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# Delay Discounting in Opioid Use Disorder: Differences between Heroin and Prescription Opioid Users

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# Abstract

**Background**—Among those with opioid use disorder, heroin use is associated with poorer prognosis relative to use of prescription opioids alone. However, relatively little is known about distinguishing features between those who use heroin relative to those who use prescription opioids. In the present study we evaluated differences in delay discounting in those with opioid use disorder based on primary opioid of use. Delay discounting is associated with a range of negative outcomes and is an important therapeutic target in this population.

**Methods**—Treatment-seeking adults with opioid dependence completed self-report measures including past-month opioid use and the Monetary Choice Questionnaire (Kirby and Marakovic, 1996; Kirby et al., 1999), a measure of delay discounting. Participants were divided into two groups based on whether they used any heroin in the past 30 days or only prescription opioids, and delay discounting scores were compared between the groups. Group differences in sociodemographic or clinical variables were included in the analysis as covariates.

**Results**—Results from a forward stepwise linear regression indicated that heroin use was associated with significantly higher delay discounting (B = -0.99,  $SE_B = 0.34$ , t = -2.88, p = .005), even when considering covariates.

**Conclusions**—Adults with opioid dependence who exclusively used prescription opioids had lower delay discounting relative to those who used heroin. This finding contributes further to the

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**Conflict of Interest** 

Dr. Weiss is has served as a consultant to Indivior, U.S. WorldMeds, and GW Pharmaceuticals. All other authors declare they have no conflicts of interest.

Contributors

Drs. Weiss, McHugh, and Griffin designed the original study. Dr. McHugh and Ms. Karakula designed this secondary analysis. Ms. Karakula and Dr. McHugh undertook the statistical analysis, and wrote the first draft of the manuscript. Dr. Weiss, Dr. Griffin, Ms. Borges, and Mr. Bailey contributed to the interpretation and presentation of study findings. All authors contributed to and have approved the final manuscript.

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literature suggesting that heroin use is associated with greater clinical severity among those with opioid use disorder.

#### Keywords

opioid use disorder; prescription opioids; heroin; delay discounting; impulsivity; addiction

# **1. INTRODUCTION**

The prevalence of heroin use and heroin use disorders has risen sharply in recent years, with the rate of past-year heroin use disorders almost doubling between 2002 and 2013 in the U.S. (Jones, 2013; Jones et al., 2015). Following years of increases in prescription opioid misuse, many new heroin users report having initiated opioid use with opioid analgesic medications (Cerda et al., 2015; Jones, 2013; Novak et al., 2016). The initiation of heroin use in this population is associated with a significantly poorer prognosis (Brands et al., 2004; Moore et al., 2007; Potter et al., 2013; Rosenblum et al., 2007; Sigmon, 2006; Weiss et al., 2011). However, little is known about mechanisms that might explain differences between prescription opioid and heroin users, or facilitate the transition from prescription opioids to heroin.

One major difference between prescription opioids and heroin is that prescription opioids can be obtained legally for medical purposes. This may foster the belief that prescription opioids are less risky than heroin (Daniulaityte et al., 2012; Lord et al., 2011). Accordingly, greater impulsivity may be a distinguishing feature of those who use heroin relative to those who use only prescription opioids. Those who are more impulsive perceive risky behaviors to be less harmful (Arria et al., 2008; Doran et al., 2011; Robbins and Bryan, 2004), and may be more likely to engage in such behaviors, even adjusting for perceived risk (Ryb et al., 2006).

The tendency to over-value immediate relative to delayed reward, i.e., delay discounting, is a common feature of substance use and other addictive disorders (Bickel et al., 2014; MacKillop et al., 2011; Reynolds, 2006). Those with higher delay discounting act quickly on current needs, while devaluing future needs. This facet of impulsivity is particularly notable in opioid use disorder; heroin users not only show a dramatically larger preference for immediate rewards relative to healthy comparison samples (Kirby et al., 1999; Madden et al., 1997), but also have higher discounting rates compared to people who abuse alcohol, and rates similar to those who abuse cocaine (Kirby and Petry, 2004). Delay discounting has also been associated with negative treatment outcome in a variety of substance use disorder populations (MacKillop and Kahler, 2009; Stanger et al., 2012; Washio et al., 2011). However, only one published study has examined delay discounting among those who are dependent upon prescription opioids (Yi et al., 2007). To date, no published studies have examined differences in delay discounting between those with opioid use disorder who use prescription opioids vs. heroin.

The objective of the present study was to compare levels of delay discounting in a sample of adults with opioid use disorder, based on the type of opioid abused in the past month. We

hypothesized that heroin users would show elevated delay discounting relative to prescription opioid users.

### 2. METHODS

#### 2.1. Participants

A sample of 151 participants seeking treatment for opioid use disorder was recruited from an inpatient detoxification unit as part of a larger study. In this sample, 139 participants completed the delay discounting measure. Eligible participants were 18 years or older, met *Diagnostic and Statistical Manual of Mental Disorders, 4th Edition* (American Psychiatric Association, 2000) criteria for opioid dependence, and were undergoing a buprenorphine stabilization and taper protocol as part of medical detoxification at the time of study participation. Exclusion criteria included presence of an acute psychiatric or medical disorder or cognitive impairment that would impede the ability to perform study procedures. Those with co-occurring substance use disorders and/or other non-acute psychiatric disorders were eligible to participate; no specific disorders were excluded.

#### 2.2. Procedure

All procedures were approved by the local Institutional Review Board. Participants who provided written informed consent completed a battery of self-report questionnaires during one 30-minute session.

Participants were categorized into one of three groups: 1) heroin users (heroin use on 4 days in the 30 days prior to hospitalization, with <4 days of prescription opioid use during that time), 2) prescription opioid users (prescription opioid use on 4 of the 30 days prior to hospitalization, with <4 days of heroin use during that time), or 3) combined heroin and prescription opioid users ( 4 days of both heroin and prescription opioid use in the 30 days prior to hospitalization).

#### 2.3. Measures

Participants self-reported demographic information. Substance use disorder and other psychiatric diagnoses were extracted from participants' medical records.

The 17-item Brief Addiction Monitor (BAM; Cacciola et al., 2013) was used to collect information about past-month substance use. The BAM has shown good test-retest reliability as well as strong predictive validity (Cacciola et al., 2013).

The Monetary Choice Questionnaire (MCQ; Kirby and Marakovic, 1996; Kirby et al., 1999) is a 27-item self-report measure that was used to characterize participants' rates of delay discounting. Each item presents a choice between two monetary rewards—one smaller, immediate reward and one larger, delayed reward (e.g., "Would you prefer \$19 today or \$25 in 53 days?"). "Discounting rates," or k values, represent the rates at which individuals devalue rewards based on their delay, and can be inferred from individuals' patterns of choices across items. Estimation of k values was performed according to standardized scoring procedures that utilize Excel-based software (Kirby, 2000; Kirby et al., 1999). Inconsistencies in the data, such as nonsystematic discounting, are accounted for by

estimating k using a participant's observed pattern of choices rather than his or her performance on a single item (Kirby, 2000). These procedures result in the k value that most likely gave rise to the observed data; these values can range from 0.00016 to 0.25, with higher k values indicating a stronger preference for smaller, immediate rewards and thus a more impulsive decision-making style. Consistent with previous literature (Kirby et al., 1999), k scores were log-transformed to approximate a normal distribution. The MCQ has demonstrated strong retest reliability (Kirby, 2009).

#### 2.4. Data Analysis

The three groups of study participants were compared on sociodemographic and drug use variables using univariate analysis of variance (ANOVA) for continuous variables and chisquare tests for categorical variables. Variables for which differences were statistically significant were added to the main study analysis.

These groups were then collapsed into two groups: prescription opioid use only and any heroin use. We then conducted a forward stepwise regression examining the association between opioid type and delay discounting (log-transformed k score). This approach was used to account for the potential confounding effects of covariates, while also mitigating concern about high overlap among independent variables.

# 3. RESULTS

Descriptive statistics for sociodemographic and drug use variables are presented in Table 1. Based on self-reported opioid use in the 30 days prior to hospital admission, 60 participants (43.2%) were defined as heroin users, 33 (23.7%) were defined as prescription opioid users, and 46 (33.1%) were defined as combined users. The groups differed significantly with respect to age, education, current cigarette smoking, and diagnosis of a stimulant use disorder (see Table 1). Mean *k* scores for heroin users, combined users, and prescription opioid users were .095, .090, and .060, respectively; higher *k* scores denote greater delay discounting (see Figure 1).

Because the heroin and combined users were similar in relation to both opioid use and sociodemographic characteristics, we collapsed these groups to reflect a single heroin-user group. This group was compared with the prescription opioid users, controlling for age, education, cigarette smoking, and diagnosis of a stimulant use disorder. The results of the regression indicated that only opioid group was associated with delay discounting (B = -0.99,  $SE_B = 0.34$ , t = -2.88, p = .005); all other variables were excluded from the model (*p*s ranged from .52 to .74). Specifically, delay discounting was significantly higher among those who used heroin relative to those who exclusively used prescription opioids.

# 4. DISCUSSION

In a sample of adults hospitalized for opioid use disorder, we found that those who used heroin (whether with or without prescription opioids) exhibited greater delay discounting than those who exclusively used prescription opioids. These differences remained statistically significant after controlling for sociodemographic and drug use variables that

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differed between groups. Delay discounting is of significant clinical concern in substanceusing populations due to its association with the initiation of use, addiction severity, increased risk of relapse, and risky behaviors (for review, see Bickel et al., 2014).

This finding adds to the growing literature showing clinically meaningful differences among those with opioid use disorder who use heroin vs. prescription opioids only. A recurring theme in this literature is that heroin use is associated with more severe baseline presentation and a more negative treatment outcome relative to prescription opioid use (Brands et al., 2004; Moore et al., 2007; Nielsen et al., 2015; Potter et al., 2013; Rosenblum et al., 2007; Sigmon, 2006).

Although our cross-sectional design cannot rule out the possibility that heroin use itself led to increases in delay discounting, there is some support in the literature that delay discounting precedes substance use. Audrain-McGovern et al. (2009) found that delay discounting is a trait-like characteristic across adolescence to young adulthood, and that greater delay discounting promotes acquisition of cigarette smoking. Longitudinal research examining this association in opioid users is needed to investigate whether these differences in delay discounting precede heroin use among nonmedical prescription opioid users.

The current findings suggest the potential relevance of targeting delay discounting in the treatment of heroin users, which could entail use of medication or behavioral interventions targeting impulsivity. Several studies have supported the benefits of contingency management for the treatment of opioid use disorders (Everly et al., 2011; Petry and Carroll, 2013; Petry and Martin, 2002), and this treatment appears to be associated with significant decreases in delay discounting (Landes et al., 2012). Other novel strategies for reducing delay discounting, such as the use of cognitive rehabilitation strategies (i.e., working memory training), have shown initial promise for other substances of abuse and are potential candidates for application with opioid use disorder (Bickel et al., 2011). Longitudinal research will help to identify whether such intervention is appropriate for preventing the initiation of heroin use in nonmedical prescription opioid users, for reducing relapse among those with current heroin use, or both.

This study has several limitations. The study was cross-sectional, so we are unable to determine the temporal relationships between heroin use and rates of delay discounting. Second, we do not have lifetime opioid use histories for our participants, and thus do not know whether opioid use was initiated with heroin or a prescription opioid, how long participants had been using opioids, or whether the prescription opioid users had a lifetime history of heroin use. Future research that investigates the lifetime course of opioid use disorder and accounts for the duration of opioid use trajectories. Third, although groups did not differ with respect to prevalence of psychiatric disorders generally, we cannot rule out the possibility that co-occurring disorders affected the findings. Fourth, in our delay discounting task, monetary choices were hypothetical and not related to any actual monetary reward outcome. However, a number of studies that investigated real versus hypothetical rewards in delay discounting procedures found no effect of reward type (Lagorio and Madden, 2005; Madden et al., 2003, 2004), somewhat mitigating this concern. Finally, although the brevity

of the MCQ is a strength, longer measures may afford more precision in estimating delay discounting (Epstein et al., 2003). Future studies also considering other impulsivity measures will help to clarify the link between impulsivity and heroin use. Other future directions may include consideration of the impact of factors such as the quantity and frequency of opioid use.

This study found statistically significant differences in delay discounting between those who used a substantial amount ( 4 days in the past month) of heroin relative to those who exclusively used prescription opioids. This finding provides further evidence for meaningful differences between those who misuse heroin vs. prescription opioids. Identifying risk factors for the initiation of heroin use is critical to early intervention to mitigate the significant hazards of heroin use.

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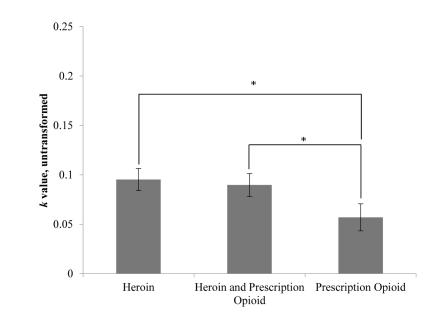
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# Highlights

• Delay discounting is the tendency to overvalue immediate vs. delayed reward.

- Adults with opioid use disorder completed a measure of delay discounting.
- Prescription opioid (PO) users had lower delay discounting vs. heroin users.
- Any use of heroin was associated with greater delay discounting than POs alone.
- Heroin use is associated with a more impulsive style of decision-making.

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#### Figure 1.

Delay Discounting Scores by Primary Opioid. \*p<.05; significance tests reflect univariate ANOVA comparing the natural log of k scores among the 3 groups

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Table 1

Participant Characteristics by Primary Opioid

	Primar	Primary opioid (N=139)		6 / 5	
	Prescription opioid (n= 33)	Heroin $(n=60)$	Combined $(n = 46)$	$\mathcal{X}_{\mathcal{A}}$	<i>p</i> -value
<u>Sociodemographic</u>					
Age (years)	38.3 (14.8)	26.1 (5.4)	28.0 (8.4)	19.01	<.001
Female, %	27.3	26.7	19.6	06.0	.639
Caucasian, % (n=138)	6.06	93.2	97.8		в
Hispanic, % (n=126)	0.0	0.0	10.0		<i>aa</i>
Beyond high school, %	6.06	63.3	58.7	10.44	.005
Employed, % (n=133)	50.0	41.4	35.6	1.55	.461
Past month opioid use, days					
Heroin use	0.0 (0.0)	24.7 (7.1)	24.9 (7.8)	183.60	<.001
Prescription opioid use	24.0 (7.9)	0.4~(0.8)	13.0 (9.4)	140.49	<.001
Other substance use disorders $b, \underline{\%}$					
Alcohol	30.3	23.3	19.6	1.23	.540
Benzodiazepines/other sedatives	15.2	15.0	26.1	2.46	.292
Cocaine/other stimulants	3.0	25.0	17.4	7.19	.027
Cannabis	18.2	20.0	30.4	2.17	.338
Current smoking status, %	45.5	86.7	82.2	21.17	<.001
Axis I psychiatric comorbidity, %	63.6	60.0	67.4	0.61	.736

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 $\boldsymbol{b}_{\mathrm{No}}$  participants met criteria for hallucinogen or inhalant use disorders