

Effects of Extended Cannabis Abstinence on Cognitive Outcomes in Cannabis Dependent Patients with Schizophrenia vs Non-Psychiatric Controls

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Cross-sectional studies of the effects of cannabis on cognition in schizophrenia have produced mixed results. Heavy and persistent cannabis use in schizophrenia is a common clinical problem, and effects of controlled abstinence from cannabis in these patients have not been carefully evaluated. The present study sought to determine the effects of cannabis abstinence on cognition in patients with schizophrenia and co-occurring cannabis dependence. We utilized a 28-day cannabis abstinence paradigm to investigate the state-dependent effects of cannabis on select cognitive outcomes in cannabis-dependent patients with schizophrenia and non-psychiatric controls. Nineteen patients and 20 non-psychiatric male cannabis-dependent participants underwent 28 days of cannabis abstinence. Cognition was assessed on day 0, 14, and 28 using a comprehensive neuropsychological battery. Clinical symptoms were assessed weekly. Abstinence was facilitated by contingency reinforcement confirmed by twice weekly urinalysis. Forty-two percent of patients and 55% of controls achieved end-point abstinence ($p = 0.53$), which was biochemically-verified (day 28 urinary THC-COOH < 20 ng/ml). In this preliminary study, schizophrenia-abstainers demonstrated improvements in Hopkins Verbal Learning Test-Revised (HVLT-R) performance over time [$F(2,14) = 4.73$, $p < 0.03$] ($d = 1.07$). Lesser improvements on HVLT-R were observed in non-psychiatric control abstainers ($d = 0.66$), and with abstinence on other cognitive test measures, in both patients and controls. Verbal memory and learning may improve in schizophrenia and control subjects with cannabis abstinence, but larger more definitive studies are needed. Our findings underscore the importance of developing effective interventions for cannabis use disorders in schizophrenia.

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INTRODUCTION

Cannabis may be a risk factor for the development of psychotic disorders such as schizophrenia (Moore *et al*, 2007). Importantly, rates of cannabis use disorders (CUD) are higher among patients with schizophrenia (~25%) compared to the general population (~2.9%; Hasin *et al*, 2015; Koskinen *et al*, 2010). Cannabis users with schizophrenia are more likely to be male, have an earlier onset of the illness and experience a more severe course of the disorder compared to non-cannabis using patients (Koskinen *et al*, 2010; Large *et al*, 2011; Zammit *et al*, 2008). Clinical evidence demonstrates that cannabis is associated with increased rates of psychotic relapse,

decreased treatment adherence, and poorer psychosocial functioning in patients with schizophrenia (Manrique-Garcia *et al*, 2014; Patel *et al*, 2016; Zammit *et al*, 2008). Accordingly, one might predict that cannabis compromises core cognitive processes in these patients. However, findings remain controversial (Coulston *et al*, 2007b; Rabin *et al*, 2011; Yucel *et al*, 2012), and warrant further investigation.

Surprisingly, several studies report that cannabis-using patients with schizophrenia have superior cognition compared to their non-using counterparts, and two recent meta-analyses examining this relationship support this notion (Rabin *et al*, 2011; Yucel *et al*, 2012). One theory proposed to explain this paradox is that patients with co-morbid CUDs represent a higher functioning subgroup with inherently superior cognition and prognosis (Dixon *et al*, 1991; Leeson *et al*, 2011; Rodriguez-Sanchez *et al*, 2010). These individuals are thought to have better premorbid adjustment, social skills, and lower levels of negative symptoms that enable them to navigate and socialize within drug scenes to maintain their habit (Arndt *et al*, 1992; Potvin *et al*, 2005).

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Other investigators have posited that these individuals possess a lower vulnerability for developing psychosis compared to patients without a history of cannabis use (Schnell *et al*, 2009; Yucel *et al*, 2012). Cannabis may interact with underlying neurobiological vulnerability factors to trigger the transition to psychosis that may not have occurred in the absence of cannabis (Caspi *et al*, 2005). Together, these theories suggest that cannabis is associated with better cognition, rather than cannabis itself leading to improved cognitive function. Thus, better cognition may reflect trait characteristics of this subgroup, a phenomenon not seen among non-psychiatric cannabis users (Broyd *et al*, 2016). Unlike studies in patient populations, there are no reported observations that cannabis using non-psychiatric controls perform better than non-users (Broyd *et al*, 2016). Thus, it is conceivable that cannabis may differentially affect cognitive function in schizophrenia compared to controls.

Evidence that cannabis has a deleterious effect on cognition is well documented (Lev-Ran *et al*, 2012; Ringen *et al*, 2010). Acute administration of tetrahydrocannabinol (THC) worsens, rather than enhances, cognition in patients with schizophrenia specifically in domains of verbal memory and attention (D'Souza *et al*, 2005). Further, these patients were more vulnerable to THC's effects compared to non-psychiatric controls. Results from a previous study from our group demonstrated that cumulative cannabis use dose-dependently impaired cognitive performance in tasks mediated by the prefrontal cortex and the hippocampus in current, but not former cannabis-dependent patients who were abstinent for at least 6 months (Rabin *et al*, 2013). Taken together, these findings suggest that cannabis exerts a state-dependent negative effect on neuropsychological performance facilitated by brain regions rich in cannabinoid type-1 receptors (CB1Rs; Mackie, 2005). Moreover, these cannabis-induced impairments may be reversible with sustained periods of abstinence.

Explanations for conflicting results on effects of cannabis on cognition in schizophrenia may be due to methodological limitations in published studies. For example, many studies have failed to control for the period of time elapsed between the last use of cannabis and neuropsychological testing. Depending on this interval, studies may be assessing the impact of acute cannabis intoxication (D'Souza *et al*, 2005), withdrawal effects (Coulston *et al*, 2007a), or the residual or longer lasting consequences of cumulative cannabis exposure (Jockers-Scherubl *et al*, 2007; Schnell *et al*, 2009). Moreover, all research studies examining the relationship between cannabis and cognition in schizophrenia have employed cross-sectional designs, making it difficult to draw firm conclusions about direct effects of cannabis on cognitive function. In fact, studies may be simultaneously assessing both the state- and trait-dependent effects of cannabis. This is problematic given the premise that they may be associated with opposite effects on cognitive function. A more reliable and robust method is to employ a longitudinal design to examine within-subject differences related to cannabis abstinence and reinstatement of use.

In the present study, we employed a 28-day cannabis abstinence paradigm to more appropriately assess the long-term state-dependent effects of cannabis on key cognitive outcomes, within regions of high CB1R density, in cannabis-dependent patients with schizophrenia and non-psychiatric

controls. More specifically, our main outcome of interest was verbal learning and memory given its high sensitivity to cannabis (D'Souza *et al*, 2005; Rabin *et al*, 2013) and documented impairment in schizophrenia (Heinrichs and Zakzanis, 1998). Secondary aims of the study examined other areas of moderate to high cannabinoid density, such as the prefrontal cortex and the cerebellum (Herkenham, 1991).

Therefore, we hypothesized that cognitive performance as assessed by the HVLT would improve over time in patients with schizophrenia who successfully remained abstinent from cannabis for the 28-day abstinent period. In addition, we expected greater magnitude of cognitive change in abstaining patients compared to abstaining controls given that patients with schizophrenia demonstrate enhanced sensitivity to the cognitive-impairing effects of cannabis (D'Souza *et al*, 2005). The 28-day abstinence period was based on the unique pharmacokinetic profile of THC (Nahas, 2001). This duration reflects the time needed for full cannabinoid elimination (ie, including residual cannabinoids) in order to achieve biochemically confirmed abstinence at a standardized cut-off of 20 ng/ml (Ellis *et al*, 1985; Smith-Kielland *et al*, 1999).

MATERIALS AND METHODS

Participants

Patients with schizophrenia were recruited through the Centre for Addiction and Mental Health (CAMH) using flyers posted around the hospital and through referrals made by outpatient clinicians. Non-psychiatric cannabis users were recruited from the community by posted ads. Study eligibility was assessed by an initial telephone screening, followed by an in-person comprehensive interview. Recruitment began in April 2012 and ended by December 2015.

Male participants between the ages of 18 and 55 were recruited for the study. All participants met criteria for current cannabis dependence based on DSM-IV-TR (APA, 2000). A positive urine test for THC-COOH (MEDTOX; Wilmington, NC) was required to confirm current cannabis use. To control for the effects of tobacco on cognition, all participants were daily cigarette smokers (≥ 5 cigarettes per day, CPD). Moreover, all participants had to achieve Full Scale Intelligent Quotient (FSIQ) scores ≥ 80 , using the Wechsler Adult Reading Test (WTAR; Wechsler, 2001). Psychiatric participants met DSM-IV-TR criteria for schizophrenia or schizoaffective disorder; we excluded subjects with schizophreniform disorder and psychosis NOS. Patients were psychiatrically stable at the time of the interview with a total score < 70 on the Positive and Negative Syndrome Scale for Schizophrenia (PANSS; Kay *et al*, 1987) and no hospitalizations in the 3 months prior to enrollment. In addition, patients had to be maintained on a stable dose of antipsychotic medication with no changes for at least 1 month. Non-psychiatric controls were excluded if they met criteria for a current or past DSM-IV Axis I diagnosis (except for major depression in remission > 1 year) or if they were taking psychotropic medications. Individuals with a current substance use disorder (SUD) or past (remission < 6 months) SUD (other than cannabis, nicotine, and caffeine) or those testing positive on urine toxicology for illicit drugs other than cannabis (ie, cocaine, opiates,

amphetamine, phencyclidine, and barbiturates) were not eligible for study participation. In addition, participants were excluded if they were actively seeking treatment for cannabis. Head injury with loss of consciousness for >30 min requiring hospitalization, or evidence of intracranial injury or neurological/medical conditions affecting cognition was also exclusionary. Written informed consent was obtained from all participants, as approved by the Research Ethics Board at CAMH.

Measures

Substance use measures. Current cannabis dependence, past alcohol and other SUD were diagnosed using DSM-IV-TR criteria. Cumulative cannabis exposure was indexed as joint-years, where one joint-year is the equivalent of using one joint per day for 1 year (Rabin *et al*, 2013). The Timeline Follow-Back (Sobell *et al*, 1988; TLFB) is a self-report of substance use frequency and was collected for cannabis in grams, tobacco cigarettes, alcoholic beverages and caffeine in the seven previous days. Level of nicotine dependence was measured using the Fagerstrom Test of Nicotine Dependence (FTND) (Heatherton *et al*, 1991) and the Alcohol Use Disorders Identification Test (AUDIT; Saunders *et al*, 1993) assessed problematic drinking behaviors. Lastly, cannabis withdrawal was assessed using the Marijuana Withdrawal Checklist (MWC; Budney *et al*, 2003).

Neuropsychological test battery. All participants completed a comprehensive neuropsychological test battery. Neuropsychological test measures included those with demonstrated sensitivity in patients with schizophrenia and/or cannabis users. Cognitive sessions took an average of 2.5 h to complete, and were administered by R.A.R. The Test of Memory Malingering (TOMM) (Tombaugh, 1997), a measure of effort and motivation (Sharland and Gfeller, 2007) was only administered at the neuropsychological training session to ensure credibility of performance across other measures. Similarly, the Wisconsin Card Sorting Test (WCST; Heaton *et al*, 1993) and Iowa Gambling Test (IGT; Bechara *et al*, 1994), measures of executive function and decision-making respectively were also only administered at the training session. Other tests, less sensitive to repeated testing, were administered at Days 0, 14, and 28 and included: Spatial Delayed Response task (SDR; Hershey *et al*, 1998), a measure of visuospatial working memory; Digit Span Forwards and Backwards (Wechsler, 1997), a measure of working memory and executive function respectively; Continuous Performance Test II (CPT-II; Conners, 2000), an assessment of sustained attention; Trail Making Test A and B (TMT; Lezak *et al*, 2004) to evaluate psychomotor speed and executive function respectively; Grooved Pegboard (Lafayette Instrument Company, 1989) to measure manual dexterity and fine motor movement; Balloon Analog Risk Task (BART; Lejuez *et al*, 2002) a computerized measure of risk taking behavior; and lastly, Kirby Delayed Discounting Test (KDDT; Kirby *et al*, 1999) to assess impulsive choices. We utilized the Hopkins Verbal Learning Test Revised (HVLT-R; Brandt and Benedict, 2001) to assess verbal memory and learning. Six alternate word lists from this test were administered in a counterbalanced order.

Clinical Measures

In patients with schizophrenia, positive and negative symptoms were assessed using the PANSS (Kay *et al*, 1987) and extrapyramidal symptoms were measured with the Simpson Angus Rating Scale (SARS) (Simpson and Angus, 1970), the Abnormal Involuntary Movement Scale (AIMS; Guy and Cleary, 1976), and the Barnes Akathisia Rating Scale (BARS; Barnes, 1989). The Calgary Depression Scale for Schizophrenia (CDSS; Addington *et al*, 1993) assessed depression exclusively in patients and the Hamilton Rating Scale for Depression (HAM-D; Hamilton, 1967) assessed mood symptoms in both patients and controls.

Laboratory Procedures

Once eligibility was confirmed, participants were invited to the lab for a neuropsychological training visit to familiarize participants with test measure procedures so as to minimize practice effects associated with repeated testing (Sacco *et al*, 2006). Participants were instructed to abstain from cannabis for 12 h prior to this visit to minimize the possibility of cannabis intoxication as well as withdrawal (Budney *et al*, 2003). Given that acute periods of abstinence cannot be biochemically confirmed, this was done through clinical observation (ie, conjunctival injection, euphoria, giggling, sedation, and lethargy (APA, 1994)). Participants were then scheduled to quit cannabis the night before coming into the lab for the day 0 (baseline) visit for a full 28-day period. Participants attended weekly study visits. Clinical measures assessing psychiatric, depressive, and withdrawal symptoms were assessed weekly, while cognitive function was evaluated biweekly, on days 0, 14, and 28. Urine was collected twice weekly and then stored in a -80°C freezer for future gas chromatography mass spectrometry (GC-MS) analysis. To encourage cannabis abstinence individual supportive therapy was given weekly (20–30 min) by trained clinical staff in the Schizophrenia Division at CAMH. Sessions included a combination of motivational interviewing, psycho-education, and coping skills. Contingency management was used as the primary reinforcer: participants who successfully abstained from cannabis for the full 28 days (MEDTOX; THC-COOH $<50\text{ ng/ml}$) were rewarded with a \$300 bonus. Four weeks later (day 56), participants attended a follow-up study visit, where both clinical and cognitive outcomes were evaluated. See Figure 1.

Abstinence Verification

Abstinence verification was based on meeting all three of the following criteria:

- (1) No self-reported cannabis use for 28 days based on TLFB.
- (2) Biochemical confirmation that no new cannabis was introduced over the abstinence period. GC-MS analysis was conducted to calculate THC-COOH/creatinine ratios on 9 urine samples collected twice weekly (baseline+2 samples per week, for 4 weeks). Urine samples were collected at the weekly study visit and the other sample was collected 3 days later. Each ratio was divided by the previously collected sample quotient (urine2/urine1). A prediction model was applied to these

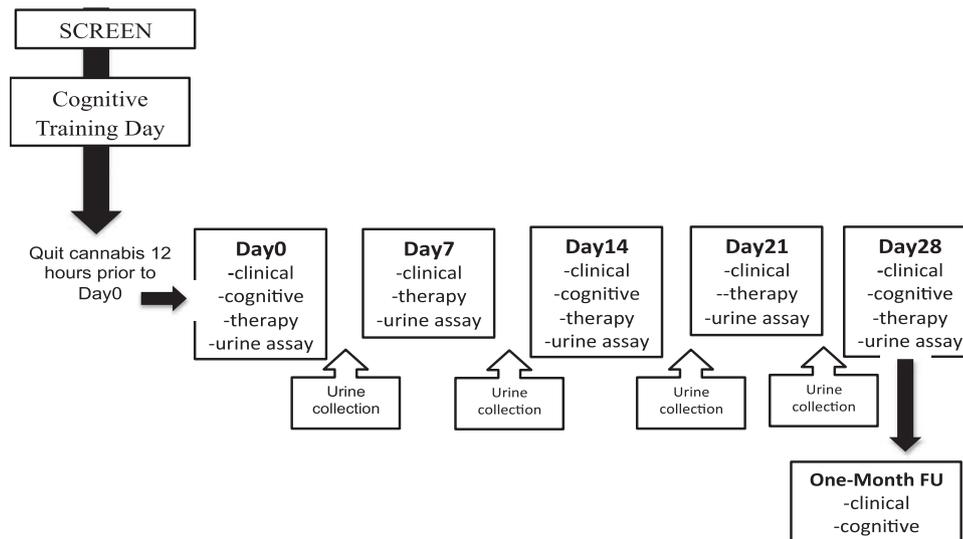


Figure 1 Study Design. Outline of the study design including screening and training visit, weekly assessments and twice weekly urine collections.

quotients to biochemically determine whether new cannabis use was introduced during the 28-day abstinence period (Schwilke *et al*, 2011).

- (3) No THC-COOH was present in urines at Day 28. A sensitivity cut-off of <20 ng/ml at day28 was employed (Ellis *et al*, 1985).

Data Analysis

Independent *t*-tests and χ^2 tests were used to analyze demographic variables between cannabis-dependent patients with schizophrenia and cannabis-dependent non-psychiatric controls. Between group differences on baseline neuropsychological measures were assessed using ANCOVAs controlling for education level and IQ. Bonferroni corrections were performed to account for multiple comparisons of cross-sectional data comparisons ($\alpha/41$, $p=0.001$).

$2 \times 2 \times 3$ repeated measures of analysis of co-variance (RM-ANCOVA) were used to assess change over time for cognitive and clinical variables in patients with schizophrenia and controls. Time was the within-subject factor, and diagnosis and abstinence status were employed as the between-subject factors. This analytic approach was adopted in order to capture time \times abstinence status interactions and time \times diagnosis \times abstinent status interaction. Time \times abstinence status evaluated potential differences in recovery in the sample as a whole and time \times diagnosis \times abstinence status evaluated. Whether abstaining and non-abstaining patients cognitively behave in a differential manner compared to non-psychiatric controls. Separate models were estimated for each cognitive and clinical outcome. When significant differences were detected, post-hoc univariate ANOVAs were conducted. Main effects of time were followed up with LSD *post hoc* tests to detect between which time-point significant change occurred. Because of group differences in education level and IQ, these were controlled for in separate analyses presented in the Supplementary Material (see Table 1). Effect sizes (Cohen's *d*) were computed to determine the magnitude of change between baseline and day 28 in patient-abstainers (see Supplementary Material,

Table 3). Difference scores in cognitive outcomes between baseline and day 28 were computed. MANCOVAs were conducted with the cognitive difference score as the dependent variable and diagnostic group and abstinence status as fixed factors to see if cognitive change differed by diagnostic group and/or abstinence status (Supplementary Material). Finally, we also conducted Linear Regression models in patients and controls separately to determine whether duration of abstinence (ie, days of sustained abstinence) predicted change in cognitive performance between baseline (day 0) and day 28. These results are now included in the Supplementary Material (Table 2).

Data were analyzed using the Statistical Program for Social Sciences (SPSS) version 24.0 (SPSS Inc., Chicago, IL). All tests were two-tailed and the level of significance was set at $p < 0.05$. Our *a priori* hypothesis based on previous studies (Rabin *et al*, 2013) was that verbal memory and learning as measured by the Hopkins Verbal Learning Test (HVLT) would be altered by cannabis abstinence. Therefore, Bonferroni corrections were applied to account for multiple time-point comparisons ($\alpha/3$, $p=0.0167$).

RESULTS

Sample Demographics

Nineteen cannabis-dependent patients with schizophrenia and 20 cannabis-dependent controls completed the study. A CONSORT diagram to outline subject disposition is provided in Figure 2. The completion rate of the 4-week abstinence period was 83%. The difference in attrition rates between patients and controls did not statistically differ $\chi^2 = 3.50$ ($df=1$), $p > 0.06$. The patient group had a 5% attrition rate, while the controls had a 25% attrition rate. Notably, the majority of controls ($n=5$) dropped out after the screening visit, before cannabis abstinence was initiated.

Table 1 Demographic, Clinical and Substance-using Characteristics

	SCZ (n = 19)	CTL (n = 20)	p-value
<i>Demographic variables:</i>			
Age (years)	31.58 ± 9.1	30.80 ± 8.1	0.78
Race (C/A/O) ^a	8/8/3	15/3/3	0.10
FSIQ	91.21 ± 8.6	101.60 ± 9.2	<0.01*
Education (years)	10.97 ± 1.8	13.35 ± 2.4	<0.01*
<i>Clinical variables:</i>			
CPZ equivalents	352.42 ± 194.0	NA	NA
Atypical/typical antipsychotics	16/2	NA	NA
PANSS positive	13.84 ± 3.9	NA	NA
PANSS negative	13.11 ± 4.0	NA	NA
PANSS General	25.8 ± 44.3	NA	NA
PANSS total	52.84 ± 9.9	NA	NA
CDSS	2.63 ± 2.8	NA	NA
HAM-D	4.37 ± 2.9	2.75 ± 3.1	0.06
<i>Substance-using variables:</i>			
Joint-years	10.10 ± 7.3	9.76 ± 6.6	0.88
GPD	1.22 ± 0.8	1.63 ± 1.2	0.21
Baseline THC-COOH:Cr	49.00 ± 47.7	100.26 ± 104.2	0.12
(\$)/Week on cannabis	43.95 ± 36.2	64.75 ± 58.6	0.19
Age of first use of cannabis	15.00 ± 2.5	15.05 ± 2.9	0.95
Age of onset of regular (weekly) cannabis use	17.05 ± 3.6	18.25 ± 5.2	0.41
MWC	10.58 ± 6.8	7.55 ± 7.2	0.19
CPD	12.56 ± 7.3	11.19 ± 10.6	0.64
CO level	19.42 ± 6.9	17.05 ± 10.7	0.42
FTND	4.95 ± 2.1	3.30 ± 2.8	<0.05*
AUDIT	5.70 ± 4.4	6.10 ± 3.7	0.78
Alcoholic drinks/week	0.50 ± 0.6	0.50 ± 0.6	0.81
Caffeinated beverages/week	2.05 ± 2.9	2.69 ± 2.2	0.44

Abbreviations: A, African; AUDIT, alcohol use identification test; C, Caucasian; CDSS, Calgary Depression Scale for Schizophrenia; CO, carbon monoxide; CPD, cigarettes per day; CPZ equivalents, chlorpromazine equivalents; FTND, Fagerstrom test of nicotine dependence; GPD, average grams of cannabis per day; HAM-D, Hamilton Depression Rating Scale; O, other race; MWC, marijuana withdrawal checklist; PANSS, Positive and Negative Syndrome Scale; THC-COOH:Cr, carboxy-tetrahydrocannabinol (THC) normalized to creatinine. Values given in mean ± SD; a, values are in numbers; *p < 0.05.

Thus, it was not cannabis abstinence itself that led to attrition. After the baseline visit (post-quit) only one patient and two controls dropped-out.

Demographic and clinical variables are presented in Table 1. According to DSM-IV, 14 patients met criteria for schizophrenia and five patients met criteria for schizoaffective disorder. (collectively designated 'schizophrenia' patients). We included patients with schizophrenia with past Axis 1 disorders [anxiety (*n* = 1) and MDD (*n* = 1)] as well as those with past SUD in remission for at least 6 months (other than cannabis). In the same respect, we included controls that had an SUD in remission for at least 6 months, and included controls that met for MDD in full remission (*n* = 2). Because of our within-subjects design, these past

diagnoses should not have influenced cognitive outcome trajectories.

Patients and the non-psychiatric control group did not differ significantly on age and race. Group differences emerged on IQ [$t(37) = 3.64$, $p < 0.01$] and years of education [$t(37) = 3.45$, $p < 0.01$], with controls demonstrating higher IQ and greater number of years of education. On the CDSS at baseline, patients with schizophrenia had mean scores of 2.6 ± 2.8 .

The majority of patients were taking atypical antipsychotics [clozapine (*n* = 1), risperidone (*n* = 7), olanzapine (*n* = 4), quetiapine (*n* = 2), and paliperidone (*n* = 3)]. One patient was taking a typical antipsychotic (flupenthixol) and one patient was on a combination of an atypical and typical medication (quetiapine+fluphenazine). Chlorpromazine equivalents are listed in Table 1. Patients were also prescribed medication for their mood [citalopram (*n* = 2), escitalopram (*n* = 1) and valproate (*n* = 1)], and one patient was prescribed a benzodiazepine (lorazepam). Patients were taking medication for hypertension (*n* = 2), Type 2 diabetes (*n* = 3) and hypercholesterolemia (*n* = 3). Among controls, only one participant was taking medication, and this was for hypercholesterolemia (rosuvastatin).

Substance Use Characteristics

Substance use characteristics are listed in Table 1. Groups were matched on cannabis-using variables such as cumulative use (joint-years), grams of cannabis used in the previous week (GPD), money spent on cannabis in the prior week and baseline THC-COOH/creatinine levels. Age of onset of weekly cannabis use was also similar between groups, as were the mean baseline scores for MWC. On FTND, patients demonstrated higher levels of nicotine dependence compared to controls [$t(37) = 1.98$, $p = 0.04$]. Interestingly, the average number of cigarettes per day did not differ between groups. Problematic alcohol use and alcoholic beverages consumed over the previous week did not differ between groups. Similarly, caffeine use was comparable between patients and controls.

Abstinence Classification

Participant abstinence rates were not significantly different between patients with schizophrenia and non-psychiatric control groups: 42.1% of patients (8/19) and 55% of controls (11/20) successfully achieved abstinence verification criteria [$\chi^2 = 0.65$ ($df = 1$), $p = 0.53$]. Notably, the non-abstaining group collectively included individuals with varying abstaining/relapse trajectories.

These included: (1) individuals who quit, but whose day 28 THC-COOH levels did not drop below 20 ng/ml (SCZ, *n* = 2; CTL, *n* = 3); (2) individuals who lapsed even just once (SCZ, *n* = 3; CTL, *n* = 2); (3) individuals who did not quit but who nevertheless cut down on their current use (SCZ, *n* = 1; CTL, *n* = 1); (4) Participants who relapsed after minimal cessation and then continued to use cannabis throughout the study period (SCZ, *n* = 5; CTL, *n* = 3).

Baseline Neurocognitive Performance

The TOMM was completed at the neuropsychological test training session to examine effort exerted and motivation. All participants scored ≥ 45 on trial 2, suggesting high

Table 2 Baseline Relationships Between Cannabis and Cognition

Cognitive measure	Subtest	SCZ (n = 19)	CTL (n = 20)	p-value
TOMM	Trial 2	49.52 ± 1.2	49.70 ± 0.8	0.97
WCST	# Trial	106.16 ± 20.3	100.15 ± 23.3	0.38
	# Correct	73.21 ± 8.6	74.15 ± 12.4	0.58
	% Error	28.68 ± 14.5	23.40 ± 14.4	0.61
	% Perseverative response	17.21 ± 10.4	12.45 ± 9.1	0.77
	% Perseverative error	15.36 ± 8.4	11.6 ± 7.4	0.83
	% Non perseverative error	13.53 ± 7.4	11.9 ± 9.4	0.30
	Categories completed	4.89 ± 1.7	5.15 ± 1.5	0.29
	Conceptual response	63.68 ± 21.1	69.75 ± 19.7	0.56
IGT	Net total	-1.89 ± 24.0	5.90 ± 28.2	0.89
	Total money	-1250.53 ± 1108.3	-670.75 ± 1301.7	0.47
CPT	% Hits	98.70 ± 1.7	99.45 ± 0.7	0.11
	% Commission	42.39 ± 24.7	30.92 ± 20.4	0.21
	Hit rate	392.19 ± 63.2	382.87 ± 57.9	0.45
	Variability	14.96 ± 14.4	11.30 ± 12.8	0.41
	Attentiveness	0.65 ± 0.49	0.93 ± 0.4	0.15
HVLТ	Trial 1	4.68 ± 2.1	6.915 ± 1.6	0.16
	Trial 2	7.21 ± 1.6	8.63 ± 1.9	0.21
	Trial 3	8.4 ± 1.9	9.94 ± 1.6	0.30
	Sum of trial 1-3	20.16 ± 5.4	24.74 ± 4.5	0.16
	Delayed recall	5.57 ± 2.6	8.79 ± 2.1	0.01
	Repetitions	1.74 ± 2.6	1.42 ± 1.6	0.89
	Intrusions	1.57 ± 1.8	0.89 ± 1.4	0.56
	True positives	10.63 ± 1.4	11.31 ± 1.2	0.31
	False positives	1.47 ± 1.8	0.47 ± 0.9	0.08
	% Retention	63.78 ± 20.8	85.11 ± 17.8	0.01
	Discrimination index	9.15 ± 2.1	10.84 ± 1.9	0.07
Digit span	Forwards	9.89 ± 2.3	11.50 ± 2.9	0.71
	Backwards	5.63 ± 1.9	8.00 ± 2.8	0.18
	Total	15.53 ± 3.5	19.50 ± 4.8	0.53
TMT	A (seconds)	31.71 ± 11.3	26.10 ± 10.2	0.50
	B (seconds)	91.97 ± 49.3	56.38 ± 20.6	0.10
	B minus A (seconds)	60.25 ± 42.2	27.65 ± 12.6	0.07
SDR	5-s delay	17.63 ± 4.8	18.26 ± 5.6	0.33
	15-s delay	25.32 ± 8.7	24.21 ± 10.2	0.66
	30-s delay	30.52 ± 10.2	30.79 ± 16.0	0.79
KDDT	Geomean	0.04 ± 0.03	0.05 ± 0.06	0.63
BART	Avg. adjusted pumps	39.68 ± 17.5	47.78 ± 21.8	0.48
Grooved pegboard	Total time	170.52 ± 46.4	142.48 ± 22.8	0.06
	Total pegs dropped	1.84 ± 2.0	0.350 ± 0.59	0.05

Abbreviations: BART, Balloon Analog Risk Task; CPT, Continuous Performance Test; HVLТ, Hopkins Verbal Learning Test; KDDT, Kirby Delay Discounting; SDR, Spatial Delay Response; TOMM, Test of Memory Malingering; TMT, Trail Making Test; WCST, Wisconsin Card Sorting Test. Values given in mean ± SD; a, values are in numbers; analyses controlled for IQ and education; p-value was set at <0.001.

motivation (Rees *et al*, 1998). In addition, no difference was observed between patients and controls; [$t(37) = 0.53$, $p = 0.60$]. Comparing baseline cognitive function between patients with schizophrenia and controls, no group differences emerged on WCST, IGT, CPT, TMT, SDR, KDDT, or the BART. Groups differed on the HVLТ and Grooved Pegboard, with controls demonstrating better performance than patients ($p < 0.05$). However, when we corrected for

multiple comparisons, these were no longer significant. All significant and non-significant relationships are presented in Table 2.

Effects of Cannabis Abstinence on Clinical Symptoms

PANSS scores remained constant across the abstinence period in both abstaining and non-abstaining patients.

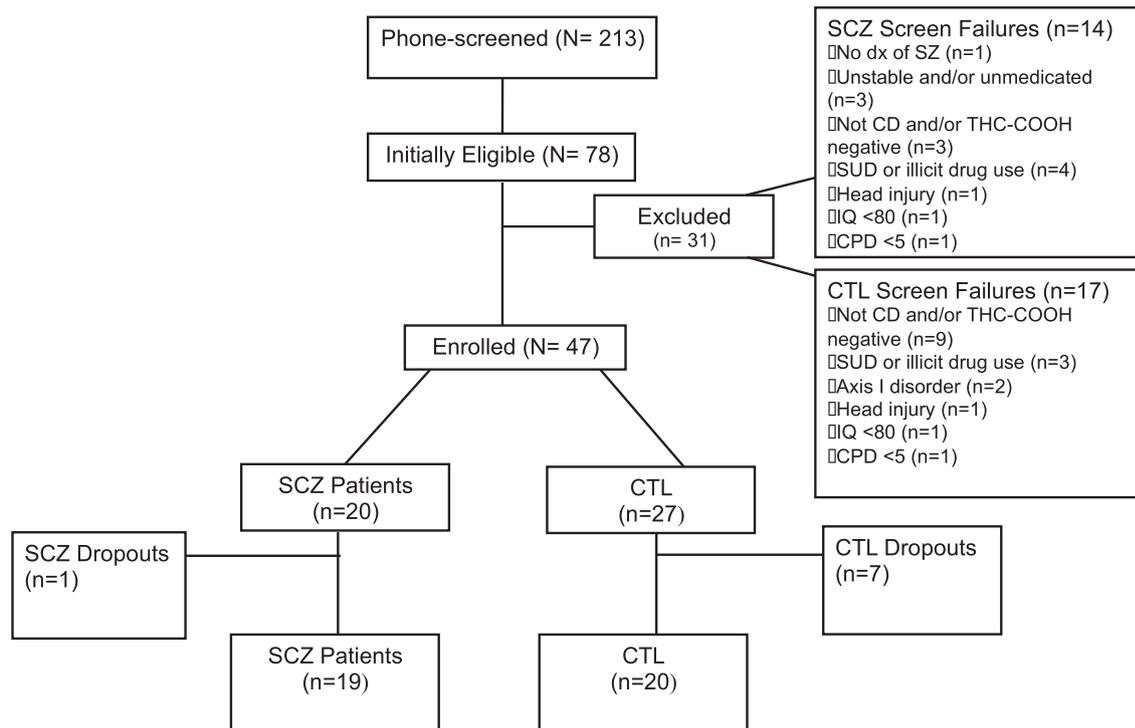


Figure 2 CONSORT Subject Flow Diagram. Details of recruitment, screening, drop-outs, and completion rates of participants for this study.

Among patient abstainers and non-abstainers, there was no change on SARS, BARS, or AIMS scores between baseline and day 28. However, for CDSS, there was a significant reduction in the total depression score over time in schizophrenia abstainers and non-abstainers [$F(4, 68) = 4.44$, $p < 0.01$]. The interaction [$F(4, 68) = 0.337$, $p = 0.88$] was not significant. In patients and control abstainers and non-abstainers, there was no significant change in the total MWC severity score over time [SCZ: $F(4, 68) = 1.61$, $p = 0.18$] [CTL: $F(4, 72) = 2.17$, $p = 0.08$].

Trajectory of Cognitive Symptoms with Abstinence

Hippocampus-mediated tasks. (i) Verbal Memory and Learning: using a $2 \times 2 \times 3$ ANOVA, there was a main effect of time [$F(2, 68) = 5.03$, $p < 0.01$] on HVLTL percent retention. The interaction between time and abstinence status was also significant [$F(2, 34) = 3.86$, $p < 0.03$]. The interactions between time \times psychiatric diagnosis, and the three-way interaction term (time \times abstinence status \times psychiatric diagnosis) were not significant. Planned contrasts, controlling for multiple pairwise comparisons (with $\alpha = 0.0167$), revealed improvement in patient abstainers [$F(2, 14) = 4.73$, $p = 0.02$] over time. The magnitude of this improvement was 39.3% ($d = 1.07$), suggesting a clinically significant change in cognitive performance (Cohen, 1988); Figure 3a. Non-abstainers had no significant change in performance over time [$F(2, 20) = 2.19$, $p = 0.14$]. Similarly, change over time was non-significant in controls; [$F(2, 34) = 0.65$, $p = 0.53$]; the magnitude of this improvement was 12.0% ($d = 0.67$); Figure 3b. Observed power for key analyses were as follows: main effects of time, 80%, $F(2, 68) = 5.034$,

$p < 0.01$; time \times abstinence status, 68.1%, $F(2, 68) = 3.86$, $p = 0.026$; time \times diagnosis \times abstinence status, 25.4%, $F(2, 68) = 1.20$, $p = 0.308$.

In exploratory analyses, since HVLTL % retention improved in patients with schizophrenia with abstinence, we determined the effects of 28 days of cannabis reinstatement. Six (of 8) abstaining patients completed the follow-up, and cannabis reinstatement occurred immediately after the abstinence period ended (day 28). Thus, we examined HVLTL performance across the 56-day period using four time-points (days 0, 14, 28, and 56). RM-ANOVAs revealed a significant change in HVLTL % retention performance over time [$F(3, 15) = 5.026$, $p < 0.02$]. LSD post-hoc analyses revealed that improvement occurred with cannabis abstinence between days 0 and 28 [$p < 0.04$], and between days 28 and 56, when cannabis was reintroduced, there was a decline in HVLTL performance [$p = 0.03$]. Among controls ($N = 4$; of 11), the re-introduction of cannabis had little effect on HVLTL performance, similar to cannabis abstinence (data not shown). RM-ANOVAs revealed no significant change in HVLTL % retention performance over time; [$F(3, 9) = 1.820$, $p = 0.21$]. Other HVLTL outcomes showed no significant change over time either in patients or controls.

Prefrontal tasks

- (i) Attention: using a $2 \times 2 \times 3$ ANOVA demonstrated there was no significant change in CPT [% Hits $F(1.5, 52.8) = 1.04$, $p = 0.85$; Hit Rate $F(2, 70) = 1.55$, $p = 0.22$] or TMT-A [$F(2, 70) = 0.45$, $p = 0.64$] outcomes over time.
- (ii) Executive function: using a $2 \times 2 \times 3$ ANOVA showed no change on the Digit Span Backwards [$F(2, 70) = 0.72$, $p = 0.49$] or the TMT-B [$F(1.5, 53.86) = 3.87$, $p = 0.04$].

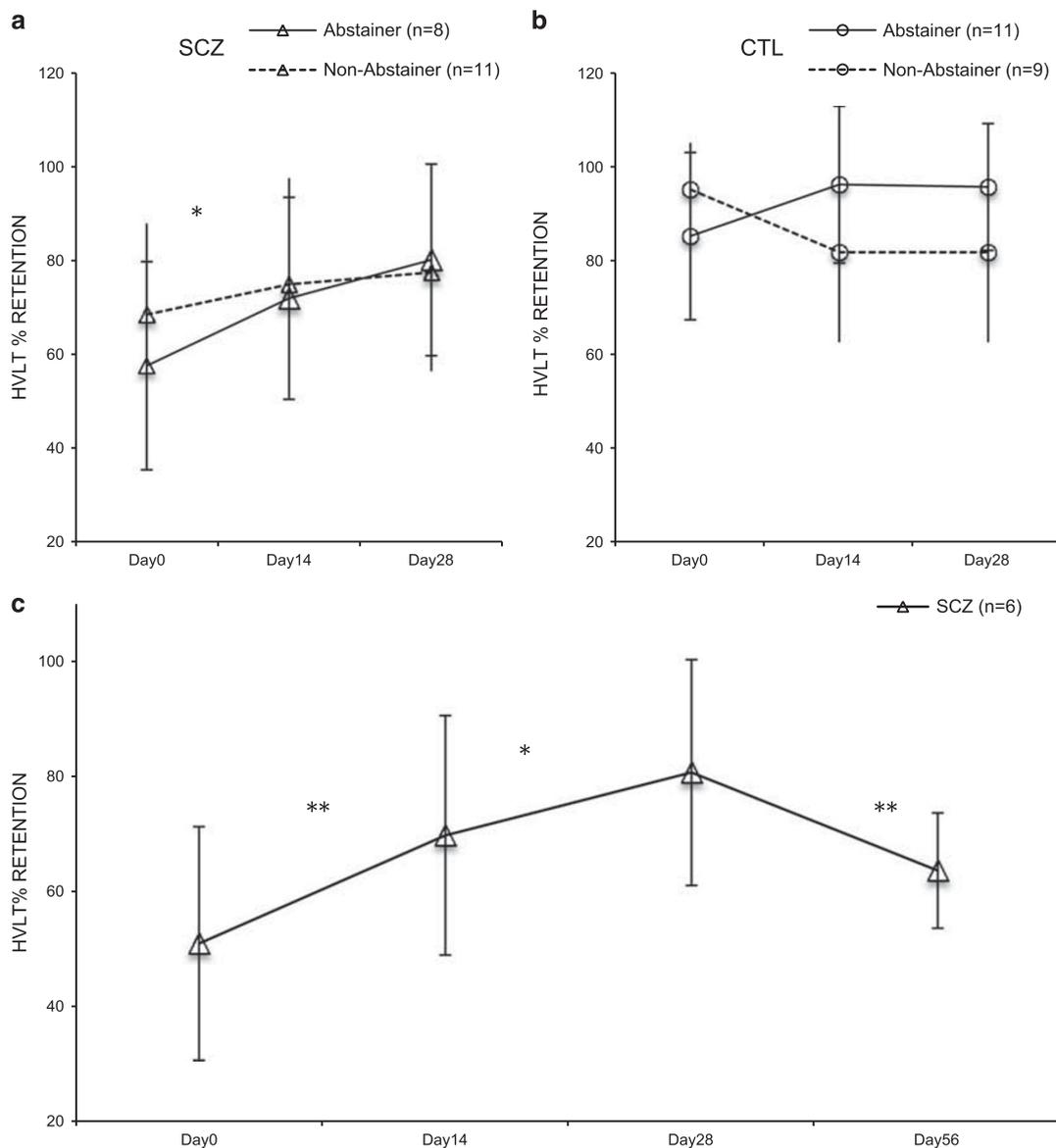


Figure 3 Trajectory of HVL T % Retention over Time. Schizophrenia abstainers demonstrated improved performance on the HVL T % retention over time, between days 0 and 28 (a). Controls showed no improvements over time in HVL T performance (b). Among patients, significant improvements in HVL T % retention performance occurred with abstinence, while cannabis relapse led to reversal of this abstinence-related cognitive change. In contrast, there was no significant change in HVL T % retention performance with abstinence, or cannabis relapse assessed at day 56 in control participants (c). Error bars reflect SD. * $p < 0.05$; ** $p < 0.01$.

- (iii) Working memory: there was a non-significant effect of time on the Digit Span Forward [$F(2, 70) = 2.61, p = 0.08$]. Performance on the SDR 5 [$F(2, 68) = 1.59, p = 0.21$], 15 [$F(2, 70) = 0.74, p = 0.48$] and 30 [$F(2, 70) = 1.67, p = 0.20$] second delay also demonstrated no change over time.
- (iv) Impulsivity: there was no significant change on measures that assess impulsivity such as the KDDT [$F(2, 70) = 0.42, p = 0.66$] and BART [$F(2, 50) = 0.36, p = 0.70$].

Cerebellar task. Motor function: RM-ANOVA yielded a significant main effect of time on the Pegboard task [$F(1.7, 59.4) = 5.61, p < 0.01$]. The interaction effect between time and diagnostic group [$F(2, 70) = 0.31, p = 0.73$] and time and abstinence status were non-significant [$F(2, 70) = 0.76, p = 0.47$].

DISCUSSION

This is the first prospective investigation of the effects of extended cannabis abstinence on cognition in cannabis-dependent patients with schizophrenia and non-psychiatric controls. We implemented a novel cannabis abstinence paradigm to isolate the state-dependent effects of cannabis on cognitive function. At study end point, 42% of patients (8/19) and 55% of non-psychiatric controls (11/20) remained cannabis free for the full 28-day period. Moreover, study retention rate was high in patients (95%) and controls (25%).

Chronic cannabis use in non-psychiatric controls generates deficits that resemble the cognitive profile of patients with schizophrenia (Solowij *et al*, 2002). However, it has been proposed that cannabis-using patients with schizophrenia may belong to a subgroup of patients who are higher

functioning, have better pre-morbid adjustment and have better cognition (eg, Schnell *et al*, 2009; Yucel *et al*, 2012). Therefore, the expected difference between cannabis-using controls and schizophrenia may be minimized when compared to this higher functioning patient subgroup. We posit that the true magnitude of impairment in cognitive performance may only become evident when cannabis is ceased.

However, over time, patient abstainers demonstrated improvements in verbal memory and learning. The magnitude of cognitive change (while not significant when correcting for multiple comparisons) may be of clinical importance, with scores improving by approximately 40% from pre- to post-abstinence ($d = 1.07$). Despite the modest sample sizes, cognitive improvement with cannabis abstinence may have been preferential for verbal memory and learning, as performance on other cognitive tests were less pronounced (see Supplementary Table 3). These results suggest that chronic cannabis use may be associated with poorer cognitive function in select cognitive domains (ie, verbal memory) in patients with schizophrenia. Importantly, these deficits may improve with 28 days of abstinence. Conversely, deficits in other cognitive domains may be insensitive to change within this abstinence timeframe.

This finding was further substantiated by exploratory data from a subset of patient abstainers. Preliminary data were collected at a 4-week follow-up visit (day 56). Six out of the eight patients (two patients were lost to follow-up) who successfully abstained from cannabis until day 28 relapsed immediately after the bonus was paid. We observed that while cannabis abstinence led to significant improvements in verbal memory and learning, performance worsened when cannabis was reintroduced; suggesting a reversal of abstinence-related improvements in schizophrenia. Further study and replication of this finding in larger samples is needed.

While some studies have attributed greater cognitive capacities of cannabis-using patients to cannabis itself (Coulston *et al*, 2007a,b), others have proposed that cannabis users belong to a subgroup of higher functioning patients (Schnell *et al*, 2009; Yucel *et al*, 2012). Our data do not support the former hypothesis, however our findings work in concert with the latter. Thus, we speculate that cannabis exerts a deleterious effect on cognition (state-effect) that is superimposed upon a higher functioning subgroup of patients (trait-effect). However, given that we did not have non-using comparison groups, this could not be empirically confirmed. Findings are in line with our previous study that reported associations between increasing cumulative cannabis use and progressive cognitive impairment in current, but not former (>6 months abstinent) cannabis-dependent patients with schizophrenia (Rabin *et al*, 2013). Thus, while continued cannabis use results in deterioration of cognitive performance, when compared to non-users cannabis-users appear to have relatively higher cognitive function.

Although selective deficits may improve with 28 days of abstinence, other cognitive domains may do so at differential rates. Thus, continued abstinence beyond 28 days may be warranted for full cognitive recovery. It is possible that specific brain regions are more vulnerable and/or resilient to cannabis compared to others. Therefore, recovery of one cognitive domain does not necessarily predict recovery in others. In other words, while verbal learning and memory

appears to improve with 28 days of abstinence, more enduring deficits may occur within other cognitive domains.

Interestingly, verbal learning and memory have been the most consistently impaired cognitive functions in studies of cannabis use (Ranganathan and D'Souza, 2006). Moreover, this domain is also the area in which patients with schizophrenia demonstrate the most significant deficits (Heinrichs and Zakzanis, 1998). A laboratory study by D'Souza *et al* (2005) showed that of all tests administered the HVLT was most sensitive to THC administration in both patients and controls (eg, verbal fluency, distractibility, and vigilance; D'Souza *et al*, 2005). Other studies are in agreement that heavy cannabis use selectively impairs verbal learning and memory tasks in patients (Nunez *et al*, 2016). Thus, the ability and/or time to cognitively recover may correlate with CB1R density in the brain region responsible for mediating that specific cognitive task. That is, improved cognitive function (with 28 days of abstinence) may first occur in tasks facilitated by areas with high CB1R concentrations, such as the hippocampus (Herkenham, 1991). Although chronic cannabis exposure may induce CB1R down-regulation and desensitization, conceivably abstinence may reverse these processes and increase CB1R availability. Notably, the magnitude of rate of down-regulation and desensitization are region-dependent (Ceccarini *et al*, 2015; Hirvonen *et al*, 2012), thus adaptations do not occur uniformly across the brain. Accordingly, tasks mediated by brain regions with lower CB1R concentrations may require longer abstinence periods for recovery of performance. This proposed underlying mechanism supports findings from the current study in that patients with schizophrenia demonstrated improvements in the HVLT, a verbal memory and learning task predominantly mediated by the hippocampus (Squire, 1992).

It is important to note that the non-abstaining group collectively included individuals with varying abstaining/relapse trajectories. The heterogeneity of these abstaining/relapsing trajectories may help to explain the upward trend of improvement in HVLT scores in non-abstaining patients with schizophrenia. It should also be emphasized that improvement in performance is unlikely related to practice effects. First, alternate HVLT forms were administered on different days. Second, at the one-month follow-up, upon the fifth administration of the HVLT, there was a decrement in performance in patients with schizophrenia who re-introduced cannabis following abstinence.

In contrast to patients, cognitive change was not observed with abstinence in our non-psychiatric controls. It has been suggested that compared to controls, patients with schizophrenia possess an enhanced sensitivity to the cognitive-impairing effects of cannabis (D'Souza *et al*, 2005) and hence may also be more susceptible for recovery of cognitive function. A dysfunctional endocannabinoid system, including reduced CB1R availability (Ranganathan and D'Souza, 2006), has been implicated in the pathophysiology of schizophrenia and may help to explain this observation (Muller-Vahl and Emrich, 2008). Thus, if hippocampus-mediated tasks are most susceptible to cognitive impairment, and patients with schizophrenia have increased susceptibility to these effects, then it follows that this brain region may also be most vulnerable to the reversal of deficits in patients. However, given our study design we cannot fully exclude the possibility that cannabis exposure might not have impaired cognition in the control group.

Importantly, cannabis cessation in patients with schizophrenia did not lead to any adverse effects. According to the self-medication hypothesis (Khantzian, 1985), cannabis may alleviate symptoms associated with the disorder. Our results do not support this theory. Psychiatric symptoms and extrapyramidal symptoms did not worsen with abstinence. Other studies have found that cannabis use is not associated with beneficial effects in patients with schizophrenia (Henquet *et al*, 2010; Ringen *et al*, 2013). The only adverse outcomes experienced with abstinence were cannabis withdrawal symptoms, which dissipated within the first 14 days of cessation (Rabin *et al*, 2017).

Recovery of cannabis-induced cognitive impairment in controls may not be as rapid as in patients. Our finding that cognitive impairment does not fully recover following 28 days of cannabis cessation is consistent with some (Bolla *et al*, 2002; Medina *et al*, 2007), but not all, prior studies (Pope *et al*, 2001). One possible explanation for this finding is that cannabis may produce irreversible neurotoxic effects (Meier *et al*, 2012; Solowij and Battisti, 2008). Results are consistent with the speculation that cannabis use during adolescence, when the brain is undergoing critical neurodevelopment processes, may produce neurobiological dysfunction (Bossong and Niesink, 2010). This has been proposed as a mechanism for the development of schizophrenia in individuals with pre-existing susceptibility factors (Caspi *et al*, 2005). However, for individuals who use cannabis during adolescence and do not develop a psychotic illness, the resulting consequences may be enduring cognitive impairment irrespective of whether or not cannabis use is sustained. In addition, given that the mean age of onset in controls was 15 and cumulative years of use was reported to be almost 10 years, this supports the idea that permanent cognitive deficits may result. While cumulative cannabis use was comparable between groups, indexed as joint-years, baseline THC-COOH was twice as high in controls *vs* schizophrenia patients, perhaps suggesting heavier use, or the use of higher potency cannabis strains. This may make the control group more vulnerable to permanent, non-reversible deficits relative to our schizophrenia group (Tait *et al*, 2011). Moreover, given that schizophrenia patients have been shown to be more sensitive to the cognitive effects of cannabis (D'Souza *et al*, 2005), this may explain why improvement more readily occurred in the patients group *vs* the control group.

Our study results must be interpreted in the context of several limitations. First, we did not include non-cannabis using control groups (eg, non-cannabis-using non-psychiatric controls or non-cannabis using patients), or cannabis-using patients and controls receiving non-contingent (yoked) control interventions. The lack of these comparison groups makes it difficult to characterize the magnitude of baseline cognitive deficits, and prospective changes in cognitive outcomes in cannabis non-using and using subjects. Future studies should include these comparative controls. Second, ceiling effects on specific cognitive tests (eg, HVLТ) at baseline may have prevented the detection of cognitive change in cannabis abstaining control participants. For example, HVLТ-R may not provide be complex enough (eg, high cognitive task load) to elicit sufficient variability in scores and error types in healthy controls (Lacritz and Cullum, 1998). Thus, we were unable to detect a large

enough magnitude of change in memory retention. It should be noted that mean HVLТ % retention for our control group at baseline was 85.11 ± 17.8 , and by week 4 scores improved to 95.61 ± 13.5 . As mentioned, this improvement did not achieve statistical significance ($p=0.27$). HVLТ % retention normative data for this age group is $91.15+13.07$ (from the HVLТ Professional manual) (Brandt and Benedict, 2001). In addition, repeated testing (four times in total by day 28) may have induced some practice effects. There is evidence to suggest that controls may experience greater practice effects from repeated testing compared to patients with schizophrenia (Szoke *et al*, 2008). Third, we set a biochemical cut-off level to distinguish abstainers from non-abstainers. This may have inadvertently posed a bias towards classifying heavier cannabis users who abstained as non-abstainers, as they may have not been able to eliminate cannabinoids to levels below 20 ng/ml within 28 days (Ellis *et al*, 1985; Goodwin *et al*, 2008). Thus, it follows that if cognitive function is most likely to show the greatest magnitude of change in heavier users then lengthier abstinence periods are warranted. However, we believe that there may be a trade-off as a longer duration of abstinence may be related to subject attrition. Perhaps with the implementation of extended duration contingency management and adequate incentives, increasing the length of abstinence might be feasible. Finally, this study exclusively studied males. Future studies should assess whether findings extend to female cannabis users. This is especially important because despite the lower rates of cannabis use among women, females are more susceptible to the development of CUDs, have more severe withdrawal symptoms, and are more likely to relapse compared to men (Cooper and Haney, 2014; Craft *et al*, 2013; Fergusson *et al*, 2006). These limitations should be addressed in future studies.

CONCLUSIONS

Our findings provide additional justification to support clinical efforts to encourage patients with schizophrenia to abstain from cannabis. While it may be thought of as a benign drug with low addiction potential, we provide evidence that cannabis might possess cognitive-impairing properties in schizophrenia. Intact cognitive function is especially critical in schizophrenia given that cognition is one of the most reliable predictors of functional outcomes in these patients (Green *et al*, 2000). Accordingly, remedying cannabis-related cognitive dysfunction may provide a critical element in the successful rehabilitation of patients with schizophrenia. Future studies should explore the cognitive effects of longer abstinence periods and functional neuroimaging techniques should be incorporated into these paradigms to monitor accompanying changes in brain activity. Understanding the degree to which the human brain may recover as a function of extended cannabis abstinence has important implications for the treatment of SUDs as well as the neurobiology of comorbid psychiatric disorders.

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