

Pilot study of the 5-HT_{2A}R agonist psilocybin in the treatment of tobacco addiction

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Abstract

Despite suggestive early findings on the therapeutic use of hallucinogens in the treatment of substance use disorders, rigorous follow-up has not been conducted. To determine the safety and feasibility of psilocybin as an adjunct to tobacco smoking cessation treatment we conducted an open-label pilot study administering moderate (20 mg/70 kg) and high (30 mg/70 kg) doses of psilocybin within a structured 15-week smoking cessation treatment protocol. Participants were 15 psychiatrically healthy nicotine-dependent smokers (10 males; mean age of 51 years), with a mean of six previous lifetime quit attempts, and smoking a mean of 19 cigarettes per day for a mean of 31 years at intake. Biomarkers assessing smoking status, and self-report measures of smoking behavior demonstrated that 12 of 15 participants (80%) showed seven-day point prevalence abstinence at 6-month follow-up. The observed smoking cessation rate substantially exceeds rates commonly reported for other behavioral and/or pharmacological therapies (typically <35%). Although the open-label design does not allow for definitive conclusions regarding the efficacy of psilocybin, these findings suggest psilocybin may be a potentially efficacious adjunct to current smoking cessation treatment models. The present study illustrates a framework for future research on the efficacy and mechanisms of hallucinogen-facilitated treatment of addiction.

Keywords

Hallucinogen, tobacco, smoking cessation, nicotine, addiction, psilocybin, psychedelic

Introduction

A promising research area from the 1950s through 1970s involved the therapeutic use of 5-HT_{2A}R agonist hallucinogens in the treatment of drug dependence, including alcohol and opioid dependence (Chwelos et al., 1959; Hollister et al., 1969; Ludwig et al., 1969; Savage and McCabe, 1973; Smart et al., 1966). Some studies did not include the rigor and controls expected of modern clinical research, thus obscuring efficacy. However, a recent meta-analysis of randomized controlled trials found that LSD-facilitated treatment of alcoholism approximately doubled the success rates of control conditions at the first follow-up (Krebs and Johansen, 2012). Despite suggestive findings, this line of investigation was abandoned due to controversy surrounding the recreational use of hallucinogens and regulatory restrictions impeding subsequent research with 5-HT_{2A}R agonists (Mangini, 1998; Nutt et al., 2013).

Recently, the 5-HT_{2A}R agonist psilocybin was found to occasion mystical-type experiences with enduring personal meaning and spiritual significance in the majority of healthy volunteers (Griffiths et al., 2006). Moreover, at 14-month follow-up, 61% of volunteers associated these experiences with moderate to extreme positive behavior change (Griffiths et al., 2008). In another study, a moderate dose of psilocybin was found to significantly decrease anxiety and depression in patients with advanced stage cancer (Grob et al., 2011). Results from these studies are atypical for pharmacotherapies in that positive effects were observed well beyond the time-course of acute drug effects. Although participants were not drug dependent, these results suggest the feasibility of a psilocybin-facilitated intervention for addiction treatment, consistent with findings that increased levels of spirituality are

associated with improved outcomes in drug dependence recovery (Cole et al., 2006; Galanter, 2006; Piderman et al., 2007; Piedmont, 2004).

Smoking-related mortalities in the USA are currently estimated at 480,000 annually (US Department of Health and Human Services, 2014), and 5 million annually worldwide (World Health Organization, 2011), highlighting the urgent need for novel treatments. Furthermore, most behavioral interventions and pharmacotherapies for smoking cessation exhibit only modest success rates at 6 months (typically <35%; Cahill et al., 2014; Mottillo et al., 2009). The present study was conducted as an open-label pilot study to determine the safety and feasibility (i.e. potential efficacy) of psilocybin-facilitated smoking cessation treatment. Although there is experimental evidence suggesting safety and efficacy of classic hallucinogens in the treatment of addiction, this is largely in relation to the use of LSD for treating alcoholism (Krebs and Johansen, 2012; Mangini, 1998). There has heretofore been no research on use of classic hallucinogens in the

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Table 1. Demographic and smoking characteristics, $N=15$.

Categories	Mean (SD)	Range
Sex ^a	10 M, 5 F	
Age (years)	51 (10.5)	26-65
Education ^b		
Some college	4 (26.7)	
Bachelor's degree	7 (46.7)	
Graduate degree	4 (26.7)	
Cigarette dependence (FTCD) ^c	5.3 (1.3)	3-8
Years smoking	31 (9.9)	10-49
Previous quit attempts	6 (3.6)	2-12
Cigarettes/day at intake	19 (2.9)	15-25
Breath CO at intake	30 (9.9)	13-53
Urine cotinine at intake	1676 (594)	841-3212

^aMean (SD) do not apply to sex.

^bMean (SD) do not apply to education. Educational achievement is represented as n (%) of total study sample.

^cFTCD=Fagerström Test for Cigarette Dependence. Possible scores on the FTCD range from 0 to 10, with higher scores indicating greater dependence.

treatment of tobacco addiction. Therefore, this pilot study was conducted without a control condition as a first step to evaluate both the safety of the approach, and whether efficacy rates would be promising enough to warrant the investment of resources necessary for a randomized trial.

Methods

This study utilized a 15-week course of smoking cessation treatment, with psilocybin administration occurring in weeks 5, 7, and 13. This study was approved by the Johns Hopkins University School of Medicine Institutional Review Board, and participants provided informed consent.

Participants

Participants were recruited through advertisements offering a novel treatment for smoking cessation involving psilocybin. In all, 323 individuals were telephone screened, and 27 of these underwent in-person medical and psychological screening. Individuals disqualified by phone were primarily seeking pay for participation, currently on medications, or did not meet minimum smoking requirements. Fifteen participants (10 males) were enrolled in the study. All of these individuals completed the study (Table 1). For inclusion in the study individuals needed to meet these criteria: smoke a minimum of 10 cigarettes per day, be healthy as determined via medical interview, electrocardiogram, blood and urinalysis laboratory tests, have multiple unsuccessful past quit attempts, and still desire to quit smoking. Furthermore, a Structured Clinical Interview for DSM-IV-TR (First et al., 2012) was performed at screening to exclude individuals with personal or family history of psychotic or bipolar disorders, and/or drug dependence (including alcohol; excluding nicotine) within the past 5 years. The study sample was relatively racially homogeneous, including 14 (93%) White individuals, and one Asian individual (7%). Ten participants reported minimal past use of hallucinogens with a mean of 8 (range 1–17) lifetime uses, with the most recent use being a mean of 27 years (range 7–43)

before study intake. Five were hallucinogen-naïve. Participants received no monetary compensation.

Procedures

Participants were assigned to a team of two or three study staff who conducted treatment and oversaw psilocybin sessions. At least one member of each team was a doctoral-level psychologist trained in delivering the study intervention and conducting psilocybin sessions. A Target Quit Date (TQD) was set to coincide with the first psilocybin session at week 5 of treatment. Participants attended four weekly meetings in which a manualized intervention consisting of cognitive behavioral therapy (CBT) for smoking cessation and preparation for psilocybin administration were delivered (Table 2). On the TQD, participants were administered a moderate dose of psilocybin (20 mg/70 kg). Participants continued meeting weekly with study staff, and received another dose of psilocybin at week 7, and optionally again at week 13. Although the default dose for sessions 2 and 3 was a high dose (30 mg/70 kg), participants were permitted to repeat the moderate dose on these sessions.

The second and third sessions were intended as additional quit opportunities for those who failed to achieve abstinence after the first session. For those who did quit on the first session, the additional psilocybin sessions were meant to support motivation for long-term abstinence. This approach was informed by studies of hallucinogen-facilitated substance dependence treatment in which investigators described an extended, time-limited, post-session period (sometimes referred to as an “afterglow”) associated with decreased substance use, elevated mood and energy, decreased anxiety, and increased capacity for close interpersonal relationships (e.g. Bowen et al., 1970; Pahnke et al., 1970). As investigators have suggested, long-term abstinence may be enhanced by including multiple sessions, which may work by extending the “afterglow” period through the time of greatest relapse risk, or by increasing the probability of a transformative mystical experience (Grinspoon and Bakalar, 1979; Halpern, 1996; Osmond et al., 1967; Savage and McCabe, 1973; Smart et al., 1966).

The doses and dosing sequence used in this study were informed by previous psilocybin dose–response research showing greater prevalence of mystical experience at 20 mg/70 kg and 30 mg/70 kg over other lower doses, and that ascending dose sequences produced significantly increased well-being, life satisfaction, and persisting positive mood at 1-month post-session as compared with descending dose sequences (Griffiths et al., 2011). Study staff met with participants the day after each psilocybin session, and weekly after the TQD to discuss session experiences and provide support for smoking cessation, for a total of 19 in-person meetings. A staff member made brief (<5 min.) daily phone calls to participants for 2 weeks after the TQD to provide encouragement for smoking abstinence. Participants were instructed to abstain from using any additional smoking cessation treatments (e.g. nicotine replacement) during the study.

Psilocybin sessions. Sessions were conducted as previously described (Griffiths et al., 2006, 2011), with the exception that participants repeated their brief motivational statement for smoking cessation before each psilocybin administration, and participated in a guided imagery exercise at the resolution of psilocybin effects on the first psilocybin session (see *Cognitive behavioral*

Table 2. Summary of treatment meetings and content.

Meeting (duration)	CBT modules	Psilocybin session-related topics
Prep. Mtg. ^a 1 (90 min)	Assign Target Quit Date (TQD) Sign contract to quit NURD ^b program Smoking diary introduction	Introduction to overall approach Discuss childhood / early family life
Prep. Mtg. 2 (90 min)	Smoking diary review Reasons to quit vs. continue smoking Develop brief motivational statement WEST-D ^c program Health effects of smoking Smoking financial costs	Discuss current and past relationships / romantic life
Prep. Mtg. 3 (90 min)	Smoking diary review Discuss previous quit attempts Weight gain and quitting smoking Withdrawal framed as "recovery"	Discuss work, hobbies, and other important activities
Prep. Mtg. 4 (90 min)	Smoking diary review Dealing with urges after TQD Preparing to quit	Discuss worldview (philosophical and/or spiritual beliefs) What to expect in psilocybin sessions
Psilocybin Ses. ^d (8 hrs)	–	–
Integration Mtg. ^e (1 hr)	–	Participant narrative of session
Support Mtg. ^f (45 min)	Support smoking abstinence Review CBT techniques	Discuss session experiences

^aPreparation meetings were held weekly for 4 weeks prior to TQD. Approximately half the duration of each meeting was spent conducting CBT, and half preparing for psilocybin sessions.

^bNURD program card was to be read each time a cigarette was smoked: "This cigarette is giving me *No* satisfaction; This is an *Unpleasant* experience; This cigarette is making me feel *Rotten*; I am losing the *Desire* to smoke."

^cWEST-D program card was to be read each time participants noticed an urge to smoke: "What's the trigger? Each time I feel like smoking, *Stop, Think, Deprogram.*"

^dPsilocybin sessions were introspective and contained no formal treatment content other than listening to a music program.

^eIntegration meetings occurred the day after each psilocybin session. Participants' experiences of the preceding day's psilocybin session were discussed.

^fSupport meetings were conducted weekly after TQD for 10 weeks.

therapy section below). Sessions followed safety guidelines for human hallucinogen research (Johnson et al., 2008). Participant blood pressure and heart rate were monitored at ≤ 60 min intervals, and at least one staff member was present throughout sessions. For each session a physician was on call, and rescue medications were available in case of adverse cardiovascular or psychological events (Johnson et al., 2008). During sessions, participants were encouraged to lie down on a couch and focus on their internal experience. Participants wore an eye mask and listened to a music program through headphones. During sessions, staff provided non-directive interpersonal support for managing psilocybin effects, but did not deliver smoking cessation-specific content. After drug effects subsided, participants were asked to write an open-ended narrative describing their session for discussion with staff the following day.

Cognitive behavioral therapy. In four weekly preparation meetings, participants received smoking cessation CBT (Marks, 1993; Perkins et al., 2007), largely based on the Quit For Life program (Table 2), shown to be effective in controlled studies (Marks and Sykes, 2002; Sykes and Marks, 2001). Sessions started with a brief (<10 min) "body-scan" meditation (Kabat-Zinn, 1990). In preparatory sessions participants developed their most important reasons to quit smoking into a brief motivational statement (e.g. "I want to be free, clean, and clear").

Study treatment also included two components of an effective group-based smoking cessation therapy (Zernig et al., 2008). First, participants smelled a scented oil during preparatory and support meetings before each exercise. This oil was provided to the participant at the TQD and the participant was encouraged to smell it when experiencing cravings. Second, brief (<10 min) guided imagery exercises were conducted during preparatory and support meetings, and at the end of the first psilocybin session.

Measures

Biological markers of smoking abstinence. Two measures of recent smoking, exhaled carbon monoxide (CO) and urinary cotinine level (Benowitz et al., 2002), were assessed at intake, weekly throughout the intervention, and at 6-month follow-up. Breath CO was measured using a Bedfont Micro III Smokerlyzer (Haddonfield, NJ) to detect smoking over approximately the past 24 hours. Urine cotinine samples were collected and sent to an independent laboratory for analysis (Friends Medical Laboratory, Baltimore, MD) to detect smoking over approximately the previous six days. Urine cotinine levels of <200 ng/mL, and breath CO of ≤ 6 ppm were considered as biological verification of non-smoking status (Bramer and Kallungal, 2003; Javors et al., 2005; Middleton and Morice, 2000). A list of measures and their schedule of administration is summarized in Table 3.

Table 3. Schedule of measures administered.

Measures	Time points assessed
Fagerström Test for Cigarette Dependence	Intake
Breath carbon monoxide	Intake, Weeks 2–15, 6-month follow-up
Urine cotinine	Intake, Weeks 2–15, 6-month follow-up
Timeline follow-back	Intake, Weeks 2–15, 6-month follow-up
Questionnaire on Smoking Urges	Intake, Week 6, Weeks 8–15, 6-month follow-up
Smoking Abstinence Self-efficacy Scale	Intake, Week 6, Weeks 8–15, 6-month follow-up
Wisconsin Smoking Withdrawal Scale	Intake, Week 6, Weeks 8–15, 6-month follow-up
Visual Effects Questionnaire	Intake, 6-month follow-up
Post-session Headache Interview	Day after psilocybin sessions
Mysticism Scale	Intake, 1-week post-psilocybin sessions 2 and 3
States of Consciousness Questionnaire	Day of psilocybin sessions (7 hours after drug administration)
Persisting Effects Questionnaire	1 week after every psilocybin session

Timeline follow-back. Participants completed a smoking timeline follow-back (TLFB) assessment at each study meeting. The TLFB is a self-report calendar completed retrospectively by participants indicating the number of cigarettes smoked each day (Sobell and Sobell, 1992).

Fagerström Test for Cigarette Dependence. The Fagerström Test for Cigarette Dependence (FTCD; formerly Fagerström Test for Nicotine Dependence) is a 6-item questionnaire widely used to characterize the level of dependence of cigarette smokers (Fagerström, 2012; Heatherton et al., 1991). FTCD data were collected at intake.

Smoking cessation-related measures. Three supplemental measures related to smoking cessation were administered at intake, weekly post-TQD until end of treatment (excluding psilocybin session weeks), and at 6-month follow-up. The Questionnaire on Smoking Urges (QSU) is a multidimensional assessment of smoking craving with demonstrated sensitivity to smoking cessation (Tiffany and Drobes, 1991). The Smoking Abstinence Self-Efficacy scale (SASE) provides a measure of smokers' confidence to abstain from smoking in 20 hypothetical situations, and temptation to smoke in those situations, and has been found to correlate with smoking cessation outcomes (DiClemente and Prochaska, 1985; DiClemente et al., 1985). The Wisconsin Smoking Withdrawal Scale (WSWS) measures severity of smoking withdrawal and exhibits good validity and reliability in smoking cessation studies (Shiffman et al., 2004; Welsch et al., 1999).

Visual Effects Questionnaire. To probe for potential Hallucinogen Persisting Perception Disorder (Halpern and Pope, 2003), participants completed this 16-item questionnaire designed to assess occurrence, duration, and severity of visual disturbances (e.g. halos, strobe-like trails) at intake and 6-month follow-up.

Post-session headache interview. The first five participants had completed their psilocybin sessions when another study found that psilocybin increased headache occurrence (Johnson et al., 2012). Therefore, the final 10 participants completed a staff-administered scale the day after each psilocybin session which assessed the occurrence, severity, and time-course of post-psilocybin headache.

Mysticism Scale. This 32-item questionnaire was designed to assess primary mystical experience across the lifetime (Hood, 1975), and has been employed extensively in research, demonstrating good validity, and cross-cultural reliability (Hood et al., 2001), as well as sensitivity to the effects of psilocybin (Griffiths et al., 2006, 2011). Participants completed the Mysticism Scale at intake, 1-week post-psilocybin session 2, and 1-week post-session 3 (when applicable).

States of Consciousness Questionnaire (SOCQ). This 100-item questionnaire has previously been used to characterize the subjective effects of psilocybin (Griffiths et al., 2006, 2011). For this study it was used to assess the occurrence of fearful or otherwise challenging experiences. Participants completed the SOCQ at the conclusion of each psilocybin session, approximately 7 hours after psilocybin administration.

Persisting Effects Questionnaire. This 145-item questionnaire was designed to measure changes in attitudes, moods, behavior, and spiritual experience, and has demonstrated sensitivity to the intermediate and long-term effects of psilocybin (Griffiths et al., 2008, 2011). In total, 140 of the items were rated on a 6-point scale ranging from 0 (*none*) to 5 (*extreme*) in six categories: Attitudes about life (26 items); Attitudes about self (22 items); Mood changes (18 items); Relationships (18 items); Behavioral changes (two items); Spirituality (43 items). Each category reflected positive and negative changes, resulting in 12 total subscales (Table 4). The questionnaire included five additional questions. The first three asked participants to rate the overall personal meaning, spiritual significance, and effects on well-being or life satisfaction attributed to their most recent psilocybin experience. The final two asked participants to endorse mechanisms by which they believed psilocybin had facilitated smoking cessation (including that psilocybin had not helped), and to rank order these items in terms of their importance (Table 5). This questionnaire was administered 1 week after each psilocybin session.

Data analysis

Safety data consisted of acute measures of cardiovascular function during psilocybin sessions, post-session States of Consciousness Questionnaire ratings of acute adverse psychological effects, next-day headache ratings, and Visual Effects Questionnaire data.

Table 4. Persisting Effects Questionnaire Ratings at 1 week after Psilocybin Sessions*.

Subscale / Item	Mean (SEM)	Mean (SEM)	Mean (SEM)
	Session 1 ^d	Session 2	Session 3
Positive attitudes about life	49.3 (6.5)	63.2 (4.2)	69.7 (6.0)
Negative attitudes about life	10.4 (3.2)	6.3 (1.4)	5.5 (1.4)
Positive attitudes about self	42.2 (6.6)	57.7 (5.2)	65.3 (5.8)
Negative attitudes about self	10.4 (2.4)	6.1 (1.3)	6.1 (1.3)
Positive mood changes	34.6 (6.0)	53.5 (5.8)	62.0 (8.4)
Negative mood changes	14.1 (5.4)	4.4 (1.9)	5.4 (4.5)
Altruistic/positive social effects	34.9 (7.8)	56.3 (6.0)	62.2 (7.1)
Antisocial/negative social effects	3.5 (1.7)	2.8 (1.7)	3.7 (2.2)
Positive behavior changes	52.9 (9.3)	65.3 (7.9)	80.0 (4.9)
Negative behavior changes	7.1 (5.0)	0.0 (0.0)	3.3 (2.3)
Increased spirituality	40.0 (7.4)	55.1 (6.0)	60.5 (7.1)
Decreased spirituality	3.4 (1.3)	1.2 (0.6)	1.0 (0.5)
How personally meaningful was the experience? (score range: 1–8) ^a	5.4 (0.5)	6.3 (0.2)	6.3 (0.3)
How spiritually significant was the experience? (score range: 1–6) ^b	3.4 (0.4)	4.2 (0.2)	4.4 (0.4)
Did the experience change your sense of well-being or life satisfaction? (score range: -3 to +3) ^c	1.4 (0.5)	2.5 (0.2)	2.7 (0.3)

*Data are mean scores with 1 SEM shown in parentheses ($N=15$); data on attitudes, mood, altruistic/social effects, and behavior changes are expressed as percentage of maximum possible score; data for the final three questions are raw scores.

^aRating scale: 1=no more than routine, everyday experiences. 2=similar to meaningful experiences that occur on average once or more a week. 3=similar to meaningful experiences that occur on average once a month. 4=similar to meaningful experiences that occur on average once a year. 5=similar to meaningful experiences that occur on average once every 5 years. 6=among the 10 most meaningful experiences of my life. 7=among the five most meaningful experiences of my life. 8=the single most meaningful experience of my life.

^bRating scale: 1=not at all. 2=slightly. 3=moderately. 4=very much. 5=among the 5 most spiritually significant experiences of my life. 6=the single most spiritually significant experience of my life.

^cRating scale: -3=decreased very much. -2=decreased moderately. -1=decreased slightly. 0=no change. 1=increased slightly. 2=increased moderately. 3=increased very much.

^dOne participant had incomplete Persisting Effects data for session 1 and was excluded from analysis.

Smoking cessation outcomes were assessed using Timeline Follow-back, and biomarker data (breath CO, urine cotinine). Changes in smoking between study intake and 6-month follow-up were examined with 2-tailed paired *t*-tests. For the Timeline Follow-back data, the *t*-test compared mean cigarettes per day between the 30 days prior to study intake and the 6 months after TQD. These *t*-tests were conducted for both the entire study sample ($N=15$) and the subsample of participants who continued smoking post-TQD ($n=3$).

Repeated measures ANOVA tested for changes in Questionnaire on Smoking Urges, Smoking Abstinence Self-Efficacy, and Wisconsin Smoking Withdrawal Scale scores across 10 time points from intake to 6-month follow-up, and tests for linear contrasts were performed to examine directionality of change over time. Lifetime Mysticism scale scores were compared between intake and 1-week post final psilocybin session using 2-tailed paired *t*-tests to assess effects of psilocybin administration on scores of mystical experience. Descriptive statistics were calculated to characterize Persisting Effects Questionnaire data regarding positive and negative effects across psilocybin sessions (Table 4), and mechanisms attributed to psilocybin for smoking cessation (Table 5).

Results

Safety data

Twelve participants completed three psilocybin sessions. Three participants did not undergo a third session, but completed the study. One participant chose a moderate dose on the second psilocybin session; all other participants chose the default

recommended dosing sequence (moderate in first session and high in subsequent sessions). During the 42 psilocybin sessions (16 moderate, 26 high dose) conducted in the course of this study, no clinically significant adverse events requiring physician or pharmacologic intervention occurred. As expected from previous research (Griffiths et al., 2011), States of Consciousness Questionnaire data showed that one participant (7%) reported extreme ratings, and five others (33%) reported strong ratings of fear, fear of insanity, or feeling trapped at some time during a session. These episodes occurred in six participants (40%), and occurred during five moderate and five high-dose sessions. They were readily managed by interpersonal support, and had resolved by the end of the sessions.

Consistent with previous findings (Griffiths et al., 2006; Grob et al., 2011), blood pressure and heart rate were elevated during drug effects, with peak values occurring on average between 1.5 and 2.5 hours post-psilocybin administration. Systolic blood pressure showed a mean (SD; range) peak of 153 (11; 134–173) mmHg (compared with baseline: 125 [10; 105–153] mmHg). Diastolic blood pressure showed a mean (SD; range) peak of 87 (11; 72–105) mmHg (compared with baseline: 71 [8; 55–89] mmHg). HR showed a mean (SD; range) peak of 87 (11; 66–120) beats/min (compared with baseline: 68 [9; 51–89] beats/min).

Visual Effects Questionnaire data showed no increase in the occurrence of clinically significant or bothersome visual effects, comparing intake and 6-month follow-up assessments.

Of the 10 participants assessed for headache, eight reported at least one post-psilocybin headache with a mean (SD; range) duration of 5.8 (2.4; 2.0–9.5) hours, onset at a mean of 6.2 (2.1;

Table 5. Mechanisms attributed to psilocybin for smoking cessation at 3 weeks post-TQD*.

Mechanism ^a	No. endorsed (%)	Mean rank order ^b
By changing how you prioritize values in life, so that reasons to smoke no longer outweighed reasons to quit	10 (67.7%)	1.9
By changing the way you orient yourself concerning the future, such that you now act in your long-term holistic benefit, rather than acting in response to immediate desire	11 (73.3%)	2.1
By reframing your quitting and staying quit as a sacrament or spiritual task	8 (53.3%)	2.4
By strengthening your belief that you have the ability to quit and stay quit	11 (73.3%)	3.2
By reducing the amount of stress involved with quitting and staying quit	7 (46.7%)	3.6
Not applicable. The sessions did not help me quit or stay quit	1 (6.7%)	
Other ^c		
"By helping me understand that my true self is and always has been a non-smoker" –Participant 1	1 (6.7%)	
"By seeing the 'so much more' than smoking and how it is interconnected" –Participant 8	1 (6.7%)	
"Unclear as to the effect of the psilocybin session on quitting smoking" –Participant 9	1 (6.7%)	
"Gain[ed] distance in thinking" –Participant 11	1 (6.7%)	
"By revealing the real issues that I was using the smoking to mask" –Participant 15	1 (6.7%)	

*TQD=Target Quit Date.

^aParticipants were asked: "If you believe the most recent psilocybin session helped you quit smoking or remain quit, then how do you believe this effect was achieved? (Check all that apply)."

^bParticipants were asked: "Please rank order the following items in terms of their importance to helping you quit smoking or remain quit." Participants did not rank order *Not applicable* or *Other* items.

^cFive participants endorsed *Other*. Each wrote in an additional mechanism in his/her own words.

3.5–11.5) hours after psilocybin administration and mean severity rating of mild, or 2.6 (1.3; 1–5) on a scale from 1=*minimal*, to 6=*excruciating*. Five reported use of over-the-counter headache medication the evening following the session to alleviate symptoms. These were considered approved medications for use during the study, as there is no evidence to suggest they would affect smoking cessation outcomes.

Smoking cessation outcomes

Based on the Timeline Follow-back and verified by CO and cotinine measures, 12 of 15 (80%) participants showed seven-day point prevalence abstinence at 6-month follow-up (Figure 1(a, b)). Eleven of these 12 self-reported quitting smoking on their TQD and demonstrated biologically verified smoking abstinence throughout the following 10 weeks of active treatment. One participant of these 12 reported quitting on the TQD and was biologically verified as abstinent at all attended meetings, but was unexpectedly required to leave the country for business, and was therefore unable to undergo a third psilocybin session, or provide CO and urine samples for weeks 6–10 post-TQD. Three of these 12 participants reported self-corrected lapses¹ (consisting of 1, 4, and 48 cigarettes) during the 16-week period between end of treatment and 6-month follow-up. Another participant reported a relapse after 13 weeks of continuous abstinence, smoking an average of 5 cigarettes/day for 14 weeks (compared with a mean of 19 cigarettes/day at intake), but resumed smoking abstinence prior to 6-month follow-up, as biologically confirmed. This participant also reported use of nicotine replacement (lozenges and gum) during the relapse. No other participant reported use of smoking cessation medications throughout the study.

Two-tailed paired *t*-tests demonstrated significant reductions in self-reported daily smoking from intake to 6-month follow-up (TLFB; $t_{14}=11.1$, $p<.001$) among the entire study sample (Figure 2(a)). Smoking biomarker data among the entire sample also demonstrated statistically significant reductions from intake to

6-month follow-up for breath CO levels ($t_{14}=3.8$, $p<.01$), and urine cotinine ($t_{14}=2.3$, $p=.04$).

Three participants tested positive for smoking at 6-month follow-up, and reported periods of 4, 11, and 22 days of smoking abstinence post-TQD, with two showing >6 days biologically verified continuous abstinence. These individuals ultimately resumed daily smoking. However, a 2-tailed paired *t*-test analysis of Timeline Follow-back data for these three participants revealed significantly reduced smoking post-TQD (Figure 2(b)), with a reported average of 20 cigarettes/day before TQD, and an average of 14 cigarettes/day afterwards ($t_2=5.3$, $p=.03$). Two-tailed paired *t*-tests found no significant differences in these three participants' smoking biomarker data from intake to 6-month follow-up for breath CO or urine cotinine levels.

Repeated measures one-way ANOVA of the entire study sample ($N=15$) showed significant differences across time points for Smoking Abstinence Self-Efficacy confidence ($F_{2,34}=24.9$, $p<.001$), Smoking Abstinence Self-Efficacy temptation ($F_{3,43}=18.5$, $p<.001$), Questionnaire on Smoking Urges ($F_{3,39}=12.7$, $p<.001$), and Wisconsin Smoking Withdrawal Scale scores ($F_{4,46}=4.0$, $p=.009$). Post-hoc testing for linear contrast found significantly increased confidence to abstain (SASE; $p<.001$) from intake to 6-month follow-up, as well as significantly decreased craving (QSU; $p<.001$) and temptation to smoke (SASE; $p<.001$) across all time points. Withdrawal scores (WSWS) fluctuated over time, peaking at 1-week post-TQD, and decreasing significantly through 6-month follow-up ($p<.001$). One participant had incomplete Wisconsin Smoking Withdrawal Scale data, and was excluded from analysis. Figure 1(c–f) illustrates mean scores on these measures.

Persisting effects

Lifetime Mysticism scale scores indicated a significant increase in mystical experience from intake to 1-week post final psilocybin session (difference $M=+54$; $t_{14}=3.5$, $p=0.004$). Participants

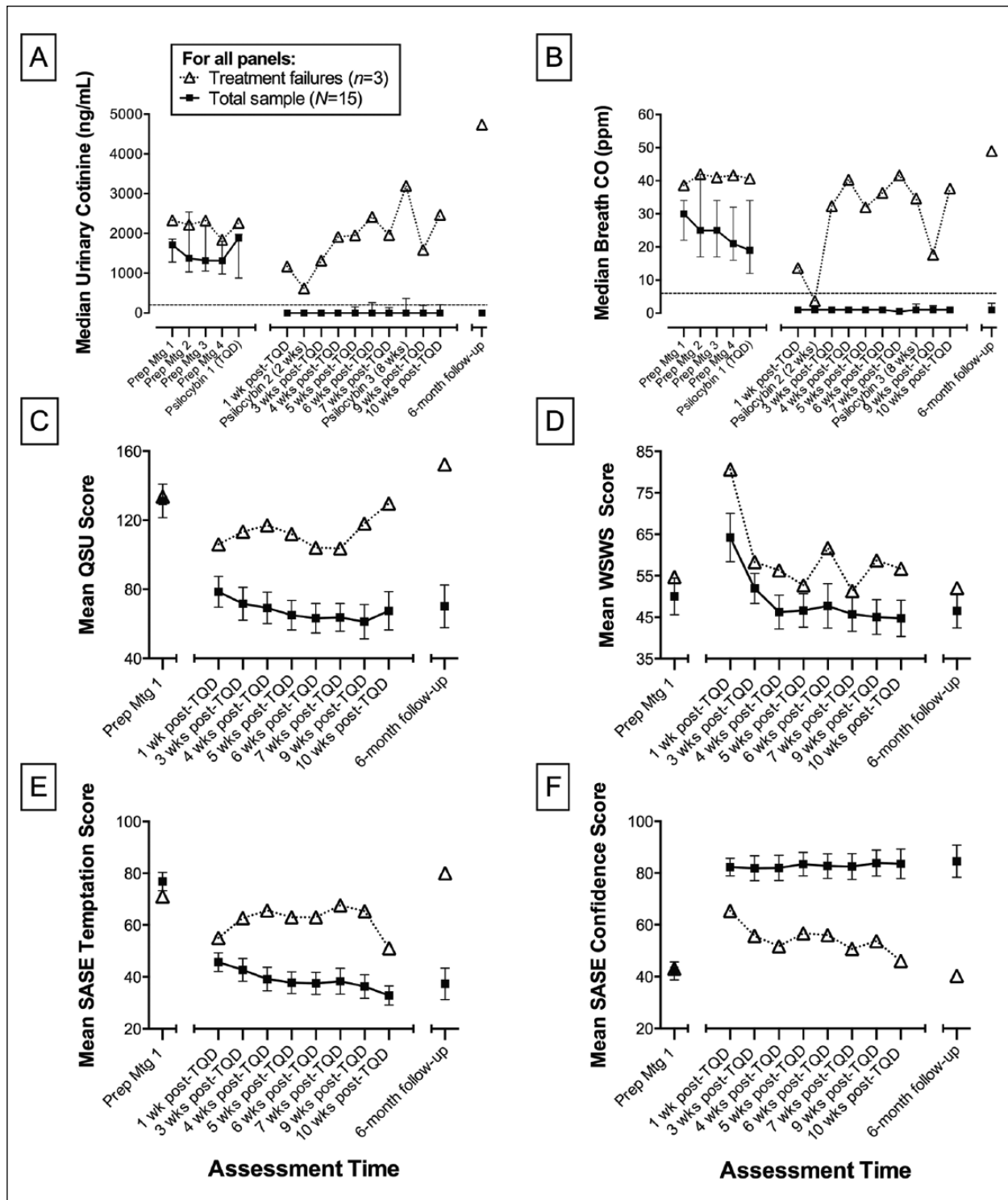


Figure 1. Smoking cessation measures. Median (interquartile range) of urinary cotinine (A) and breath CO (B) at all time points are shown. Prep Mtg=Preparation Meeting. TQD=Target Quit Date. Threshold values for determining non-smoking status are indicated at dotted line (<200 ng/mL cotinine; \leq 6ppm breath CO). Mean (SEM) of Questionnaire on Smoking Urges (QSU) scores (C), Wisconsin Smoking Withdrawal Scale (WSWS) scores (D), Smoking Abstinence Self-Efficacy (SASE) temptation (E) and confidence (F) scores at Prep Mtg 1, at 1, 3, 4, 5, 6, 7, 9, and 10 weeks post-TQD, and at 6-month follow-up are shown. C, The QSU contains 32 items (e.g. "Smoking would make me feel happier now"), rated on a 7-point scale ranging from *strongly disagree*, to *strongly agree* (Range=32–224). D, The WSWS contains 28 items (e.g. "I have been tense or anxious") rated on a 5-point scale ranging from *strongly disagree*, to *strongly agree* (Range=28–140). E, The SASE assesses temptation to smoke and confidence in smoking abstinence (F) in 20 hypothetical situations (e.g. "At a bar or cocktail lounge having a drink") rated on a 5-point scale ranging from *not at all*, to *extremely* (Range=20–100).

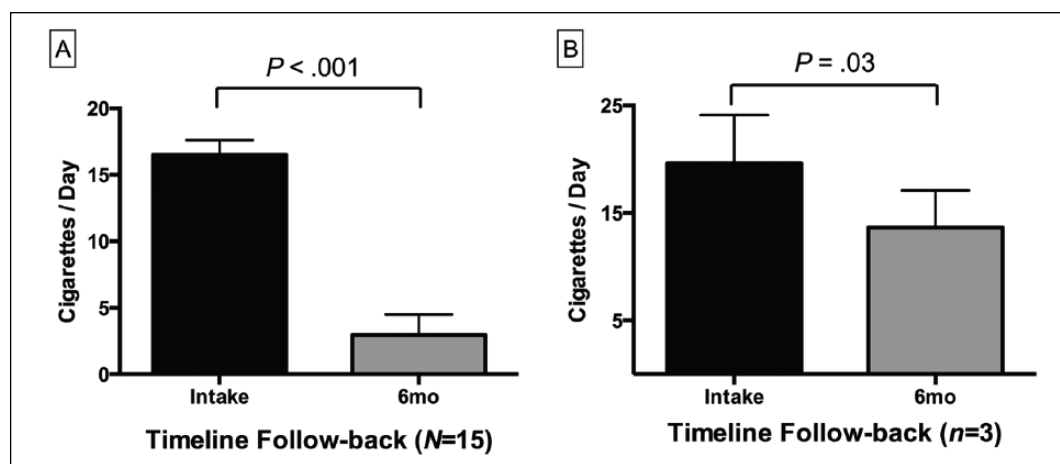


Figure 2. Smoking self-report data. Mean (SEM) of Timeline Follow-back data at intake and 6-month follow-up for the entire study sample, $N=15$ (A), and for participants who tested positive for smoking at 6-month follow-up, $n=3$ (B). 6mo=6-month follow-up. Results shown are for 2-tailed paired t -tests comparing average daily smoking.

reported greater positive than negative effects across all Persisting Effects Questionnaire subscales, with a mean (SD) score of 5.3% (3.6) across negative subscales, and a mean (SD) score of 55.8% (12.1) across positive subscales (scores expressed as percentage of maximum possible score; Table 4).

Participants attributed substantial personal meaning to their psilocybin experiences, with 13 (87%) rating at least one psilocybin session among the 10 most meaningful experiences of their lives. Eleven (73%) rated at least one psilocybin session among the five most spiritually significant experiences of their lives, and 13 (87%) reported that their personal well-being or life satisfaction had increased very much as a result of at least one psilocybin session.

Participants were asked to rate several mechanisms by which psilocybin may have helped in quitting smoking at 3-weeks post-TQD (Table 5). The most commonly endorsed items (paraphrased) included: changing orientation toward the future, so that long-term benefits outweighed immediate desires (73%); strengthening participants' belief in their ability to quit (73%); and changing life priorities/values, such that smoking was no longer more important than quitting (68%). Only one participant (7%), who exhibited the shortest duration of abstinence (4 days) among the study sample, responded that psilocybin had not helped in smoking cessation.

Discussion

This is the first study to provide preliminary data on the safety and feasibility of psilocybin as an adjunct to smoking cessation treatment. The present results are consistent with previous studies examining 5-HT_{2A}R agonists in the treatment of drug dependence (Chwelow et al., 1959; Hollister et al., 1969; Krebs and Johansen, 2012; Savage and McCabe, 1973), suggesting both safety and feasibility, which are discussed in turn.

Our results show promise regarding the safety of psilocybin as an adjunct to smoking cessation treatment. Adverse effects associated with psilocybin consisted of modest acute increases in blood pressure, heart rate, dysphoric subjective effects (e.g. anxiety, fear; <7 hours), and headaches (<24 hours). Consistent with

previous research administering psilocybin in controlled settings, these effects were readily managed (Griffiths et al., 2006, 2011; Grob et al., 2011; Johnson et al., 2008, 2012). Such time-limited effects stand in contrast to adverse effects (e.g. nausea, insomnia, abnormal dreams) associated with approved smoking cessation medications requiring daily administration (e.g. bupropion, varenicline; Gonzales et al., 2006; Jorenby et al., 2006). An advantage of the temporally confined adverse effects of psilocybin is that staff and medical personnel can readily respond with appropriate support and treatment.

Results of the present pilot study also support the feasibility of the approach, as 80% of participants were abstinent at 6-month follow-up. Results should be interpreted with caution given the small N and open-label design. Therefore, no definitive conclusions can be drawn about the causal role of psilocybin per se. However, abstinence rates were substantially higher than typical. For example, when paired with 12 brief weekly counseling meetings, pharmacotherapies have shown seven-day point prevalence abstinence rates of 24.9% to 26.3% (bupropion) and 33.5% to 35.2% (varenicline) at approximately 6 months post-TQD (Gonzales et al., 2006; Jorenby et al., 2006). Furthermore, a randomized controlled trial of the Quit for Life CBT program that provided the primary foundation for the manualized intervention used in this study found a 17.2% abstinence rate at 6-month follow-up (Sykes and Marks, 2001), although participants in our study received substantially more contact with study staff.

Our study provided higher levels of psychosocial support than typical in smoking cessation treatment. However, efficacy rates were higher than observed in studies utilizing similarly extensive CBT-based support. For example, in two trials of extended smoking cessation treatments using a combination of bupropion, nicotine replacement, and CBT ranging from 5 to 12 months in duration, participants showed 45% to 59% seven-day point prevalence abstinence at approximately 6 months (Hall et al., 2009; Killen et al., 2008). Another issue is the relative racial homogeneity and high education levels of the present study sample, which may have impacted the outcomes. Future studies would benefit from more diverse samples.

The study design was unable to discern differential benefits of moderate-dose (20 mg/70 kg) and high-dose (30 mg/70 kg) sessions. All participants who quit smoking ($n=12$) did so after their initial moderate-dose session, and those who did not quit ($n=3$) were unable to do so even after their subsequent high-dose sessions.

One potential concern is the use of an abused drug (psilocybin) in the treatment of dependence on another drug (tobacco). However, 5-HT_{2A}R agonists do not engender compulsive drug seeking (National Institute on Drug Abuse, 2001, 2005), consistent with evidence that they are not reliably self-administered in animals (Fantegrossi et al., 2004; Griffiths et al., 1980; Poling and Bryceland, 1979). Furthermore, the observation that two participants voluntarily declined a third psilocybin session suggests a lack of psilocybin seeking in the context of the present study.

Several plausible mechanisms of hallucinogen-facilitated treatments have been proposed (Bogenschutz and Pommy, 2012; Carhart-Harris et al., 2012, 2014; Ross, 2012; Vollenweider and Kometer, 2010). However, the mechanistic role of psilocybin in smoking cessation remains unclear. Participant responses in the present study suggest that increased temporal horizon, increased self-efficacy, and altered life priorities may be involved. The present results regarding tobacco addiction, combined with previous studies showing efficacy of 5-HT_{2A}R agonists for treatment of alcoholism and opioid dependence (Chwelow et al., 1959; Hollister et al., 1969; Krebs and Johansen, 2012; Savage and McCabe, 1973), suggest higher-order psychological and/or biological mechanisms related to addiction are involved. This contrasts with conventional pharmacotherapies for drug dependence, which are typically specific to the pharmacology of a particular drug class. Valuable directions for future research include qualitative study of participants' accounts regarding potential psychological mechanisms, and neuroimaging studies to inform biological mechanisms.

This study is the first to examine a 5-HT_{2A}R agonist in the treatment of tobacco addiction, and illustrates a viable framework for psilocybin-based addiction treatment interventions. An estimated 5 million worldwide deaths per year are caused by tobacco use, and those numbers are projected to rise to over 8 million deaths annually by 2030 (World Health Organization, 2011). Given the global scope of smoking-related mortality, and the modest success rates of approved smoking cessation treatments (Cahill et al., 2014; Gonzales et al., 2006; Jorenby et al., 2006; Mottillo et al., 2009), the novel approach presented here warrants further investigation with a randomized controlled trial.

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Conflict of interest

Dr. Griffiths is on the board of directors of the Heffter Research Institute, a funding sponsor of the study.

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Note

1. Smoking lapses were defined as any discrete instances of smoking (even a puff of a cigarette) post-TQD, while relapses were defined as smoking on seven or more consecutive days after TQD (Hughes et al., 2003).

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