

The cognitive effects of 5-Methoxy-*N,N*-Dimethyltryptamine (5-MeO-DMT) are associated with improvements in depression and anxiety conditions

Sara So^{1,2}, Rafael Lancelotta³, Joseph P Barsuglia⁴, Roland R Griffiths², Alan K. Davis²,

¹Johns Hopkins University, Bloomberg School of Public Health, ²Johns Hopkins University School of Medicine, Psychedelic Research Unit

³University of Wyoming, Counseling Department, ⁴New School Research



JOHNS HOPKINS
MEDICINE

Introduction

5-methoxy-*N,N*-dimethyltryptamine (5-MeO-DMT), is a short-acting (30-90 minutes) tryptamine found in the venom and skin of *Bufo alvarius* toads and can also be synthetically produced (1, 2, 3). According to a recent epidemiological survey study, 5-MeO-DMT is used infrequently, primarily for spiritual exploration, has a safe profile of use and low potential for psychiatric or biomedical consequences, and might have psychotherapeutic effects (4). More specifically, there have been reports of spontaneous and unintended symptom improvements in anxiety and depression (4).

Study Aim

The primary aim of this current analysis is to examine whether use of 5-MeO-DMT is associated with spontaneous and unintended improvements in depression and anxiety among people who have used 5-MeO-DMT in the US with procedures that guide the source, dose, and administration of 5-MeO-DMT, and the preparation of and support during/following sessions. The second aim of this study is to examine factors associated with improvement in depression and anxiety.

Method & Data Analysis

Using an email distribution list of people in the US that use 5-MeO-DMT in a specific group setting, we recruited English-speaking adults to complete an anonymous web-based survey. The primary survey used for this study included an extensive series of questions about the patterns of use, acute subjective effects, and potential consequences and benefits of using 5-MeO-DMT in this group setting. Depression and Anxiety measures, the Mystical Experiences Questionnaire, the Challenging Experiences Questionnaire, and Persisting Effects Questionnaire was included.

Sample Characteristics

Sample

362 respondents completed the survey

The sample was comprised as follows:

- Mean age = 48 (*SD*=13)
- White/Caucasian = 84%
- Heterosexual = 79%
- College Graduates = 75%
- Females = 45%

Sample after 5-MeO-DMT use

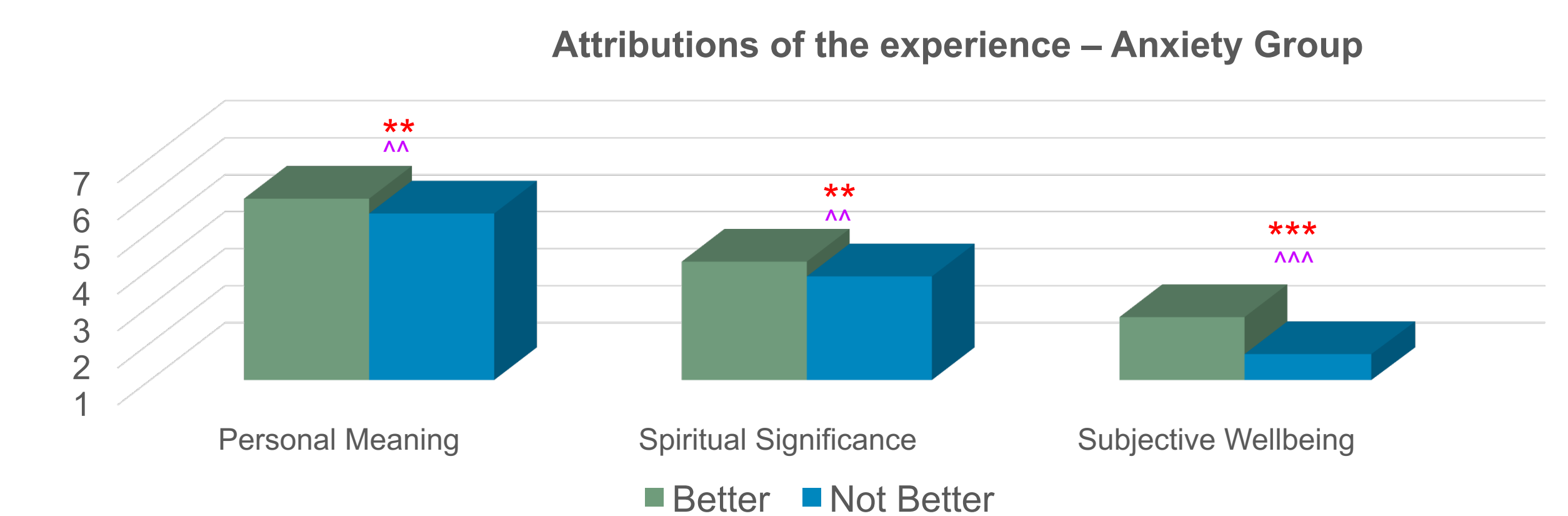
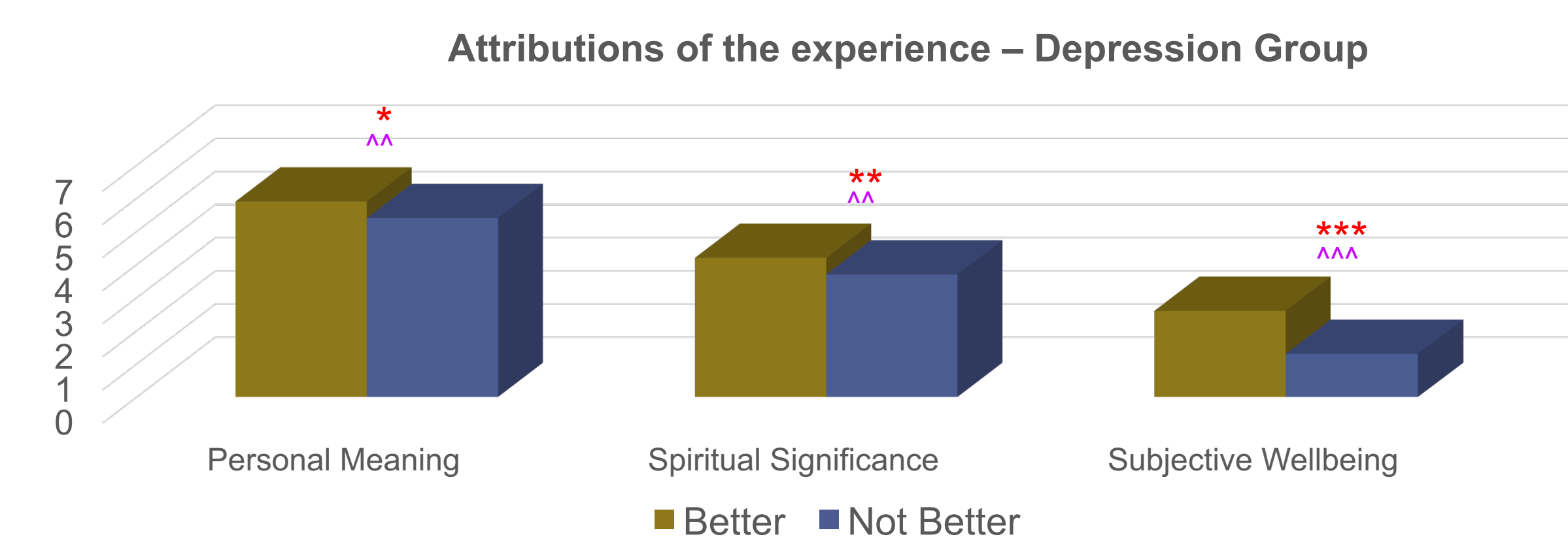
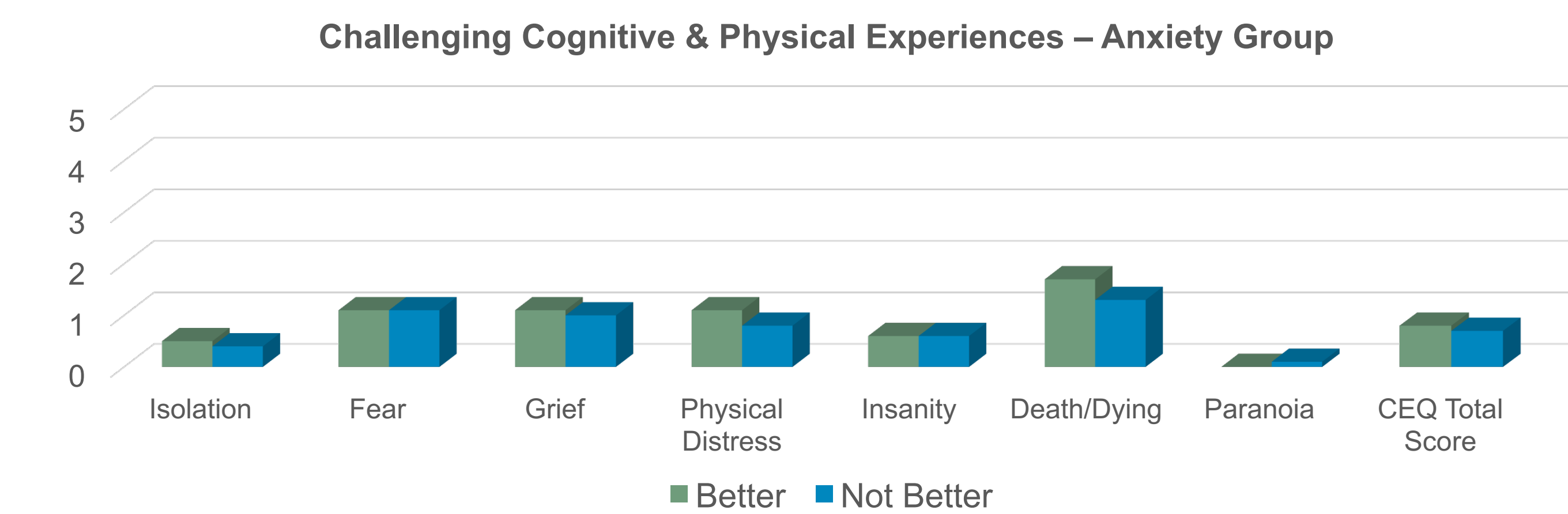
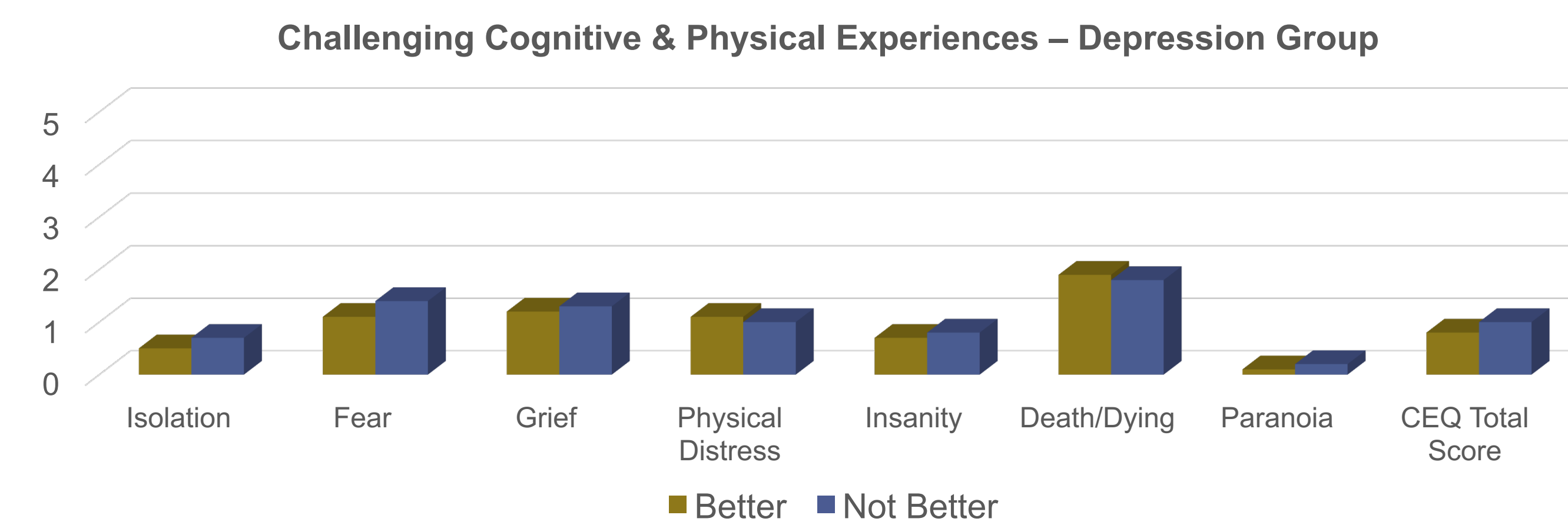
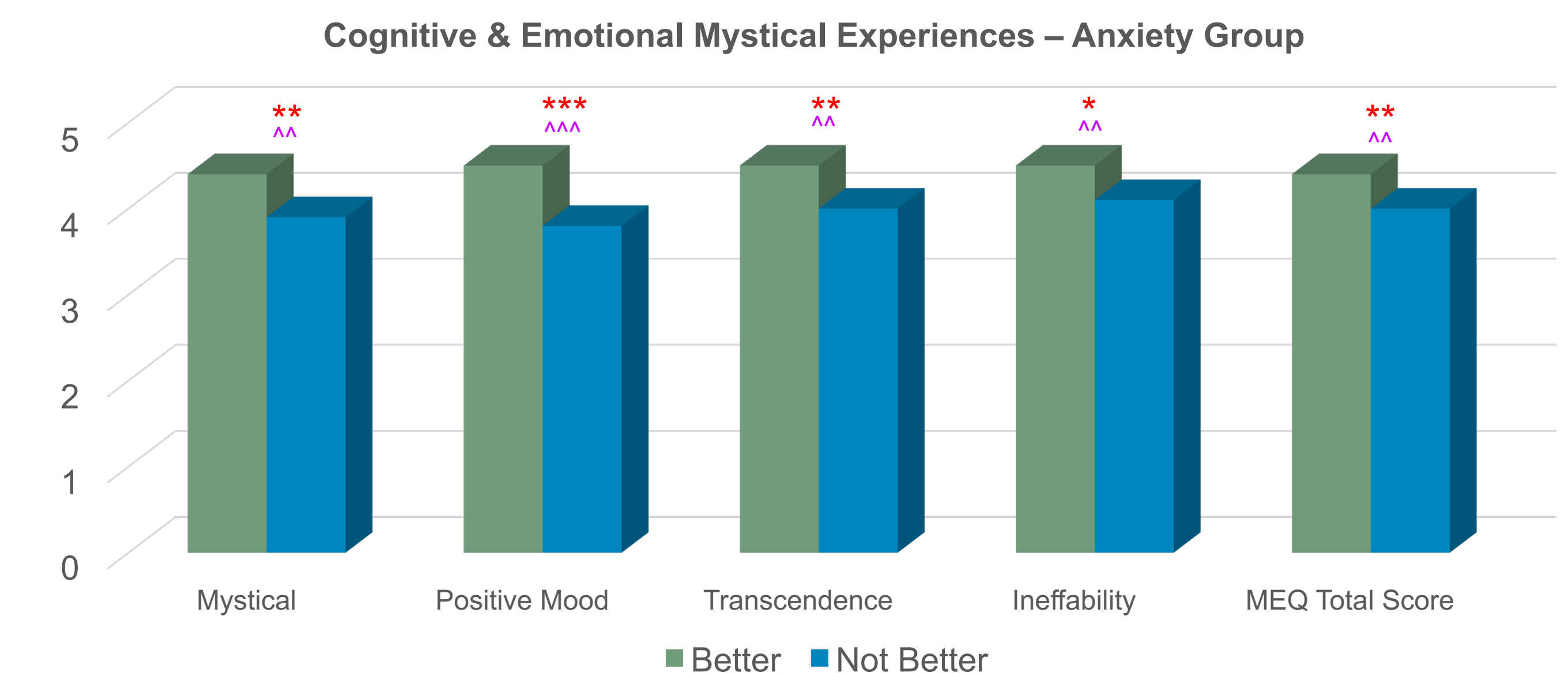
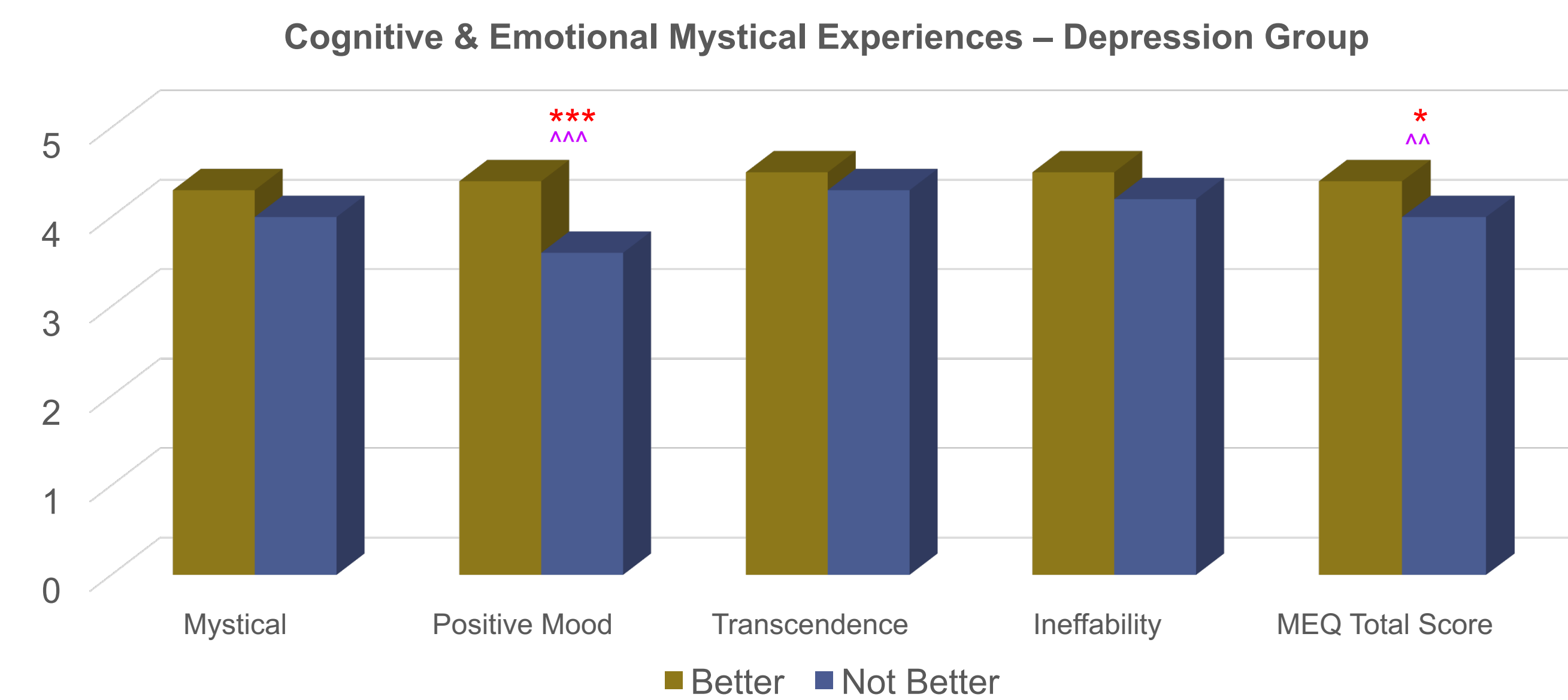
Rates of Anxiety (n=173)

- Improved: 79%
- Stayed the same/worsened: 21%

Rates of Depression (n=149)

- Improved: 81%
- Stayed the same/worsened: 19%

Results



* $p < .05$; ** $p < .01$; *** $p < .001$ ^ Small effect; ^^ Medium effect; ^^^ Large effect

Conclusions

When administered in a naturalistic group setting, 5-MeO-DMT appears to be associated with spontaneous and unintended improvements in self-reported depression and anxiety (approximately 80%), which were related to more intense acute mystical effects and increases in ratings of the personal meaning and spiritual significance of the 5-MeO-DMT session, as well as higher ratings of the degree to which the session contributed to improved well-being and life satisfaction. These results are consistent with laboratory studies that found positive psychotherapeutic effects of tryptamines as an adjunct to supportive psychotherapy (5-8) and suggests the importance of the acute mystical effects of psychedelic substances as one of the mechanisms by which they exert psychotherapeutic effects (8-11).

References

1. Araújo AM, Carvalho F, Bastos MD, Pinho PG, Carvalho M. The hallucinogenic world of tryptamines: an updated review. Arch Toxicol. 2015;89(8):1151–73.
2. Ott J. Pharmepéna-Psychonautics: Human Intranasal, Sublingual and Oral Pharmacology of 5-Methoxy-N, N-Dimethyl-Tryptamine. J Psychoactive Drugs. 2001;33(4):403–7.
3. Shulgin AT, Shulgin A. Tihkal: the continuation. Berkeley, CA: Transform Press; 1997.
4. Davis AK, Barsuglia JP, Lancelotta R, Grant R and Renn E. The epidemiology of 5-Methoxy-N,N-Dimethyltryptamine (5-MeO-DMT) use: Benefits, consequences, patterns of use, subjective effects, and reasons for consumption. J Psychopharmacol. 2016;32(7):779–92.
5. Johnson MW, Griffiths RR. Potential Therapeutic Effects of Psilocybin. Neurotherapeutics. 2017;14(3): 734–40.
6. Carhart-Harris RL, Roseman L, Bolstridge M, Demetriou L, Pannekoek JN, Wall MB, Tanner M, Kaelen M, McGonigle J, Nutt DJ. Psilocybin for treatment-resistant depression: FMRI-measured brain mechanisms. Sci Rep. 2017;7(1).
7. Ross S, Bossis T, Guss J, Agin-Lieb G, Malone T, Cohen B, Mennenga S, Belser A, Kalliontzis K, Babb J, et al. Rapid and sustained symptom reduction following psilocybin treatment for anxiety and depression in patients with life-threatening cancer: A randomized controlled trial. J Psychopharmacol. 2016;30(12):1165–80.
8. Griffiths RR, Johnson MW, Carducci MA, Umbricht A, Richards WA, Richards BD, Cosimano MP, Klinedinst MA. Psilocybin produces substantial and sustained decreases in depression and anxiety in patients with life-threatening cancer: A randomized double-blind trial. J Psychopharmacol. 2016;30(12):1161–9.
9. Griffiths RR, Richards W, Johnson M, McCann U, Jesse R. Mystical-type experiences occasioned by psilocybin mediate the attribution of personal meaning and spiritual significance 14 months later. J Psychopharmacol. 2008;22(6):621–32.
10. Griffiths RR, Johnson MW, Richards WA, Richards BD, McCann U, Jesse R. Psilocybin occasioned mystical-type experiences: Immediate and persisting dose-related effects. Psychopharmacology. 2011;218(4):649–65.
11. Garcia-Romeu A, Griffiths RR, Johnson MW. Psilocybin-occasioned mystical experiences in the treatment of tobacco addiction. Curr Drug Abuse Rev. 2014;7(3):157–64.

Funding

AKD was supported by NIDA (DA007209). RL was supported by Source Research Foundation to provide administrative support on this study. RL, JPB and AKD are on the board of directors at SRF. The funding source had no role in study design, data analysis, or interpretation.

Contact email for corresponding author: adavi157@jhmi.edu