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Predictors of methamphetamine psychosis: History of ADHD-relevant childhood behaviors and drug exposure

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Abstract

The goal of this study was to extend our previous research that reported a significant association between Attention Deficit Hyperactivity Disorder (ADHD)-relevant childhood behaviors and the frequency of methamphetamine (MA)-induced psychotic symptoms in an expanded sample. 190 participants who met DSM-IV criteria for MA dependence were administered the Methamphetamine Experience Questionnaire that assessed MA-induced psychosis. Data related to MA exposure, comorbid drug use, education, familial psychiatric history and assessments of ADHD-relevant childhood behaviors as measured by the Wender Utah Rating Scale (WURS) were collected. Although WURS scores did not differ between 145 MAP+ and 45 MAP- subjects, MAP+ subjects with higher WURS scores were significantly more likely to report more frequent psychosis. Although mean daily MA dosage did not differ between the MAP+ and MAP- subjects, MAP+ who consumed larger doses of MA were significantly more likely to experience frequent psychosis. These data suggest that ADHD-relevant childhood behaviors may interact with MA exposure to reflect a neurobiological vulnerability related to the emergence of frequent MA-induced psychotic symptoms. These results may elucidate factors that contribute to the psychiatric sequelae of MA abuse.

Keywords

methamphetamine; psychosis; attention; ADHD; prefrontal cortex; substance abuse; predictors

1. Introduction

Worldwide use of methamphetamine (MA) is now estimated to be at 51 million users (Roehr, 2005; Degenhardt et al., 2008; Nations, 2008, 2009), with global abuse of amphetamine/methamphetamines now surpassing that of cocaine and opiates combined (United Nations, 2009). Approximately 5% of the adult population in the United States has used MA on at

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least one occasion and Emergency Department admissions related to MA use have doubled during the period of 1994 to 2002 (SAMSHA, 2004). A subset of individuals who chronically abuse MA also develop severe psychotic symptoms, which are often associated with high levels of psychiatric hospitalization and serious social dysfunction (Sato, 1992; Chen et al., 2003; Dore and Sweeting, 2006; McKetin et al., 2006; Pasic et al., 2007). In 2010, the nation's Drug Abuse Warning Network estimated that 66,308 drug-related Emergency Department visits involved amphetamines and methamphetamine use (SAMSHA, 2010) and that patients with MA-related psychotic disorders were 33% more likely than patients with cocaine disorders to be transferred to the inpatient ward (Leamon et al., 2002). The highly addictive nature of MA, as well as its ability to produce psychotic symptoms, makes this drug a major public health concern in the 21st century (United Nations, 2009).

Approximately 60% of MA abusers report a history of paranoia, delusions and hallucinations while under the influence of the drug (Dore and Sweeting, 2006; McKetin et al., 2006; Schuckit, 2006; Degenhardt et al., 2008; Salo et al., 2008). Early symptom descriptions of psychosis induced by amphetamines that are still relevant today include the following: 1) paranoid delusions with ideas of reference; 2) persecutory delusions; and 3) auditory and visual hallucinations that occur in a state of clear consciousness (Connell, 1958). Although negative symptoms and thought disorder have also been reported following long-term MA and amphetamine use, they are rare (Siomopoulos, 1976; Srisurapanont et al., 2003). One large study of 309 MA users reported that after adjusting for comorbid psychiatric disorders (e.g., schizophrenia), MA abusers were 13 times more likely to develop psychotic symptoms compared to the general population (McKetin et al., 2006). Two other studies also reported high prevalence rates of psychosis in MA abusers, with one study reporting psychotic symptoms in 75% of their sample (Sato, 1992) and another reporting auditory hallucinations and persecutory delusions in 44.6 and 22.8%, respectively (Srisurapanont et al., 2003). Pre-morbid risk factors for MA psychosis include a familial history of psychiatric illness and age of first MA use (Chen et al., 2005) as well as genetic risk factors (Harano et al., 2004; Suzuki et al., 2006; Bousman et al., 2009).

Although it is well known that MA is highly neurotoxic to the brain, causing damage to multiple transmitter systems (Ricaurte et al., 1984; Schmidt et al., 1985; Seiden and Ricaurte, 1987; Axt and Molliver, 1991; Bowyer and Holson, 1995; Seiden and Sabol, 1996; Davidson et al., 2001; Thompson et al., 2004; Meredith et al., 2005; Yamamoto and Bankson, 2005; Quinton and Yamamoto, 2006), less is known about why MA abusers are at increased risk for psychotic episodes.

Similarities between amphetamine/methamphetamine induced psychosis and psychotic symptoms associated with schizophrenia have long been observed (Angrist and Gershon, 1970; Angrist et al., 1974). Studies that have administered large doses of amphetamine to non-schizophrenia individuals reported symptoms similar to those observed in schizophrenia patients including: delusions, hallucinations and to a lesser degree thought disorder (Angrist et al., 1974). Acute amphetamine administration has also been reported to produce or enhance psychotic symptoms in schizophrenia patients that are ineffective in non-substance-using controls (Curran et al., 2004). Following amphetamine challenge in acutely ill schizophrenia patients, increased dopamine release was observed within the striatum (Laruelle et al., 1996; Laruelle et al., 1999; Abi-Dargham et al., 2003), with levels of dopamine release correlating both with positive symptoms (Abi-Dargham et al., 2000) as well as with response to dopamine blockers (Laruelle et al., 1999). Collectively these findings suggest substantial overlap between schizophrenia psychosis and amphetamine/methamphetamine-induced psychosis.

Different models have emerged to describe the mechanisms underlying drug-induced psychoses (Buckley, 1998; Chambers et al., 2001). One model is that drug use alone can cause psychosis. If this were true then one would expect 100% of MA users to experience drug-induced psychosis after using this powerful stimulant drug and the facts simply do not support this. Statistics across studies report that on average only about 60% of MA abusers report a history of paranoia, delusions and hallucinations while under the influence of the drug (Dore and Sweeting, 2006; McKetin et al., 2006; Schuckit, 2006; Degenhardt et al., 2008; Salo et al., 2008). A second widely held model is a version of the “self medication hypothesis”, which proposes that individuals with psychiatric disorders abuse substances to alleviate adverse disease symptoms or medication side-effects” (Dalack et al., 1998). This hypothesis has been disputed in recent years due to numerous factors: 1) the prevalence rate of major psychiatric disorders (e.g., schizophrenia) has remained constant while the incidence of substance abuse in these patients has increased exponentially (Chambers et al., 2001); 2) drugs of abuse have wide ranging effects in psychiatric patients, some of which alleviate symptoms and some that exacerbate them (DeQuardo et al., 1994; Addington and Duchak, 1997); and 3) drug abuse often occurs before the onset of the disease (Berti, 1994). A third model is the primary addiction hypothesis that proposes the existence of common neural circuits underlying the neuropathology of both schizophrenia and addiction (Chambers et al., 2001). These neural circuits include a dysregulation of a wide neural network that involves mesolimbic dopamine in the nucleus accumbens along with cortical and hippocampal inputs (Jentsch and Taylor, 1999). And finally, there is the stress-vulnerability model that proposes that individuals who are at pre-existing risk for developing psychosis, develop psychosis after using drugs (Bramness et al., 2012).

In the current study we sought to replicate and extend our previous research that reported a significant association between Attention Deficit Hyperactivity Disorder (ADHD) relevant childhood behaviors as measured by the Wender Utah Rating Scale (WURS) and the frequency of MA-induced psychotic symptoms in a small sample of 39 MA abusers (Salo et al., 2008). Recent studies have documented a high-level of ADHD symptomatology in stimulant-dependent individuals (Kaye et al., 2012), while other studies have also reported increased WURS scores in chronic substance abusers (Clure et al., 1999; Lynskey and Hall, 2001; Matsumoto et al., 2005a; Matsumoto et al., 2005b), including individuals dependent on methamphetamine (Simon et al., 2000) and cocaine (Boutros et al., 2002). However, this is one of the first studies to link measures of ADHD-relevant childhood behaviors to MA-induced psychoses.

The main goal of the current study was to examine whether the presence of MA psychosis was associated with any risk factors (MAP+ vs. MAP- comparison) and conditional on having MA psychosis, whether or not increased frequency of psychosis was associated with any predictors. Furthermore, as cannabis use has been linked to non-substance induced psychosis (Broome et al., 2005), we wanted to examine cannabis use patterns as well as the prevalence of family history of psychiatric disorders in our sample.

2. Method

2.1. Participants

The MA-dependent subjects comprised 99 men and 91 women meeting criteria for lifetime MA dependence according to DSM-IV criteria. Data from thirty nine of the subjects were reported as part of our previous study (Salo et al., 2008). The MA abusers were recruited from substance abuse treatment centers and residential housing programs in the Sacramento area. Our study criteria required that our sample was/had: 1) MA abstinent at the time of study; 2) no current or lifetime comorbid non-substance induced Axis I disorder; 3) free of

any neurological medical issues. 3) no substance dependence (other than MA and nicotine) within the past five years; and 4) no self-reported history of a seropositive test for HIV.

Among the 190 subjects, 91 were currently in treatment and 99 were in non-treatment settings. It should be emphasized however that most of our sample had received structured substance abuse treatment at some point in their addiction. All subjects had been drug abstinent for a minimum period of three weeks (range 3 weeks to 10 years) by self report and random urine drug screens performed at referring sites. All subjects were literate and completed a standardized measure of verbal IQ (National Adult Reading Test; NART) (Nelson, 1982). All subjects signed informed consent approved by the University of California Davis Institutional Review Board and were paid a modest stipend for study participation.

2.2. Assessments and semi-structured interviews

2.2.1. Structured clinical interview for DSM-IV—All participants were interviewed by PhD level research personnel using the Structured Clinical Interview for DSM-IV (SCID) and a consensus diagnosis was obtained for each participant. The SCID was used as an adjunctive measurement to assess the presence and frequency of psychotic episodes associated with MA use.

2.2.2. Methamphetamine Experience Questionnaire—All participants were interviewed using the Methamphetamine Experience Questionnaire (MEQ) which is an interview based on the Cocaine Experience Questionnaire (Gelernter et al., 1994; Leamon et al., 2010). The MEQ is designed to assess the lifetime frequency of psychotic episodes associated with MA use, conditions in which psychotic episodes occur, as well as the persistence of these symptoms.

Sample MEQ questions include:

1. How often have you had paranoid experiences while using methamphetamine, using a 0 to 5 Likert Scale? (Where 0 is never and 5 is always)
2. Were you more likely to get paranoid when you used greater amounts of methamphetamine?

2.2.3. Wender Utah Rating Scale (WURS)—All participants completed the WURS which retrospectively assesses Attention Deficit Hyperactivity Disorder (ADHD)-relevant childhood behaviors and symptoms in adults (Wender, 1985; Ward et al., 1993). The version of the WURS used in the current study contained 25 items that assessed symptoms which occurred between five and twelve years of age.

2.3. Statistical analyses

Statistical analyses were conducted using the SAS Institute SAS Version 9.3 (SAS Institute, 2002-2010) and included descriptive statistics for all categorical and continuous variables. First, differences between MA users with (MAP+) and without (MAP-) a history of psychosis were assessed using chi-squared test for the categorical variables, two sample t-tests for the continuous variables that were normally distributed and nonparametric Wilcoxon two-sample tests for those continuous variables that violated the assumption of normality. Second, for the MAP+ users, proportional-odds models (McCullagh, 1980) were used to examine the association between demographic and clinical characteristics and the frequency of psychosis. These models can be thought of as an extension of the logistic regression model for dichotomous dependent variables, allowing for more than two ordered response categories and have the advantage of being able to capture information from the

entire spectrum of psychosis (measured on an ordinal Likert scale). The odds ratio (OR) is calculated for each cut-off across the frequency psychosis (for example, 1 versus 2-5, then 1-2 versus 2-5, and so on), and then a summary OR is calculated from the individual ORs, under the assumption that the individual OR are constant for all categories. When the testable proportional odds assumption is met, the odds ratio provided by the model does not reflect a simple odds ratio, but rather an average across multiple thresholds of classification of the outcome and can be interpreted as the odds of having “more frequent” psychosis across the whole range of the psychosis. We first examined unadjusted models (with one predictor at the time), followed by adjusted models to assess if the significant effect of the predictors from the univariate analyses could be an artifact of confounding. The validity of the proportional odds assumption was tested using Score tests and was deemed adequate for all models.

3. Results

Table 1 presents summary measures (means and standard deviations for the continuous variables and frequencies and percentages for the categorical ones) for all demographic and clinical characteristics, stratified on the history of paranoia. Out of the 190 participants in the sample, 145 (76%: 79 men and 66 women) reported symptoms associated with MA psychosis that included paranoid delusions as well as visual, auditory and tactile hallucinations (MAP+), and 45 (24%: 20 men, 25 women) did not (MAP-). The mean age in the entire sample was 38 years (standard deviation = 8, range 19 – 55 years), with the average participant having completed high school. The two groups did not differ significantly in age, gender composition, education, years of parental education, estimates of pre-morbid intelligence as assessed by the NART (Nelson, 1982), or WURS scores (see Table 1). An analysis of the methamphetamine use patterns revealed that on average the MAP+ and MAP- subjects had similar ages of first use, duration of drug use, mean daily MA dose consumed and months drug abstinent (see Table 1).

We had information on history of psychiatric illness in first-degree relatives from 98% of our sample. Four subjects were adopted at an early age and thus were unable to provide family history. In the whole MA abusers sample, 54 (28%) reported a family history of psychiatric illness. The proportion of participants with a history of psychiatric illness was 31% in the MAP+ group and 21% in the MAP- group, but this difference did not reach statistical significance (see Table 1).

3.1. Comorbid drug use

In any sample of illicit drug users it is common that most abusers will have used/abused more than one drug during their lifetime. Among the 190 MA abusers, several met criteria for lifetime, but not current, abuse/dependence for the following substances: Alcohol: 56%, Cocaine: 26%; Cannabis: 76%, Tobacco: 83%. Of those who had lifetime comorbid disorders, inclusion criteria required any that comorbid dependence on any substance (other than tobacco) must have occurred more than 5 years prior to study. Comorbid abuse was allowed if it did not occur within the past year. Median length of abstinence from the comorbid drugs was: Alcohol: 8 years, Cocaine: 15 years, Cannabis: 5 years.

Comorbid drug use patterns were similar in the two groups, for alcohol and cocaine, as well as tobacco use (see Table 1). Planned analysis between the MAP+ and the MAP-groups revealed no difference in the proportion of abuse/dependence users, age of first cannabis use, or years of cannabis use (see Table 1). There was however a difference in the abstinence times, with MAP+ participants having longer cannabis abstinence than the MAP- participants ($p = 0.04$, see Table 1).

73% of the MAP+ subjects experienced both auditory and visual hallucinations, 17% reported hallucinations in the visual modality only, 8% in the auditory modality only, and 2% reported delusions with no experience of hallucinations in either sensory modality. None of the MAP- subjects had experienced delusions or hallucinations. Among the 145 participants with reported history of psychosis, 23 (16%) had the lowest frequency (1), 27 (19%) had frequency 2, 38 (26%) frequency 3, 33 (23%) frequency 4 and 24 (17%) reported the highest levels (5).

Table 2 summarizes the results of the univariate unadjusted analyses (each predictor entered the model by itself, without any adjustment for other variables) examining the associations between demographic and clinical characteristics and the frequency of psychosis. Neither gender nor any of the other demographic characteristics were associated with frequency of psychosis in univariate analyses. Although WURS scores did not differ between the MAP+ and the MAP- subjects, MAP+ subjects with higher WURS scores were more likely to report being in a category with more frequent psychosis (OR = 1.04, 95% CI 1.02 – 1.05, $p < 0.0001$). There was a 4 percent increase in the odds of a more frequent psychosis per 1 point increase in WURS score. This interpretation holds across the entire range of psychosis (from 1 to 5). Among the MA use variables, the mean daily MA dosage was associated with frequency of psychosis. MA abusers who consumed larger quantities of MA on a daily basis had higher likelihood of being in a more frequent psychosis category (OR = 1.61, 95% CI 1.17 – 2.20, $p = 0.003$). None of the comorbid drug use variables were associated with frequency of psychosis, although there was a trend for the tobacco use and cannabis abuse/dependence to be associated with higher frequency of psychosis ($p = 0.08$ and 0.06 , respectively) in unadjusted analyses. We then fitted an adjusted model in which the two main predictors associated with frequency of psychosis in univariate analysis (WURS and mean daily MA dosage) were included as predictors of the frequency of psychosis simultaneously, to examine if they would be significant after controlling for the effect of the other. The significance of WURS remained unchanged, but the coefficient for daily MA dosage decreased (OR = 1.46, 95% CI 1.06 – 2.00, $p = 0.02$).

3.2. Gender differences

This dataset was well balanced on gender (males = 99; females = 91), thus we were in a position to examine gender as a moderator of MA psychosis. Across the entire sample of 190 MA abusers, the males and females did not differ in age (Wilcoxon two-sample $z = 1.26$, $p = 0.21$), years of education ($z = -0.37$, $p = 0.71$), years of parental education ($z = -0.55$, $p = 0.58$), or estimates of pre-morbid intelligence as assessed by the NART ($t_{183} = -0.01$, $p = 0.99$). An examination of the drug use patterns revealed that male and female MA abusers did not differ in duration of drug use ($z = 0.96$, $p = 0.34$), age of first use ($z = 0.46$, $p = 0.64$), mean daily dose of MA consumed ($z = -0.81$, $p = 0.42$), or the number of months for which they were MA abstinent at the time of study ($z = -1.16$, $p = 0.25$). No gender differences among the MAP+ participants reached significance.

4. Discussion

The data in the current study extend our previous findings of a significant positive correlation between frequency of MA-related psychotic episodes in the MAP+ subjects and scores on the WURS, a scale that retrospectively measures childhood attention function and hyperactivity (Salo et al., 2008). Although WURS scores did not differ between the MAP+ and the MAP- subjects, those MAP+ subjects who reported a higher lifetime frequency of MA induced psychotic episodes were those with the highest WURS scores reflecting ADHD-relevant childhood behaviors. In this new study we also examined mean daily MA dose consumed and found that although the mean daily dose consumed did not differ between the MAP+ and the MAP- subjects, among the MAP+ subjects there was a

correlation between the amount of MA consumed and the frequency of MA psychotic episodes. These results appear to be consistent with other published studies (Batki and Harris, 2004; McKetin et al., 2013).

We did not replicate our previous finding that a familial history of psychiatric illness may further increase the risk of developing MA-induced psychotic symptoms (Chen et al., 2005; Salo et al., 2008). In this dataset, there was no statistically significant difference in either the prevalence or the frequency of MA induced psychotic episodes between those 54 MA Abusers with a family history of psychiatric illness, versus those who had no family history. We also failed to detect gender differences related to psychosis in this sample. These findings are inconsistent with others who have reported that MA dependent women were more likely than their male counterparts to report experiencing various psychotic symptoms (Mahoney et al., 2010). Additional studies in a larger community based sample are needed to explore these issues further.

Within the context of existing models that propose frameworks relating drug use to psychosis (Buckley, 1998; Chambers et al., 2001; Bramness et al., 2012), our results are perhaps the most consistent with the stress-vulnerability model. Our data suggest there is a degree of dose-response relationship between meth exposure and frequency of psychotic symptoms among individuals vulnerable to psychosis and that a pre-existing history of ADHD-relevant childhood behaviors is yet another contributing factor. If one accepts this model that preexisting vulnerabilities put one at risk for developing MA-induced psychosis then this can also explain why some individuals develop psychotic symptoms when exposed to methamphetamine and others do not. However, the one issue that remains unresolved is why WURS scores do not differ between the MAP+ and MAP- subjects. One explanation might be that early attention problems represent only one of several factors that predispose individuals to MAP and that although ADHD-relevant childhood behaviors may contribute to the severity (i.e., frequency) of psychotic episodes experienced, it is not a sufficient factor on its own to produce the symptoms.

As recent epidemiological studies have shown that early substance use (i.e., cannabis) among individuals who are at genetic risk may increase the likelihood of developing psychotic symptoms in adulthood (Luzi et al., 2008; Di Forti et al., 2009; Fisher et al., 2009; Smith et al., 2009; Barkus and Murray, 2010; Mazzoncini et al., 2010; Stilo and Murray, 2010; Casadio et al., 2011; Paparelli et al., 2011; Donoghue et al., 2012), we also examined the contribution of comorbid drug use to MA psychosis, specifically cannabis. In this data set the prevalence of cannabis use did not differ between the MAP+ and MAP-subjects and there was no significant relationship between the frequency of MA psychosis and the age at which they first used cannabis. There was however a statistical trend towards increased frequency of MA-induced psychotic episodes among those MAP+ subjects with a history of cannabis use. However, as the MAP+ participants had longer cannabis abstinence than the MAP- participants, it is unclear how strong the relationship is between cannabis use and psychosis in this sample. A recent meta-analysis of studies that examined the role of cannabis use on psychosis suggests that the reported link between cannabis use and psychotic symptoms is inconsistent (Zammit et al., 2008).

Although MA exposure does not appear to differ between the MAP+ and the MAP- subjects, mean daily lifetime dosage does appear to correlate with the frequency of which the psychotic symptom occurs. This might suggest that pharmacological exposure alone may not be the sole factor that produces MA psychosis, but might interact with other factors that contribute to an increased frequency of psychotic symptoms. If MA exposure alone were sufficient to produce MAP, then why do approximately 40% of chronic MA abusers never

experience MA-induced psychotic episodes? This paradox points to the need to search for other factors.

Although genetic risk factors clearly play a role (Bousman et al., 2009), further studies are needed to determine if the development and recurrence of MA psychosis is linked to abnormal brain function in regions such as the prefrontal and temporal cortices, both of which are often implicated in the emergence of non-substance induced psychotic symptoms (e.g., schizophrenia) (Fusar-Poli et al., 2011a; Fusar-Poli et al., 2011b; Fusar-Poli et al., 2012). ADHD is associated with dysfunction in cortical circuits supporting attention and executive function including frontal parietal attentional networks and the anterior cingulate cortex. These circuits are also disrupted in MA abusing individuals. In rodent models (Pycock et al., 1980), as well as in individuals with schizophrenia, impaired prefrontal function is associated with increased dopamine release in the striatum (Tost et al., 2010) and excessive DA release in the striatum in schizophrenia is associated with clinical symptoms of psychosis (Laruelle et al., 1999; Abi-Dargham et al., 2000). In the context of this literature the present results suggest that childhood ADHD, with its associated deficits on cortical attention circuitry that persists throughout adolescence and into adulthood, may render MA abusing individuals more vulnerable to psychosis by limiting their ability to modulate excessive sub-cortical dopamine release. Future studies comparing the function of the neural circuitry modulating attention and the function of sub-cortical dopamine system will enable a test of these hypotheses and inform potential treatments (Grelotti et al., 2010).

4.1. Limitations

There are several limitations with the use of retrospective measures, especially in clinical populations. They include but are not limited to: 1) memory impairments; 2) report bias; and 3) subjective nature of the instrument. It is possible that a subgroup of MA abusers may have overestimated both psychotic symptoms and ADHD-relevant childhood behaviors in childhood, and others may underestimated them. Such response patterns could explain the correlations observed in the present study. It is also possible that substance abusers may be limited in their ability to accurately recollect childhood attention behaviors. Although we believe that such a recollection bias is unlikely to have played a role in our results, such concerns can be addressed by future studies designed specifically to control for such confounds. If such a bias would be present, it is unclear why it would manifest in a unique correlation with severity of psychosis and not with the other self-report variables that were measured.

As our participants were recruited from treatment and residential housing programs, it may limit the generalizability of our findings to the entire community of individuals with methamphetamine dependence. Finally, although the correlational patterns reported within the current study cannot imply causality, the data nonetheless reveal an interesting relationship between both the quantity of MA consumed and measures of early attention/hyperactivity and the emergence of frequent psychotic episodes in individuals who abuse MA heavily in adulthood and reach criteria for dependence.

4.2. Conclusion

The results suggest the existence of possible behavioral markers reflecting an early cognitive vulnerability to the development of frequent MA-induced psychotic symptoms. These behavioral markers may interact with drug exposure in individuals with other pre-existing genetic and cortical vulnerabilities. Understanding and identifying these cortical risk factors for MA psychosis will provide insight into the general mechanisms of psychosis as well as serve as a foundation to detect MAP susceptibility and the development of targeted clinical interventions.

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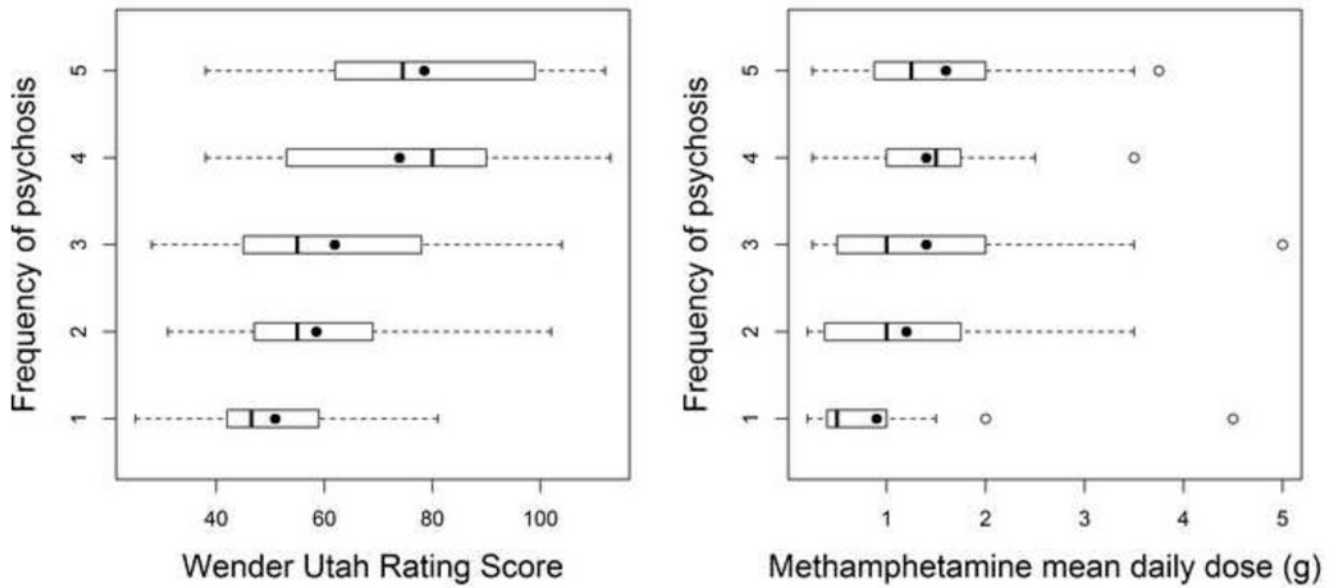


Figure 1.

Association between Wender Utah Rating Scale of childhood attention and methamphetamine daily dose with frequency of psychosis measured on a Likert scale. Box plots define the values for median, range, 25th and 75th percentiles. Means are represented by filled circles. The proportional odds model was used to calculate the odds (OR) of developing more frequent psychosis for higher Wender Utah scores (OR = 1.04, $p < 0.0001$) and larger methamphetamine daily dose (OR = 1.61, $p = 0.003$), respectively.

Table 1
Summary of the demographic and clinical characteristics of the methamphetamine abusers with psychosis (MAP+) and without psychosis (MAP-)

	MAP+ (n = 145)	MAP- (n = 45)	Test Statistic	p-value
<i>Demographic Characteristics</i>				
Age	37.6 (8.2)	37.7 (8.4)	$z = -0.06$	0.95
Females	66 (46%)	25 (56%)	$\chi^2 = 1.39$	0.24
Education (years)	12.2 (2.0)	12.6 (1.8)	$z = 1.30$	0.20
Parental Education (years) ¹	13.0 (2.6)	13.2 (2.9)	$z = 0.46$	0.65
NART ²	105.6 (5.6)	107.3 (6.4)	$t_{183} = 1.71$	0.09
Family History of Psychiatric Disorder ³	45 (31%)	9 (21%)	$\chi^2 = 1.78$	0.18
<i>Clinical Characteristics</i>				
WURS score ¹	64.9 (21.8)	59.4 (20.7)	$z = -1.41$	0.16
Methamphetamine use				
Age of first use	18.8 (6.0)	18.8 (6.1)	$z = -0.11$	0.91
Years of use	14.1 (6.7)	14.2 (7.2)	$z = 0.06$	0.95
Mean Dose (g)	1.3 (1.0)	1.2 (0.9)	$z = -1.17$	0.24
Months Abstinent	18.2 (24.2)	14.5 (23.0)	$z = -1.72$	0.09
Alcohol use				
Alcohol Abuse/Dependence	81 (56%)	26 (58%)	$\chi^2 = 0.05$	0.82
Years Alcohol clear	10.1 (7.9)	10.2 (8.3)	$z = 0.09$	0.93
Tobacco use				
Tobacco smokers	121 (83%)	37 (82%)	$\chi^2 = 0.04$	0.85
Pack years	14.8 (11.3)	16.0 (9.9)	$z = 0.89$	0.37
Cocaine use				
Cocaine Abuse/Dependence	41 (28%)	8 (18%)	$\chi^2 = 1.98$	0.16
Years Cocaine clear	15.7 (8.0)	18.3 (7.6)	$t_{46} = 0.97$	0.34
Cannabis use				
Cannabis Abuse/Dependence	113 (78%)	32 (71%)	$\chi^2 = 0.88$	0.35
Age of first use	14.2 (2.6)	14.7 (4.7)	$z = -0.77$	0.44
Years of Cannabis use	12.2 (9.0)	15.1 (9.9)	$z = 1.55$	0.12
Years Cannabis clear	9.7 (9.6)	6.5 (8.5)	$z = -2.04$	0.04

Note: continuous variables are summarized as mean (SD) and categorical ones as frequency (percent). Group differences were assessed using Chi-squared (χ^2) tests for categorical variables, two-sample t-test the continuous variables whose distribution was normal, and Wilcoxon two-sample (z) tests for the continuous variables who violated the assumption of normality.

¹Frequency missing = 3 in MAP+ sample

²Frequency missing = 3 in MAP+ sample and 2 in the MAP- sample

³Frequency missing = 2 in MAP+ sample and 2 in the MAP- sample

Table 2
Results of the unadjusted proportional odds models examining the association between demographic and clinical characteristics of the 145 MAP+ methamphetamine abusers with the frequency of psychotic episodes

	OR (95% CI)	<i>p</i> -value
<i>Demographic Characteristics</i>		
Age (years)	1.02 (0.98 – 1.06)	0.28
Female Gender	0.93 (0.52 – 1.66)	0.81
Education (years)	0.99 (0.86 – 1.15)	0.94
Parental Education (years) ¹	0.93 (0.83 – 1.05)	0.24
NART	0.99 (0.94 – 1.05)	0.82
Family History of Psychiatric Disorder ²	1.16 (0.62 – 2.17)	0.64
<i>Clinical Characteristics</i>		
WURS score ¹	1.04 (1.02 – 1.05)	< 0.0001
Methamphetamine use		
Age of first use	1.01 (0.96 – 1.05)	0.84
Years of use	1.02 (0.98 – 1.06)	0.40
Mean Dose (g)	1.61 (1.17 – 2.20)	0.003
Months Abstinent	1.00 (0.99 – 1.01)	0.61
Alcohol Abuse/Dependence	1.01 (0.57 – 1.80)	0.98
Tobacco use	2.00 (0.92 – 4.39)	0.08
Cocaine Abuse/Dependence	0.94 (0.49 – 1.78)	0.84
Cannabis Abuse/Dependence	2.00 (0.99 – 4.04)	0.06

Note: unadjusted models with frequency of psychosis as a dependent variable were fit for each of the demographic and clinical characteristics

¹Frequency missing = 3

²Frequency missing = 2

Table 3
Frequency of Psychosis measured on a Likert Scale, mean Wender Utah score and mean daily methamphetamine (MA) dose across the range of frequency of psychosis

Frequency of Psychosis in Likert Units	<i>n</i>	Wender-Utah (WURS)	Mean daily MA dose
1	23	50.9	0.88
2	27	58.5	1.16
3	38	61.9	1.43
4	33	73.9	1.40
5	24	78.5	1.64