra-VTA injection, maximal expression of all viral-encoding constructs is observed 24 h after injection. Behavioral tests were carried out 3 d after injection and then dissipates and is not detectable by day 7, as reported previously (Carlezon et al., 1997; Barrot et al., 2002). This time course is independent of the particular transgene expressed.

SurgerY. Surgery was performed as described previously (Barrot et al., 2002). Animals were anesthetized and received unilateral injections into the VTA [rats: 2 μl of virus stock; anteroposterior (AP), −4.9 to −6.2 mm caudal to bregma; lateral (L), +2.1 mm from the midline; arm set at a 10° angle toward the midline; dorsoventral (DV), −7.3 mm from dura; mice: 0.5 μl of virus stock; AP, −2.8 mm caudal to bregma; L, +1.0 mm from midline; arm set at a 7° angle toward midline; DV, −4.6 mm from dura] or the lateral substantia nigra pars compacta [rats only; AP, −5.3 mm caudal to bregma; L, +3.1 mm from midline; arm set at a 10° angle toward midline; DV, −6.5 mm from dura], which served as an anatomical control. For sham surgery, the syringe was lowered 1.0 mm below the dura, but no injection was made. We used unilateral injections, because we found that bilateral injections in the VTA offer no greater biological effect than unilateral injections, and the latter involve less stress to the animal and less labor on the part of the experimenter (Carlezon et al., 1997; Bolanos et al., 2003).

Behavior. Using a balanced and unbiased paradigm, place conditioning occurred in a three-compartment apparatus as described previously (Kelz et al., 1999). This apparatus consisted of a central small shuttle chamber between two larger conditioning chambers, which differed in
lighting, walls, and floors. Before surgery, animals were placed into the central compartment and allowed to freely explore the entire apparatus for 30 min. Animals that did not show a baseline preference (>18 min) or a preference for the central chamber (>6 min) were admitted into the study; >80% of all animals met these criteria.

After surgery, animals were allowed either 1 d of recovery and subsequently conditioned for 3 d consecutively or were allowed 2 d of recovery and conditioned for 3 d consecutively. No significant difference between the two conditioning protocols was observed. These time points were chosen based on the fact that maximal HSV-transgene expression occurs on days 1–4 after surgery (Barrot et al., 2002). On conditioning days, two pairing sessions were given. On the first trial of each day, animals received a volume of saline equivalent to the volume of drug they would receive and were restricted to one of the conditioning chambers for 1 h. After the first pairing session, animals were returned to their home cages and allowed to rest for a minimum of 3 h. On the second trial of each day, animals received a dose of drug and were restricted to the other conditioning chamber for 1 h. Depending on the group, drugs were given as follows: 1.25 mg/kg cocaine, i.p., or 0.125, 0.25, and 0.5 mg/kg morphine, s.c. After the final training day, animals were tested for conditioned place preference (CPP) by placing them in the central chamber and allowing them to freely explore the entire apparatus for 30 min. Drug and saline pairings were given at the same time on each conditioning day, and animals were tested for preference at the same time as the drug pairing on the previous day. After testing (or for light microscopy, 3 d after intra-ventral tegmental area (VTA), isolated as described above, were homogenized in 1 ml of TRIzol reagent (Invitrogen, San Diego, CA) and extracted following the protocol of the manufacturer. RNA was resuspended in DEPC-treated water, quantified from A260 measurements, and assessed for quality by visual inspection of samples electrophoresed on agarose gels. The RNA was treated with 2 U of DNase using the DNA-free kit (Ambion, Austin, TX) and then heated at 65°C for 10 min to inactivate the enzyme. cDNA synthesis was performed using the Superscript First-Strand Synthesis System for reverse transcription (RT)-PCR (Invitrogen) as follows: 1 μg of RNA was preincubated with 1 μl of Oligo(dT) 12–18 (0.5 μg/μl) and 1 μl of 10 μM deoxyribonucleoside triphosphates at 65°C for 5 min. After chilling on ice for 1 min, 4 μl of First Strand buffer (5X), 2 μl of DTT (0.1 M), 1 μl of RNase-OUT, and 1 μl (50 U) of Superscript II reverse transcriptase were added for a final volume of 20 μl. The mixture was incubated at 42°C for 50 min and then 70°C for 15 min to terminate the reaction. Each sample was then incubated at 37°C for 20 min with 2 U of RNase H.

Real-time RT-PCR and data analysis. PCRs were performed using the LightCycler-FastStart DNA Master SYBR Green I kit (Roche Products, Welwyn Garden City, UK) as follows. A total of 2 μl of cDNA diluted 1:10 in nuclease-free water was added to 2.5 μl of FastStart reaction mix, 2 μl of 25 mM MgCl2 (final MgCl2 concentration, 3 mM), and water to a final volume of 25 μl. Thermocycling and detection of the SYBR green fluorescent dye were performed using the LightCycler System (Cepheid). Thermocycling parameters were as follows: 95°C for 10 min (DNA polymerase activation) followed by 18 cycles of denaturation at 95°C for 10 s, annealing at 58°C for 10 s, and extension at 72°C for 10 s. Optical detection of the fluorescent signal occurred during the extension phase. Each experiment was performed in duplicate, and the results were averaged for data analysis. The cycle threshold (Ct) was calculated by software supplied by the manufacturer as the cycle during which the second derivative of the fluorescent signal reached maximum value. Data for all genes of interest were normalized to the expression levels of two housekeeping genes (B-actin, GAPDH) by the following formula: ΔCt = Ct(gene of interest) – Ct (GAPDH or B-actin).

Results

Chronic morphine exposure increases CRE-mediated transcription in the VTA in a topographically specific manner

We mapped the ability of chronic morphine administration to activate CRE-mediated transcription in the VTA using CRE-LacZ transgenic mice. Because the VTA is a heterogeneous collection of cell groups, we characterized β-gal induction by dividing the VTA into two subregions (rostral and caudal VTA) based on functional differences between these subregions reported previously in the rat (Arnt and Scheel-Kruger, 1979; Ikemoto et al., 1997b, 1998; McBride et al., 1999; Ikemoto and Wise, 2002; Zangen et al., 2002; Bolanos et al., 2003). In both rostral and caudal VTA, there was a significant increase in CRE-LacZ-positive cells after chronic morphine exposure (Fig. 1A).

Because the VTA is known to be populated by both GABAergic and dopaminergic neurons, we next sought to characterize the neuronal type in which this upregulation occurs by colabeling with an antibody to TH, a marker of dopamine neurons. In rostral-VTA...
and caudal VTA, we observed a significant upregulation of CRE-LacZ in dopaminergic neurons (Fig. 1B). However, in rostral VTA, a significant number of these β-gal-positive neurons were TH negative (Fig. 1C) (presumably GABAergic), indicating that CREB activation in this region may occur in a more heterogeneous population of neurons than in caudal VTA, a supposition supported by work presented below (see Fig. 6). Thus, these findings suggest that chronic morphine exposure induces CREB activity in partly distinct populations of neurons in rostral versus caudal VTA.

Viral-mediated expression of CREB-GFP or mCREB-GFP in the VTA alters conditioned place preference for cocaine in an anatomically distinct manner

Based on our anatomical data that drug induction of CRE activity differs between rostral and caudal VTA and on previously reported data for functional differences between these regions (see the Introduction), we considered the possibility that CREB may differentially regulate the effects of drugs of abuse in the rostral versus caudal VTA. To test this hypothesis, we expressed CREB-GFP or mCREB-GFP in different areas of the rat VTA using HSV vectors. The GFP-tagged fusion proteins made it possible to distinguish transgenic CREB (or mCREB) from endogenous CREB. Cre-GFP (a bacterial protein, Cre recombinase, tagged with GFP) was expressed as a control. We assessed the rewarding effects of cocaine using the place-conditioning paradigm, in which animals learn to prefer a drug-paired environment. In an attempt to map different functional subregions of the VTA, we plotted place-preference scores as a function of the placement of viral infection in each rat. The results are shown in Figure 2. From these data, we identified two functionally different subregions in rat VTA. In rostral VTA, which spans the supramammillary area to the rostral pole of the interpeduncular nucleus (−4.9 to −5.4 mm bregma), expression of CREB-GFP made a threshold dose of cocaine highly rewarding, whereas mCREB-GFP made this dose of cocaine aversive. Conversely, in caudal VTA, which spans the rostral pole of the interpeduncular nucleus to the caudal end of the VTA (−5.5 to −6.2 mm bregma), CREB-GFP expression made the threshold dose of cocaine aversive, whereas mCREB-GFP made this cocaine dose highly rewarding (Fig. 2A, B). These two anatomical areas correlate well with previous studies describing two functionally
different subregions of the VTA (Ikemoto et al., 1997b, 1998; Carlezon et al., 2000; Ikemoto and Wise, 2002; Zangen et al., 2002; Bolanos et al., 2003; Rodd et al., 2005) and with our data demonstrating differential induction of CRE activity within non-dopaminergic cells of these subregions in CRE-LacZ mice (Fig. 1). Subsequently, all experimental animals were divided into two groups with injections into either rostral or caudal VTA. As found in several previous studies (Neve and Geller, 1995; Carlezon et al., 1997, 1998, 2000; Neve et al., 1997), injection of HSV vectors caused minimal damage to the infected area, which was indistinguishable from that seen with vehicle injections alone (our unpublished observations).

To confirm the effectiveness of CREB-GFP and mCREB-GFP in regulating CRE-mediated transcription in the VTA in vivo, we injected HSV-CREB-GFP, HSV-mCREB-GFP, or HSV-Cre-GFP throughout the VTA of CRE-LacZ transgenic mice. Expression of CREB-GFP significantly increased CRE-LacZ expression in both rostral and caudal VTA relative to control animals, whereas expression of mCREB-GFP significantly decreased CRE-LacZ expression levels relative to control animals injected with HSV-Cre-recombinase-GFP. Data are expressed as cell numbers per section ± SEM (n = 3–4). *p < 0.05; t test.

Viral-mediated expression of CREB-GFP or mCREB-GFP in the VTA alters conditioned place preference for morphine

We next tested whether altered CREB activity in the VTA also affects the rewarding effects of a threshold dose of morphine. Results equivalent to those seen for cocaine were obtained. Animals that received HSV-CREB-GFP in rostral VTA showed a strong preference for a threshold dose of morphine, whereas those that received HSV-mCREB-GFP into rostral VTA showed a mild aversion. In contrast, animals that received HSV-CREB-GFP in caudal VTA showed a mild aversion for this morphine dose, whereas those that received HSV-mCREB-GFP in caudal VTA exhibited strong place conditioning (Fig. 4A). In contrast, injection of the HSV vectors into the nearby substantia nigra had no effect on morphine reward (data not shown), indicating the regional specificity of CREB action within the VTA.

To better understand these effects of morphine, place conditioning was studied over a full-dose range of the drug. At higher doses of morphine, the effects of HSV-mediated changes in CREB activity in rostral and caudal VTA disappeared, presumably because higher doses of drug overcome the inhibitory effects of HSV-mCREB-GFP in rostral VTA and of HSV-CREB-GFP in caudal VTA. However, a higher dose of morphine (0.5 mg/kg)
was apparently necessary to overcome the effect of HSV-mCREB-GFP in rostral VTA compared with HSV-CREB-GFP in caudal VTA (0.25 mg/kg) (Fig. 4B, C).

GluR1 and TH are putative target genes for CREB in the VTA

CREB affects molecular, cellular, and behavioral processes by regulating the transcription of numerous genes. As a first step in understanding the mechanism by which expression of CREB-GFP and mCREB-GFP within the VTA alters CPP to drugs of abuse, we used real-time RT-PCR to measure changes in message levels of two CREB-regulated genes implicated in drug reward. Previous studies suggest that both GluR1, an AMPA glutamate receptor subunit that regulates drug reward via its actions in the VTA (Carlezon et al., 1997, 2000; Carlezon and Nestler, 2002), and TH, the rate-limiting enzyme in dopamine biosynthesis induced by drugs of abuse in the VTA (Nestler, 2001), are regulated by CRE sites in their respective promoters (Joh et al., 1978; Lewis et al., 1987; Hiremagalur et al., 1993; Kim et al., 1993; Boundy et al., 1998; Myers et al., 1999; Borges and Dingledine, 2001). We found that expression of CREB-GFP in the VTA increased local GluR1 and TH message levels, whereas expression of mCREB-GFP had the opposite effect (Fig. 5A). The relative difference in message levels between CREB-GFP and mCREB-GFP-treated animals was nearly twofold (Fig. 5A). Additional evidence for CREB regulation of these genes was obtained by Western blot analysis. As shown in Figure 5B, injection of HSV-CREB-GFP into the VTA significantly increased levels of TH and GluR1 immunoreactivity in this brain region.

Different neuronal populations are infected in rostral versus caudal VTA

The VTA is thought to contain two primary types of neurons: dopaminergic neurons, which comprise approximately two-thirds of the neurons in this region, and GABAergic neurons, which comprise the remainder. However, the number of dopaminergic neurons is reportedly greater in more caudal subregions of the VTA (Swanson, 1982; German and Manaye, 1993). We therefore hypothesized that our HSV vectors might be infecting different proportions of dopaminergic and GABAergic neurons in caudal VTA versus rostral VTA, and that such differences could contribute to the different behavioral effects observed after vector injections into these subregions. Using double-label immunohistochemistry for GFP (to mark viral infection) and for TH or GAD (a marker of GABAergic neurons), we characterized the types of neurons infected with HSV-CREB-GFP in rostral versus caudal VTA. As shown in Figure 6A, the percentage of infected neurons colabeled with TH increased in an approximately linear manner from rostral to caudal VTA. When we examined the number of HSV-infected neurons that expressed GAD, we saw the opposite pattern; the percentage of infected neurons colabeled with GAD decreased in an approximately linear manner from rostral to caudal VTA. Similar results were obtained for HSV-LacZ (data not shown), a control viral vector without behavioral effects, consistent with previous studies demonstrating that efficiency of HSV infection is independent of neuronal type or viral-encoded transgene.

When we pooled the neurons counted into rostral and caudal VTA groups based on the anatomical boundaries used in our behavioral experiments, we found that approximately equal proportions of GABAergic and dopaminergic neurons were infected in rostral VTA. However, in caudal VTA, by far, more infected neurons were dopaminergic than GABAergic (Fig. 6B).

More dopaminergic neurons in caudal VTA project to the NAc than in rostral VTA

Previous studies have shown that dopamine neurons in the VTA, particularly caudal regions, project to the NAc (Swanson, 1982), a brain region also strongly implicated in drug and natural reward. To determine whether the neurons that we were infecting in caudal VTA project to the NAc, we injected a retrograde tracer (HSV) into the NAc shell and HSV-LacZ into both rostral and caudal VTA. We saw no difference in the number of β-gal-positive neurons between rostral and caudal VTA (Fig. 7A). However, consistent with previous studies, we found more retrogradely labeled neurons in caudal than rostral VTA, indicating that more neurons in this subregion project to the NAc. We next examined the projections of these rostral and caudal VTA neurons by colabeling for β-gal and HS. Of the neurons expressing β-gal, significantly more neurons in the caudal VTA compared with the rostral VTA were colabeled with HS. Thus, a significantly greater number of infected neurons in caudal VTA than in rostral VTA project to the NAc. In addition, triple labeling for β-gal, HS, and TH demonstrated that nearly all of these infected projection neurons, in either rostral or caudal VTA, were dopaminergic, a finding consistent with the fact that the majority of VTA projections to the NAc are dopaminergic (Swanson, 1982) (Fig. 7A). These results, combined with those presented above, suggest that the population of HSV transgene-expressing neurons in rostral VTA differs significantly from that in caudal VTA not only in terms of cell type but also in efferent projections.

Discussion

Using CRE-LacZ reporter mice, we show that CREB activity is upregulated throughout the VTA by chronic morphine administration. We then used viral-mediated expression of CREB-GFP or mCREB-GFP in the VTA to assess the contribution of CREB activity to drug reward. Our study defines two functionally distinct subregions of the VTA (rostral and caudal), which differ dramatically in their responses to CREB-GFP and mCREB-GFP expression. In rostral VTA, CREB-GFP expression made mor-
also in the types of neurons that populate them, as well as in their regions. Differences observed in this and previous studies. These subregions could underlie the functional differences in rostral versus caudal VTA that could underlie the functional effects of CREB-GFP and mCREB-GFP expression in both subregions. This study is the first to demonstrate that CREB-activated transcription in response to chronic drug administration occurs throughout the VTA, and that this induction occurs in different populations of neurons in rostral versus caudal VTA.

Using anatomical boundaries of rostral and caudal VTA described previously, we found that CRE-LacZ expression was increased in rostral and caudal subregions of VTA after chronic drug exposure, consistent with the recent finding that phospho-CREB levels are increased in this region after chronic exposure to drugs of abuse (Walters et al., 2003). CRE-LacZ expression was induced in TH-positive neurons in both VTA subregions, whereas induction in TH-negative neurons, presumably GABAergic, was seen in rostral VTA only. This study is the first to demonstrate that CREB-activated transcription in response to chronic drug administration occurs throughout the VTA, and that this induction occurs in different populations of neurons in rostral versus caudal subregions.

Using viral vectors, we next showed that CREB activation in rostral versus caudal VTA has opposite effects on morphine and cocaine reward. Greater doses of morphine can overcome the effects of CREB-GFP and mCREB-GFP expression in both subregions. This may be a result of the activation of other brain regions involved in morphine reward. Another possibility is that, at higher doses, other transgene-negative VTA neurons are recruited. It is also interesting to note that a higher dose of morphine is required to overcome these effects in rostral rather than caudal VTA. This may reflect the differences in the sensitivity of these two neuronal populations to the effects of either CREB or morphine.

As a transcription factor, CREB produces its effects by regulating the expression of other genes. We examined the expression of two candidate genes that may contribute to CREB action in the VTA: GluR1 and TH. Both genes have functional CREs in their promoters (Lewis et al., 1987; Hiremagalur et al., 1993; Kim et al., 1993; Boundy et al., 1998; Myers et al., 1999; Borge and Duling, 2001), and both are upregulated in the VTA by chronic drug exposure (Fitzgerald et al., 1996; Lane-Ladd et al., 1997). Furthermore, upregulation of GluR1 expression, like CREB, has opposite effects on drug reward in rostral versus caudal VTA (Carlezon et al., 2000). Although we were not able to distinguish rostral and caudal VTA in these experiments because of technical limitations (see Materials and Methods), we did find that CREB-GFP expression increases GluR1 and TH mRNA and protein levels in the VTA, and that mCREB-GFP has the opposite effect.

We hypothesized that the functional differences between the effects of CREB in rostral and caudal VTA and similar rostrocaudal differences reported previously may be attributable to neuroanatomical differences between these two subregions. To address this possibility directly, we characterized the relative abundance of two types of VTA neurons, GABAergic and dopaminergic, in rostral and caudal VTA. Whereas approximately equal percentages of GABAergic and dopaminergic neurons were infected in rostral VTA, a much larger percentage of dopaminergic neurons was infected in caudal VTA. Because HSV vectors do not preferentially infect one neuronal cell type more than another, our results presumably reflect the different cellular make up of these two subregions, consistent with previous studies reporting a proportionately greater number of dopamine neurons in caudal VTA (Swanson, 1982; German and Manaye, 1993). In addition, we present evidence that rostral and caudal VTA neurons differ in their efferent projections: many more caudal VTA neurons project to the NAc. Using triple-labeling techniques, we found that, in both rostral and caudal subregions, the large majority of NAc-projecting neurons are dopaminergic. In the present study, retrograde tracer was injected into the mid-rostrocaudal shell of the NAc. Additional studies are needed to determine whether this topographical organization of VTA dopamine projections holds for other NAc subregions.

These rostrocaudal differences in neuronal cell type and projections could explain why the two subregions influence drug reward mechanisms so divergently. The explanation of such differences remains highly speculative, particularly because we do not yet know what effect CREB activity has on the excitability of
rostral and caudal VTA neurons. Nevertheless, a working hypothesis can be proposed.

Previous studies have shown that expression of GluR1 or administration of picrotoxin, a GABA<sub>A</sub> receptor antagonist, into rostral VTA is rewarding. Both of these treatments could amplify the excitability of rostral VTA neurons either by increasing their sensitivity to glutamate or by disinhibiting them via blockade of GABA<sub>A</sub> receptors. Ikemoto et al. (1997a) suggested that this excitation promotes reward by increasing mesolimbic (i.e., VTA to NAc) dopamine neurotransmission, a hypothesis supported by their finding that administration of picrotoxin into rostral VTA increases dopamine release in the NAc. A similar increase in neuronal activity via CREB-induced GluR1 expression could occur in rostral VTA mesolimbic dopaminergic neurons. Mesolimbic dopamine release could be further augmented by an increase in TH expression and, consequently, dopamine levels.

Another mechanism that could account for the effects of CREB in rostral VTA is a similar excitation of mesocortical neurons, which predominate in this subregion (Swanson, 1982; Loughlin and Fallon, 1984). Both GABAergic and dopaminergic VTA neurons project to the prefrontal cortex, and activation of both neuronal types inhibits prefrontal cortex neurons (Pirot et al., 1992). This inhibition could lead to decreased corticlimbic excitatory drive, a mechanism proposed to promote reward (Carlezon and Wise, 1996a,b; Wise, 1996, 2002). This hypothesis is supported by our recent finding that injection of HSV vectors into rostral VTA causes transgene expression in mesocortical GABA neurons (V. Olson and E. J. Nestler, unpublished observations), which make up ~70% of the mesocortical projection (Swanson, 1982). It is also possible that these two mechanisms act synergistically to mediate the effects of CREB in rostral VTA on reward.

Why upregulation of CREB in caudal VTA antagonizes drug reward is less straightforward, but one possible explanation is that elevated CREB levels in this region sensitize an animal to the aversive properties of drugs. Because most neurons in this region are dopaminergic, this effect would appear to be mediated almost exclusively by this cell type. CREB induction of GluR1 and TH in these neurons could increase their excitability and, therefore, local dopaminergic tone. An increase in local dopamine release could activate inhibitory autoreceptors on mesolimbic dopamine neurons (Falkenburger et al., 2001; Beckstead et al., 2004) and could also activate D<sub>1</sub> receptors on GABA nerve terminals arising from the NAc, increasing local GABA release and further attenuating mesolimbic dopamine neuron firing (Mansour et al., 1991; Sugita et al., 1992; Cameron and Williams, 1993). Such activation of local autoreceptors may also explain why lower, but not higher, doses of cocaine are self-administered into the caudal VTA (Rodd et al., 2005). Moreover, mesocortical dopamine neurons would be impervious to these changes because they do not express inhibitory autoreceptors (Bunney et al., 1973; Aghajanian and Bunney, 1977; Bannon et al., 1981; White and Wang, 1984). A final consideration is that caudal VTA may be a direct target of drugs of abuse, whereas rostral VTA may influence drug reward through indirect, circuit-level interactions among the major reward regions of the brain. Additional studies are needed to clarify these important issues.

The studies reported here support a fundamental reconsideration of the role of the VTA in brain reward circuitry. Our findings contribute to a growing body of evidence suggesting that the VTA is a heterogeneous brain region, with at least two functionally discrete subregions that can modify drug reward in a distinct and independent manner, neither of which fit the conventional model of drug reward circuitry. We contend that differences in rostral versus caudal VTA, by analogy with differences in the shell versus core subregions of the NAc, should be taken into account in studies of VTA function. We provide two mechanisms for why...
these VTA subregions differ, including a difference in the proportion of dopaminergic versus GABAergic neurons in rostral and caudal VTA, and a difference in the efferent projections of these neurons. Our findings support a distinct role for these VTA subregions in drug reward and provide new insight into the molecular neuropathological changes in this region that contribute to drug addiction.

References
Ikemoto S, Murphy JM, McBride WJ (1997b) Self-infusion of GABA(A) antagonists directly into the ventral tegmental area and adjacent regions. Behav Neurosci 111:369–380.


