

Advancing preclinical models to accelerate the discovery of novel psychedelic therapeutics

*Dr Jack Prenderville
Neuroscience Ireland Meeting
10th September 2021*



The oldest and least understood psychopharmacological agent

Psychedelics are powerful psychoactive substances that alter perception and mood and affect numerous cognitive processes (Nichols, 2016)

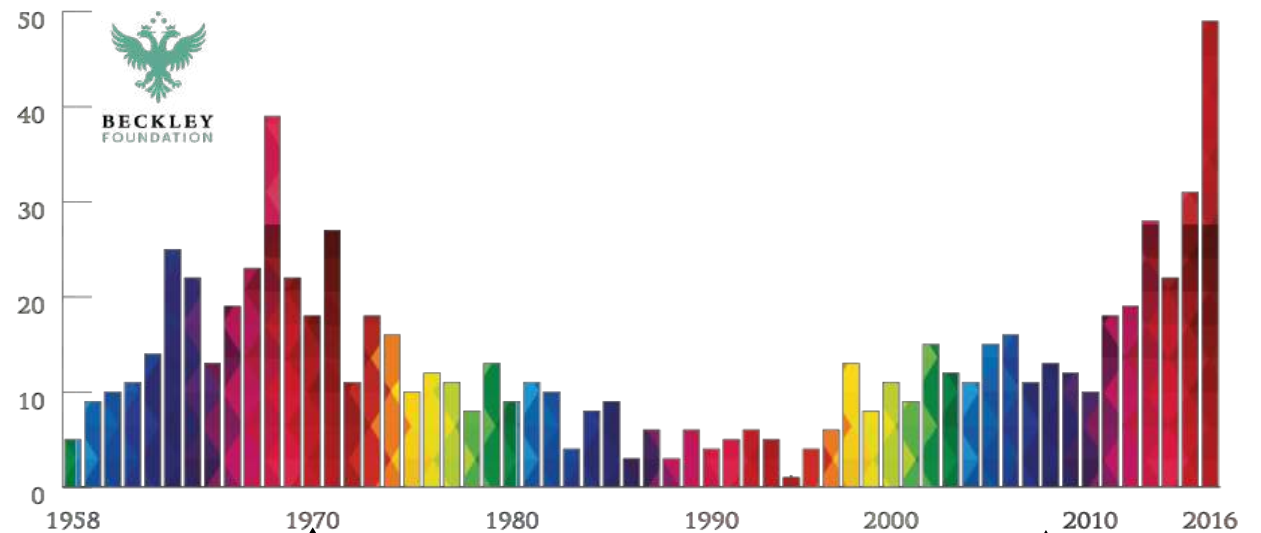
"...was seized by a peculiar sensation of vertigo and restlessness... In a dreamlike state... With my eyes closed, fantastic pictures of extraordinary plasticity and intensive colour seemed to surge towards me."

Albert Hofmann, on his accidental exposure to LSD, 1943

"I lost all control of time: space and time became more and more disorganised and I was overcome with fears that I was going crazy... Occasionally I felt as being outside my body. I thought I had died. My 'ego' was suspended somewhere in space and I saw my body lying dead on the sofa."

Albert Hofmann, on his deliberate exposure to LSD, 1943

Number of scientific articles published about psilocybin



USA passes Controlled Substances Act
Psychedelic drugs - Schedule 1
(most restrictive category)

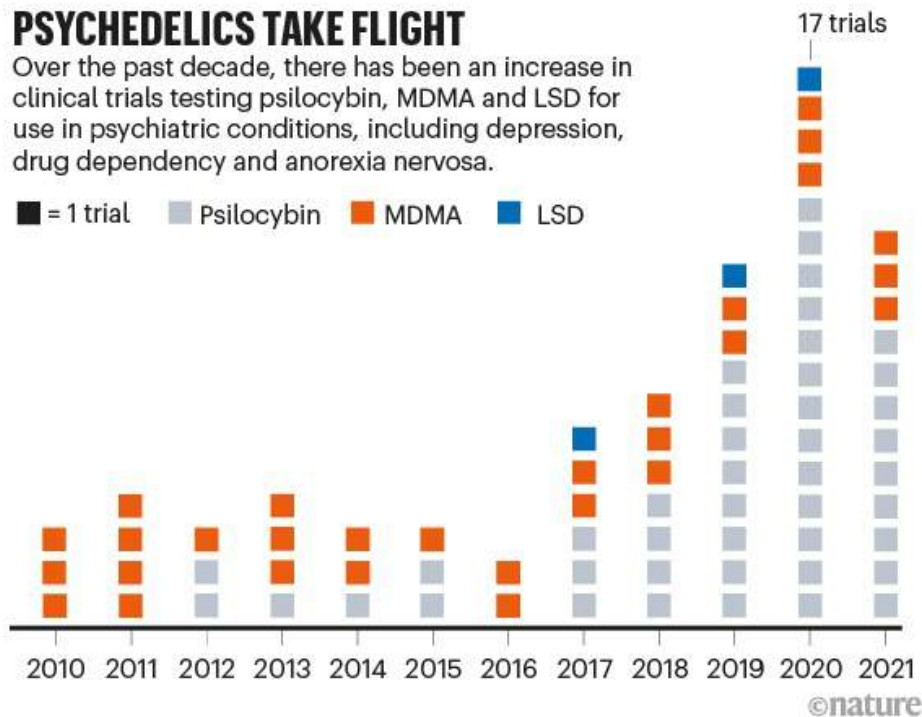
Therapeutic effect of psychedelics
End-of-life anxiety

Psychedelics in the clinic

PSYCHEDELICS TAKE FLIGHT

Over the past decade, there has been an increase in clinical trials testing psilocybin, MDMA and LSD for use in psychiatric conditions, including depression, drug dependency and anorexia nervosa.

■ = 1 trial ■ Psilocybin ■ MDMA ■ LSD



The NEW ENGLAND JOURNAL of MEDICINE

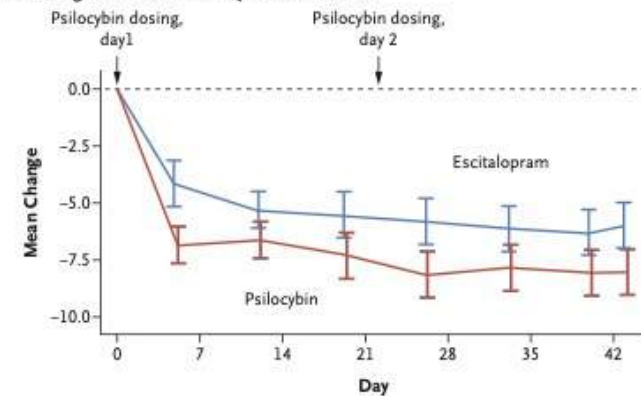
ORIGINAL ARTICLE

Trial of Psilocybin versus Escitalopram for Depression

N Engl J Med 2021;384:1402-11.
DOI: 10.1056/NEJMoa2032994
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Michelle Baker-Jones, B.A., Ashleigh Murphy-Beiner, M.Sc.,
Roberta Murphy, M.D., Jonny Martell, M.D., Allan Blemings, M.Sc.,
David Erritzoe, M.D., and David J. Nutt, M.D.

A Change from Baseline in QIDS-SR-16 Score



B Change from Baseline in WEMWBS Score

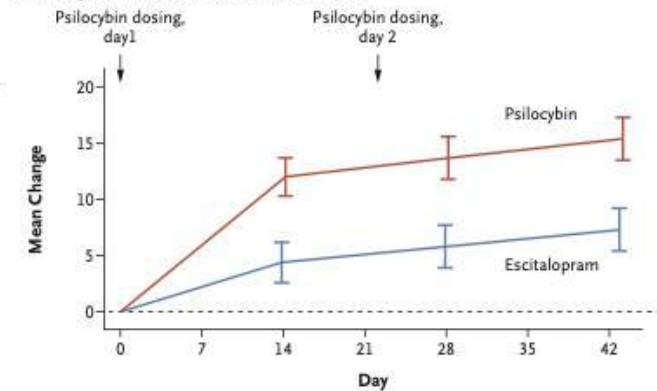
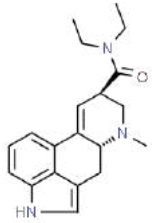


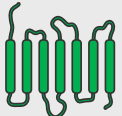
Figure 1. Change in Depression Severity and in Well-Being over 6 Weeks.

Psychedelics (classic serotonergic 'hallucinogens')

Agonist or partial agonist



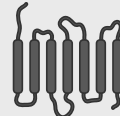
LSD



5-HT_{2A}



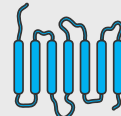
5-HT_{2C}



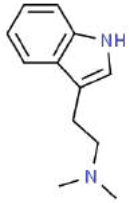
5-HT_{1A}



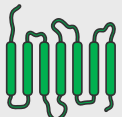
α₂



D₂



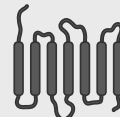
DMT



5-HT_{2A}



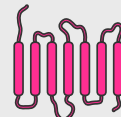
5-HT_{2C}



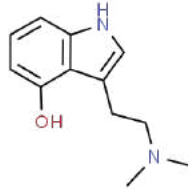
5-HT_{1A}



Sigma-1



TAAR₁



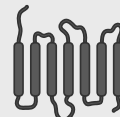
Psilocin
(Psilocybin)



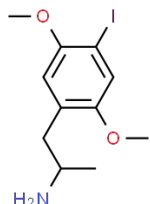
5-HT_{2A}



5-HT_{2C}



5-HT_{1A}



DOI

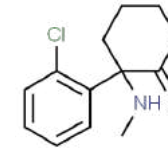


5-HT_{2A}



5-HT_{2C}

Dissociatives



Ketamine

Antagonist

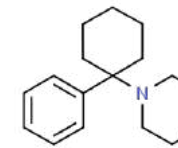


NMDA

Agonist



D₂



PCP



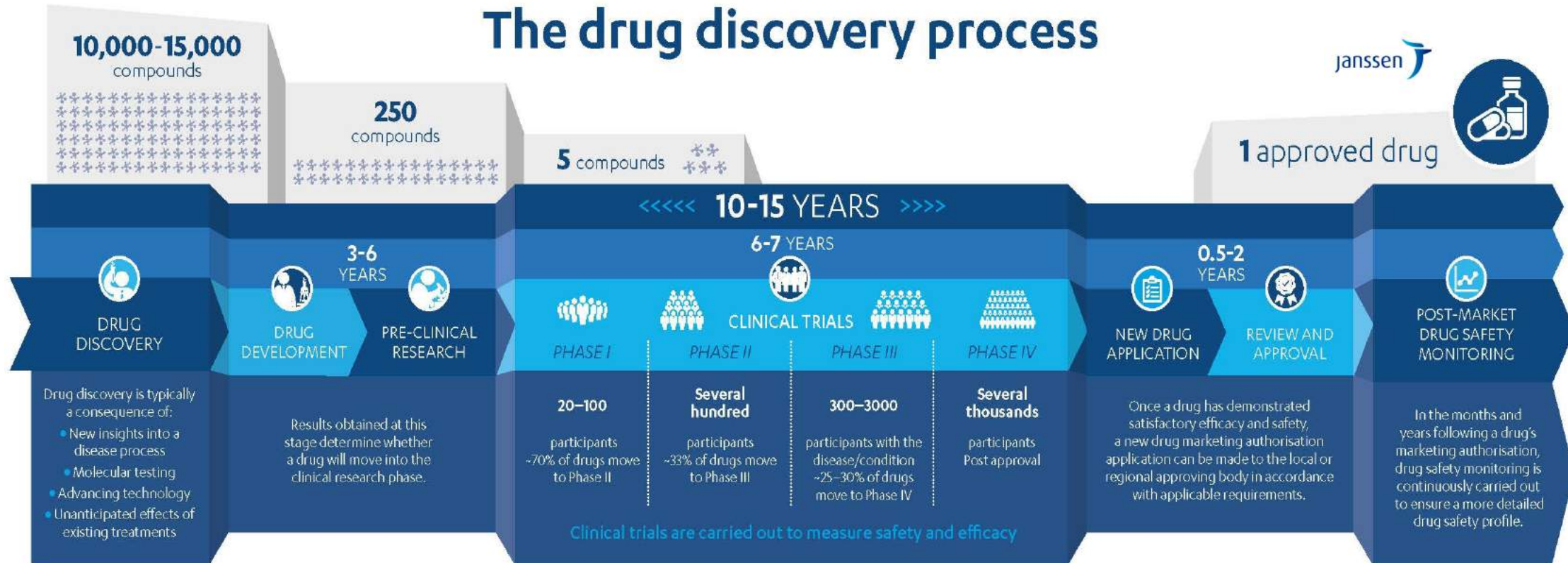
NMDA



D₂

How can preclinical models support (psychedelic) discovery?

The drug discovery process



I. Target engagement

II. Preclinical efficacy testing

III. Safety pharmacology

IV. Mechanism of action / biomarkers

5HT-2A receptor target engagement – Head Twitch Response

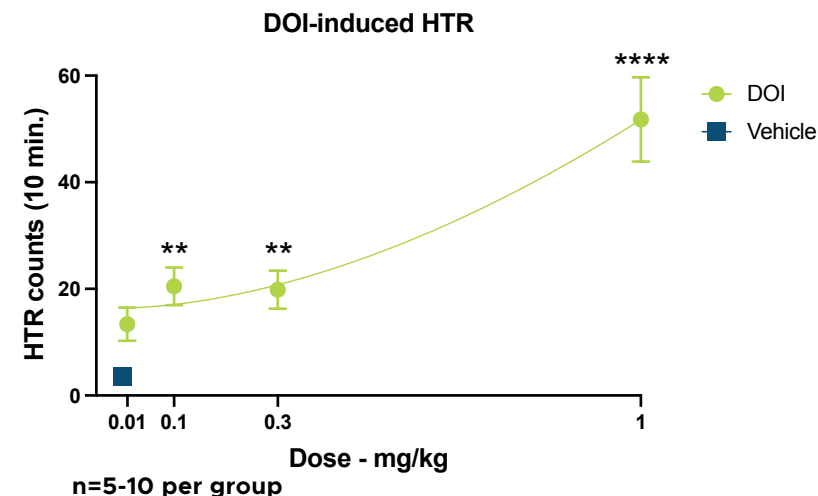
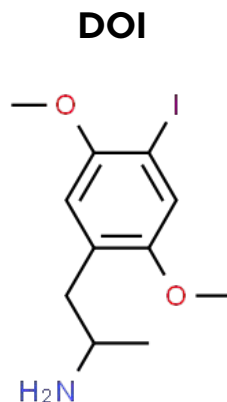
Science 8 September 1956 “Permanent” Alteration of Behavior in Mice by Chemical and Psychological Means

DORIS L. KELLER

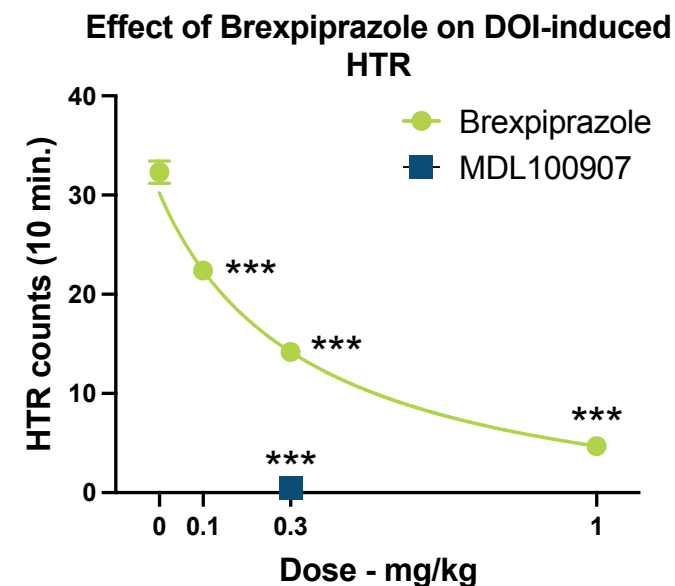
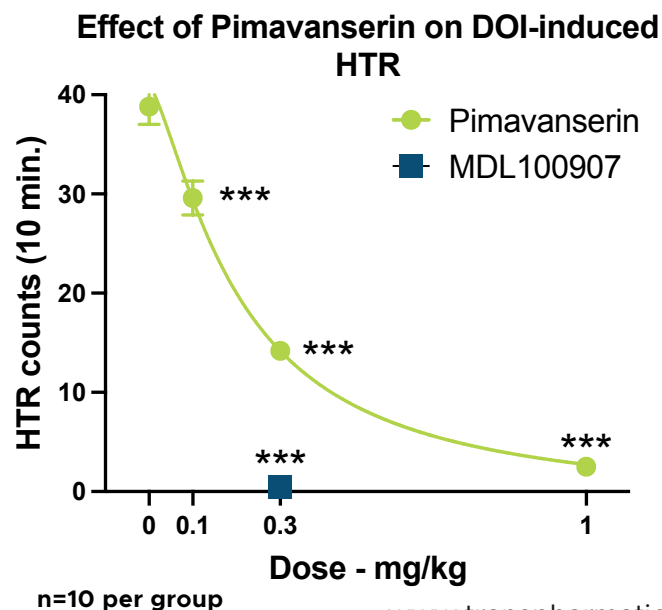
WAYNE W. UMBREIT

Merck Institute for Therapeutic
Research, Rahway, New Jersey

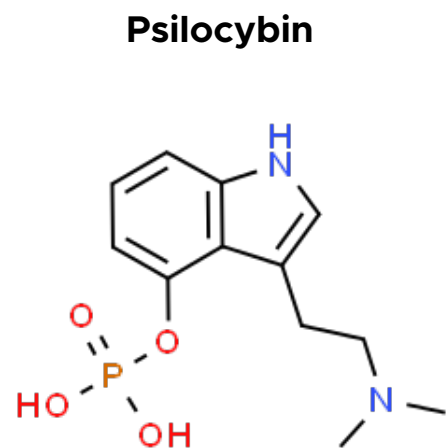
The response consists of a rapid and violent head shaking when any area about the back of the head is touched very lightly with a small stick or pencil point. The head-twitch response does not occur in normal mice, and with a little experience the response is easy to detect. It is only rarely that one is uncertain whether a particular animal possesses the head twitch or not, and



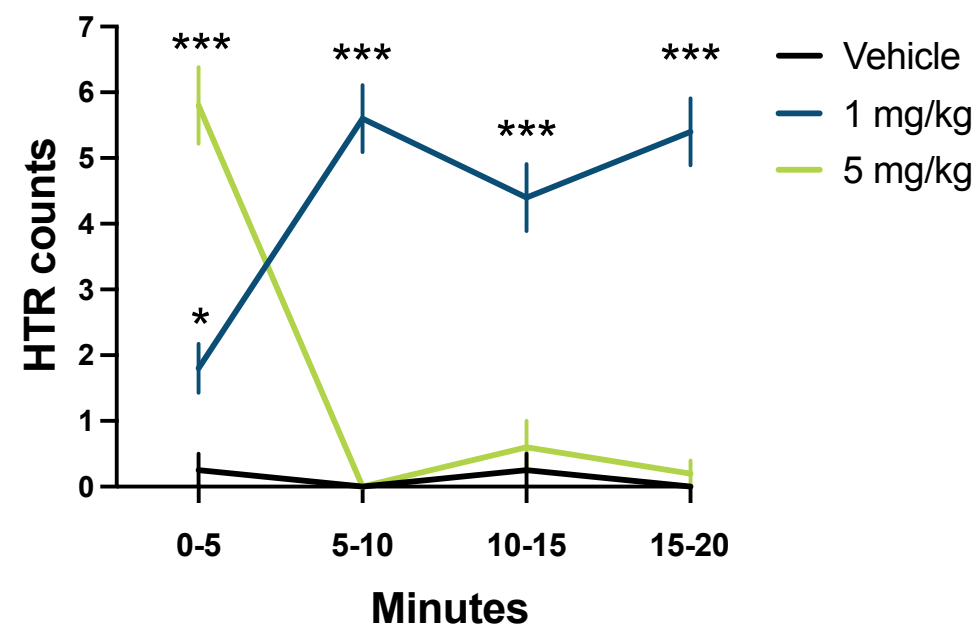
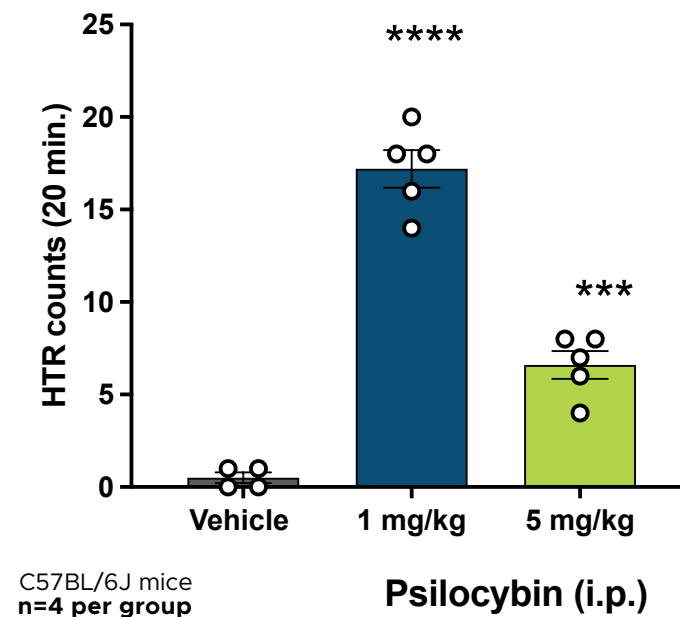
DOI-induced HTR: 5-HT_{2A} receptor antagonists



Target engagement - HTR

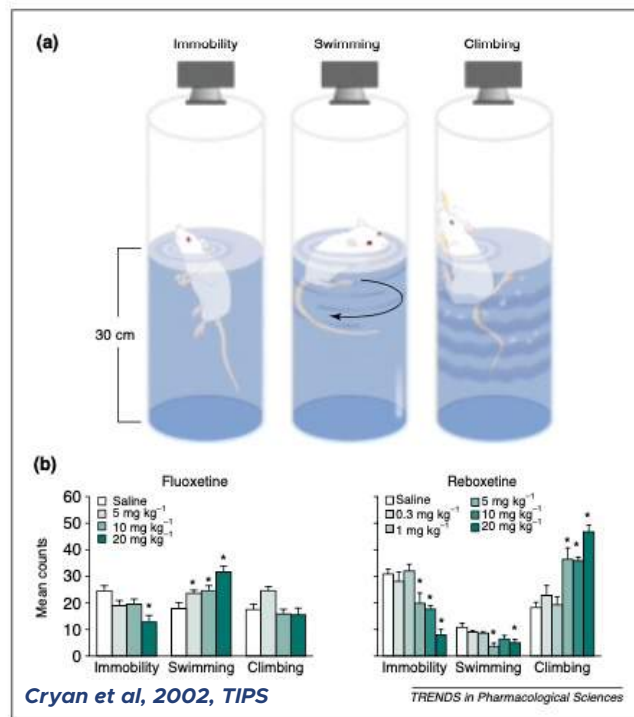


Psilocybin-induced HTR



Preclinical Efficacy testing - Depression

The forced swim test (FST): monoaminergic antidepressants



The Rodent Forced Swim Test Measures Stress-Coping Strategy, Not Depression-like Behavior

Kathryn G. Commons,^{*} Aram B. Cholanians, Jessica A. Babb, and Daniel G. Ehlinger

Immobility in the forced swim test is adaptive and does not reflect depression

Marc L. Molendijk^{a,*,1}, E. Ronald de Kloet^{b,1}

Depression researchers rethink mouse swim test

Animal-rights group campaigns to end test that some scientists say is overused.

Too Depressed to Swim or Too Afraid to Stop? A Reinterpretation of the Forced Swim Test as a Measure of Anxiety-Like Behavior

Jeffrey Anyan¹ and Shimon Amir^{2,1}

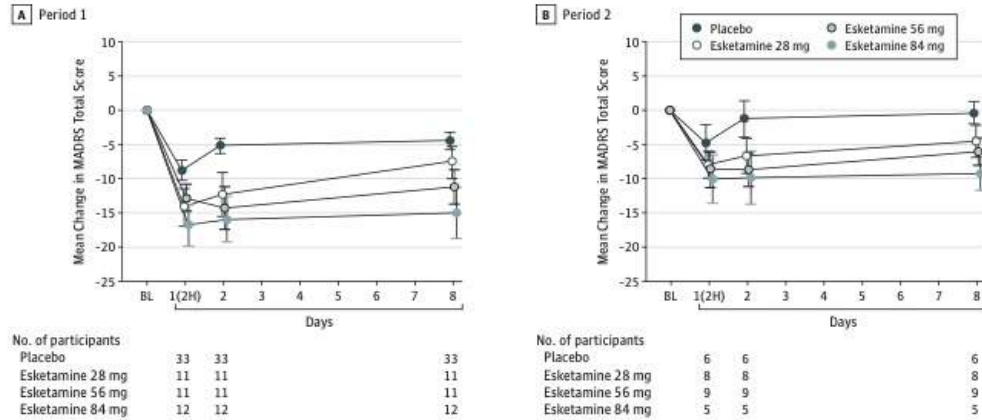
¹Center for Studies in Behavioral Neurobiology, Department of Psychology, Concordia University, Montreal, Quebec, Canada

Drug	Mechanism of action	Effect on FST immobility (rat)
Fluoxetine	SSRI	↓
Sertraline	SSRI	↓
Paroxetine	SSRI	↓
Desipramine	NRI	↓
Maprotiline	NRI	↓
Reboxetine	NRI	↓
Bupropion	NDRI (?)	↓
Venlafaxine	SNRI	↓
Milnacipran	SNRI	↓
Duloxetine	SNRI	↓
Mianserin	NaSSA (?)	↓

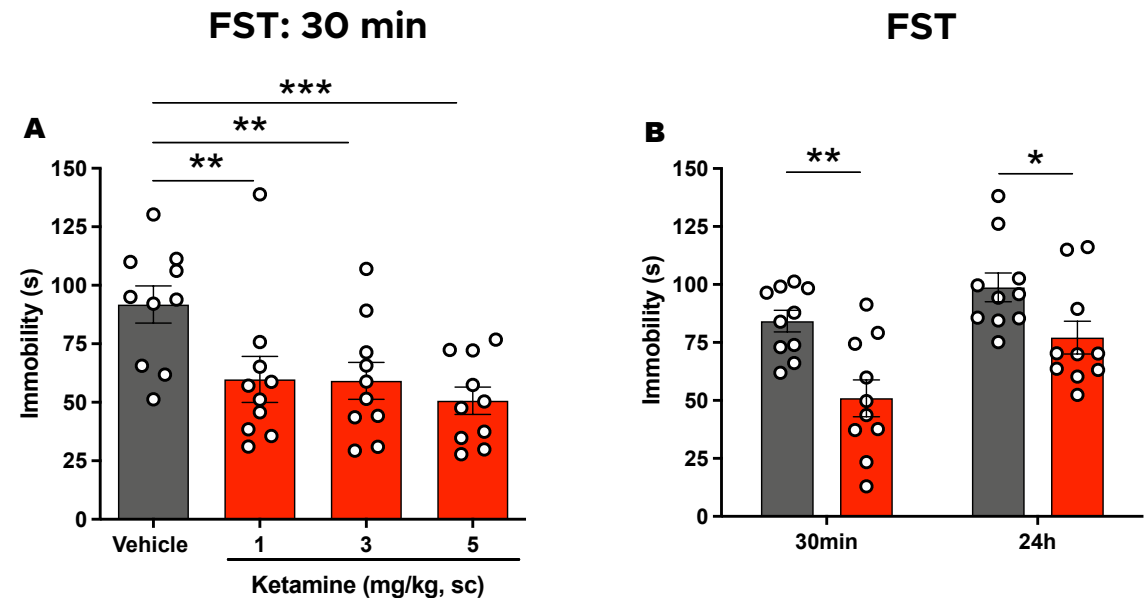
Adapted from Cryan et al, 2005, *Neurosci. Biobehav. Rev.*

Novel antidepressants and the FST: (es)ketamine

Daly et al, 2018, JAMA Psychiatry



The FST: glutamatergic antidepressants - ketamine
(McDonnell et al, 2021, Frontiers in Psychiatry)



Wistar K yoto rats
n=9-10 per group

* Significant effect compared to Vehicle. One-way ANOVA with Fishers LSD.

PHARMACEUTICAL NEWS

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FDA APPROVES NEW NASAL SPRAY MEDICATION FOR TREATMENT-RESISTANT DEPRESSION

Spravato[™]
(esketamine) [®]
nasal spray

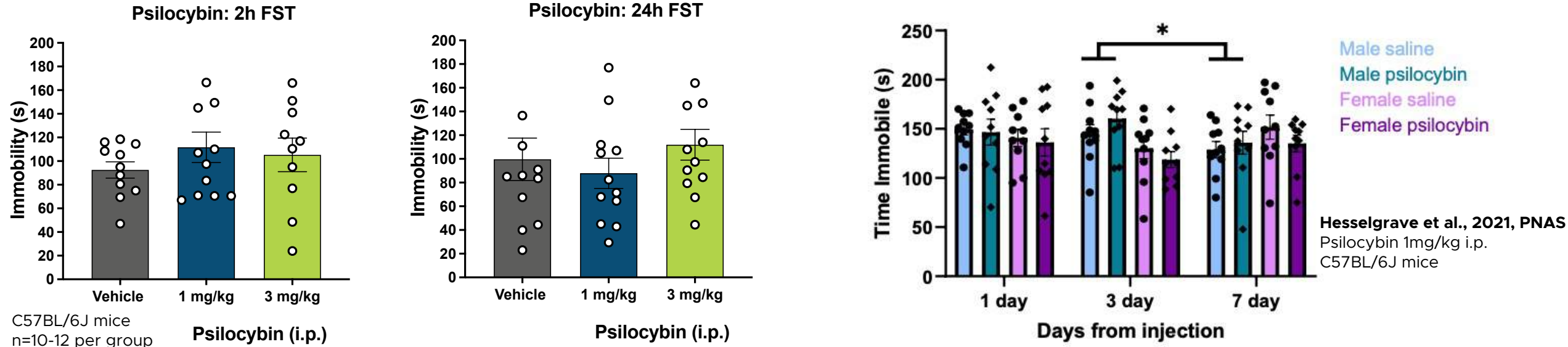
What is SPRAVATO[™]?
SPRAVATO[™] is a prescription medicine, used along with an antidepressant taken by mouth, for treatment-resistant depression (TRD) in adults.

Esketamine

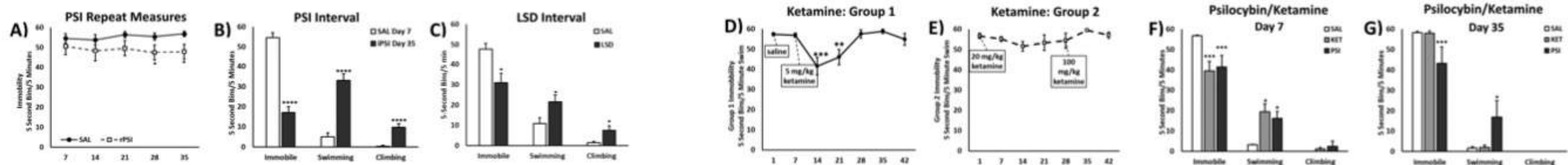
Esketamine (2S)-2-(2-chlorophenyl)-2-(methylamino)cyclohexan-1-one is the S- (more active) enantiomer of ketamine.

Key question: is the FST a suitable tool for psychedelic research?

Psilocybin and the forced swim test



Psilocybin (and LSD) produce persistent antidepressant-like effects in the FST



Hibicke et al., 2020, ACS Chem. Neuro.
Psilocybin 1mg/kg i.p.; LSD 0.15mg/kg i.p.
WKY rats

Measuring rodent motivation: the progressive ratio task

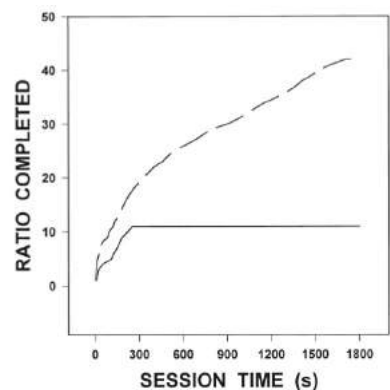
Anhedonia: Inability to feel pleasure in normally pleasurable activities

‘Markedly diminished interest or pleasure in all, or almost all activities most of the day, nearly every day’
(symptom of depression as in DSM-5)

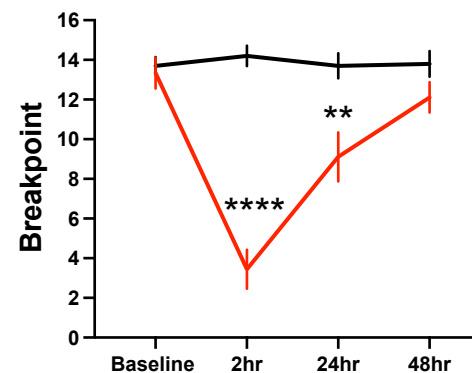
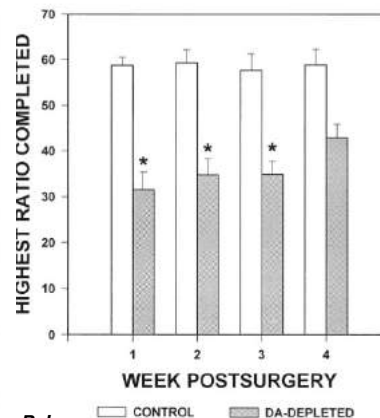
Motivational anhedonia: loss of interest or absence of anticipatory pleasure

Consummatory anhedonia: loss of pleasure in response to previously rewarding stimuli

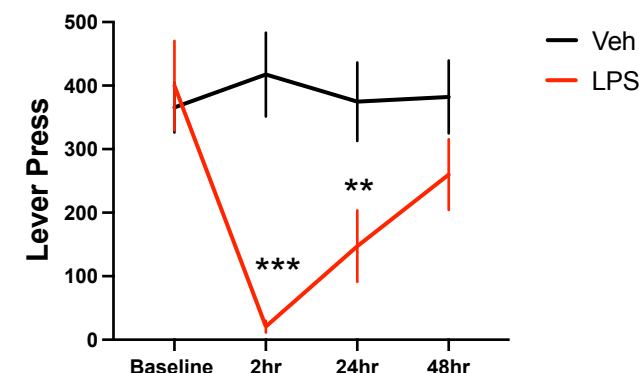
Rat progressive ratio (PR) task



Aberman et al., 1998, Pharmacol Biochem Behav



* Significant effect compared to baseline. One-way repeated measures ANOVA with Fishers LSD post-hoc (n=9-10 per group)



- Responses for successive reinforcers (sucrose pellet) increases according to the progression:
2, 4, 6, 9, 12, 15, 20, etc.

- Endpoints: **Lever Press** and **Breakpoint**

Micro doses of psilocybin enhance motivation



Low Doses of Psilocybin and Ketamine Enhance Motivation and Attention in Poor Performing Rats: Evidence for an Antidepressant Property

Guy A. Higgins^{1,2*}, Nicole K. Carroll¹, Matt Brown¹, Cam MacMillan¹, Leo B. Silenies¹, Sandy Thevarkunne¹, Julia Izhakova³, Lilia Magomedova², Ines DeLannoy^{3,4} and Edward M. Sellers^{2,5}

¹InterVivo Solutions Inc., Fergus, ON, Canada, ²Department of Pharmacology and Toxicology, University of Toronto, Toronto, ON, Canada, ³InterVivo Solutions Inc., Mississauga, ON, Canada, ⁴Leslie Dan Faculty of Pharmacy, University of Toronto, Toronto, ON, Canada, ⁵DL Global Partners Inc., Toronto, ON, Canada

OPEN ACCESS

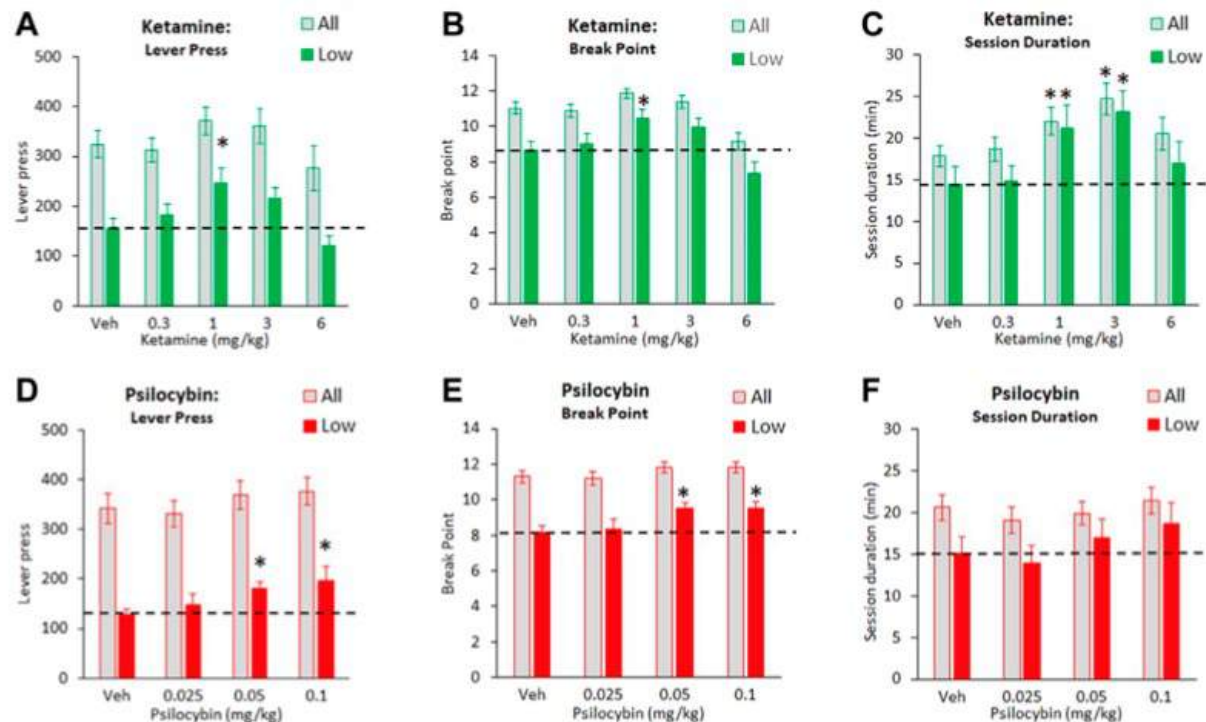
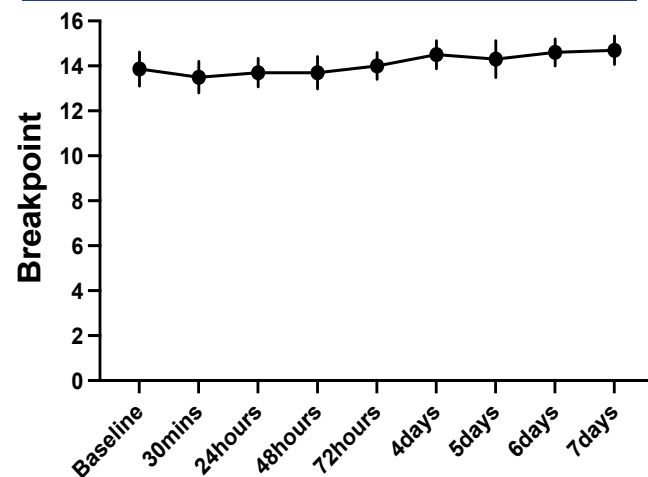


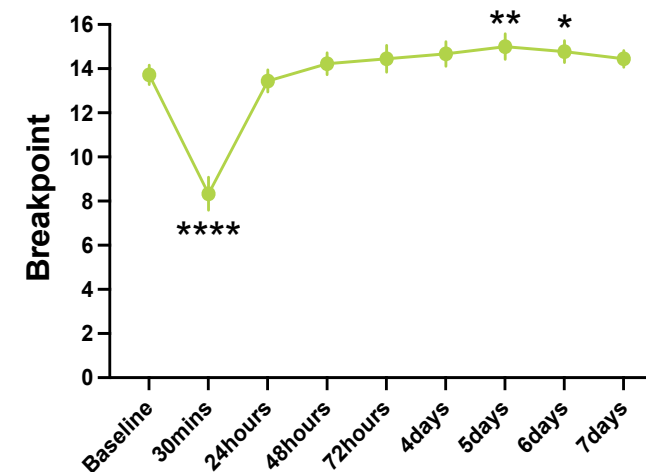
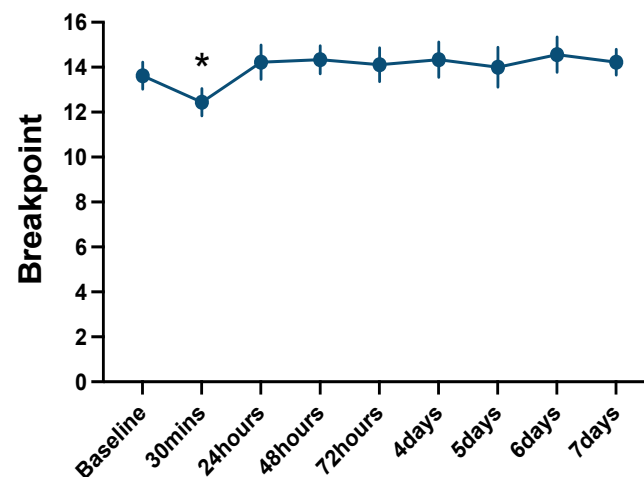
FIGURE 2 | Characterisation of ketamine (0.3–6 mg/kg IP) and psilocybin (0.025–0.1 mg/kg SC) on food responding made available under a progressive schedule of reinforcement. Data is presented for both drugs as total number of lever presses recorded during the test session (**A and D**), final break point (**B and E**), and total session duration (**C and F**). Data for each drug is presented both as all test subjects (ketamine: $N = 68$; psilocybin: $N = 72$), and subjects characterized as “low performers” based on having the lowest tertile on lever presses/break point based on performance measured over 7 days prior to onset of drug testing (ketamine: $N = 23$; psilocybin: $N = 24$). The hashed line is to highlight the level of the “low performer” subgroup following vehicle pretreatment. * $p < 0.05$ vs. vehicle control (Dunnett’s test following significant ANOVA).

DOI and motivation

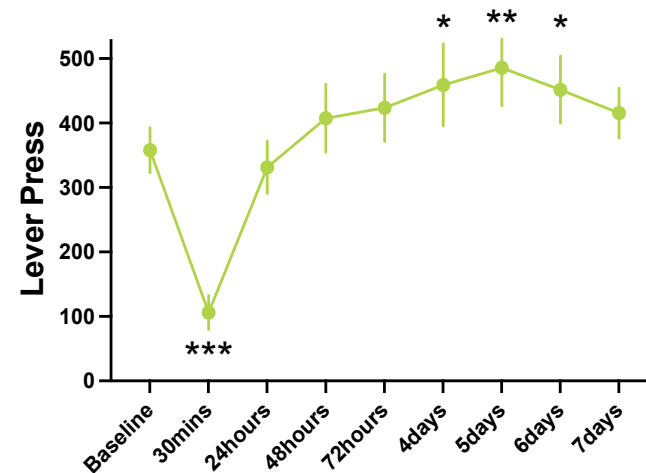
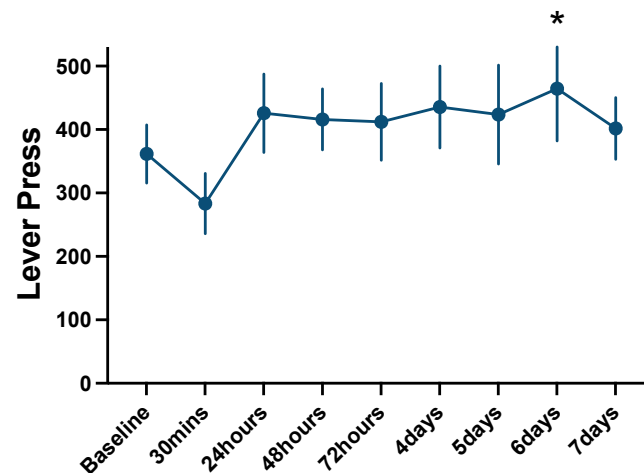
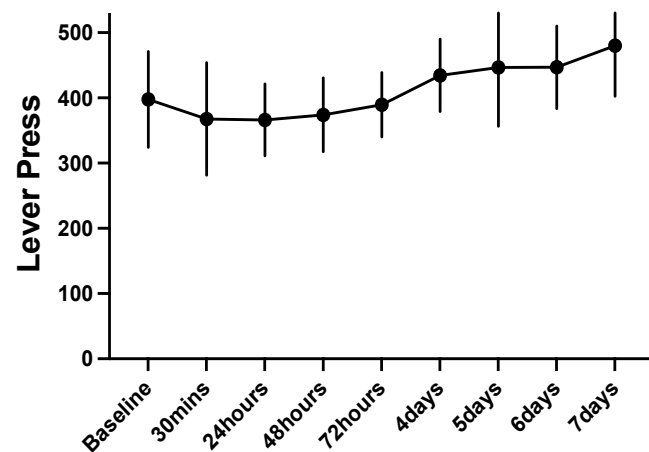
Breakpoint



● Vehicle ● DOI 0.2mg/kg SC ● DOI 1mg/kg SC



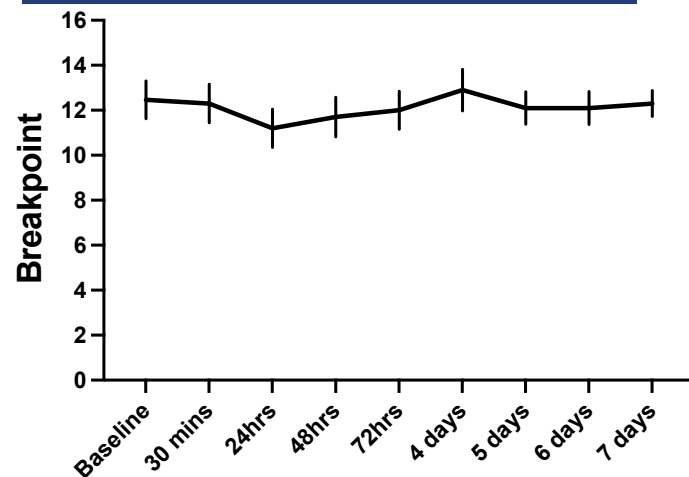
Lever Presses



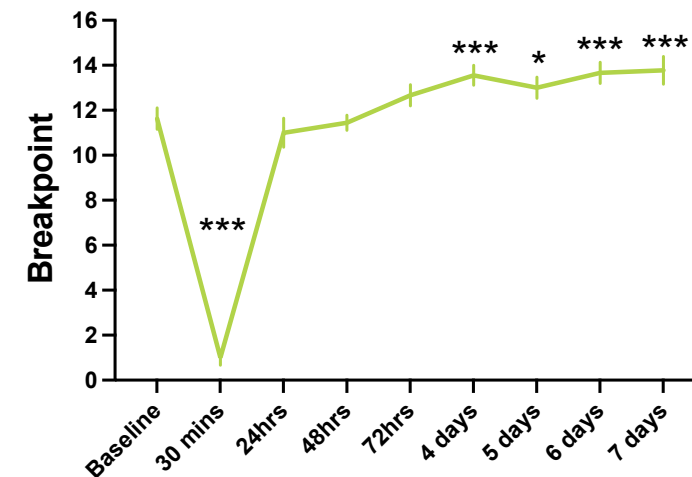
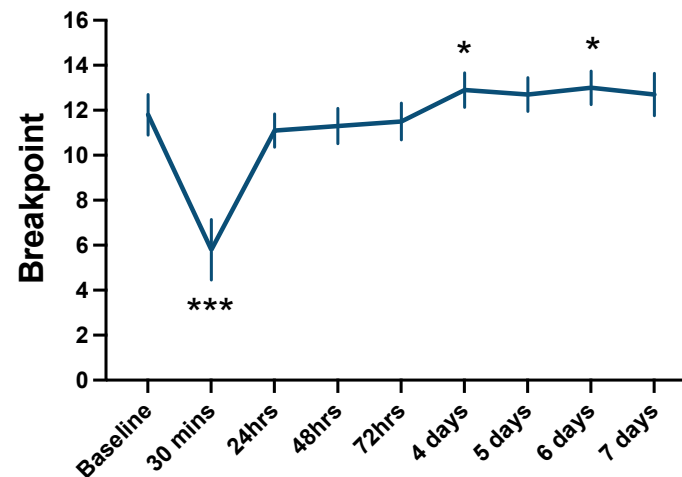
* Significant effect compared to baseline. One-way repeated measures ANOVA with Fishers LSD post-hoc (n=9-10 per group). Drugs administered by subcutaneous (SC) injection.

Psilocybin and motivation

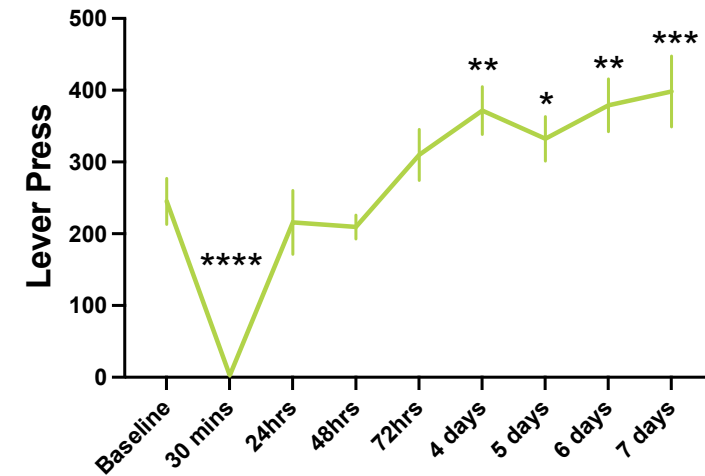
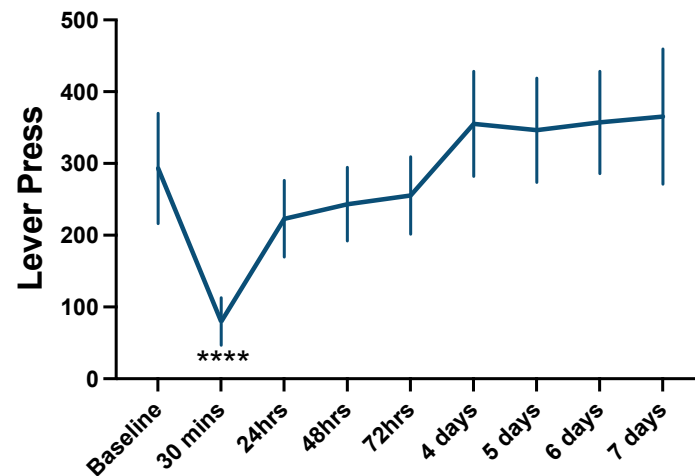
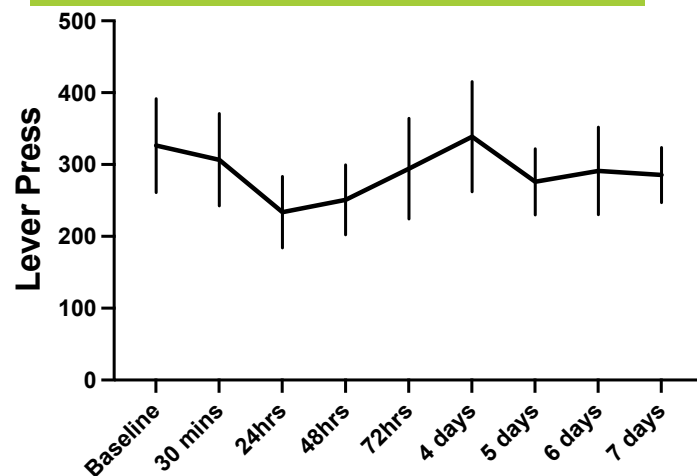
Breakpoint



— Vehicle — Psilocybin 1mg/kg SC — Psilocybin 3mg/kg SC



Lever Presses

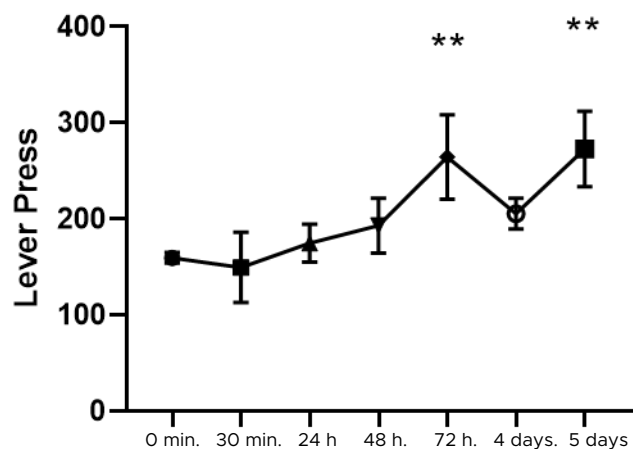


* Significant effect compared to baseline. One-way repeated measures ANOVA with Fishers LSD post-hoc (n=9-10 per group). Drugs administered by subcutaneous (SC) injection.

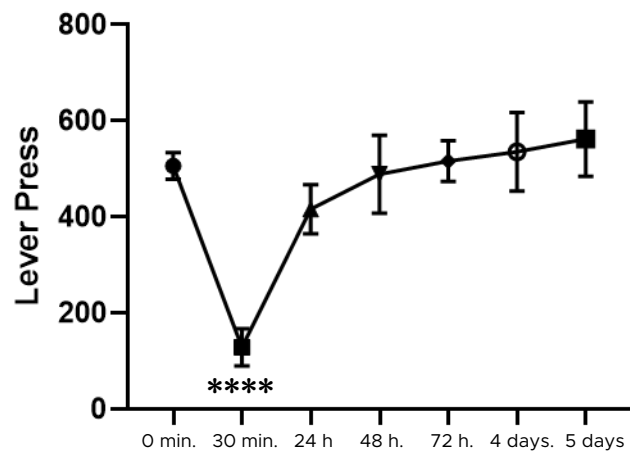
DOI (1mg/kg sc) and motivation

Lever Presses

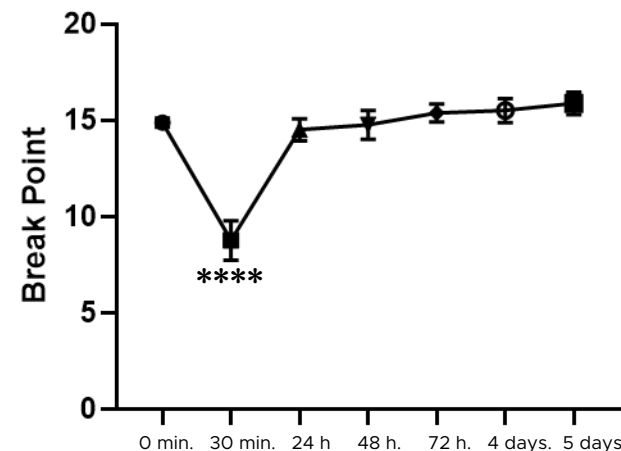
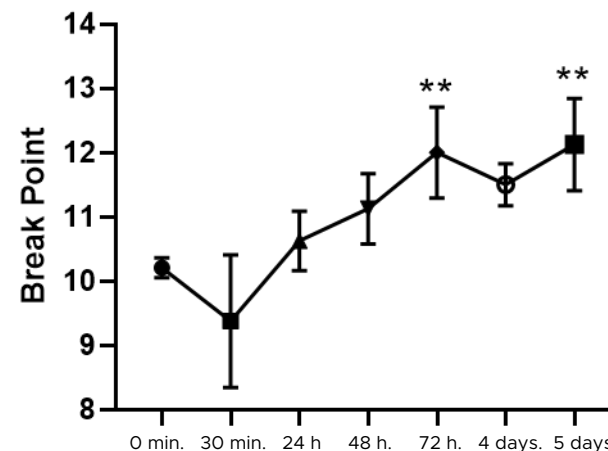
Low responders




High responders



Breakpoint



* Significant compared to Baseline group. One-way repeated measures ANOVA with Fishers LSD post-hoc (n=8 per group)



“It gave me an inner joy, an open mindedness, a gratefulness, open eyes and an internal sensitivity for the miracles of creation.”

Albert Hofmann, on LSD

Acknowledgements

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Dr. Mark Duxon

