Up-Regulation of AGS3 during Morphine Withdrawal Promotes cAMP Superactivation via Adenylyl Cyclase 5 and 7 in Rat Nucleus Accumbens/Striatal Neurons

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Received May 15, 2009; accepted June 23, 2009

ABSTRACT

Effective medical treatment of opiate addiction is limited by a high relapse rate in abstinent addicts. Opiate withdrawal causes cAMP superactivation, but the underlying molecular mechanisms are not clear. Recent evidence implicates an activator of G-protein signaling 3 (AGS3) in opiate addiction. We found previously that during a 10-min activation of opioid receptors, AGS3 binds Ga,-GDP to promote free Gβγ stimulation of adenylyl cyclase (AC) 2 and 4, and/or inactivate Ga inhibitory function, thereby transiently enhancing cAMP-dependent protein kinase A (PKA) activity. In contrast, we report here that in nucleus accumbens/striatal neurons, morphine withdrawal induces cAMP superactivation, which requires up-regulation of AGS3. cAMP increases as a function of withdrawal time, by approximately 20% at 10 min and 75% at 5 h. However, cAMP superactivation does not require Gβγ. Instead, adenosine A2A receptor activation of Ga,-GDP seems to initiate cAMP superactivation and promote AGS3 up-regulation. Elevated AGS3 binds to Ga, to prevent its inhibition on AC activation. Moreover, withdrawal-induced increases in cAMP/PKA activate phospholipase C and e protein kinase C to further stimulate AC5 and AC7, causing cAMP superactivation. Our findings identify a critical role for AC 5 and 7 and A2A receptors for up-regulation of AGS3 in morphine withdrawal-induced cAMP superactivation.

Opiate addiction is a worldwide public health problem, but effective treatment is limited by high rates of relapse during abstinence (O’Brien, 2005). Sharma et al. (1975) developed model neuronal cell systems expressing opiate receptors to identify molecular mechanisms associated with exposure to and withdrawal from opiates. They discovered that activation of opioid receptors in NG108-15 cells produced an initial reduction in cAMP followed by a compensatory up-regulation during continued exposure to morphine. It is noteworthy that cAMP increased even further after opiate receptor blockade with antagonists or by morphine withdrawal, a phenomenon described as cAMP overshoot, superactivation, supersensitization, or heterologous sensitization of adenylyl cyclase (AC) (Watts, 2002). Increased cAMP activities cAMP-dependent protein kinase A (PKA) and cAMP response element (CRE)-dependent gene transcription, thought to regulate the development of tolerance and dependence (Chao and Nestler, 2004). Opiates exert their actions by binding to the 6-opioid receptor (DOR), the μ-opioid receptor (MOR), and the κ-opioid receptor. Morphine, an opiate agonist, acts primarily on the MOR to promote opiate seeking behavior (Narita et al., 2001). Opiate receptors belong to a superfamily of pertussis toxin (PTX)-sensitive G-protein coupled receptors. Activation of MOR releases Ga and Gβγ. Ga inhibits adenylyl cyclase to decrease cAMP production. On the other hand, Gβγ can stimulate AC and also activate several other down-stream signaling molecules, including phospholipase C (PLC), potassium channels, and so forth. (Gautam et al., 1998). Emerging evidence suggests that cAMP superactivation consists of enhanced Ga,-receptor coupling, G-protein dissociation, and Ga,-adenyl cyclase interaction (Watts and Neve, 2005; Chakrabarti and Gintzler, 2007). However, it is unclear how...
G-protein dissociation could occur in the absence of morphine during withdrawal and how Gαs can be involved in G2-coupled MOR signaling. Recent evidence suggests that signaling via G-proteins can be regulated by receptor-independent activators (Takesono et al., 1999). A family of such regulatory accessory proteins includes an activator of G-protein signaling 3 (AGS3) (Blumer et al., 2007). AGS3 binds to Gαi3-GDP (De Vries et al., 2000), enhances unbound free Gβγ stimulation of AC2 and AC4 (Yao et al., 2005), and/or diminishes Gαi2-GTP inhibition of AC (Takesono et al., 1999; Kimple et al., 2002). We have shown that knockdown of AGS3 or expressing a Gβγ inhibitor blocks morphine-induced cAMP/PKA signaling in primary nucleus accumbens/striatal neurons (Yao et al., 2005). Inhibition of AGS3 expression in rat nucleus accumbens (NAC) core or rat prefrontal cortex also eliminates withdrawal-induced release of heroin-, cocaine-, and ethanol-seeking behavior (Bowers et al., 2004; Yao et al., 2005). Moreover, withdrawal from cocaine or ethanol increases AGS3 expression in rat prefrontal cortex and the core of nucleus accumbens (Bowers et al., 2004, 2008). These findings suggest that up-regulation of AGS3 may be a decisive molecular mechanism underlying opiate withdrawal-induced cAMP superactivation and relapse.

Materials and Methods

Materials. All reagents were purchased from Sigma (St. Louis, MO), except where indicated. Rp-cAMPs was obtained from BioLog (La Jolla, CA). Bisindolylmaleimide I (GF109203X) was purchased from Calbiochem (San Diego, CA). cAMP assay kit and [γ-32P]ATP (6000 Ci/mmol) were purchased from GE Healthcare (Chalfont St. Giles, Buckinghamshire, UK). Complete protease inhibitors tablets were purchased from Roche Molecular Biochemicals (Indianapolis, IN).

Cell Culture and Transfection. Primary nucleus accumbens (NAC)/striatal neuron preparations and culture were carried out as described previously (Yao et al., 2002). In brief, NAC/striatal cells from Sprague-Dawley rat newborn brain (postnatal day 0) were plated at 5 × 10^4 cells/cm^2 and grown in Neurobasal A medium (Invitrogen, Carlsbad, CA) supplemented with 1× B27 supplement (Invitrogen). Half the medium was changed 1 day after plating and weekly thereafter. Experiments were carried out on day 10. For siRNA transfection, cells were incubated with 100 pmol of siRNA for AC2, AC4, AC5, or AC7 (Santa Cruz Biotechnology, Santa Cruz, CA) plus 5 μL of Lipofectamine 2000 (Invitrogen). Transfection was performed as described by the manufacturer. For adenovirus transfection, cells were incubated with Ad5βARK1 (Yao et al., 2002) or Ad5AG53AS (Yao et al., 2005) at a multiplicity of infection of 100.

cAMP Assay. NAC/striatal neurons were treated with 100 nM morphine or not treated for 18 h. To withdraw, cells were completely washed three times with culture medium and continued incubation for the indicated time in the absence of morphine. Cells were then harvested, suspended in 1 ml of Opti-MEM I (Invitrogen) containing 1 unit/ml adenosine deaminase and 10 μM rolipram, and incubated at 37°C for 10 min. cAMP was assayed using HitHunter cAMP XS Assay kit (GE Healthcare) as described by the manufacturer. Data are mean ± S.E.M. obtained from three to six independent experiments, with each treatment carried out in triplicate.

Immunocytochemistry and Microscopy. Cells were fixed on ice-cold PBS, harvested in 1 ml of lysis buffer containing 20 mM Tris, pH 7.5, 2 mM EDTA, 10 mM EGTA, 0.25 M sucrose, and one tablet of protease inhibitor/10 ml. Cells were homogenized by 10 passes through a 26-gauge needle and centrifuged at 3000 rpm for 5 min at 4°C. The supernatant was centrifuged for 8 min at 70,000 rpm at 4°C in a TLA 120 rotor (Beckman Coulter, Fullerton, CA) to separate the membrane pellet from the cytosol. The supernatant was saved as the cytosolic fraction. The remaining pellets were suspended in 1 ml of lysis buffer containing 1% Triton X-100, titrated, and incubated on ice for 20 min. This suspension was centrifuged as described above, and the Triton-soluble material was collected as the original particulate fraction. Each experiment was repeated three times with each treatment in triplicates.

Immunoprecipitation and Western Blot. Cells in 100-mm dishes (2 × 10^6 cells/dish) were washed with ice-cold PBS, harvested in 1 ml of lysis buffer, and lysed on ice for 20 min. The lysates were centrifuged at 14,000 rpm at 4°C in an Eppendorf centrifuge and the supernatant was subject to immunoprecipitation. Five micrograms of pan PKC polyclonal immunoglobulin G antibody or AGS3 polyclonal immunoglobulin G antibody was incubated with 50 μL of protein A/G beads (Santa Cruz Biotechnology, Santa Cruz, CA) overnight at 4°C, washed three times with PBS, and coverslipped with Vectashield mounting medium (Vector Laboratories, Burlingame, CA). The cells were then washed three times with PBS, incubated at room temperature with goat anti-mouse IgM (Cappel, Aurora, OH) (diluted 1:1000) and anti-NeuN (Millipore Bioscience Research Reagents, Temecula, CA), washed three times with PBS, and overlaid with Vectashield mounting medium (Vector Laboratories, Burlingame, CA). Cells were imaged using a Carl Zeiss LSM PASCAL (Jena, Germany) laser scanning microscope equipped with an argon laser attached to an Axiosvert 200 M microscope. Images were collected as z-series. Collected data were processed using NIH Image (http://rsb.info.nih.gov/nih-image/) and Photoshop software (Adobe Systems, Mountain View, CA). All images shown were obtained under 40× magnification from individual middle sections of the projected z-series. To view PKA Co translocation, random fields on each slide were selected, and cells were examined for PKA Co staining in the Golgi, nucleus, cytoplasm, and neurites.

Cell Fractionation. Cells in 100-mm dishes (2 × 10^6 cells/dish) were washed with ice-cold PBS and lysed in 1 ml of lysis buffer containing 20 mM Tris, pH 7.5, 2 mM EDTA, 10 mM EGTA, 0.25 M sucrose, and one tablet of protease inhibitor/10 ml. Cells were homogenized by 10 passes through a 26-gauge needle and centrifuged at 3000 rpm for 5 min at 4°C. The supernatant was centrifuged for 8 min at 70,000 rpm at 4°C in a TLA 120 rotor (Beckman Coulter, Fullerton, CA) to separate the membrane pellet from the cytosol. The supernatant was saved as the cytosolic fraction. The remaining pellets were suspended in 1 ml of lysis buffer containing 1% Triton X-100, titrated, and incubated on ice for 20 min. This suspension was centrifuged as described above, and the Triton-soluble material was collected as the original particulate fraction. Each experiment was repeated three times with each treatment in triplicates.

PKC Activity Assay. Cells in 100-mm dishes (2 × 10^6 cells/dish) were washed with ice-cold PBS, harvested in 1 ml of lysis buffer, and lysed on ice for 20 min. The lysates were centrifuged at 14,000 rpm at 4°C in an Eppendorf centrifuge and the supernatant was subject to immunoprecipitation. Five micrograms of pan PKC polyclonal immunoglobulin G antibody or AGS3 polyclonal immunoglobulin G antibody was incubated with 50 μL of protein A/G beads (Santa Cruz Biotechnology, Santa Cruz, CA) overnight at 4°C, washed three times with PBS, and bound material was collected as the original particulate fraction. Each experiment was repeated three times with each treatment in triplicates.

Results

Morphine Withdrawal Induces AGS3 Up-Regulation. Recent studies suggest that increased AGS3 expression seems to be required for cocaine- and ethanol-induced re-in-
Morphine withdrawal increases cAMP and activates PKA. Because PKA seems to be required for AGS3 up-regulation during morphine withdrawal, we sought to determine whether morphine withdrawal increases cAMP in NAc/striatal neurons. Withdrawal from morphine induced cAMP production as a function of time, increasing by approximately 20% at 10 min and reaching 75% at 5 h (Fig. 2A). Increasing morphine exposure time from 18 to 24 or 48 h, or increasing morphine concentrations from 100 nM to 300 nM or 1 μM, did not further increase morphine withdrawal-induced cAMP production (data not shown).

The NAc/striatum expresses DOR and MOR. To determine which receptor is involved in cAMP superactivation, cells were pretreated with opioid receptor antagonists for 30 min and continued incubation with 100 nM morphine for 18 h, followed by 5-h withdrawal. Withdrawal-induced cAMP superactivation was attenuated by the DOR antagonist naltrexone (1 μM) (Fig. 2B). In addition, pretreatment with PTX also blocked the cAMP increase (Fig. 2B), suggesting that Goαq is required for cAMP superactivation. In control studies, opioid antagonists or PTX did not affect basal cAMP levels (data not shown).

Increased cAMP activates PKA by binding to its regulatory subunit and releasing the catalytic subunit (Ca), which translocates from the Golgi to other cell compartments to phosphorylate its substrates (Yao et al., 2006). Confocal microscopy shows that 10 min after morphine withdrawal, PKA Ca was translocated from the Golgi to the cytoplasm and the nucleus, similar to the effect of a 10-min treatment with the AC activator, forskolin (Fig. 2C). Western blots confirmed Ca translocation from the membrane to the cytosolic fraction (Fig. 2D). This finding is consistent with an opiate withdrawal-induced increase in cAMP at 10 min, suggesting that a 20% increase in cAMP is sufficient to activate PKA. By contrast, long-term exposure to morphine did not cause PKA Ca translocation (Fig. 2, C and D).

Adenosine A2A Receptors Promote cAMP Superactivation via AC5 and AC7. Increasing evidence suggests that cAMP superactivation involves receptor-Goα coupling and enhanced Goα sensitization (Chakrabarti and Gintzler, 2007). The NAc/striatum is enriched in Goα5/6-coupled adenosine A2A receptors (A2A), which stimulate cAMP production in response to released adenosine (Yao et al., 2006) and modulate the neuronal responses to heroin (Yao et al., 2006). Thus, we first determined whether A2A is involved in the initiation of withdrawal-induced cAMP superactivation by eliminating adenosine or blocking A2A before withdrawal. Indeed, adenosine deaminase, which degrades adenosine, or

Fig. 1. Morphine withdrawal increases AGS3 expression. A, representative Western blot showing AGS3 expression as a function of time. NAc/striatal neurons were treated with morphine (100 nM) for 18 h. To withdraw, cells were washed with growth medium for three times and continue cultured for the indicated time. Cells were then lysed for Western analysis of AGS3 expression. B, relative intensity of individual bands from A was quantitated using http://rsbweb.nih.gov/ij/ software. Blots were normalized to GAPDH. Data are the mean ± S.E.M. of three independent experiments. C, representative Western blot showing PKA-dependent AGS3 up-regulation. Morphine withdrawal (W.D.) was performed as above, and cells were then incubated for 5 h with or without Rp-cAMPS (Rp, 20 μM). D, relative intensity of individual bands from C was quantitated using ImageJ software. Blots were normalized to GAPDH. Data are the mean ± S.E.M. of three independent experiments. **, p < 0.01 compared with control (one-way ANOVA and Dunnett’s test).
the A2A antagonist KW-6002 inhibited morphine withdrawal-induced cAMP superactivation, respectively (Table 1). Then, we determined whether A2A plays a role in the sustained cAMP superactivation by treating cells with adenosine deaminase or KW-6002 at 3 h after withdrawal. We found that A2A was not involved in sustaining cAMP super-activation (Table 2), suggesting that A2A is required only for the initial stimulation of cAMP superactivation. Next, we determined which ACs are involved in morphine withdrawal-induced cAMP superactivation by knocking down each AC isoform with a sequence-specific siRNA. Each siRNA suppressed its corresponding AC expression by more than 80% but had no significant effect on basal cAMP levels (data not shown). However, siRNA for AC5 or AC7 alone significantly attenuated morphine withdrawal-induced cAMP superactivation; when the siRNA were added together, morphine withdrawal-induced cAMP superactivation was completely blocked. It is noteworthy that siRNA for AC2, AC4, and other ACs had no effect (Table 1, or data not shown). Activation of A2A releases Gs/olf, which should stimulate all AC isoforms (Tang and Gilman, 1991). The selective requirement for AC5 and AC7 in opiate-withdrawal induced superactivation suggests that other signaling molecules are likely to be involved.

**εPKC Is Required for cAMP Superactivation and AGS3 Expression.** It has been reported that protein kinase C (PKC) stimulates AC2, AC5, and AC7 in neuronal and non-neuronal cell lines and is implicated in morphine tolerance (Narita et al., 1994). We next sought to determine whether cAMP superactivation requires PKC. GF109203X, a PKC inhibitor, attenuated cAMP superactivation (Fig. 3A). This observation led us to determine whether morphine withdrawal activates PKC. We found that morphine withdrawal increased PKC activity by approximately 125%; this increase in PKC activity was blocked by the PKA inhibitors, Rp-cAMPS, or the PLC inhibitor Et-18-OCH3 (edelfosine), respectively (Fig. 3B), consistent with evidence for cross-talk between PKA and PKC signaling (Yao et al., 2008). We next identified a specific PKC isoform activated during morphine withdrawal. Figure 3C shows that the εPKC-specific peptide inhibitor εV1-2 blocked PKC activation, but there was no effect of the δPKC peptide inhibitor δV1-1, or the classic PKC peptide inhibitor βC2-4. These findings suggest that morphine withdrawal seems to activate εPKC. Indeed, Western blots show that during morphine withdrawal, activated εPKC translocates from membrane to cytoplasm as a function of time, starting at 2 h and peaking at 5 h (Fig. 3, D and E). Because AGS3 up-regulation requires PKA (Fig. 1, C and D), and cAMP superactivation requires εPKC (Fig. 3A), AC5, and AC7 (Table 1), we next sought to determine whether inhibition of PKC, AC5, or AC7 prevents AGS3 up-regulation. Figure 3F shows that inhibition of εPKC either by GF109203X or by εV1-2 reduced AGS3 up-regulation and that knockdown of AC5 and AC7 prevented AGS3 up-regulation. These findings suggest that PKA and εPKC seem to interact with each other to mediate AGS3 up-regulation and cAMP superactivation during morphine withdrawal.

![Fig. 2. Morphine withdrawal induces cAMP/PKA signaling. A, cAMP increases as a function of time. NAc/striatal neuron culture and morphine withdrawal (W.D.) were conducted as Fig. 1A. Cells were harvested at the indicated time, and cAMP was determined as described under Materials and Methods. Data are the mean ± S.E.M. of four independent experiments with each treatment carried out in triplicates. B, the role of MOR and DOR in cAMP superactivation. Cells were preincubated with the DOR antagonist naltrindole (NLT, 100 nM), the MOR antagonist CTOP (0.1 μM, CTOP 0.1; 0.3 μM, CTOP 0.3; or 1 μM, CTOP 1), or the nonselective opioid receptor antagonist naltrexone (NLX, 1 μM) for 30 min, or PTX (50 ng/ml) overnight, respectively. Cells were then treated with 100 nM morphine for 18 h. cAMP was assayed 5 h after withdrawal. Data are the mean ± S.E.M. of six independent experiments with each treatment carried out in triplicates. C, representative confocal microscopic image showing PKA Ca translocation from the Golgi to the cytoplasm determined 10 min after withdrawal (W.D.). Forskolin (1 μM for 10 min) was used as a positive control. Green indicates Ca, and red indicates nucleus stained with NeuN. D, representative Western blot showing PKA Ca translocation from particulate fraction (P) to cytosolic fraction (C) determined 10 min after withdrawal. *, p < 0.05; **, p < 0.01 compared with control (one-way ANOVA and Dunnett’s test).](image-url)


Up-Regulation of AGS3 Contributes to cAMP Superactivation. To determine the role of AGS3 in the cAMP superactivation, we transfected cells with adenovirus carrying AGS3 antisense before long-term morphine exposure. We found that AGS3 antisense largely prevented morphine withdrawal-induced cAMP superactivation. In addition, the role of AGS3 in the superactivation was confirmed by treating cells with AGS3 inhibitor peptide Tat-GPR (Bowers et al., 2004) (Fig. 3A). Increased AGS3 expression is thought to drive the formation of AGS3-G_{i3}, thus reducing the amount of G_{i3} available to inhibit AC (Takesono et al., 1999; Kimpile et al., 2002). To test this possibility, we sought to determine whether G_{i3} is coimmunoprecipitated with AGS3 using AGS3 antibody. Withdrawal from morphine promoted the formation of AGS3-G_{i3} (Fig. 4, A and B). This increase was blocked by the PKC inhibitor GF109203X or the PKA inhibitor Rp-cAMPS, respectively. Moreover, increased formation of AGS3-G_{i3} is not due to a concomitantly increased expression of G_{i3}, because withdrawal did not affect G_{i3} expression (Fig. 4, C and D).

Discussion

The major findings in this study are that morphine withdrawal induces up-regulation of AGS3, which in turn seems to regulate morphine withdrawal-induced cAMP superactivation. Elevated AGS3 binds to G_{i3}, preventing inhibitory G_{i3} from restraining cAMP signaling. Thus, inactivation of G_{i3} signaling facilitates activation of PKA. Increased PKA, known to increase PKC activity via PLC (Yao et al., 2008), promotes PKC stimulation of AC5 and AC7 activity, boosting cAMP production further, contributing to superactivation. We know that AGS3 binding to G_{i3}-GDP, liberates free G_{i3} and have reported that unbound G_{i3} subunits stimulate AC2 and AC4 (Yao et al., 2005). However, during morphine withdrawal, inhibition of G_{i3} function did not inhibit withdrawal-induced cAMP superactivation. Thus, in contrast to short-term morphine-induced increases in cAMP via G_{i3} subunits, morphine withdrawal-induced cAMP superactivation does not seem to require G_{i3} function but does require up-regulation of AGS3 as a central molecular event. Overall, the mechanism of action of morphine withdrawal-induced cAMP superactivation requires up-regulation of AGS3, A2A receptors, AC5, AC7, PKA, and ePKC. A schematic presentation of these molecular mechanisms is presented in Fig. 5.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>cAMP Increase</th>
</tr>
</thead>
<tbody>
<tr>
<td>W.D.</td>
<td>78 ± 9</td>
</tr>
<tr>
<td>W.D. + ADA</td>
<td>2 ± 5*</td>
</tr>
<tr>
<td>W.D. + KW-6002</td>
<td>1 ± 3*</td>
</tr>
<tr>
<td>W.D. + Control SiRNA</td>
<td>71 ± 5</td>
</tr>
<tr>
<td>W.D. + AC2 SiRNA</td>
<td>69 ± 12</td>
</tr>
<tr>
<td>W.D. + AC4 SiRNA</td>
<td>72 ± 10</td>
</tr>
<tr>
<td>W.D. + AC5 SiRNA</td>
<td>24 ± 4*</td>
</tr>
<tr>
<td>W.D. + AC7 SiRNA</td>
<td>49 ± 7*</td>
</tr>
<tr>
<td>W.D. + (AC5 + AC7) SiRNA</td>
<td>11 ± 4*</td>
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* P < 0.05 compared with withdrawal control (one-way ANOVA and Dunnett’s test).

Table 2

<table>
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<tr>
<th>Treatment</th>
<th>cAMP Increase</th>
</tr>
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<tbody>
<tr>
<td>W.D.</td>
<td>75 ± 7</td>
</tr>
<tr>
<td>W.D. + ADA</td>
<td>79 ± 11</td>
</tr>
<tr>
<td>W.D. + KW-6002</td>
<td>70 ± 6</td>
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The role of adenosine A2A receptors in cAMP superactivation 3 h after withdrawal.

NAc/striatal neurons were treated with 100 nM morphine for 18 h. Three hours after withdrawal (W.D.), cells were then treated with or without adenosine deaminase (ADA, 1 unit/ml) or the A2A antagonist KW-6002 (100 nM) for 2 h. Data are the mean ± S.E.M. of at least three experiments with each treatment carried out in triplicates.
PKA activation caused both by short-term exposure to morphine and by withdrawal after long-term exposure to morphine. Thus, knockdown of AGS3 expression completely blocks the increase in cAMP/PKA signaling in both conditions. AGS3 selectively interacts with the GDP-bound conformation of α-subunits of the Gαi/o family, enabling free Gβγ to stimulate AC2 and AC4 (Yao et al., 2005). However, G-protein dissociation is not expected to occur in the absence of morphine. Therefore, free Gβγ released from Gαi/o stimulating AC cannot explain cAMP superactivation during withdrawal from long-term exposure. Indeed, we found that cAMP superactivation, which requires AC5 and AC7, did not require Gβγ function. AGS3 also binds to and inactivates Gαi/o, preventing Gαi/o from restraining AC activation (Takesono et al., 1999; Kimple et al., 2002; Peterson et al., 2002). Because superactivation requires AGS3, it is possible that
up-regulation of AGS3 selectively diminishes signaling through Gαi. Indeed, we found that up-regulation of AGS3 may contribute to morphine withdrawal-induced cAMP superactivation apparently by binding with the Gαi-GDP complex, thereby preventing free Gαi from inhibiting AC.

cAMP activates PKA and causes its translocation from the Golgi to the nucleus, where PKA phosphorylates the cAMP response element binding protein to activate CRE-mediated gene transcription (Chao and Nestler, 2004). It is possible that up-regulation of AGS3 could occur as a result of increased gene transcription. It is noteworthy that Rp-cAMPS, a PKA inhibitor, prevented up-regulation of AGS3, suggesting the possibility that the AGS3 gene may contain CREs regulated by cAMP/PKA, although this is unknown. Despite this limitation, however, it seems that up-regulation of AGS3 by drugs of abuse seems to be another common determinant in addiction biology, possibly related to drug-seeking behavior and locomotor sensitization. Thus, AGS3 antisense prevents cocaine-induced behavioral sensitization and cocaine-, heroin-, and ethanol-seeking behavior (Bowers et al., 2003, 2008; Yao et al., 2005).

Adenosine A2A Receptors. Adenosine has been implicated in responses to several addictive agents such as ethanol (Arolfo et al., 2004), opiates (Brundege and Williams, 2002), and cannabinoids (Soria et al., 2004). The striatum is enriched in postsynaptic adenosine A2A receptors that couple to the stimulatory G-protein Gαs in most cells and primarily Gαolf in brain (Kull et al., 2000). Long-term morphine exposure can increase extracellular adenosine in striatum by up-regulating adenosine transporter binding sites (Kaplan and Leite-Morris, 1997), and adenosine seems to act on A2A. We found that A2A is required for initiating but not sustaining morphine withdrawal-induced cAMP superactivation. The sustained cAMP superactivation seems to require AGS3 up-regulation because treatment with adenosine deaminase or KW-6002 at 3 h after withdrawal, when AGS3 has been up-regulated, did not inhibit cAMP superactivation. This finding is consistent with a role for A2A in contribution to morphine-seeking behaviors. Thus, adenosine receptor antagonists attenuate development of morphine sensitization in mice (Weisberg and Kaplan, 1999) and inhibit morphine self-administration (Sahraei et al., 1999) and heroin-induced reinstatement in rats (Yao et al., 2006).

Adenylyl Cyclases. Nine different AC isoforms have been identified that are regulated by different effectors, such as Gα and Gβγ subunits and protein kinases. AC2, AC4, and AC7 are conditionally stimulated by Gβγ subunits; AC2, AC5, and AC7 are activated by PKC (Watts and Neve, 2005). We previously reported that AC2 and AC4 are stimulated by Gβγ and involved in short-term neural responses to morphine (Yao et al., 2005). By contrast, we show here that AC5 and AC7, not AC2 and AC4, are required for morphine with-
drawal-induced cAMP superactivation. Thus, opiate withdrawal seems to require specific AC isozymes (Avidor-Reiss et al., 1997). AC5 is highly enriched in the striatum (Defer et al., 2000). AC5 knockouts show a dramatic reduction in morphine-induced physical dependence and withdrawal symptoms (Kim et al., 2006). PKC has been implicated in the mechanism of action of addictive drugs (Olive and Messing, 2004). PKC phosphorylates AC5 and enhances basal and forskolin- and Goβγ-stimulated cAMP accumulation (Kawabe et al., 1994). Furthermore, PKC can be activated by PKA via a cross-talk mechanism (Yao et al., 2008). Consistent with these reports, we found that morphine withdrawal activates and translocates εPKC. It is noteworthy that cAMP superactivation is blocked by PKC inhibitors and attenuated by siRNA for AC5 or AC7, suggesting that activation of εPKC may enhance AC5 and AC7 activity contributing to cAMP superactivation during withdrawal.

Figure 5 schematically illustrates a putative model of AGS3 up-regulation and cAMP superactivation. During long-term morphine exposure, both MOR/DOR and A2A modulate basal cAMP levels. Morphine decreases cAMP via Goβγ, whereas adenosine increases cAMP via GoβγH9251. Withdrawal removes the inhibitory effect of morphine and enables A2A-stimulated increases in cAMP to rise above basal levels. This increase in cAMP is sufficient to activate PKA and increase AGS3 expression. Increased AGS3 binds Goβγ/DNP and inactivates Goβ, signaling, further increasing cAMP production. Unrestrained enhancement of cAMP/PKA signaling activates the PLC/PKC system to stimulate AC5 and AC7 and boost cAMP production, ultimately leading to cAMP superactivation. Taken together, up-regulation of AGS3 seems to play a central role in the cooperation of these diverse pathways, facilitating morphine withdrawal-induced cAMP superactivation and, perhaps, underlying some drug-seeking behaviors associated with opiate withdrawal and relapse. It is possible that agents that interfere with AGS3 function might be potential therapies for opiate and other drug addictions.

Acknowledgments

We thank Dr. Daria Mochely-Rosen (Stanford University, Stanford, CA) for PKC peptide inhibitors.

References


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