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Dysregulation of AMPA receptor transmission in the nucleus accumbens in animal models of cocaine addiction

Marina E. Wolf

Department of Neuroscience, Rosalind Franklin University of Medicine and Science, 3333 Green Bay Road, North Chicago, IL 60064-3095 marina.wolf@rosalindfranklin.edu

Abstract

Plasticity of glutamate transmission in neuronal circuits involving the nucleus accumbens (NAc) is now recognized to play a critical role in cocaine addiction. NAc neurons are excited primarily by AMPA-type glutamate receptors (AMPA) and this is required for cocaine seeking. This review will briefly describe AMPAR properties and trafficking, with a focus on studies in NAc neurons, and then consider mechanisms by which cocaine may alter AMPAR transmission. Two examples will be discussed that may be important in two different stages of addiction: learning about drugs and drug-related cues during the period of drug exposure, and persistent vulnerability to craving and relapse after abstinence is achieved. The first example is drawn from studies of cultured NAc neurons. Elevation of DA levels (as would occur following cocaine exposure) facilitates activity-dependent strengthening of excitatory synapses onto medium spiny neurons, the main cell type and projection neuron of the NAc. This occurs because activation of D1-class receptors primes AMPAR for synaptic insertion, creating a temporal window in which stimuli related to cocaine-taking are more efficacious at eliciting synaptic plasticity and thus being encoded into memory. The second example involves rat models of cocaine addiction. Cell surface and synaptic expression of AMPAR on NAc neurons is persistently increased after withdrawal from repeated cocaine exposure. We hypothesize that this increases the reactivity of NAc neurons to glutamate inputs from cortex and limbic structures, facilitating the ability of these inputs to trigger cocaine seeking and thus contributing to the persistent vulnerability to relapse that characterizes addiction.

Keywords

addiction; AMPA receptor; dopamine; glutamate; nucleus accumbens; receptor trafficking; synaptic plasticity

1. Introduction to AMPA-type glutamate receptors and plasticity

1.1 Scope of the review

It is now well accepted that drug addiction involves activity-dependent plasticity at glutamate synapses within neuronal circuits important for motivated behaviors (Wolf et al., 2004; Kauer and Malenka, 2007). This conclusion is based on evidence that drugs of abuse can influence fundamental cellular processes that regulate synaptic strength. Much of this work has focused on the nucleus accumbens (NAc), which occupies a key position in the neural circuitry related to motivation and drug addiction (Kelley, 2004; Kalivas and Volkow, 2005). The majority of NAc neurons (~90%) are medium spiny GABA neurons (MSN; Meredith and Totterdell, 1999). From both anatomical and functional perspectives, the NAc MSN serve as an interface between limbic and cortical brain regions important for regulating motivated behaviors (including drug seeking) and motor regions important for the execution of motivated behaviors (Mogenson, 1987; Groenewegen et al, 1999; Kelley, 1999). Indeed, considerable evidence indicates that activation of NAc MSN, by glutamate

inputs originating from limbic and cortical regions, is the final common pathway for drug seeking in many animal models of addiction (Kalivas and Volkow, 2005) although motor circuitry involving the dorsal striatum becomes more important as drug use becomes habitual (Everitt and Robbins, 2005).

The activation of NAc MSN by glutamate inputs is mediated primarily via AMPAR (Pennartz et al., 1990; Hu and White, 1996). Therefore it is not surprising that drug seeking, elicited by cocaine or cues previously paired with cocaine, depends upon AMPAR transmission in the NAc (Cornish et al., 1999; Cornish and Kalivas, 2000; Di Ciano and Everitt, 2001; Di Ciano and Everitt, 2004; Suto et al., 2004; Kruzich and Xi, 2006; Bäckström and Hyttiä, 2007; Conrad et al., 2008; Famous et al., 2008; Ping et al., 2008). Furthermore, several groups have shown that enhanced AMPAR transmission in the NAc is associated with enhanced drug seeking (Suto et al., 2004; Anderson et al., 2008; Conrad et al., 2008). These results and others suggest that the strength of AMPAR transmission in the NAc represents a critical determinant of the intensity of drug seeking (for a comprehensive review, see Wolf and Ferrario, 2010).

The present review focuses on specific examples of how cocaine, which acts initially by elevating extracellular dopamine (DA) levels, may enhance AMPAR transmission in the NAc. The examples are drawn from our own work (presented at the 2009 Congress of the Fundación Cerebro y Mente) and work from other laboratories. Section 1 will briefly review the role of AMPAR in synaptic plasticity. Most studies on this topic have been conducted in hippocampal neurons, but similar results have been obtained in NAc neurons. In Sections 2 and 3, the focus will be cocaine- or DA-mediated mechanisms that may be important in two different stages of addiction: 1) learning about drugs and drug-related cues during the period of drug exposure, and 2) persistent vulnerability to craving and relapse after abstinence is achieved. Specifically, Section 2 will discuss how cocaine administration, by elevating DA levels, may facilitate synaptic plasticity in NAc neurons. This may help explain why drug-related learning is exceptionally powerful and persistent. Section 3 will discuss studies of in vivo AMPAR plasticity after withdrawal from repeated cocaine exposure. Based on these studies, we hypothesize that upregulation of AMPAR at excitatory synapses onto NAc neurons enhances their response to glutamate inputs that trigger drug craving and relapse. This in turn may contribute to persistent vulnerability to relapse even after long periods of abstinence.

In light of the central role of AMPAR in many forms of experience-dependent plasticity (Kessels and Malinow, 2009), cocaine-induced alterations in AMPAR function are expected to profoundly alter the responsiveness of NAc neurons and their subsequent plasticity. However, it should be emphasized that many other neuroadaptations are critically important in regulating the excitability and output of NAc MSN after cocaine withdrawal. We touch upon a few of these, but the reader is referred to recent reviews for more comprehensive coverage (Kauer and Malenka, 2007; Thomas et al., 2008; Goto and Grace, 2008; Kalivas, 2009; Wheeler and Carelli, 2009).

1.2 AMPAR trafficking and synaptic plasticity

The importance of AMPAR for plasticity was first demonstrated approximately 10 years ago by studies showing that trafficking of AMPAR in and out of synapses determines the level of excitatory synaptic strength and contributes to LTP and LTD (Malinow and Malenka, 2002). It is now appreciated that AMPAR trafficking involves receptor endocytosis and exocytosis, occurring in specialized regions outside of the synapse, coupled with receptor lateral diffusion along the surface of the plasma membrane between synaptic and extrasynaptic compartments (Groc and Choquet, 2006).

Functional AMPAR are tetramers (dimers of dimers) (Mano and Teichberg 1998; Rosenmund et al., 1998; Greger and Esteban, 2007). They are composed of GluA1-4 subunits (formerly termed GluR1-4; see Collingridge et al., 2009). Early information about AMPAR subunit composition was provided by landmark co-immunoprecipitation (co-IP) studies using adult rat hippocampus (CA1/CA2) as starting material (Wentholt et al., 1996). These studies showed that about 80% of GluA1 was pulled down with GluA2/A3 antibody. Together with other results, this indicated that GluA1A2 receptors are an important population. About 30% of the GluA2 was not pulled down by GluA1 antibody. This could represent GluA2 present in GluA2A3 receptors (Wentholt et al., 1996). However, subsequent studies have shown that GluA2 in cultured neurons and rat brain is largely retained in the endoplasmic reticulum (ER) in dimer form whereas GluA1 is mainly found in tetrameric receptors in post-ER compartments (Greger et al., 2003). These findings suggest that “left-over” GluA2 may represent partially assembled receptors, rather than a significant GluA2A3 population. If this is the case, the co-IP results (Wentholt et al., 1996) indicate a predominant role for GluA1A2 receptors, which is consistent with more recent results obtained using a single cell-genetic approach combined with electrophysiological recordings (Lu et al., 2009). This latter study, conducted in hippocampal slices from young mice, found that GluA1A2 receptors comprise ~81% of AMPAR in synapses on CA1 pyramidal neurons and ~95% of receptors in extrasynaptic regions of their soma, whereas GluA2A3 receptors represented ~16% of the synaptic pool and were absent from the extrasynaptic somatic pool.

Subunit composition plays a critical role in determining biophysical properties of the AMPAR (Dingledine et al., 1999; Palmer et al., 2005). For example, absence of the GluA2 subunit results in Ca^{2+} permeability of the AMPAR channel (see below). Subunit composition is also important in determining “rules” that govern AMPAR trafficking. According to the leading theory, AMPAR comprised of subunits with short C-termini (GluA2 or GluA3) cycle constitutively in and out of synapses, whereas AMPAR containing a subunit with a long C-terminus (GluA1 or GluA4; GluA4 is important mainly during early development) are inserted into synapses in an activity-dependent manner, e.g., during LTP (Malinow, 2003). This may reflect differences in regulatory phosphorylation events and protein-protein interactions between subunits with long and short C-terminal tails (Song and Hugarir, 2002; Derkach et al., 2007). AMPAR gating and trafficking are also regulated by AMPAR auxiliary subunits (transmembrane AMPA receptor regulatory proteins, or TARPs, and cornichon proteins) (Milstein and Nicoll, 2008; Schwenk et al., 2009; Tigaret and Choquet, 2009).

Synaptic strength can also be regulated by switching between AMPAR containing the GluA2 subunit (e.g., GluA1A2) and AMPAR lacking the GluA2 subunit (homomeric GluA1, homomeric GluA3 or GluA1A3) (Cull-Candy et al., 2006; Thiagarajan et al., 2007; Liu and Zukin, 2007; Isaac et al., 2007). GluA2-lacking AMPAR are permeable to Ca^{2+} and therefore have higher conductance than GluA2-containing AMPAR. Their higher conductance means that insertion or removal of only a small number of these receptors markedly alters synaptic strength (Liu and Cull-Candy, 2000; Plant et al., 2006; Guire et al., 2008), whereas their Ca^{2+} permeability permits AMPAR transmission to be directly coupled to the many important Ca^{2+} -dependent signaling pathways that govern subsequent plasticity (Thiagarajan et al., 2007). GluA2-lacking AMPAR can be identified experimentally based on inward rectification due to voltage-dependent block by polyamines as well as sensitivity to antagonists that do not block GluA2-containing AMPAR.

Although the cellular mechanisms underlying LTP and LTD vary at different synapses, the best studied form of LTP requires strong synaptic activation, Ca^{2+} entry through the NMDA receptor (NMDAR), and activation of protein kinase dependent cascades that recruit additional AMPAR to the synapse. Synaptic activation leading to LTD is associated with

less Ca^{2+} entry and selective activation of protein phosphatases, leading to AMPAR removal from synapses (Shepherd and Huganir, 2007; Pi and Lisman, 2008). These events occur in the early phase of LTP. There is also a late phase of LTP in which protein synthesis is required to stabilize the increase in synaptic strength (see Abraham and Williams, 2008 for a recent review).

Recently, there has been considerable interest in a homeostatic form of plasticity termed synaptic scaling, in which prolonged activity blockade (typically 1-3 days) leads to augmented excitatory synaptic transmission, whereas prolonged increases in activity produce opposite effects (Turrigiano and Nelson, 2004; Turrigiano, 2008). Synaptic scaling is believed to stabilize activity of neurons and neuronal circuits. For example, scaling up would prevent rundown of synaptic strength after repeated LTD. Although many mechanisms contribute to the expression of synaptic scaling, including presynaptic changes in transmitter release, the major postsynaptic mechanism involves increases in synaptic AMPAR levels in response to activity blockade and decreased AMPAR levels after prolonged increases in activity. Synaptic scaling is a promising candidate mechanism for slowly developing plasticity that occurs after drug withdrawal (Sun and Wolf, 2009; see Section 2.3).

1.3 AMPAR in the NAc

Excitatory synapses in the NAc undergo NMDAR-dependent LTP and several forms of LTD, a topic that has been reviewed recently (Kauer and Malenka, 2007) and therefore will not be addressed here. Instead, this review will focus on studies of AMPAR subunit composition and trafficking in NAc neurons.

To determine the subunit composition of NAc AMPAR, we conducted quantitative co-IP studies identical to those of Wenthold et al. (1996). This section will discuss data obtained using NAc membrane preparations from untreated adult rats as starting material (Reimers et al., submitted), but similar results were found in the NAc of rats with saline self-administration experience (Conrad et al., 2008; see also Boudreau et al., 2007) and in other brain regions (Reimers et al., submitted). In the NAc of untreated rats, we found that ~90% of the GluA1 is physically associated with GluA2 or GluA3, and the majority of this represents GluA1A2. GluA1A3 complexes also exist, but this is a minor population (only ~6% of the GluA3 is pulled down by GluA1 antibody). It is possible that there is also a very small population of homomeric GluA1 receptors (Reimers et al., submitted). These findings are consistent with earlier studies suggesting that most GluA1-containing AMPAR in the striatum and other forebrain regions also contain GluA2 or GluA3 (Bernard et al., 1997; Gold et al., 1997). Turning to the question of GluA2A3 receptors, we found that ~50% of GluA2 was not physically associated with GluA1 and therefore could represent GluA2A3. However, using blue native polyacrylamide gel electrophoresis to assess AMPAR assembly state, we found that a substantial portion of GluA2 was present in dimers or monomers, whereas relatively more GluA1 was present in tetramers (Reimers et al., submitted). These findings are consistent with prior work in cultured neurons and rat brain (Greger et al., 2003; see Section 1.2) and argue that most of the “left-over” GluA2 probably represents partially assembled receptors. Overall, our results suggest a predominant role for GluA1A2 receptors in the NAc (Reimers et al., submitted) and thus fall into line with single-cell genetic/electrophysiological assessments of AMPAR subunit composition in hippocampus (Lu et al., 2009). However, a role for GluA2A3 or GluA1A3 receptors in the NAc should not be ruled out. GluA3 is expressed on the cell surface in the adult rat NAc (Boudreau et al., 2007) and this measure is altered after cocaine self-administration (Conrad et al., 2008). GluA4 is not present in medium spiny neurons (Bernard et al., 1997; Stefani et al., 1998).

Although the co-IP studies described above suggest that a small pool of GluA2-lacking AMPAR is present in NAc neurons (Reimers et al., submitted; Conrad et al., 2008; Boudreau et al., 2007), electrophysiological studies indicate that they play a minimal role in synaptic transmission. Thus, whole-cell patch clamp recordings in the NAc core of adult rats with previous saline self-administration experience found linear AMPAR current-voltage relationships and only a small reduction (~5%) in the evoked excitatory postsynaptic current (EPSC) amplitude after bath application of 1-naphthylacetylsperimine (Naspm), a selective antagonist of GluR2-lacking AMPAR (Conrad et al., 2008). Linear AMPAR current-voltage relationships have also been found in the NAc shell of young mice given saline injections (Kourrich et al., 2007, P24-28; Mameli et al., 2009; P16-35), although a small contribution of GluA2-lacking AMPAR to synaptic transmission in the NAc shell was found in untreated adult mice (Campioni et al., 2009; P56-70).

To study AMPAR trafficking in the NAc, we have used primary cultures prepared from the NAc of postnatal day 1 rats and co-cultures in which prefrontal cortex (PFC) neurons are included to restore excitatory synapses onto NAc MSN. Despite the fact that NAc MSN are GABA neurons, AMPAR trafficking was similar to what has been described in hippocampal pyramidal neurons. Thus, brief exposure to glutamate agonists produces a decrease in AMPAR surface expression in cultured NAc neurons (Mangiavacchi and Wolf, 2004b) akin to that implicated in LTD in other brain regions (Carroll et al., 2001). We have also described the activity-dependent synaptic insertion of GluA1-containing AMPAR in cultured NAc neurons. This occurs through a two-step process that involves externalization onto the cell surface at extrasynaptic sites followed by translocation into the synapse (Fig. 1). The first step is accelerated by protein kinase A (PKA), most likely via phosphorylation of GluA1 at serine 845 (a PKA phosphorylation site), whereas the second step requires NMDAR stimulation (Chao et al., 2002a,b; Mangiavacchi and Wolf, 2004a; Sun et al., 2008). A similar two-step mechanism occurs in other cell types (Passafaro et al., 2001; Esteban et al., 2003; Sun et al., 2005; Gao et al., 2006; Oh et al., 2006; Man et al., 2007; Yudowski et al., 2007). Recent studies have provided additional information about this process. Petrini et al. (2009) demonstrated that AMPAR recycling via postsynaptic endocytic zones is crucial for maintaining a mobile pool of extrasynaptic AMPAR that can be inserted into synapses in an activity-dependent manner to increase synaptic strength. Furthermore, in addition to the important priming role noted above for PKA phosphorylation of serine 845, it has been demonstrated that protein kinase C phosphorylation of serine 816 and serine 818 of GluA1 enhances the interaction with its binding partner 4.1N and thus enhances insertion of GluA1 into surface extrasynaptic pools (Lin et al., 2009).

While most studies of synaptic plasticity in NAc neurons have focused on LTP or LTD, we showed recently that NAc MSN also exhibit bidirectional synaptic scaling. Prolonged activity blockade [1-3 days of incubation with tetrodotoxin (TTX) or the AMPAR antagonist CNQX] leads to increased levels of GluA1A2-containing AMPAR on the surface of MSN, both at synaptic and extrasynaptic locations, whereas prolonged increases in activity (produced by incubating with the GABA_A receptor antagonist bicuculline) lead to opposite effects. Increased AMPAR surface expression after prolonged activity blockade requires protein synthesis and is occluded by inhibition of the ubiquitin-proteasome system, perhaps indicating a requirement for new receptors as well as slowing of the turnover of existing receptors (Sun and Wolf, 2009). It is possible that synaptic scaling contributes to the increased AMPAR surface and synaptic expression that is observed in the rodent NAc after withdrawal from repeated cocaine exposure (see Section 3). Imaging studies in humans and primates have demonstrated persistent metabolic hypoactivity after cocaine exposure in cortical areas that send glutamate projections to the NAc (Goldstein and Volkow, 2002; Porrino et al., 2007), which could translate into decreased excitatory transmission onto NAc neurons after cocaine withdrawal. We have speculated that this leads to synaptic scaling in

NAc neurons, resulting in the accumulation of AMPAR at excitatory synapses onto these neurons and explaining in vivo results indicating AMPAR upregulation after cocaine withdrawal (Section 3). For discussion of rodent data that support this hypothesis, see Sun and Wolf (2009).

2. Facilitation of AMPAR synaptic insertion by DA may contribute to the strength of drug-related learning

2.1 D1-class receptors prime AMPAR for synaptic insertion

The insertion of new AMPAR into synapses is responsible for increasing synaptic strength in many forms of experience-dependent plasticity and learning (Kessels and Malinow, 2009). As discussed in more detail in Section 2.2, one important role of DA is to facilitate learning about natural rewards and drugs of abuse. A very simple way to accomplish this would be to directly facilitate AMPAR synaptic insertion at relevant synapses. This mechanism is plausible in several brain regions - including the dorsal striatum, NAc (ventral striatum), PFC and hippocampus - where dendrites of the principal neurons receive convergent DA and glutamate inputs, forming a synaptic triad (Sesack et al., 2003; Fig. 1). Within the triad, DA receptors are well-positioned to modulate plasticity mechanisms, including AMPAR trafficking, at nearby glutamate synapses.

We tested this idea in cultured neurons. Our initial studies used primary cultures prepared from postnatal NAc neurons. They revealed that stimulation of D1-class receptors, which are positively coupled to adenylyl cyclase, increase the rate of externalization of GluA1-containing AMPAR to the cell surface, in concert with stimulation of GluA1 phosphorylation at serine 845 (Chao et al., 2002a,b; Mangiavacchi and Wolf, 2004a; we are using the term D1-class because pharmacological agents used in these studies do not distinguish between D1 and D5 receptors). We could not study AMPAR synaptic targeting in “pure” NAc cultures, because the NAc does not contain intrinsic glutamate neurons and thus cultures prepared from the NAc do not contain glutamate synapses. To overcome this, we restored excitatory synapses onto the NAc neurons by co-culturing them with PFC neurons (Sun et al., 2008). We also studied primary cultures prepared from postnatal rat hippocampus (Gao et al., 2006) and PFC (Sun et al., 2005), brain regions that contain glutamate neurons and therefore yield cultures containing glutamate synapses. In all neurons studied (NAc MSN, hippocampal pyramidal neurons, and PFC pyramidal neurons), we found that brief (5-15 min) stimulation of D1-class receptors increased GluA1 surface expression through a mechanism requiring PKA activation (Sun et al., 2005; Gao et al., 2006; Sun et al., 2008) and the secretory pathway (Gao et al., 2006). Protein synthesis was not required in our studies (Gao et al., 2006; Sun et al., 2008), although D1-class agonists can also increase synaptic GluA1 levels in hippocampal neurons through a mechanism that involves stimulation of dendritic protein synthesis (Smith et al., 2005). Interestingly, in all cases, we found that D1-class receptor stimulation increased AMPAR surface expression exclusively at extrasynaptic sites; no changes in synaptic AMPAR levels were observed (Sun et al., 2005; Gao et al., 2006; Sun et al., 2008). These results indicated that stimulation of D1-class receptor/PKA signaling is not sufficient for AMPAR synaptic insertion, at least not under our experimental conditions (see next paragraph). However, we hypothesized that D1-class receptor stimulation, by increasing extrasynaptic AMPAR levels, might prime AMPAR for synaptic insertion. To test this, we took advantage of the ability of glycine, an obligatory co-agonist at the NMDAR, to elicit AMPAR synaptic insertion and LTP (Lu et al., 2001). We identified a concentration of glycine that was subthreshold for this effect. If cultures were incubated briefly with D1 agonist followed immediately by 3 min incubation with glycine, we observed that this subthreshold concentration of glycine was able to produce AMPAR synaptic incorporation. These results demonstrate that D1-class receptors

can facilitate activity-dependent synaptic plasticity (Sun et al., 2005; Gao et al., 2006; Sun et al., 2005).

An implication of these results is that D1-class receptor agonists will only facilitate AMPAR synaptic insertion under conditions of sufficient NMDAR tone. This may explain some electrophysiological results in NAc neurons that do not support this mechanism. For example, Nicola et al. (1996) found no evidence for D1-class receptor mediated enhancement of postsynaptic AMPAR transmission, but their recordings were conducted in brain slices at membrane potentials that prohibit NMDAR transmission. In contrast, when NAc are activated by excitatory inputs in vivo, it is likely that that NMDAR tone would be sufficient to observe the priming effect. However, many factors are no doubt important in determining the in vivo consequences of D1-class receptor stimulation, including effects on voltage-gated conductances (e.g., Nicola et al., 2000; Zhang et al., 1998; Zhang et al., 2002) and the existence of presynaptic D1-class receptors in the NAc that dampen excitatory synaptic transmission (e.g., Pennartz et al., 1992; Harvey and Lacey, 1996; Nicola et al., 1996).

In conclusion, our results in cultured NAc, PFC and hippocampal neurons provided some of the first direct support for the two-step process of GluA1 synaptic incorporation described in Section 1.3 and demonstrated that D1-class receptors prime AMPAR for synaptic insertion by accelerating the first step (PKA-dependent externalization onto extrasynaptic regions of the cell surface) and thus increasing the number of AMPAR available for translocation into synapses when NMDAR are subsequently stimulated (see Fig. 1 for summary). Facilitation of AMPAR synaptic insertion by D1-class receptors was blocked by KN-62, an inhibitor of Ca^{2+} -calmodulin-dependent protein kinases (Gao et al., 2006). This is consistent with the fact that AMPAR synaptic insertion during LTP ultimately requires activation of Ca^{2+} -calmodulin-dependent protein kinase II (CaMKII) (Lisman et al., 2002). Overall, our results indicate that the D1-class receptor/PKA pathway works cooperatively with the NMDAR/CaMKII pathway to induce AMPAR synaptic incorporation, thereby facilitating early phases of synaptic plasticity. Consistent with this, activation of D1-class receptors has been shown to augment the magnitude of early LTP in CA1 slices (Otmakhova and Lisman, 1996). This role of D1-class receptor/PKA signaling is distinct, but complementary, to its well established role in the late protein synthesis-dependent phase of LTP (Frey et al., 1990, 1991, 1993; Huang and Kandel, 1995; Matthies et al., 1997; Swanson-Park et al., 1999; Duffy and Nguyen, 2003; Morris et al., 2003).

D1-class receptor stimulation also increased GluR1 synaptic incorporation in VTA DA neurons co-cultured with PFC neurons to restore excitatory synapses, although the mechanism differed from that described above for NAc, PFC and hippocampus (Gao and Wolf, 2007). The VTA DA neurons responded to acute PKA activation with increased GluR1 surface expression, as expected from our prior results (above). However, D1-class receptors are not present on their surface (at least in our culture system), and thus D1-class receptor agonists did not reproduce this effect of PKA activation. Instead, we found that brief stimulation with a D1-class agonist (10 min) increased GluA1 and GluA2 surface expression through a glutamate receptor-dependent pathway, apparently by acting on D1-class receptors on the PFC neurons in the co-culture and thereby altering their excitatory transmission onto VTA DA neurons. We also tested the effect of a longer incubation with DA (1 hr), to mimic the duration of elevated DA levels produced by systemic cocaine administration. In this case, we observed an increase in surface and synaptic GluA1 but not GluA2 on VTA DA neurons, suggesting an increase in GluA2-lacking AMPAR. Surface GluA3 and the area of GluA1A3 colocalization were also increased (Gao and Wolf, 2007). Due to the higher conductance of GluA2-lacking AMPAR, their addition to VTA DA neurons would be expected to significantly enhance DA neuronal activation by glutamate

inputs. These studies provide an *in vitro* model for the cocaine-induced potentiation of excitatory transmission onto VTA DA neurons that contributes to the initiation of behavioral sensitization (Ungless et al., 2001). However, although there is evidence that GluA2-lacking AMPAR contribute to this potentiation (Bellone and Lüscher, 2006), the *in vivo* phenomenon is complex and cannot be fully explained by synaptic addition of GluA2-lacking AMPAR (see Kauer and Malenka, 2007).

2.2 Significance of the priming effect of D1-class receptors for motivated behavior

This section addresses the possible behavioral significance of the ability of D1-class receptors to prime AMPAR for synaptic insertion in principal neurons of the PFC (Sun et al., 2005), hippocampus (Gao et al., 2006) and NAc (Sun et al., 2008). Although the role of DA in motivation and reward is very complex, one idea is that DA transmission facilitates associative learning about natural rewards and drugs of abuse that is important for shaping future behavior (Di Chiara, 1998; Berke and Hyman, 2000; Schultz, 2000; Kelly and Berridge, 2002; Lisman and Grace, 2005; Wheeler and Carelli, 2009). Our results suggest that DA may enhance learning about reward-related stimuli at least in part by priming AMPAR for synaptic insertion and thereby facilitating synaptic plasticity. Thus, when a stimulus is presented within a temporal window in which DA levels are elevated, that stimulus is more likely to be encoded as an increase in the strength of excitatory synapses. This mechanism may help explain many results obtained in NAc, PFC and hippocampal neurons indicating that D1-class receptors facilitate LTP and learning (Jay, 2003; O'Donnell, 2003; Lisman and Grace, 2005). For example, Li et al. (2003) found that brief exposure to a novel environment, leading to D1 receptor-PKA signaling, enhanced the ability of a weak tetanus to induce LTP in hippocampal CA1 neurons. This observation is readily explained by a priming effect of D1-class receptor stimulation (Fig. 1), although it is important to note that DA also modulates other cellular mechanisms that regulate neuronal excitability and plasticity, including voltage-gated ion channels (e.g., Seamans and Yang, 2004) and NMDAR transmission (Cepeda and Levine, 2006) (see Section 2.1 for more specific examples related to D1-class receptors in the NAc).

A similar priming effect has been demonstrated after stimulating hippocampal β -adrenergic receptors (which like D1-class receptors are positively coupled to adenylyl cyclase) with norepinephrine, a transmitter important for stress responses. Norepinephrine produced a robust increase in GluA1 phosphorylation at serine 845 (PKA phosphorylation site), and to a lesser extent at serine 831 (CaMKII or protein kinase C phosphorylation site), an effect that was reproduced by emotional stress (Hu et al., 2007). Through this mechanism, norepinephrine facilitated GluA1 synaptic delivery and LTP (Hu et al., 2007). This is proposed to contribute to stress-related learning (Krugers and Hoogenraad, 2009). More broadly, the ability to promote plasticity by facilitating AMPAR synaptic delivery may be a shared property of neuromodulator receptors that are positively coupled to adenylyl cyclase (see Seol et al., 2007).

In contrast to natural rewards, psychomotor stimulants like cocaine, by blocking the DA transporter, produce robust and unregulated increases in extracellular DA levels. During repeated cocaine exposure, unregulated DA receptor signaling may lead to inappropriate modulation of AMPAR trafficking and abnormal synaptic plasticity. These processes may contribute to the rewiring of neuronal circuits that underlies the transition to compulsive drug use. More specifically, we speculate that unregulated DA release promotes very strong learning about events that occur under the influence of the drug, when DA levels are high, including cues and context that are associated with the drug-taking experience. As cues are important triggers for drug craving, this pathologically strong learning may explain the persistent vulnerability to relapse that makes addiction so difficult to treat. It is interesting to

speculate that learning related to the negative consequences that follow the act of drug-taking (e.g., family disapproval or loss of employment) is much less effective because DA levels are low when the negative consequences are experienced.

2.3 Long-term effects of DA receptor stimulation

The studies described in Section 2.2 focused on acute effects of DA receptor stimulation on AMPAR trafficking. We wondered whether long-lasting effects also occur. This is difficult to address using primary neuronal cultures because there is a limited time window for experimentation. Typically, at least a week *in vitro* is required for synaptic connections to be established, but cultures deteriorate after about 3 weeks (although this depends on cell density). Within this window, we attempted to mimic repeated cocaine exposure by treating NAc/PFC co-cultures repeatedly with DA (30 min per day on days 7, 9 and 11 *in vitro*). Two interesting effects were observed 4 days after the last DA exposure (day 15). First, D1-class agonist treatment no longer increased AMPAR surface expression. This refractoriness was associated with decreased D1 receptor surface expression (Sun et al., 2008). Second, upregulation of GluA1 and GluA2 surface expression on NAc MSN was detected on day 15 (Sun et al., 2008; Sun and Wolf, 2009). This effect required CaMK activity during the 4 day “withdrawal” period and was associated with CaMKII activation (Sun et al., 2008). Obviously cultures do not reproduce the circuitry of the brain and are prepared from young neurons, which differ from adult neurons in many respects. For example, unlike adult rodent NAc neurons (Section 1.3), cultured NAc neurons, prepared from P1 rats, express a substantial number of GluA2-lacking AMPAR under basal conditions (Sun and Wolf, 2009). Therefore, caution must be used in extrapolating these and other *in vitro* results to the adult brain. However, our results in NAc/PFC co-cultures indicate that repeated DA treatment has effects on plasticity mechanisms that outlast the period in which DA levels are elevated, perhaps via changes in excitatory activity that are triggered by prior DA receptor overstimulation. Specifically, these results are reminiscent of the increased GluA1 and GluA2 surface expression that we have observed in the NAc of cocaine-sensitized rats (see Section 3.2). These rats show a transient elevation of CaMKII levels in the NAc on withdrawal day 7 (Boudreau et al., 2009), suggesting another parallel to the *in vitro* findings discussed here.

Interestingly, we found that the upregulation of surface GluA1 and GluA2 in cultured NAc neurons produced by repeated DA treatment occluded subsequent synaptic scaling induced by prolonged activity blockade, i.e., no further increase in AMPAR surface expression was produced by activity blockade when it was preceded by repeated DA treatment (Sun and Wolf, 2009). If this occlusion occurs *in vivo*, it would argue against the hypothesis that synaptic scaling is responsible for AMPAR upregulation in the NAc after withdrawal from repeated *in vivo* cocaine exposure (see Section 1.3). On the other hand, if the occlusion of scaling occurs *in vivo* but is transient, it could help explain the delayed onset of AMPAR upregulation after cocaine withdrawal (see Section 3). Interestingly, a different kind of homeostatic plasticity, in which MSN adjust their membrane excitability to functionally compensate for changes in the level of excitatory synaptic input, shows a prolonged impairment in the NAc of cocaine-sensitized rats (Ishikawa et al., 2009).

3. AMPAR upregulation after cocaine withdrawal may contribute to incentive sensitization

3.1 Overview

AMPA transmission in the NAc has been studied after both non-contingent and contingent cocaine administration. When cocaine is administered by the experimenter, this is referred to as “non-contingent” because drug administration is not dependent upon the animal's

behavior. In cocaine self-administration procedures, drug administration is dependent on the rat's response, such as a nose poke or lever press, and is therefore "contingent." As described below, we and others have observed increases in AMPAR surface and synaptic expression in the NAc following withdrawal from repeated non-contingent cocaine injections leading to behavioral sensitization (Section 3.2) and cocaine self-administration leading to the incubation of cocaine craving (Section 3.3), although the AMPAR subtype involved differs in these two situations. It is important to note that these increases do not occur during cocaine exposure but require a withdrawal period to become evident. Thus, they do not result from direct effects of elevated DA levels on mechanisms that control AMPAR trafficking. Instead, just as changes in the level of presynaptic activity can trigger LTP, LTD or synaptic scaling (Section 1.2), we speculate that changes in the activity of glutamate pathways terminating in the NAc are responsible for this withdrawal-dependent plasticity. At present, the nature of such changes is not known. Candidate mechanisms include altered presynaptic glutamate transmission due to impaired cystine-glutamate transport (Kalivas, 2009) or hypoactivity of cortical areas that send glutamate projections to the NAc (Goldstein and Volkow, 2002; Porrino et al., 2007). Another factor that cannot be neglected is "whole-cell plasticity" related to drug-induced changes in voltage-gated ion channels (Zhang et al., 1998, 2002; Hu et al., 2004, 2005; Dong et al., 2006; Ishikawa et al., 2009; Kourrich and Thomas, 2009; Mu et al., 2010). For discussion of possible interactions between these phenomena and postsynaptic AMPAR plasticity, see Wolf and Ferrario (2010).

3.2 AMPAR plasticity in behavioral sensitization

Increased AMPAR surface expression in the NAc after withdrawal from repeated cocaine exposure was first observed in the behavioral sensitization model. Behavioral sensitization refers to the progressive augmentation of behavioral responses to a drug that develops during repeated drug treatment and then persists long after drug exposure is discontinued. Sensitization of locomotor stimulatory effects is commonly studied. In addition, sensitization develops to the incentive-motivational properties of drugs, in other words, the properties that make them "wanted" (Robinson and Berridge, 2008). Thus, rats previously sensitized to psychomotor stimulants will subsequently show enhanced motivation to obtain drugs compared to control rats (Vezina, 2004), just as an addict will continue to seek drugs even in the face of adverse consequences.

Our studies were conducted using adult rats and initially employed a cocaine regimen that produces locomotor sensitization in about half the cocaine-treated rats. Using a protein crosslinking assay that can distinguish between cell surface and intracellular pools of AMPAR subunits (Boudreau and Wolf, 2005), we found that sensitized rats, but not cocaine-treated rats that failed to sensitize, exhibit a withdrawal-dependent increase in AMPAR surface expression in the NAc. This was detected after 7, 14 or 21 days of withdrawal, but not on the first day of withdrawal, indicating that AMPAR cell surface expression increases sometime during the first week of withdrawal (Boudreau and Wolf, 2005; Boudreau et al., 2007). More recently, we have used a cocaine regimen that produces sensitization in all rats, and found that all rats treated with this regimen exhibit increased AMPAR surface expression in the NAc on withdrawal day 14 (Ferrario et al., 2010).

Evidence for AMPAR upregulation has also been obtained with other approaches. Electrophysiological studies found an increased AMPA/NMDA ratio in the NAc of cocaine-sensitized young mice after 14 days of withdrawal (Kourrich et al., 2007). A more recent study found that synaptic strength was increased on withdrawal day 35 due to the addition of GluA2-lacking AMPAR (Mameli et al., 2009; see next paragraph for more discussion). Biochemical studies found increased GluA1 and GluA2 levels in synaptosomal membrane fractions prepared from the NAc of sensitized rats after 21 days but not 1 day of withdrawal (Ghasemzadeh et al., 2009a; Schumann and Yaka, 2009), while increased GluA1 in tissue

homogenates was reported after 1-3 weeks but not 1 day of withdrawal (Churchill et al., 1999; Scheggi et al., 2002). Finally, the locomotor response elicited by infusion of AMPA into the NAc is enhanced after withdrawal from a sensitizing cocaine regimen (Pierce et al., 1996; Bell and Kalivas, 1996; Bachtell and Self, 2008). The cellular mechanisms that trigger and maintain AMPAR upregulation in cocaine-sensitized rats remain unclear, but NMDAR stimulation and ERK activation during withdrawal are implicated (Boudreau et al., 2007; Boudreau et al., 2009; Schumann and Yaka, 2009; Huang et al., 1999). CaMKII and PKA may also be involved (Boudreau et al., 2009). Interestingly, despite the fact that both cocaine and amphetamine elevate DA transmission in the NAc, sensitization to these drugs is associated with different effects on postsynaptic AMPAR transmission (see Ferrario and Wolf, 2010).

The NAc is divided into two subregions, core and shell, that can be distinguished based on morphology, connectivity, and functional role (Meredith et al., 2008). While our studies and others (Schumann and Yaka, 2009) used a combined core/shell dissection, other studies examined these subregions separately and obtained results supporting the idea that AMPAR upregulation occurs in both subregions (Pierce et al., 1996; Bell and Kalivas, 1996; Kourrich et al., 2007; Mameli et al., 2009; Ghasemzadeh et al., 2009a). Differences and similarities between core and shell subregions are beyond the scope of this review, although AMPAR transmission in both regions is implicated in drug seeking behavior in cocaine-experienced rats (see Wolf and Ferrario, 2010).

Most results suggest that GluA1A2-containing AMPAR, rather than GluA2-lacking AMPAR, increase after withdrawal from non-contingent cocaine treatment leading to sensitization. Thus, both GluA1 and GluA2 subunits show increased surface expression (Boudreau and Wolf, 2005; Boudreau et al., 2007) and enrichment in synaptosomal membrane fractions (Ghasemzadeh et al., 2009a) on withdrawal days 14-21, and electrophysiological studies found no evidence for GluA2-lacking AMPAR in the shell of cocaine-sensitized mice on withdrawal day 14 (Kourrich et al., 2007). This is in contrast to results in the incubation model (see Section 3.3). In this model, we observed addition of GluA2-lacking AMPAR to NAc synapses in the core subregion after 42-47 days of withdrawal from extended access cocaine self-administration, and further showed that this underlies the withdrawal-dependent increase in cue-induced cocaine seeking (Conrad et al., 2008). Why did GluA1A2 receptors increase in the sensitization model while GluA2-lacking AMPAR increased in the incubation model? We hypothesized that this was due to different patterns of neuronal activity in pathways projecting to the NAc as a result of non-contingent (sensitization) versus contingent (incubation) cocaine exposure. However, Mameli et al. (2009) found evidence for GluA2-lacking AMPAR in the NAc of cocaine-treated mice after 35 days of withdrawal regardless of whether they received 10 daily i.p. cocaine injections or 8 days of cocaine self-administration (4 h/day). This suggests that a long withdrawal period leads to synaptic incorporation of GluA2-lacking AMPAR regardless of whether cocaine administration is contingent or non-contingent, and raises the possibility that GluA1A2-containing AMPAR increase during the first weeks of withdrawal and are then replaced by GluA2-lacking AMPAR. Alternatively, the results of Mameli et al. (2009) in the sensitization model may be related to the age of the mice when cocaine treatment began (P16-35). This is an important point, since there are developmental changes in the prevalence of GluA2-lacking AMPAR (e.g., Ho et al., 2007) and plasticity mechanisms (McCutcheon and Marinelli, 2009).

What is the functional significance of AMPAR upregulation in the NAc of cocaine-sensitized rats? Some results indicate that enhanced AMPAR transmission in the NAc mediates the expression of locomotor sensitization (Pierce et al., 1996; Bell et al., 2000). On the other hand, several dissociations have been observed between the level of AMPAR

transmission and the expression of sensitization (Bachtell and Self, 2008; Bachtell et al., 2008; Ferrario et al., 2010). For example, locomotor sensitization is expressed on withdrawal day 1, yet AMPAR surface expression on withdrawal day 1 does not differ between cocaine-sensitized and saline-treated rats (Boudreau and Wolf, 2005). Furthermore, if cocaine-sensitized rodents are administered a challenge injection of cocaine, AMPAR surface and synaptic expression are decreased 24 h later (Thomas et al., 2001; Boudreau et al., 2007; Kourrich et al., 2007). In other words, the AMPAR upregulation that occurs after withdrawal is reversed by cocaine re-exposure, although both behavioral and protein crosslinking results suggest that upregulation is restored within 7 days (Bachtell and Self, 2008; Ferrario et al., 2010). However, if a second cocaine challenge is administered 24 h after the first, that is, when AMPAR surface and synaptic expression have been decreased by the first cocaine challenge, the magnitude of locomotor sensitization is not decreased (Ferrario et al., 2010). In fact, Bachtell and Self (2008) observed that it was enhanced. These and other results suggest that treatments which decrease AMPAR transmission in the NAc do not necessarily decrease the magnitude of the expression of locomotor sensitization to cocaine. Finally, results obtained after transient viral-mediated over-expression of wild-type GluA1 or a pore-dead GluA1 mutant suggest that the magnitude of sensitization varies inversely with the strength of AMPAR transmission (Bachtell et al., 2008). Related to this, one theory holds that rapid AMPAR internalization following drug challenge mediates the expression of locomotor sensitization (Brebner et al., 2005), although this is based on studies of amphetamine, which has very different effects on NAc AMPAR compared to cocaine (Nelson et al., 2009; Wolf and Ferrario, 2010). Arguing against this theory as it applies to cocaine, we have not observed any change in AMPAR surface expression in the NAc 30 min after a cocaine challenge injection is administered to sensitized rats (Ferrario et al., 2010). In conclusion, some evidence suggests that enhanced AMPAR transmission in the NAc underlies locomotor sensitization, but other studies suggest no clear relationship or even the opposite relationship.

In contrast, nearly all evidence supports the idea that cocaine seeking behavior, measured in a number of different animal models, requires activation of AMPAR on NAc neurons by glutamate afferents originating from PFC or limbic regions (Cornish and Kalivas, 2000; Vorel et al., 2001; Di Ciano and Everitt, 2001; 2004; Park et al., 2002; McFarland et al., 2003; Hayes et al., 2003; Ito et al., 2004; Di Ciano et al., 2007; Bäckström & Hyttiä, 2007; Conrad et al., 2008; Famous et al., 2008; Ping et al., 2008; Suto et al., 2009; Sari et al., 2009). Furthermore, several studies have observed a positive correlation between the magnitude of drug seeking behavior and the level of AMPAR transmission in the NAc (Suto et al., 2004; Conrad et al., 2008; Anderson et al., 2008; see Section 3.3 for more discussion). As noted above, it is well established that rats administered non-contingent psychomotor stimulant treatment leading to locomotor sensitization will subsequently show enhanced acquisition of drug self-administration and will work harder to obtain the drug in progressive ratio experiments (Vezina, 2004). Given that cocaine seeking seems to depend on level of AMPAR transmission, we propose that prior sensitization, leading to AMPAR upregulation in the NAc, is responsible for this subsequent enhancement of drug taking and seeking. In other words, it is possible that AMPAR upregulation after non-contingent cocaine exposure is more directly related to incentive sensitization than to locomotor sensitization (see Wolf and Ferrario, 2010 for more discussion).

There are studies that do not support a positive relationship between drug seeking and AMPAR transmission in the NAc. These involve experiments in which GluA1 has been over-expressed or deleted (Mead et al., 2007; Bachtell et al., 2008), situations in which compensatory changes may affect the outcome, or experiments in which extinction training followed cocaine self-administration (Sutton et al., 2003). Extinction training involves new learning (Self et al., 2004) and therefore, not surprisingly, different AMPAR changes are

observed in the NAc if cocaine self-administration is followed by extinction training than if it is followed by withdrawal in home cages (Sutton et al., 2003; Ghasemzadeh et al., 2009b). Recognition of this difference may resolve at least some of the apparent discrepancies in this literature (Wolf and Ferrario, 2010).

Our discussion has not attempted to incorporate literature on the dorsal striatum. However, this region is also important for locomotor sensitization and shares some sensitization-related neuroadaptations with the NAc (Patrick et al., 1991; Robinson and Kolb, 2004; Jedynak et al., 2007). Yet, cocaine sensitization is associated with different and in some respects opposite changes in AMPAR distribution in the dorsolateral striatum versus the NAc (Ferrario et al., 2010). This may argue further for dissociation between locomotor sensitization and the AMPAR redistribution observed in the NAc. However, AMPAR plasticity, albeit different from that induced by cocaine, may nevertheless contribute to amphetamine sensitization (e.g., Loweth et al., 2010).

3.3 AMPAR plasticity in the incubation model

After extended access cocaine self-administration (e.g., 6 hrs/day), there is a progressive increase in cue-induced cocaine seeking over the first weeks to months of withdrawal that has been termed “incubation” (Neisewander et al., 2000; Grimm et al., 2001; Lu et al., 2004a,b; Sorge & Stewart, 2005). We showed that GluA2-lacking AMPAR are added to NAc synapses in association with the incubation of cue-induced cocaine seeking (Conrad et al., 2008). Thus, they were not found in NAc synapses of drug-naïve rats or cocaine-experienced rats evaluated on withdrawal day 1, when cue-induced craving is low. However, on withdrawal days 42-47, when cue-induced craving is high, GluA2-lacking AMPAR were present. This was demonstrated using a variety of approaches, including protein crosslinking studies showing selective increases in surface GluA1, quantitative co-IP studies showing an increase in the portion of GluA1 not physically associated with GluA2, and electrophysiological studies showing inwardly rectifying and Nasp-sensitive AMPAR EPSC (Conrad et al., 2008). Consistent with our results, inwardly rectifying AMPAR EPSC were demonstrated in the mouse NAc shell on withdrawal day 35 from cocaine self-administration (4 h/day for 8 days; Mameli et al., 2009). This represents a dramatic change in the synaptic function of MSN due to the Ca^{2+} permeability and resulting higher conductance of GluA2-lacking AMPAR (Section 1.2). Finally, we showed that blocking GluA2-lacking AMPAR, by injecting Nasp into the NAc core, prevented the expression of incubated cue-induced cocaine seeking on withdrawal day 45 (Conrad et al., 2008). Together, these results may suggest GluA2-lacking AMPAR as a target for the design of anti-craving medications.

Cocaine seeking requires activation of glutamate inputs to the NAc and a resultant increase in AMPAR transmission onto MSN (see Section 3.2). Thus, the results described above (Conrad et al., 2008; Mameli et al., 2009) suggest that MSN become more responsive to these glutamate inputs after prolonged withdrawal from cocaine self-administration due to the synaptic incorporation of high conductance GluA2-lacking AMPAR. Thus, when a cue previously paired with cocaine is presented and glutamate is released in the NAc, MSN respond more strongly and cue-induced craving is enhanced. These ideas are consistent with electrophysiological data showing that incubation of cue-induced cocaine seeking is accompanied by an increase in the portion of NAc neurons exhibiting phasic activation in response to cocaine self-administration or cocaine-associated cues, as well as an increase in the strength of neuronal activation (Hollander and Carelli, 2005, 2007). In addition to enabling stronger depolarization of NAc neurons due to their higher conductance, the synaptic incorporation of GluA2-lacking AMPAR after prolonged withdrawal from cocaine self-administration would be predicted to qualitatively alter synaptic transmission and subsequent plasticity at NAc synapses due to their ability to pass Ca^{2+} ions and thus activate

Ca²⁺-dependent signaling pathways. It should be noted that the full circuitry responsible for the expression of incubated cocaine craving has not been worked out. We have focused in this review on NAc, but prior work has established a critical role for excitatory transmission in the central amygdala (Lu et al., 2005, 2007).

Results in the incubation model show that AMPAR transmission in the NAc is enhanced after withdrawal from extended access cocaine self-administration. What about limited access cocaine self-administration (e.g., 2 h/day)? Although AMPAR transmission in the NAc is required for cocaine seeking after such regimens (e.g., Cornish and Kalivas, 2000), cell surface or synaptic AMPAR levels have not been measured. It will be important to determine if AMPAR upregulation occurs and, if so, what type of AMPAR is involved. Interestingly, withdrawal from limited access cocaine self-administration is associated with potentiation of extracellular field potentials in the NA core evoked by *in vivo* stimulation of the PFC, which could indicate AMPAR upregulation (Moussawi et al., 2009). Furthermore, there is evidence that a rapid increase in NAc cell surface levels of GluAR1-containing AMPAR may contribute to cocaine-induced reinstatement of drug seeking following limited access cocaine self-administration and extinction training. D1 receptors, L-type Ca²⁺ channels, and CaMKII activation were implicated in this effect (Anderson et al., 2008). These findings could be related to our demonstration that D1 receptor stimulation facilitates AMPAR synaptic insertion in cultured NAc neurons (Sun et al., 2008; see Section 2.1).

Another important challenge is to understand the relationship between altered AMPAR trafficking and observations of altered LTP or LTD at NAc synapses after cocaine withdrawal (Thomas et al., 2001; Yao et al., 2004; Goto and Grace, 2005; Martin et al., 2006; Mameli et al., 2009; Moussawi et al., 2009). Overall, these studies indicate that repeated cocaine exposure impairs normal plasticity mechanisms in the NAc. However, some divergent results have been reported, probably because these studies differ with respect to whether drug exposure is contingent or non-contingent, whether extinction training has occurred, and withdrawal time (see Wolf and Ferrario, 2010 for more discussion).

3.4 Global versus ensemble-specific neuroadaptations

The upregulation of NAc AMPAR after sensitization and incubation is likely to occur in many or all NAc neurons, based on the ability to detect changes in tissue homogenates (e.g., Boudreau and Wolf, 2005; Conrad et al., 2008). Furthermore, in our electrophysiological studies in the incubation model, we detected inwardly rectifying AMPAR-EPSC in all MSN recorded from cocaine-experienced rats after 30-47 days of withdrawal (Conrad et al., 2008 and unpublished findings of Kuei-Yuan Tseng and Marina Wolf). Other cocaine-induced neuroadaptations that influence MSN output are also likely to affect a large number of MSN, such as decreased extracellular glutamate levels (Kalivas, 2009) and decreased intrinsic excitability of MSN (Zhang et al., 1998, 2002; Hu et al., 2004, 2005; Dong et al., 2006; Ishikawa et al., 2009; Kourrich and Thomas, 2009; Mu et al., 2010). In contrast, many results demonstrate the importance of functional ensembles of NAc neurons in mediating responses to natural rewards and drugs (e.g., Peoples et al., 2007; Koya et al., 2009; Wheeler and Carelli, 2009). This is probably due in large part to selective activation or modulation of MSN by various presynaptic inputs. For example, DA is known to gate the activation of MSN by excitatory inputs (Nicola et al., 2000; O'Donnell, 2003; Wheeler and Carelli, 2009). Perhaps DA and other inputs select the population of MSN that is activated in a particular behavioral context (e.g., Owesson-White et al., 2009), and then global adaptations such as AMPAR upregulation influence the strength of activation. A challenge for future studies is to understand the relationship between presynaptic and postsynaptic changes over the course of cocaine withdrawal.

4. Conclusions

Research from our lab and others has established that psychomotor stimulants produce both transient and long-lasting changes in the strength of excitatory synapses in brain circuits that control motivation and reward. We have focused on two examples in this review, but the number of mechanistically distinct forms of drug-induced plasticity suggested by recent studies is impressive. The challenge of relating drug-induced synaptic changes to addiction-related behavioral changes is daunting, but progress is being made. Indeed, due in part to its strong animal models and relatively well characterized underlying circuitry, addiction research is emerging as a major front in ongoing efforts to characterize the cellular mechanisms that link synaptic and behavioral plasticity.

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Figure 1.

AMPA enter synapses through a two-step process, indicated by red boxes 1 and 2 in the figure: 1) exocytosis onto the extrasynaptic cell surface (this step is accelerated by PKA), and 2) translocation into the synapse following activation of NMDAR and CaMKII. In the NAc and several other regions (e.g., PFC and hippocampus), spines of principal neurons receive convergent inputs from glutamate and DA terminals; thus, DA receptors are well positioned to modulate AMPAR trafficking at glutamate synapses. In primary cultures from these brain regions, we have found that brief (5-15 min) treatment with a D1-class receptor agonist, leading to PKA activation, increases the rate of AMPAR externalization at extrasynaptic sites. This in turn primes AMPAR for synaptic insertion in response to subsequent NMDAR stimulation. This schematic is based on data presented in Sun et al. (2005), Gao et al. (2006), and Sun et al. (2008).