



# Decriminalize Nature Berkeley

We are suffering under misinformation about traditional entheogenic plants. These are traditional healing plants, respectfully used for religious and healing purposes for millennia, around the world. Although currently illegal outside of limited clinical and religious contexts, these plants aren't addicting, socially destabilizing drugs.

"Entheogenic plants" are different from, for example, amphetamine, cocaine, heroin, alcohol, tobacco. Contrary to political messaging, the science shows that these plants are not addictive - in fact, they help alleviate addiction, depression, and, when used with respect and care, can be deeply healing to individuals and communities. Here are a few facts about these plants, with their scientific references listed at the bottom of the page.

Entheogenic plants and fungi have co-evolved with humans for millennia, first used, possibly, to enhance hunting ability through heightened sensory awareness allowing greater success during hunting and gathering. From there, the plants and fungi began their co-evolution with humans, creating a unique synergistic relationship between these entheogens and human communities via shamanic integration.

Eventually, the sensory and consciousness benefits, including the social cohesion and expansion of consciousness, were brought into the villages and communities. From there, the relationship between the entheogens and humans grew and are thought to be the genesis of the mystical experiences that led to growth of religion, and to expanded capacity of humans to contextualize themselves within the greater cosmos and understand their surroundings as a fully integrated ecosystem and to create a workable narrative of their own existence within this ecosystem.

These plants and fungi have been revered throughout the world, but have become limited to humanity by millennia of colonization, authoritarianism, industrialization, and the disenfranchisement of humans from their ecological roots.

In addition, countless indigenous groups throughout the world, including Native Americans, the Olmec, Zapotec, Maya, and Aztec used peyote and psilocybe mushrooms. Mushrooms, stones, and art date back to 3000 BC in the Americas and as far back as 11,000 BC in northern Africa. The Aztecs would drink chocolate and eat mushrooms called "Teonanácatl" in Nahuatl (literally "god mushroom"—compound of the words teo(tl) (god) and nanácatl

(mushroom)) with honey. The act of ingesting the mushroom sacrament is known as monanacahuia (to mushroom oneself).

Entheogenic plants and fungi are known to increase an individual's respect for the earth and the ecosystem and to heighten human awareness of their own condition within society, a fact that may have led to these plants and fungi being placed on the Federal Schedule 1 list of banned substances.

A top Nixon aide, John Ehrlichman, admitted: "You want to know what this was really all about. The Nixon campaign in 1968, and the Nixon White House after that, had two enemies: the antiwar left and black people. You understand what I'm saying. We knew we couldn't make it illegal to be either against the war or black, but by getting the public to associate the hippies with marijuana and blacks with heroin, and then criminalizing both heavily, we could disrupt those communities. We could arrest their leaders, raid their homes, break up their meetings, and vilify them night after night on the evening news. Did we know we were lying about the drugs? Of course we did."

Wall Street investors are already investing in psilocybin companies which are developing synthetic versions of nature that will cost \$7,000- \$10,000 a gram and potentially reduce access for our community, rather than increase access.

Individual and community empowerment comes from being able to know where our entheogens come from, from our garden, or from our community. We should not have to rely upon synthetically made substances from the pharmaceutical industry that distances us from our relationship to Earth.



## Ayahuasca

- **Substance use disorder and alcohol dependence**

Winkelman, M. (2014). Psychedelics as Medicines for Substance Abuse Rehabilitation: Evaluating Treatments with LSD, Peyote, Ibogaine and Ayahuasca. *Current Drug Abuse Reviews* 7, pp. 101-116.

*Winkelman (2014) describes a successful addiction treatment center called Takiwasi. "For more than 20 years, Jacques Mabit and the Takiwasi center in Peru has been a pioneer in the use of ayahuasca treatments for addiction. The Takiwasi program incorporates ayahuasca in ritual treatments for addicts in remote settings in the Peruvian Amazon. The Takiwasi program is known for its work with cocaine and cocaine paste addicts, as well as a variety of polysubstance users.*

Loizaga-Velder, A. and R. Verres (2014). Therapeutic effects of ritual ayahuasca use in the treatment of substance dependence—qualitative results. *Journal of Psychoactive Drugs* 46(1), pp. 63-72.

*Loizaga-Velder and Verres (2014) qualitative research findings indicate that ayahuasca can serve as a valuable therapeutic tool that, in carefully structured settings, can catalyze neurobiological and psychological processes that support recovery from substance dependencies and the prevention of relapse.*

- **Treatment-resistant depression**

Palhano-Fontes, F., Barreto, D., Onias, H., Andrade, K., Novaes, M., Pessoa, J., . . . Araújo, D. (2019). Rapid antidepressant effects of the psychedelic ayahuasca in treatment-resistant depression: A randomized placebo-controlled trial. *Psychological Medicine*, 49(4), 655-663. doi:10.1017/S0033291718001356

*Palhano-Fontes et al. (2019) randomized placebo controlled trial found significant decreased depression severity, which persisted two and seven days after the trial. For example, on Day 7 the Hamilton Depression Rating scale remission rate was 43% in ayahuasca versus 13% in placebo, and the MADRS remission rate showed 36% ayahuasca and 7% placebo.*

- **Recurrent depression**

de L. Osório, F., et al. (2015). Antidepressant effects of a single dose of ayahuasca in patients with recurrent depression: a preliminary report. *Revista Brasileira de Psiquiatria* 37(1), pp. 13-20.

*de L. Osório, F. et al. (2015) preliminary report on the antidepressant effects of a single dose of ayahuasca suggest fast-acting anxiolytic and antidepressant effects in patients with a depressive disorder.*

- **Grief**

González, D., et al. (2017). Potential Use of Ayahuasca in Grief Therapy. *OMEGA—Journal of Death and Dying*, pp. 1-26.

*Gonzalez et al. (2017) comparison between 6 (avg.) ayahuasca experiences compared to 12 months of peer support found 83.3% (25/30) believed their ayahuasca-drinking experience had a very positive influence on their grieving process 16.7% (5/30) felt it had a positive influence.*

- **Neuroprotective and Parkinson's**

Samoylenko, V., et al. (2010). Banisteriopsis caapi, a unique combination of MAO inhibitory and antioxidative constituents for the activities relevant to neurodegenerative disorders and Parkinson's disease. *Journal of Ethnopharmacology*, 127 (2), pp. 357–367. doi:10.1016/j.jep.2009.10.030.

*University of Mississippi: Samoylenko et al. (2010) study shows inhibition of MAO-B activity by  $\beta$ -carbolines harmine (7) and harmaline (6), in addition to potent MAO-A inhibition responsible for antidepressant activity, provide protection against neurodegeneration, and has a potential therapeutic value for the treatment of Parkinson's diseases.*

Djamshidian, A. , Bernschneider-Reif, S. , Poewe, W. and Lees, A. J. (2016), *Banisteriopsis caapi*, a Forgotten Potential Therapy for Parkinson's Disease?. *Mov Disord Clin Pract*, 3: 19-26. doi:[10.1002/mdc3.12242](https://doi.org/10.1002/mdc3.12242)

*Djamshidian's (2015) research proposes that harmine should be reconsidered as a potential rapidly acting anti-Parkinsonian agent.*

- **Evidence of health and safety**

*Halpern et al. (2008) study shows physical exam and test scores revealed healthy subjects. Members claimed psychological and physical benefits from ayahuasca. 19 subjects met lifetime criteria for a psychiatric disorder, with 6 in partial remission, 13 in full remission, and 8 reporting induction of remission through Church participation. 24 subjects had drug or alcohol abuse or dependence histories with 22 in full remission, and*

*all 5 with prior alcohol dependence describing ayahuasca-based church participation as the turning point in their recovery.*

Harris, R., and Gurel, L. (2012). A Study of Ayahuasca Use in North America. *Journal of Psychoactive Drugs* 44(3): 209-215.

*Harris and Gurel (2012) qualitative study reveals that ayahuasca users reduced their alcohol intake, ate healthier diets, enjoyed improved mood and greater self-acceptance and felt more loving and compassionate in their relationships.*

- **Diabetes**

Wang, P. et al., (2015). A high-throughput chemical screen reveals that harmine-mediated inhibition of DYRK1A increases human pancreatic beta cell replication. *Nature Medicine* 21, pp. 383–388.

*Wang et al. (2015) study using three different mouse and human islet in vivo–based models, we show that harmine is able to induce beta cell proliferation, increase islet mass and improve glycemic control. These observations suggest that harmine analogs may have unique therapeutic promise for human diabetes therapy.*

- **Recidivism**

*In 2013, volunteer therapists working with Acuda, a prisoner’s rights group based in Port Velho, Brazil, began integrating yoga, reiki, and ayahuasca ceremonies as part of a wide-scale rehabilitation effort to help the half-million-plus inmates scattered across the nation.*



## Iboga

Table 1. Open-label case series and clinical trials describing the antiaddictive effects of ibogaine

References	Design/setting	Subjects	Dose	Main results	Safety
Sheppard (1994) <sup>a</sup>	Open-label Non-medical Amsterdam, The Netherland	7 (heroin, methadone, codeine, and ethanol) 5 men, mean age 29.28 years	Single doses 11.7 25 mg/kg	Absence of withdrawal symptoms after 24 38 hr for all subjects, two relapsed within 48 hr, two after several weeks, one reverted to intermittent heroin use, and three remained drug-free >14 weeks	No medical screening or monitoring No serious adverse reactions
Luciano (1998)	Open-label Setting (?)	3 (cocaine, opiates/opioids, and ethanol) Gender, age (?)	Single doses 20 25 mg/kg	Absence of withdrawal and craving symptoms after 24 hr for all subjects Abstinence (?)	Medical screening and monitoring No serious adverse reactions
Alper et al. (1999) <sup>a</sup>	Open-label Non-medical United States and The Netherlands	33 (heroin, methadone, and cocaine) 22 men, mean age 27.3 years	Single doses 6 29 mg/kg	Absence of withdrawal symptoms and drug use for 25 (76%) subjects after 72 hr, four did not report symptoms but used drugs in <72 hr, two reported attenuated symptoms and no drug use in <72 hr, and one reported absence of effects	Medical screening and monitoring one fatality, probably caused by concomitant heroin use
Mash et al. (2000) <sup>b</sup>	Open-label Private clinic St. Kitts, West Indies	27 (heroin, methadone [?] and cocaine) 23 men, mean age 34.6 (opiates/opioids) and 37.5 (cocaine) years	Single doses 500, 600, and 800 mg	Significant ( $p < .005$ ) reductions in all HCQN-29 subscales <sup>c</sup> and in two CCQN-45 subscales <sup>d</sup> after 36 hr and 14 days, and significant ( $p < .0005$ ) reductions in BDI scores after one month Abstinence (?)	Medical screening and monitoring No serious adverse reactions
Mash et al. (2001) <sup>b</sup>	Open-label Private clinic St. Kitts, West Indies	32 (heroin and methadone) 23 men, mean age 33.6 years	Single dose 800 mg	Significant ( $p < .05$ ) reductions in OOWS scores after 12 24 hr and in OP-SCL scores after <72 hr and 6 9 days, and months of abstinence in many subjects (although data were not presented)	Medical screening and monitoring No serious adverse reactions
Alper (2001) <sup>a</sup>	Open-label Non-medical United States and The Netherlands	41 (heroin, methadone, cocaine, sedatives, ethanol) Gender, age (?)	Single doses 6 29 mg/kg	15 (29%) subjects reported abstinence for <2 months, 15 (29%) for $\geq 2$ months/<6 months, 7 (13%) for $\geq 6$ months/<1 year, 10 (19%) for >1 year, and 5 (10%) the outcomes could not be determined	Medical screening and monitoring No serious adverse reactions
Schenberg et al. (2014)	Open-label Private hospital Santa Cruz do Rio Pardo, Brazil	75 (ethanol, cannabis, cocaine, and crack-cocaine) 67 men and 8 women, mean age 34.16 (men) and 29.50 (women) years	Single and multiple doses; mean number of sessions 3.83 (men) and 5.40 (women) 17 20 mg/kg	Significant ( $p < .001$ ) increases in abstinence duration, and 61% of subjects were abstinent after single (median 5.5 months) and multiple (median 8.4 months) doses	Medical screening and monitoring No serious adverse reactions
Glue, Cape, Tunnicliff, Lockhart, Lam, Gray, et al. (2016)	Randomized, double-blind, placebo-controlled, single ascending-dose clinical trial Research unit Dunedin, New Zealand	27 (methadone) 21 men, mean age 41.2 years	Single doses (noribogaine) 60, 120, and 180 mg	Non-significant effects on SOWS, OOWS, COWS, and time to resumption of MST	Medical screening and monitoring No serious adverse reactions Significant QT interval prolongation

Note. BDI, Beck Depression Inventory; CCQN-45, Cocaine Craving Questionnaire; COWS, Clinical Opioid Withdrawal Scale; HCQN-29, Heroin Craving Questionnaire; OOWS, Objective Opiate Withdrawal Scale; OP-SCL, Opiate-Symptom Checklist; MST, methadone substitution treatment; SOWS, Subjective Opioid Withdrawal Scale. Interrogation: not informed.

<sup>a</sup>Included subjects from the same sample.

<sup>b</sup>Included subjects from the same sample.

<sup>c</sup>Desire to Use, Intention to Use, Anticipation of Positive Outcomes, Relief of Negative States, and Lack of Control.

<sup>d</sup>Relief of Negative States and Lack of Control.



## Psilocybe mushrooms

### Psilocybin from Psilocybe mushrooms

- **Smoking Cessation**

Johnson, M. et al. (2017). An online survey of tobacco smoking cessation associated with naturalistic psychedelic use. *Journal of Psychopharmacology* 31 (7), pp. 841-850.

*Johnson et al. (2014) open-label, proof of concept study found the clinical outcome of successful smoking cessation in 80% of the sample (12/15 participants), with biologically verified abstinence 6 months after each participant's planned quit date. 73% of participants rated at least one of their sessions as among the top five most spiritually significant experiences of their lives.*

- **Alcohol dependence**

Bogenschutz, M., et al. (2015). Psilocybin-assisted treatment for alcohol dependence: A proof-of-concept study. *Journal of Psychopharmacology* 29(3), pp. 289-299.

*Bogenschutz et al. (2015) proof of concept study found patients in the study exhibited a significant improvement in drinking after their first psilocybin session and scores on ratings of psilocybin-occasioned mystical experience correlated strongly with change in drinking behavior.*

- **Treatment resistant depression**

Lyons, T. and Carhart-Harris, R. (2018). Increased nature relatedness and decreased authoritarian political views after psilocybin for treatment-resistant depression. *Journal of Psychopharmacology*, 32(7), pp. 811–819.

*Centre for Psychiatry, Department of Medicine, Imperial College London: Lyon et al (2018) study on depressive symptoms: One week after psilocybin treatment, depressive symptoms were significantly reduced to levels more comparable with controls. The*

*patients' depressive symptoms remained significantly reduced at the 7–12-months follow-up.*

- **End-of-life anxiety**

Ross, S., et al. (2016). Rapid and sustained symptom reduction following psilocybin treatment for anxiety and depression in patients with life-threatening cancer: a randomized controlled trial. *Journal of Psychopharmacology*, 30(12), pp. 1165-1180.

*New York University: Ross et al. (2016) double-blind, placebo-controlled, crossover trial, on 29 patients with cancer-related anxiety and depression found “immediate, substantial, and sustained improvements in anxiety and depression and led to decreases in cancer-related demoralization and hopelessness, improved spiritual well being, and increased quality of life.” At the 6.5-month follow-up “approximately 60–80% of participants continued with clinically significant reductions in depression or anxiety.”*

- **Openness**

MacLean, K., et al. (2011). Mystical experiences occasioned by the hallucinogen psilocybin lead to increases in the personality domain of openness. *Journal of Psychopharmacology*, 25(11) 1453–1461.

*Johns Hopkins Medical School: MacLean et al. (2011) combined two double blind studies with 52 participants to show significant increases in Openness after a high dose psilocybin session with mystical experiences that were larger in magnitude than changes in personality typically observed in healthy adults over decades of life experience. Openness remained significantly higher than baseline more than 1 year after the session.*

- **Nature-Relatedness**

Lyons, T. and Carhart-Harris, R. (2018). Increased nature relatedness and decreased authoritarian political views after psilocybin for treatment-resistant depression. *Journal of Psychopharmacology*, 32(7), pp. 811–819.

*Centre for Psychiatry, Department of Medicine, Imperial College London: Lyons et al. (2018) open-label pilot study with a mixed-model design on treatment resistant depression found patients reported being significantly more connected to nature 1 week and 7–12 months after psilocybin treatment compared with baseline.*

- **Decreased Authoritarian**

Lyons, T. and Carhart-Harris, R. (2018). Increased nature relatedness and decreased authoritarian political views after psilocybin for treatment-resistant depression. *Journal of Psychopharmacology*, 32(7), pp. 811–819.

*Centre for Psychiatry, Department of Medicine, Imperial College London: Lyons and colleagues (2018) open-label pilot study with a mixed-model design on treatment resistant depression found patients were significantly less authoritarian 1 week after*

*psilocybin treatment and a trend-level decrease was found at 7–12 months compared with baseline.*

- **Most Meaningful**

Griffiths, R., Richards, W., Johnson, M., McCann, U., & Jesse, R. (2008). Mystical-type experiences occasioned by psilocybin mediate the attribution of personal meaning and spiritual significance 14 months later. *Journal of psychopharmacology (Oxford, England)*, 22(6), 621–632. doi:10.1177/0269881108094300

*Johns Hopkins Medical School: Griffiths et al. (2006, 2008) 67% of the volunteers rated the experience with psilocybin to be either the single most meaningful experience of his or her life or among the top five most meaningful experiences of his or her life, which continued at the 14-month follow-up. 79% of the volunteers rated that the psilocybin experience increased their current sense of personal well-being or life satisfaction “moderately” (50%) or “very much” (29%) Lifetime mystical experience and spiritual transcendence scores were significantly higher in the group that received psilocybin. These results were relatively the same after a 14-month follow-up.*

- **Mystical experiences**

Griffiths, R., Richards, W., Johnson, M., McCann, U., & Jesse, R. (2008). Mystical-type experiences occasioned by psilocybin mediate the attribution of personal meaning and spiritual significance 14 months later. *Journal of psychopharmacology (Oxford, England)*, 22(6), 621–632. doi:10.1177/0269881108094300

*Johns Hopkins Medical School: Griffiths et al. (2006, 2008) 22 of the total group of 36 volunteers had a “complete” mystical experience after psilocybin, staying relatively the same after a 14-month follow-up.*

## General Benefits Indoleamine compounds

- **Reduced suicidality**

Hendricks, P., et al. (2015). Psilocybin, psychological distress, and suicidality. *Journal of Psychopharmacology*, 29(9), pp. 1041–1043.

*Hendricks et al. (2015) found that having ever used any classic psychedelic substance—namely, dimethyltryptamine (DMT), ayahuasca... mescaline, peyote or San Pedro, or psilocybin—was associated with a significantly reduced likelihood of past month psychological distress, past year suicidal thinking, past year suicidal planning, and past year suicide attempt in the United States adult population.*

- **Cluster Headaches**

Schindler, E. et al., (2015) Indoleamine Hallucinogens in Cluster Headache: Results of the Clusterbusters Medication Use Survey, *Journal of Psychoactive Drugs*, 47(5), pp. 372-381. DOI:10.1080/02791072.2015.1107664

*Schindler et al. (2015) survey from Cluster Buster Headaches 496 respondents with diagnosis of cluster headache verified by a neurologist or headache specialist. "Psilocybin... along with another hallucinogen, DMT, were used daily to weekly for abortive purposes (n = 23). For prevention, they were used every few weeks to twice yearly (n = 80). The word "single" or "once" to indicate one dose of psilocybin or LSD was clearly written by eight responders. In contrast to these conventional medications, the current study shows that psilocybin and LSD provided over 70% of those who tried them with at least moderate protection from attacks. Complete preventive efficacy was about 40% for each drug, which is greater than that reported for any other conventional medication."*

- **Intimate Partner Violence & Recidivism**

Walsh, Z. , et al. (2016). Hallucinogen use and intimate partner violence: Prospective evidence consistent with protective effects among men with histories of problematic substance use. *Journal of Psychopharmacology*, pp. 1-7. DOI:10.1177/0269881116642538.

*Walsh et al. (2016) study of 302 inmates at a US county jail found that any lifetime use of hallucinogens was associated with lower rates of Intimate Partner Violence; 26.79% of the hallucinogen-use group were arrested for later IPV (mean survival time=62.76 months) compared with 41.79% of the group that reported no hallucinogen use (mean survival time=54.85 months). 13.64% of the lifetime hallucinogen-use disorder group were arrested for IPV (mean survival time=68.82 months) compared with 35% of the group that reported no lifetime hallucinogen use disorder (mean survival time=58.50 months).*

- **Recidivism**

Hendricks, P. S., Clark, C. B., Johnson, M. W., Fontaine, K. R., & Cropsey, K. L. (2014). Hallucinogen use predicts reduced recidivism among substance-involved offenders under community corrections supervision. *Journal of Psychopharmacology*, 28(1), 62–66. <https://doi.org/10.1177/0269881113513851>

*University of Alabama Observational Study: Hendricks (2014) collected data from 25,622 individuals charged with a felony and under community corrections supervision for individuals with a history of substance involvement. Hallucinogen use disorder was associated with a decreased probability of supervision failure. This stands in contrast to any cannabis, cocaine, alcohol, opiate, and amphetamine use disorder, each of which was associated with an increased probability of supervision failure. Any hallucinogen use disorder was the third strongest predictor of supervision outcome among all predictors, trailing only any cocaine use disorder and any cannabis use disorder (both predicting failure).*

### **Additional Resources**

- Mushrooms: <https://www.iceers.org/psilocybin-mushrooms-basic-info/>
- Cacti containing mescaline: <https://www.iceers.org/san-pedro-basic-info/>
- Ayahuasca: <https://www.iceers.org/ayahuasca-basic-info/>
- Iboga: <https://www.iceers.org/iboga-basic-info/>