



# Atypical anti-schizophrenic drugs prevent changes in cortical N-methyl-D-aspartate receptors and behavior following sub-chronic phencyclidine administration in developing rat pups

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## ABSTRACT

We sought to determine the relationship between phencyclidine (PCP)-induced alterations in behavior and NMDAR expression in the cortex by examining the effect of anti-schizophrenic drug treatment on both. Sprague–Dawley rat pups were pretreated with risperidone or olanzapine prior to treatment with PCP on postnatal day 7 (PN7) or sub-chronically on PN7, 9, and 11. Pre-pulse inhibition (PPI) of acoustic startle was measured on PN24–26 and following a challenge dose of 4 mg/kg PCP, locomotor activity was measured on PN28–35. PCP treatment on PN7 did not cause a deficit in PPI, but did cause locomotor sensitization. This was prevented by both antipsychotics. PCP treatment on PN7 caused an up-regulation of NR1 and NR2B, which was not affected by either anti-schizophrenic drug. PCP treatment on PN7, 9, and 11 caused a deficit in PPI and a sensitized locomotor response to PCP challenge as well as an up-regulation of NR1 and NR2A, all of which were prevented by both atypical anti-schizophrenic drugs. These data support the hypothesis that sub-chronic, but not single injection PCP treatment in developing rats results in behavioral alterations that are sensitive to antipsychotic drugs and these behavioral changes observed could be related to up-regulation of cortical NR1/NR2A receptors.

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## 1. Introduction

Schizophrenia is a severe neuropsychiatric disorder which afflicts approximately 1% of the population worldwide and shows strong genetic tendencies, with symptoms first presenting in early adulthood (Bromet and Fennig, 1999; Lewis and Lieberman, 2000). The disease is characterized by the presence of both positive, e.g. paranoia, hallucinations, delusions, and negative symptoms, including but not limited to emotional and social withdrawal, anhedonia, and depression, as well as cognitive impairments such as memory and attention deficits (McGlashan, 1996).

Phencyclidine (PCP) intoxication in humans has been shown to mimic both the positive and negative symptoms of schizophrenia as well as exacerbate psychosis in schizophrenics. These psychotomimetic properties led researchers to examine the effects of PCP in animals (Javitt and Zukin, 1991; Luby et al., 1962). PCP administration to rats results in increased locomotor activity, stereotypy, ataxia, head-weaving, and circling and walking backwards (Braff and Geyer, 1990; Castellani and Adams, 1981; Geyer and Ellenbroek, 2003; Martinez

et al., 2000; Steinpreis, 1996). In general, alterations in motoric behaviors such as these are thought to model the positive symptoms of schizophrenia.

N-methyl-D-aspartate (NMDA) receptor antagonists (PCP and MK-801) have been shown to reproducibly disrupt pre-pulse inhibition (PPI) of acoustic startle in animals and are routinely used to model the sensorimotor gating deficits of schizophrenia (Geyer et al., 2001; Rasmussen et al., 2007). It is a measure of the reduction of the startle response when a smaller non-startling acoustic stimulus (pre-pulse) is presented 80–120 ms prior to the startling stimulus (pulse) (Swerdlow et al., 1994). PPI of acoustic startle is a measure of information processing in specific pathways that are known to be abnormal in schizophrenia (Adams and Moghaddam, 1998; Bunney, 2000; Castellani and Adams, 1981; Ellenbroek and Cools, 2000; Wang et al., 2001). This deficit may contribute to the thought disorder and cognitive fragmentation characteristic of this disease (Braff and Geyer, 1990).

The etiology of schizophrenia has been described as a neurodevelopmental disorder (Pilowsky et al., 1993; Weinberger, 1987, 1996). Neurodegeneration during early stages of development has been shown following PCP or MK-801 treatment in the cortex, hippocampus, and striatum, all of which are regions of the brain implicated in schizophrenia (Ikonomidou et al., 1999; Wang and Johnson, 2005, 2007). In addition, PCP treatment on postnatal day (PN) 7, results in neurodegeneration (positive silver staining) in the frontal cortex,

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striatum, and hippocampus within 9 h of treatment (Wang and Johnson, 2005, 2007). Accompanying increases in caspase-3 immunoreactivity and terminal dUTP nick-end labeling (TUNEL) of broken DNA show this degeneration to be apoptotic in nature. Furthermore, administration of MK-801 and/or PCP to the neonate has been shown to produce aberrant behaviors later in development (Beninger et al., 2002; Wang et al., 2001) specifically, locomotor sensitization and deficits in PPI of acoustic startle were reported in postnatal (PN24–28) rats treated with PCP on PN7, 9, and 11 (Wang et al., 2001).

While it is generally acknowledged that the schizophrenic “break” does not often occur before puberty, there is evidence that this does occur in some cases (Nicolson et al., 2000; Ross et al., 2006b). Further, when examined retrospectively, there is ample evidence of abnormalities such as poor academic performance and poor social interaction skills in children who are eventually diagnosed as schizophrenic. This laboratory and others have demonstrated that this model (PCP treatment on PN7, 9, and 11) has face validity in that it results in behavioral deficits in rats similar to those seen in schizophrenia, including deficits in spatial learning, PPI of acoustic startle and social behavior (Harich et al., 2007; Rasmussen et al., 2007; Wang et al., 2001; Wiley et al., 2003a,b). Further, pre-treatment with olanzapine was able to prevent both behavioral and neurotoxic indices of postnatal PCP administration, suggesting that this treatment paradigm is a suitable model of schizophrenia in rats (Wang et al., 2001).

Up-regulation of the NR1 subunit of the NMDA receptor has been reported to be associated with PCP-induced behavioral changes (Wang et al., 2001); however, it is not known whether this is accompanied by commiserate increases in NR2 subunits, which would be necessary for altered receptor function (Kutsuwada et al., 1992; Monyer et al., 1994; Paoletti and Neyton, 2007). Further, it is not known whether behavioral changes following single dose PCP treatment on PN7 administration are associated with NMDAR up-regulation, and if so, whether these changes can also be modulated by anti-schizophrenic drugs. Therefore, in this study we determined the effect of PN7 and PN7, 9, 11 PCP administration on two behaviors thought to model different aspects of schizophrenia as well as on the levels of the three most abundant cortical and striatal NMDA receptor subunits (NR1, NR2A and NR2B). Finally, the relevance of the changes in behavior and NMDA receptor subunit protein as a model of schizophrenia was challenged by determining the effect of pre-treatment with two anti-schizophrenic drugs, risperidone and olanzapine.

## 2. Materials and methods

### 2.1. Animals

Timed, day 14 pregnant female Sprague–Dawley rats were obtained from Charles River Laboratories (Wilmington, MA). The dams were housed individually with a regular 12 h light–dark cycle (lights on 0700, off at 1900) with food and water ad libitum. Following parturition, male and female pups from four dams were combined and randomly cross-fostered to one of the four lactating dams. Each litter consisted of ten to twelve pups with approximately equal numbers of each gender. All experiments were conducted in accordance with the NIH and the University of Texas Medical Branch at Galveston Institutional Animal Care and Use Committee.

### 2.2. Drugs

Phencyclidine was acquired from the National Institute on Drug Abuse (NIDA, Rockville, MD) and dissolved in 0.9% NaCl. Risperidone was obtained as a solution from the UTMB Pharmacy (Janssen Pharmaceutica, Titusville, NJ) and dissolved in 0.9% NaCl. Olanzapine was obtained from the UTMB Pharmacy in the powdered form (Eli Lilly and Company, Indianapolis, IN) and was reconstituted with 2.1 mL sterile water. Doses were chosen based on prior experiments that

addressed PCP-induced regulation of the NMDAR (Wang et al., 2001) and our own preliminary experiments.

### 2.3. Experimental design

Male and female rat pups were treated on either PN7 or on PN 7, 9, and 11 (sub-chronic) with 10 mg/kg PCP or saline vehicle (s.c.). Olanzapine (1 mg/kg, s.c.) or risperidone (0.25 mg/kg, s.c.) were administered 30 min prior to PCP or saline administration on PN7 or on PN 7, 9, 11. Pups were sacrificed by decapitation on PN7 at 24 h following saline/vehicle (control, time=0 h), PCP/vehicle, antagonist/vehicle, or antagonist/PCP treatment. In a separate experiment, pups were sacrificed by decapitation 24 h following the last injection of the aforementioned drug regimens on PN7, 9 and 11 (on PN12). For biochemical studies, the frontal cortex or striatum was dissected as described below and used for Western blot analysis. We chose to investigate the frontal cortex and striatum because of the prominent role these two brain regions play in the pathophysiology of schizophrenia. Furthermore, the frontal cortex and striatum are part of the brain circuitry involved in the expression of both PPI (Ellenbroek et al., 1996; Koch and Bubser, 1994; Swerdlow et al., 1995) and locomotor sensitization (Pierce and Kalivas, 1997). In behavioral experiments, animals were assessed for PPI of acoustic startle on PN24–26 and then tested for locomotor activity following a 4 mg/kg PCP (i.p.) challenge on PN28–35.

### 2.4. Sub-cellular fractionation

Protein extracts were prepared from the frontal cortex or striatum as previously described with some modifications (Anastasio and Johnson, 2008; Wang et al., 2001). Briefly, 2 mm sections corresponding to 4.7 to 2.7 mm anterior to Bregma for the frontal cortex and 0.7 mm to –1.3 mm for the striatum (Paxinos and Watson, 1986) were cut with the aid of an aluminum brain mold. Cortical brain sections were homogenized in 500  $\mu$ L of lysis buffer with the aid of an automatic tissue grinder (Kontes Pellet Pestle Motor, Kimble/Kontes, Vineland, New Jersey). The lysis buffer consisted of 10 mM HEPES (pH 7.4), 1 mM EDTA, 2 mM EGTA, and 500  $\mu$ M DTT. Just prior to use, protease inhibitor cocktail [4-(2-aminoethyl) benzenesulfonyl fluoride (AEBSF), pepstatin A, E-64, bestatin, leupeptin, and aprotinin without metal chelators (Sigma-Aldrich, St Louis, MO)] at a concentration of 10  $\mu$ L/mL was added to the lysis buffer. The homogenate was then centrifuged at 1000  $\times$ g at 4 °C for 10 min to pellet the nuclear protein fraction (P1). The supernatant (S1) was collected and centrifuged at 8000  $\times$ g at 4 °C for 30 min to pellet the membrane protein fraction (P2). The membrane fraction was re-suspended in homogenization buffer and centrifuged at 20,000  $\times$ g at 4 °C for 30 min; the resultant pellet was re-suspended in lysis buffer+1% SDS, boiled for 10 min and stored at –80 °C.

### 2.5. Western blot analysis

Equal amounts of protein were separated on 10% Bis–Tris gels (Invitrogen, NY) using SDS-PAGE with a MES-SDS running buffer system, pH 7.4. Following electrophoresis (110 V for 2 h), proteins were transferred to polyvinylidene difluoride (PVDF) membranes (0.2  $\mu$ m) in a Mini Electroltransfer Unit (Bio-Rad, Hercules, CA) overnight. The membrane was blocked in 5% nonfat milk, followed by incubation with the primary antibody in 1% milk for 2 h at room temperature. Following washes (3  $\times$  10 min) in TBS+0.1% Tween 20 (TBST), the membrane was incubated with horseradish peroxidase conjugated secondary antibodies for 1 h at room temperature. Analysis was carried out using the enhanced chemiluminescence (ECL) plus Western blotting detection reagents (Amersham Biosciences, Piscataway, NJ). The bands corresponding to the various proteins of interest were scanned and densitometrically analyzed by using an automatic imaging analysis system (Alpha Innotech

Table 1

	Weight (g)						
	PN7	PN8	PN9	PN10	PN11	PN12	PN28
Saline female	14.3±0.7	19.7±0.2	17.4±1.2	23.7±0.5	20.9±1.5	29.1±0.7	81.1±2.8
PCP female	14.7±0.8	17.3±0.6 <sup>a</sup>	16.5±0.8	21.4±0.4 <sup>a</sup>	19.9±1.3	24.9±1.5	79.2±4
Saline male	16.2±0.6	21.2±0.4	20.7±0.8	26.4±0.6	25.5±1.1	31.5±0.6	91.2±3.5
PCP male	15.9±0.8	17.7±0.5 <sup>b</sup>	17.5±0.8 <sup>b</sup>	21.9±0.7 <sup>b</sup>	20.6±1.1 <sup>b</sup>	23.8±0.3 <sup>b</sup>	85.6±4.6

Data presented as mean (g)±SEM.

<sup>a</sup>  $p < 0.05$  vs. saline female  $N = 6-11$ /group (Student's  $t$ -test).

<sup>b</sup>  $p < 0.05$  vs. saline male  $N = 6-13$ /group (Student's  $t$ -test).

Corporation, San Leandro, CA). All quantitative analyses were normalized to  $\beta$ -actin (after stripping [Reblot mild, Chemicon International, Temecula, CA]).

## 2.6. Antibodies

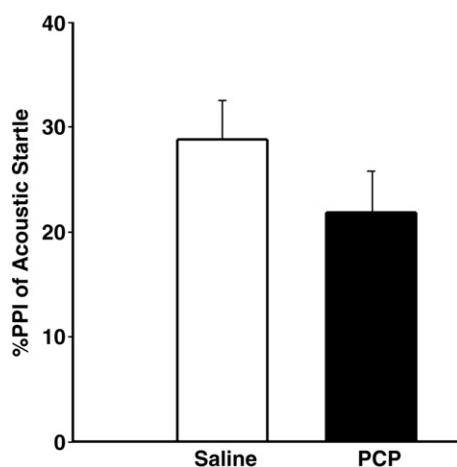
The monoclonal anti-NR1, anti-NR2A, and anti-NR2B were purchased from BD Biosciences (San Jose, CA). Anti-actin antibody was obtained from Chemicon International (Temecula, CA). Primary antibody dilution was 1:500–1:1000. Secondary antibodies were purchased from Zymed (Invitrogen Corporation, Carlsbad, CA) and used at a concentration of 1:5000.

## 2.7. Pre-pulse inhibition (PPI) of acoustic startle

Measurement of PPI of acoustic startle was performed according to previously published procedures with minor modifications (Wang et al., 2001, 2003). Testing was performed between 0900 and 1600 h as described below. Male and female rat pups (PN24–26) were transferred into a small sound-attenuated, dedicated behavior room on the day of testing and allowed to acclimate to the room for 20 min. Animals were then placed into one of three startle chambers (SR-Lab, San Diego Instruments, San Diego, CA) with a background noise level of 65 dB. Following a 10 minute acclimation period, rats were exposed to three randomly administered stimuli: no stimulus, a 73 dB 20 ms pre-pulse 100 ms prior to a 120 dB pulse, or a 120 dB 40 ms pulse alone with a variable inter-trial interval (5–20 s) for a total of 63 trials (21 no stimulus, 21 pulse alone, and 21 pre-pulse+pulse). % PPI of acoustic startle was calculated as the  $[\text{pulse} - (\text{pre-pulse} + \text{pulse})] / \text{pulse} \times 100$ .

## 2.8. Locomotor activity

On the day of testing animals were placed in locomotor chamber boxes and allowed to habituate for 30 min prior to a 4 mg/kg challenge

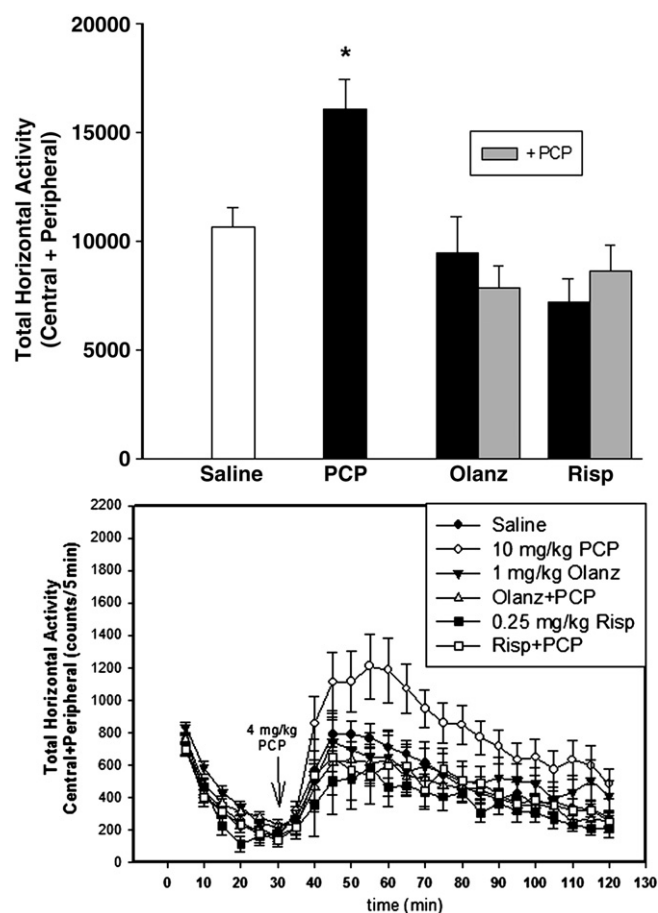


**Fig. 1.** Effects of PCP administration (PN7 only) on PPI of acoustic startle. PCP (10 mg/kg) treatment on PN7 does not produce a significant deficit in PPI of acoustic startle as measured on PN24–26.  $N = 8$  in both control and experimental groups.

dose of PCP (i.p.). Locomotor activity was measured for an additional 90 min via an open-field activity system (San Diego Instruments, San Diego, CA) which consisted of a square enclosure with Plexiglas walls (40×40×40 cm). Horizontal activity was measured with a 16×16 photobeam matrix which recorded both central and peripheral activity in 5 min bins.

## 2.9. Statistical analysis

Group comparisons were specifically defined before the beginning of each experiment; therefore, planned comparisons were performed instead of an overall  $F$  test in a multifactorial ANOVA (Keppel, 1982). Statistical comparisons for each experiment were conducted using a



**Fig. 2.** Effects of PCP administration (PN7 only) on the development of locomotor sensitization. Locomotor sensitization induced by a 4 mg/kg challenge to PCP treatment on PN7 treated animals as measured on PN28–35. Pre-treatment on PN7 with olanzapine (1 mg/kg) or risperidone (0.25 mg/kg) inhibits the development of locomotor sensitization  $N = 8$ /group (top). Time course of locomotor activity (bottom)  $*p < 0.05$  vs. saline (one-way ANOVA with Bonferroni's post hoc test).  $N = 8$  in both control and experimental groups. ●—saline; ○—10 mg/kg PCP; ▼—1 mg/kg olanzapine; △—olanzapine+PCP; ■—0.25 mg/kg risperidone; □—risperidone+PCP.

one-way ANOVA. All values are presented as mean  $\pm$  SEM. The null hypothesis was rejected at  $p < 0.05$ .

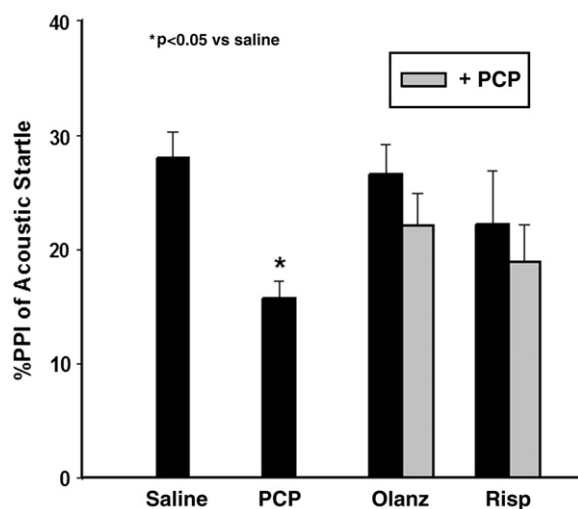
### 3. Results

PCP administration in adult rats causes substantial impairments in normal behavior (Geyer et al., 2001; Hanania et al., 1999; Phillips et al., 2001). In perinatal rats, 10 mg/kg PCP results in significant reduction of normal behaviors, most notably, a reduction of suckling behavior. This lasts approximately 6–8 h and causes a concomitant weight loss, especially in male pups. While females regain the lost weight within 24 h, the males do not; however, at the time of behavioral testing, there is no significant difference in weight between saline and PCP treated animals of either gender (Table 1). All experimental groups consisted of equal numbers of males and females.

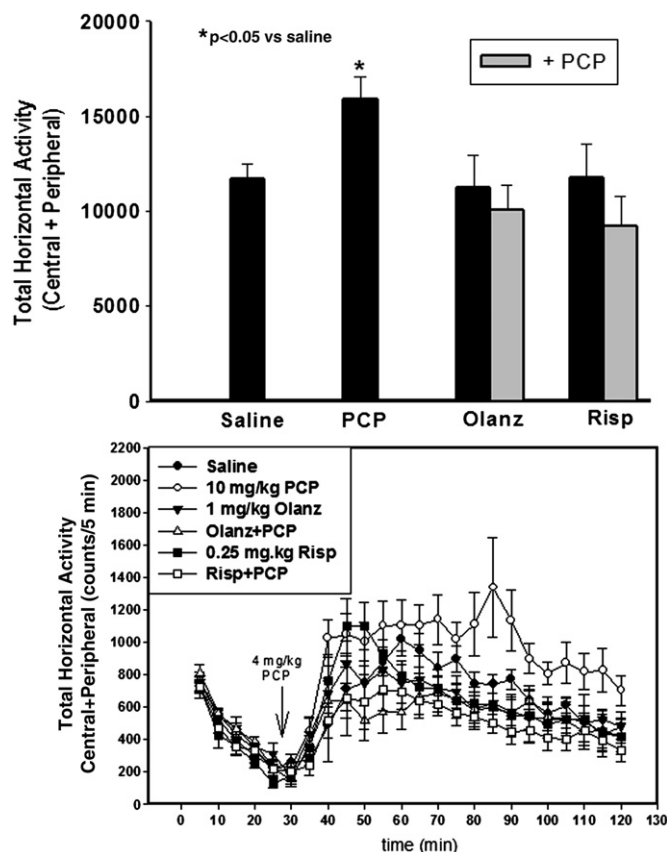
#### 3.1. Effects of PCP treatment on the development of locomotor sensitization and deficits in PPI

PCP treatment on PN7 had no significant effect on PPI ( $F_{3,31} = 2.264$ , NS) (Fig. 1). Further, PCP treatment on PN7 had no effect on the startle amplitude (data not shown). However, we found that PCP treatment on PN7 caused a significantly sensitized locomotor response to PCP challenge on PN28–35 that was prevented by either olanzapine ( $F_{3,31} = 8.071$ ,  $p < 0.05$ ) or risperidone ( $F_{3,31} = 10.001$ ,  $p < 0.05$ ) (Fig. 2).

As with PCP treatment on PN7, treatment on PN7, 9, and 11 produced no significant effect on startle amplitude (data not shown). On the other hand, PCP treatment on PN7, 9, and 11 resulted in a significant deficit in PPI of acoustic startle measured on PN24–26, which was blocked by pre-treatment with either olanzapine ( $F_{3,31} = 4.194$ ,  $p < 0.05$ ) or risperidone ( $F_{3,31} = 2.852$ ,  $p < 0.05$ ) (Fig. 3). Previous studies from this laboratory reported that the atypical antipsychotic drug, olanzapine, prevented sensitization to the locomotor activating effects of PCP challenge with 2 mg/kg on PN42 as well as the inhibition of baseline pre-pulse inhibition of acoustic startle on PN24–28 in female pups (Wang et al., 2001). This experiment was repeated using the present design which also included male pups in order to exclude any gender related differences. Further, this design was extended to risperidone, which has some atypical properties at low doses. For these studies, animals treated



**Fig. 3.** Effects of sub-chronic PCP administration (PN7, 9 and 11) on PPI of acoustic startle. PCP treatment (10 mg/kg) treatment on PN7, 9, and 11 results in a deficit in PPI of acoustic startle as measured on PN24–26. Pre-treatment on PN7, 9, and 11 with olanzapine (1 mg/kg) or risperidone (0.25 mg/kg) inhibits the PCP-induced deficit in PPI of acoustic startle. \* $p < 0.05$  vs. saline (one-way ANOVA with Bonferroni's post hoc test).  $N = 8$  in both control and experimental groups.



**Fig. 4.** Effects of sub-chronic PCP administration (PN7, 9 and 11) on the development of locomotor sensitization. Locomotor sensitization induced by a 4 mg/kg challenge to PCP treatment on PN7, 9, and 11 treated animals as measured on PN28–35. Pre-treatment on PN7, 9, and 11 with olanzapine (1 mg/kg) or risperidone (0.25 mg/kg) inhibits the development of locomotor sensitization. (top) Time course of locomotor activity (bottom) \* $p < 0.05$  vs. saline (one-way ANOVA with Bonferroni's post hoc test).  $N = 8$  in both control and experimental groups. ●—saline; ○—10 mg/kg PCP; ▼—1 mg/kg olanzapine; △—olanzapine + PCP; ■—0.25 mg/kg risperidone; □—risperidone + PCP.

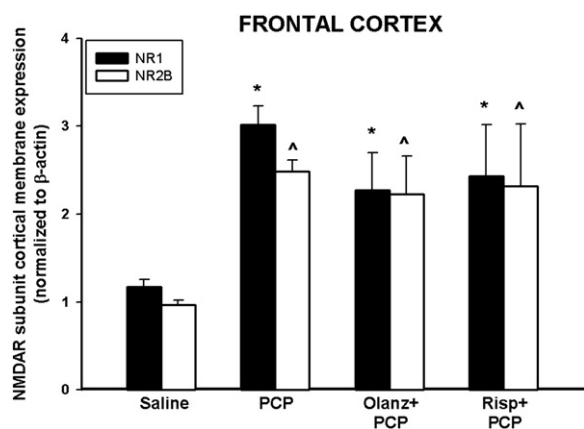
sub-chronically with PCP and then challenged with 4 mg/kg PCP on PN28–35, showed enhanced locomotor activity compared to saline treated animals (Fig. 4). Both olanzapine ( $F_{3,31} = 5.920$ ,  $p < 0.05$ ) and risperidone ( $F_{3,31} = 5.044$ ,  $p < 0.05$ ) pre-treatment on PN7, 9, and 11 prevented PCP-induced locomotor sensitization measured on PN28–35 with no effect of their own (Fig. 4).

#### 3.2. Effects of olanzapine and risperidone on PCP-mediated regulation of NR1/2A/2B

In order to delineate one possible biochemical mechanism underlying the above mentioned behavioral changes, we determined the effects of PCP treatment on PN7 or on PN7, 9, and 11 on the density of NR1, NR2A, and NR2B in the synaptosomal membrane fraction of the frontal cortex and striatum.

Quantitative western analysis revealed that PCP treatment on PN7 produced a 3-fold increase in membrane bound NR1 and NR2B subunits in the frontal cortex 24 h after PCP administration, while no effect on NR2A was observed (Anastasio and Johnson, 2008). In another experiment, perinatal animals were treated with either olanzapine (1 mg/kg) or risperidone (0.25 mg/kg) 30 min prior to PCP treatment on PN7. Twenty-four hours after the PCP injection, the rats were killed and the density of NMDA receptors was determined in the cortical membrane fraction. Pre-treatment with either olanzapine or risperidone did not inhibit the up-regulation of membrane cortical NR1 or NR2B induced by PCP treatment on PN7 (Fig. 5). Furthermore, treatment with olanzapine or risperidone alone had no effect on the



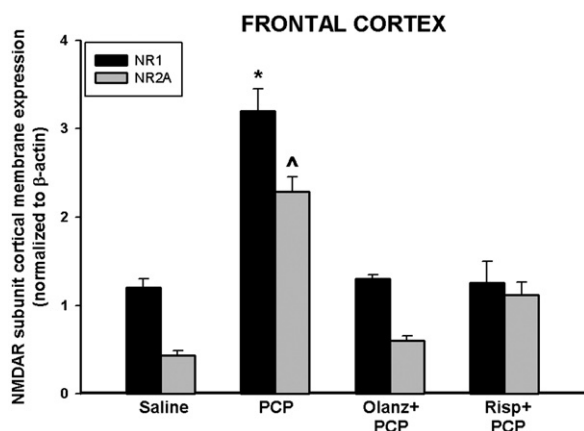


**Fig. 5.** Pharmacological antagonism of PCP treatment on PN7 effects on NR1 and NR2B in the frontal cortex. Quantitative analysis reveals that antipsychotics pre-treatment has no effect on PCP treatment on PN7-induced up-regulation of NR1 or NR2B protein levels in the frontal cortex 24 h following PCP (10 mg/kg,  $N=5-15$ /treatment). \* $p<0.05$  vs. SAL NR1 (one-way ANOVA with Bonferroni's post hoc test) ^ $p<0.05$  vs. SAL NR2B (one-way ANOVA with Bonferroni's post hoc test).

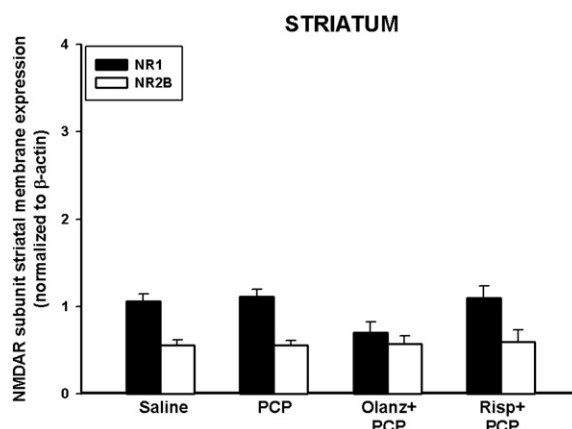
concentration of membrane NR1 or NR2B in the frontal cortex compared to saline controls (data not shown).

PCP treatment on PN7, 9, and 11 treatment caused a 3-fold increase in membrane NR1 and a 10-fold increase in NR2A protein levels in the frontal cortex with no effect on NR2B protein expression (Anastasio and Johnson, 2008). To analyze the actions of atypical antipsychotics on PCP-induced changes in the membrane protein levels of NMDAR subunits in the frontal cortex, animals were treated with olanzapine or risperidone 30 min prior to PCP on PN7, 9, and 11 and sacrificed 24 h after the last of 3 injections. Each of the antipsychotics investigated inhibited PCP-induced up-regulation of the NR1 subunit in the membrane protein fraction of the frontal cortex (Fig. 6), while having no effect on NR1 subunit expression on their own (data not shown). Pre-treatment with either olanzapine or risperidone also was able to completely block the increase in membrane cortical NR2A protein levels caused by PCP treatment on PN7, 9, and 11 treatment (Fig. 6) with no effect on their own (data not shown).

PCP treatment on PN7 had no significant effect on NR1 or NR2B protein expression levels in the striatum at 24 h following treatment (Fig. 7). The effects of pre-treatment with the anti-



**Fig. 6.** Pharmacological antagonism of PCP treatment on PN7, 9, and 11-induced up-regulation on NR1 and NR2A in the frontal cortex. Quantitative analysis reveals that pre-treatment with antipsychotics prevents PCP-induced (10 mg/kg,  $N=5-15$ /treatment) up-regulation of NR1 or NR2A protein in the frontal cortex. \* $p<0.05$  vs. SAL NR1 (one-way ANOVA with Bonferroni's post hoc test) ^ $p<0.05$  vs. SAL NR2A (one-way ANOVA with Bonferroni's post hoc test).

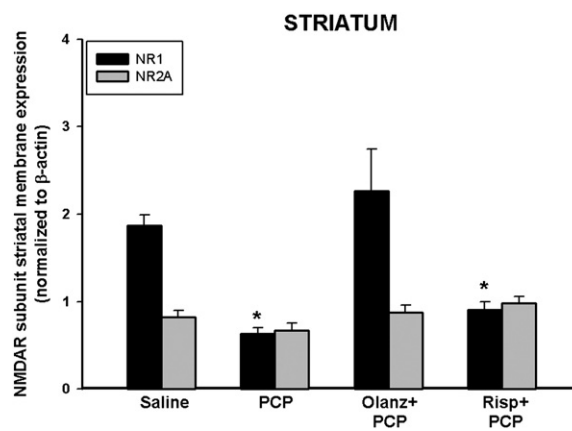


**Fig. 7.** Effects of a single injection of PCP on PN7 on the membrane protein expression of NR1 and NR2B in the striatum. Quantitative analysis reveals that PCP treatment on PN7 has no effects on the expression of striatal NR1 or NR2B (10 mg/kg,  $N=5-15$ /treatment).

psychotics on NR1 and NR2B protein levels in the striatum were also investigated following a single injection of PCP. Treatment with olanzapine or risperidone did not significantly alter expression of NR1 or NR2B protein levels in the striatum prior to PCP on PN7 at 24 h following treatment (Fig. 7) or when administered alone (data not shown). Similarly, treatment with either PCP or atypical antipsychotics did not alter striatal NR2A protein expression levels (data not shown).

Unlike the up-regulation of NR1 observed in the frontal cortex, Western blot analysis of membrane bound protein showed that levels of striatal NR1 protein were significantly decreased following sub-chronic PCP treatment on PN7, 9, and 11 (Fig. 8), with no effect on regulation of striatal NR2A (Fig. 8) or NR2B (data not shown). Pre-treatment with the anti-schizophrenic drug olanzapine, but not risperidone, prevented the down-regulation of NR1 protein expression in the striatum caused by sub-chronic PCP administration (Fig. 8). Administration of these drugs alone had no effect on NR1 or NR2A protein levels in the striatum (data not shown).

We also measured the levels of NR1, NR2A, and NR2B in the frontal cortex of animals following PCP challenge on PN28–35. No alterations in the protein levels of any of the subunits in the frontal cortex were evident in the animals treated sub-chronically with PCP (data not shown).



**Fig. 8.** Antagonism of PCP treatment on PN7, 9, and 11-induced down-regulation on NR1 in the striatum. Quantitative analysis reveals that pre-treatment with olanzapine prevents PCP-induced (10 mg/kg,  $N=5-15$ /treatment) down-regulation of NR1 in the striatum. PCP treatment had no effect on protein levels of NR2A \* $p<0.05$  vs. SAL NR1 (one-way ANOVA with Bonferroni's post hoc test).

#### 4. Discussion

It is well known that acute PCP treatment produces a disruption in PPI in adult rats, similar to that seen in schizophrenic patients (e.g. (Mansbach and Geyer, 1989; Martinez et al., 2000)). Typical antipsychotics, such as haloperidol, are not able to reverse deficits in PPI caused by acute PCP treatment in adult rats (Geyer et al., 2001) or in pre-pubertal rats (Martinez et al., 2002), but in adult rats, they can reverse the effects of dopamine agonists (Geyer et al., 2001). Atypical antipsychotics, including clozapine, olanzapine, and quetiapine, are effective at alleviating acute PCP inhibition of PPI in adult rats (Ballmaier et al., 2001; Geyer et al., 2001; Johansson et al., 1994, 1995; Martinez et al., 2002), but not in pups (PN16–19) or pre-pubertal (PN45) rats (Martinez et al., 2002). However, olanzapine is effective at preventing the deficits observed in PPI in PN24–28 pups following PCP treatment on PN7, 9 and 11 (Wang et al., 2001). In addition, both olanzapine and risperidone are able to increase PPI in NR1<sup>-/-</sup> mice (Duncan et al., 2006). The current study shows that both olanzapine and risperidone pre-treatment also blocks the PPI deficit observed in both male and female PN24–26 pups following PCP pre-treatment on PN7, 9 and 11. In contrast, PCP treatment on PN7 administration did not produce a deficit in PPI in developing rat pups. It is then reasonable to postulate that PCP treatment on PN7, 9, and 11 produces a chronic deficit in NMDA receptor function compared to a single injection of PCP and that this more closely models the disease and the developmental NMDA hypofunction theory of schizophrenia (Duncan et al., 2006).

Like other psychomotor stimulants, repeated administration of PCP causes a progressive augmentation of locomotor activity (Xu and Domino, 1994), referred to as sensitization. The neuroadaptations associated with sensitization may be linked to the mechanisms underlying addiction (Robinson and Berridge, 1993). Sensitization is also thought to be an important index related to psychosis as well as movement and thought disorders in schizophrenia (Robbins, 1990). PCP-induced sensitization is blocked by haloperidol and risperidone (Kitaichi et al., 1995) and cross-sensitizes with MK-801 (Pechnick and Hiramatsu, 1994), but not with amphetamine (Balster, 1989, 1986). In the current study, PCP treatment on PN7 or on PN7, 9, and 11 produced locomotor sensitization in rats at PN28–35 that was blocked by pre-treatment with either olanzapine or risperidone.

Our group and others have demonstrated that a single administration of PCP, MK-801 or ketamine to PN7 pups induces widespread neuronal apoptosis (Ikonomidou et al., 1999; Scallet et al., 2004; Wang and Johnson, 2005, 2007; Young et al., 2005). Other laboratories have also reported that transient NMDAR blockade by acute PCP, MK-801, ketamine, and ethanol to rodents during development causes behavioral, structural, and molecular abnormalities in adulthood (Fredriksson and Archer, 2003, 2004; Fredriksson et al., 2004; Harris et al., 2003; Wozniak et al., 2004). For example, Harris et al. (2003) found that PN7 female rat pups administered MK-801 (0.5 mg/kg, twice, 8 h apart, s.c.) showed PPI deficits and increased locomotor activity accompanied by a reduction of brain volume and neuronal number within the hippocampus and altered hippocampal NR1 subunit expression. Furthermore, the neurotoxicity evident in the developing frontal cortex following either a single injection or multiple injections suggests a role for altered cortical function in the development of locomotor sensitization (Wang et al., 2001; Wang and Johnson, 2005, 2007).

Several studies from this laboratory have investigated a possible mechanism by which PCP may elicit its neurotoxic effects and produce alterations in behavior in rats. In developing rats, PCP treatment on PN7, 9, and 11 of pups resulted in increased expression of NR1 mRNA in the frontal cortex, striatum, nucleus accumbens and olfactory cortex that was inhibited by pre-treatment with the atypical antipsychotic olanzapine (Wang et al., 2001). These data are consistent with our previous report demonstrating that chronic PCP treatment resulted in

increased NR1 immunoreactivity in the frontal cortex and striatum of adult rats treated chronically with PCP (Hanania et al., 1999). In addition, recently we reported that PCP treatment on PN7 produces an up-regulation of cortical NR1 and NR2B subunits via an increase in trafficking from intracellular compartments to the membrane, while sub-chronic PCP treatment results in an up-regulation of cortical NR1 and NR2A subunits by increasing new protein synthesis of these subunits (Anastasio and Johnson, 2008). We extended these experiments in this study and show that olanzapine and risperidone pre-treatment is unable to prevent the up-regulation of cortical NR1 and NR2B subunits following PCP treatment on PN7 administration. These data suggest that NMDAR up-regulation in the cortex and subsequent locomotor sensitization following a single injection of PCP are independent of each other. This suggests that expression of sensitized behavior is dependent on mechanisms requiring activation of dopamine (DA) D2 and/or serotonin (5-HT) 5-HT<sub>2A/2C</sub> receptors. Thus, even though a sensitized behavioral response to acute PCP challenge appears to involve activation of D2 and/or 5-HT<sub>2</sub> receptors, the increase in NMDAR trafficking appears to be independent of these receptors. Furthermore, the DA and 5-HT mechanisms involved in locomotor sensitization may be downstream of receptor regulation. If so, antipsychotics would be ineffective at preventing PCP-induced changes in receptor trafficking, but still capable of preventing PCP-induced sensitization and other behaviors. In contrast, both olanzapine and risperidone are able to prevent up-regulation of NR1 and NR2A subunits in the frontal cortex following sub-chronic PCP treatment. These data suggest that up-regulation of NR1 and NR2A subunits (rather than NR1 and NR2B) may underlie the development of deficits in PPI following PCP treatment on PN7, 9, and 11, though the relative role of NMDAR subunits in locomotor behavior and PPI are not completely understood. Several studies have reported that blockade of NR2B subunits with selective antagonists (eliprodil and Ro63-1908) does not disrupt PPI (Depoortere et al., 1999; Higgins et al., 2003; Wiley, 1998) and that it is the combined blockade of NR2A and NR2B subunits which is necessary to induce deficits in PPI as well as hyperactivity in rodents (Spooren et al., 2004). Administration of the selective NR2A antagonist, NVP-AAM007, to adult rats did not disrupt PPI and resulted in hypolocomotor activity, while the NR2B antagonist, Ro 25-6981 disrupted PPI and produced a dose-dependent hyperlocomotion (Chaperon et al., 2003). Thus, the role of NR2A and NR2B subunits in the detrimental behavioral effects of PCP merits further investigation.

Neither the mechanisms underlying PCP-induced alterations in behavior nor NMDA receptor up-regulation nor those responsible for the ability of olanzapine and risperidone to prevent these alterations is clear. Clozapine, olanzapine, and M100907, but not haloperidol, have been reported to prevent the blockade of NMDA responses in the medial prefrontal cortex caused by acute PCP administration (Arvanov and Wang, 1999; Wang and Liang, 1998). These data suggest a prominent role for 5-HT<sub>2A</sub> receptors in preventing the acute effect of PCP. Furthermore, clozapine has been shown to inhibit hypersensitive responses to NMDA following sub-chronic PCP (Arvanov and Wang, 1999), suggesting that the expected increase in NMDAR function following PCP up-regulation of NR1 and NR2A may also be prevented by olanzapine treatment. It is possible that the ability of olanzapine to inhibit PCP-induced neurotoxicity, NMDAR up-regulation and deficits in PPI following PCP treatment stems from its blockade of both DA and 5-HT<sub>2A</sub> receptors. Additionally, if PCP-induced neurotoxicity and the subsequent loss of these cortical neurons results in a hypo-glutamatergic state, then the ability of antipsychotics to prevent this loss of glutamatergic tone through blockade of DA and 5-HT receptors may be related to their effectiveness in alleviating both positive and negative symptoms of schizophrenia (Jardemark et al., 2000). Understanding the mechanism by which olanzapine and risperidone prevent PCP-induced neuronal apoptosis could provide insights into the molecular and cellular mechanisms involved in the behavioral

effects of PCP in rats, which in turn could provide insight into the etiology and pharmacotherapy of schizophrenia.

PN7–11 is a critical stage in brain development, with ample evidence of neurite growth and synapse formation, and is referred to as the brain growth spurt (Olney et al., 2002). Thus, it is reasonable that even temporary changes during this critical period could result in behavioral changes as these animals mature. No variation in the expression of cortical NR1, NR2A, or NR2B was observed in either saline or sub-chronically PCP treated pups as measured immediately following PCP challenge and behavior measurement on PN28–35. Thus, it could be argued that the behavioral effects observed at this time are independent of the effects of PCP challenge on the expression of the NMDA receptor and that are more likely due to changes in NMDA receptor expression during an earlier, critical period of development and the subsequent maturation of the brain. Additionally, these results are consistent with the neurodevelopment theory of schizophrenia, which hypothesizes that damage during this early, critical stage of brain development may account for the later manifestation of mental disorders (du Bois and Huang, 2007).

PCP treatment on PN7 had no effect on the expression of either PSD-95 (data not shown) or any of the NMDA receptor subunits in the striatum. PSD-95 interacts with the intracellular tail of NR2 subunits and is thought to be responsible for anchoring the functional NR1/NR2 receptor complex in the membrane (Wenthold et al., 2003). Thus, it is possible that the lack of an observable effect on NMDA receptor expression is related to the lack of effect on PSD-95. This suggests that the striatal neurotoxicity previously reported (Wang and Johnson, 2005) is unrelated to alterations in the expression of NMDA receptor protein. Similarly, the locomotor sensitization measured following a single injection of PCP must also be due to mechanisms other than changes in striatal NMDAR expression levels. However, this does not rule the possible role that altered NMDAR expression in the frontal cortex may play in the sensitization response.

Pre-treatment with olanzapine was able to prevent the down-regulation of striatal NR1 polypeptide that was caused by sub-chronic PCP administration; however, pre-treatment with risperidone did not. We have recently observed that neither SCH23390 (selective D1 antagonist), sulpiride (selective D2 antagonist), nor M100907 (selective 5-HT<sub>2A</sub> antagonist) were able to prevent the down-regulation of NR1 in the striatum (Anastasio and Johnson, unpublished observations). Thus, this effect of olanzapine may require blockade of both DA and 5-HT receptors. In addition to affinity for DA and 5-HT receptors, olanzapine possesses high affinity for muscarinic ACh receptors ( $K_i$ =1.89 nM), H1 histamine receptors ( $K_i$ =7.14 nM), as well as  $\alpha$ 1 noradrenergic receptors ( $K_i$ =19 nM) (Arnt and Skarsfeldt, 1998; Raggi et al., 2004); therefore, it is possible that in the striatum, an area rich in muscarinic ACh receptors, olanzapine's effect of blocking down-regulation of NR1 caused by sub-chronic PCP administration to postnatal rats could also involve an action at these receptors.

Studies of postmortem schizophrenic brains have provided conflicting data on the expression of NMDA receptor subunit expression. The NR1 protein has been shown to be up-regulated in the prefrontal, parietal, and medial temporal cortices in schizophrenia (Chen et al., 1998), while radioligand binding studies showed no significant differences in NR1 in healthy vs. schizophrenic brain tissue (Gao et al., 2000). Another study using in situ hybridization reported no difference between control and schizophrenic brains in either NR1 or NR2 mRNA levels (Akbarian et al., 1996), but NR2B mRNA was reportedly increased in the hippocampus while NR1 mRNA was diminished in the medial temporal cortex, superior temporal cortex, and frontal cortex (Akbarian et al., 1996; Meador-Woodruff and Healy, 2000). Although these are quite complex, this does not rule out a role for altered NMDAR function early in development in this disease. In fact, a number of genes related to glutamatergic function have been discovered that are significantly associated with schizophrenia, though in as yet undefined ways. Included in this list is *NRG1*

(neuregulin-1), which is known to regulate NMDAR expression (Chong et al., 2008; Craddock et al., 2005). *DTNBP1* (dystrobrevin binding protein 1) has been shown to modulate glutamate function through up-regulation of presynaptic proteins and neurotrophic effects mediated by the Akt signaling pathway (Duan et al., 2007; Numakawa et al., 2004). DAOA (D-amino acid oxidase activator) activates NMDA receptors through a series of reactions involving the glycine-like molecule, D-serine (Korostishevsky et al., 2004; Ross et al., 2006a). RGS4 (regulator of G-protein signaling 4) is under-expressed in schizophrenic prefrontal cortex in postmortem microarray studies and has been reported to regulate G proteins in glutamate neurons, thereby dampening the effects of neurotransmitter interactions at G-protein coupled receptors (Chowdari et al., 2008, 2002). In addition to the glutamate related genes above found with linkage analysis, several other genes putatively associated with the NMDA synapse have been discovered to be associated with schizophrenia, including GRM3 (mGluR3 a member of the group II metabotropic receptor family, which also includes mGluR2 (Egan et al., 2004); mGluR3 is known to modulate glutamate release. Thus, genetic evidence supports a role for alterations in various glutamatergic genes including those specific to NMDA receptor function in schizophrenia. Importantly, these data are strongly supported by pharmacological data that also imply a role for altered NMDAR function. However, the complexities of these data do not allow a direct comparison between the data in this model to what may occur in the human brain in schizophrenia. Our view is that this model may reveal the gross structural features underlying behavioral alterations in rats that are similar to schizophrenia and possibly point to novel pharmacological approaches that may be helpful in treatment of the disease.

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