# Patient perception of brain stimulation in the treatment of severe pediatric chronic pain

Salma Farag,<sup>1</sup> Elias Abou-Assaly, BA<sup>2,4</sup> Spencer Epp, BSc,<sup>2</sup> Nivez Rasic, MD FRCPC<sup>2,4</sup> Frank P. MacMaster, PhD<sup>3,8,9</sup> Adam Kirton, MD<sup>4,8</sup> Catherine Lebel, PhD<sup>5,8</sup> Laura Rayner, RN<sup>7</sup> Melanie Noel, PhD<sup>6-8</sup>, Jillian Vinall Miller, PhD<sup>2,7,8</sup>

<sup>1</sup>Neuroscience, <sup>2</sup>Anesthesiology, Perioperative & Pain Medicine, <sup>3</sup>Psychiatry, <sup>4</sup>Pediatrics, <sup>5</sup>Radiology, <sup>6</sup>Psychology, University of Calgary, Calgary AB, Canada; <sup>7</sup>Vi Riddell Children's Pain & Rehabilitation Centre, Alberta Children's Hospital, Calgary AB, Canada; <sup>8</sup>Child Brain & Mental Health Program, Alberta Children's Hospital Research Institute, Calgary, AB, Canada, <sup>9</sup>Provincial Addiction and Mental Health Portfolio, Calgary, AB, Canada

#### Introduction

- **Severe chronic pain** is characterized by pain lasting  $\geq 3$  months, interfering with daily functioning (1)
- The Intensive Pain Rehabilitation Program (IPRP) at Alberta Children's Hospital is a 3-week, interdisciplinary day-treatment program that strives to return affected youth to normal daily activities.
- Scans of IPRP patients revealed a correlation between decreased activity in the dorsolateral prefrontal cortex (DLPFC) and improved functional outcomes, which is consistent with the DLPFC's role in pain modulation (2)

#### Methods

- Youth (n=6) aged 12-18 years (83% female) with severe chronic pain were  $\bullet$ recruited from the IPRP
- rTMS was applied every weekday for three weeks to the left DLPFC (x=-30, y=36, z=42) at 10 Hz.
- Participants were monitored for adverse events and tolerability using a Pediatric TMS Safety and Tolerability Measure on days 1, 6, and 11



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- In October 2020, open label repetitive transcranial magnetic stimulation (rTMS) was added to the IPRP to investigate whether stimulating the DLPFC could enhance patient outcomes
- Youth and their parents ranked the three most and least helpful IPRP interventions before starting the IPRP (baseline) and at follow-up (end of program and 3-month follow-up). There was a total of 13 IPRP interventions to choose from.

### **Background/Goals**

We examined youth and parent perspectives of rTMS relative to other IPRP interventions, and the tolerability, and adverse events reported.

# **Hypothesis**

We hypothesized youth with severe chronic pain would demonstrate tolerability of rTMS, and families would perceive it as a helpful intervention, when combined with standard IPRP.

## Results

• At baseline, youth predicted rTMS would be the most helpful IPRP intervention. However, from baseline to follow-up the proportion of youth ranking rTMS among the top 3 interventions decreased from 75% to 33%, making it the 4<sup>th</sup> most helpful



- among youth at follow-up.
- At baseline, only 25% of parents included brain stimulation in their top 3 interventions, while at follow-up, it was the 3<sup>rd</sup> most helpful intervention according to parents, with 50% of the ranking it among their top 3.
- Pre- and post-rTMS, 33-50% of patients reported mild to moderate headaches and neck pain, as compared to 17-33% of patients by week three.
- One person experienced severe nausea and neck pain on day one, post-rTMS.
- Tolerability was considered favorable among youth, with most comparing their experience to non-noxious activities, like watching TV.

days 1 and 11 of treatment.

#### Discussion

- Preliminary data revealed that despite having high expectations, youth viewed rTMS as less favorable, post-treatment. Meanwhile, the opposite trend was seen in parents, who had initially lower expectations, followed by favorable views of the treatment post-IPRP.
- As reported in previous studies, extreme adverse effects occur rarely, while mild to moderate adverse effects may also be experienced in some patients (3).
- Mild to moderate headaches were the most reported adverse effect, however past studies have also observed similar rates of this symptomology in patients receiving sham rTMS (4)
- Similar to previous studies, symptoms appeared to decrease between week one and week three of treatment (5)
- More research is needed to further evaluate rTMS' safety and tolerability for severe pediatric chronic pain.

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#### References

[1] Treede RD et al. (2019) Chronic pain as a symptom or a disease: the IASP Classification of Chronic Pain for the International Classification of Diseases (ICD-11). Pain (Amsterdam) 160:19–27.

[2] Seminowicz, DA, Moayedi M (2017) The Dorsolateral Prefrontal Cortex in Acute and Chronic Pain. J Pain 18:1027-1035.

[3] Burt T, Lisanby SH, Sackeim HA (2002) Neuropsychiatric applications of transcranial magnetic stimulation: a meta analysis. Int J *Neuropsychopharmacol* 5:73-103.

[4] Kaster TS et al. (2018). Efficacy, tolerability, and cognitive effects of deep transcranial magnetic stimulation for late-life depression: a prospective randomized controlled trial. *Neuropsychopharmacology*, 43:2231-2238.

[5] Humaira A, Gao S, Wu L, Downar J, Blumberger D, Vila-Rodriguez F (2019) Side effects trajectories in rTMS treatment for depression: 10 Hz vs. intermittent theta-burst stimulation. Brain Stimulation 12:478–478.