

### Background

Despite the 2019 NIH campaign to address the opioid crisis, over 140 million opioids were prescribed and over 150,000 people died from an overdose in 2020. Since opioids are most commonly prescribed to treat severe pain, it is incredibly important to identify new, readily available interventions that may be used to target chronic pain and opioid use. In the last few years, the United States has witnessed enormous changes concerning public acceptance of cannabinoids (CBD), with survey studies indicating that many individuals are substituting CBD for opiates to alleviate pain. Results from recent RCTs and survey studies have concluded that CBD is an effective tool in the management of chronic pain, opioid craving and opioid use.

### Methods

- Individuals with chronic pain enrolled in a randomized, double blind, phase 2 clinical trial assessing the efficacy of CBD in reducing pain and/or opioid medication were recruited and randomly assigned to one of three conditions:
- **Broad-spectrum CBD** [bsCBD, n=46, Mean age (M)= 50.65, Standard Deviation (SD)=14.76, 80% Female]
- **Full-spectrum CBD** [fsCBD, n=43, *M*=42.56, *SD*=13.85, 79% Female]
- **Placebo** [PL, n=36, *M*=39.00, SD=13.91, 91% Female]
- Participants ingested 210 mg daily oral doses over a 12-week trial period. Pain was assessed at three timepoints via the self-report Pain Catastrophizing Scale.
- Blood cannabinoid levels were collected at each timepoint to assess adherence to trial medication.
- A mixed ANOVA with repeated measures was used to evaluate pain catastrophizing scores across timepoints. Our within-subject factor was time, with repeated measures (Week 0, Week 6, and Week 12). Our between-subject factors was condition, with three groups (bsCBD, fsCBD, PL).

### References

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## **Cannabidiol Reduces Experience of Pain in Individuals with Chronic Pain.**

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### Results

Although the omnibus test was not significant, pairwise comparisons revealed there was a significant simple effect of time

- PCS scores significantly decreased within the bsCBD condition from baseline to week 6 (p=.005) and from baseline to week 12
- PCS scores significantly decreased within the fsCBD condition from baseline to week 6 (p=.001) and from baseline to week 12
- PCS scores did not significantly decrease within the placebo condition from baseline to week 6. However, PCS scores did significantly decrease within the placebo condition from week 6 to week 12 (p=.006) and from baseline to week 12 (p=.001)

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condition bsCBD fsCBD Placebo