Pilot feasibility study with randomised control of dual site Theta burst Transcranial Magnetic Stimulation (TMS) for Methamphetamine use disorder: Preliminary results of the TARTAN study

<u>Buddhima Lokuge^{1,2}</u>, Tarun Yadav^{1,2,10}, Melissa A. Jackson^{1,3,9}, Amanda L. Brown^{1,2,10}, Bryan Paton³, Marcia Sequeira¹, Martin Nean¹, Llewllyn Mills^{4,5}, Emma K. Austin⁸, Paul B Fitzgerald^{6,7}, Adrian J. Dunlop^{1,2,9}

¹Drug and Alcohol Clinical Services, Hunter New England Local Health District, ²School of Medicine and Public Health, University of Newcastle., ³School of Psychological Sciences, University of Newcastle; Hunter Medical Research Institute., ⁴Discipline of Addiction Medicine, Central Clinical School, University of Sydney & South Eastern Sydney Local Health District, Drug and Alcohol Services, ⁵The Langton Centre., ⁶School of Medicine and Psychology, College of Health & Medicine, Australian National University., ⁷Monarch Mental Health Group, ⁸School of Environmental and Life Sciences, University of Newcastle, Newcastle NSW, Australia, ⁹NSW Drug & Alcohol Clinical Research & Improvement Network, St Leonards, Australia, ¹⁰Hunter Medical Research Institute, Brain Neuromodulation Research Program

Presenter's email: buddhima.lokuge@health.nsw.gov.au

Introduction: The feasibility of Transcranial Magnetic Stimulation (TMS) for moderate to severe methamphetamine (MA) use disorder in a public outpatient Drug and Alcohol (D&A) setting is examined. TMS is a non-invasive means to stimulate neurons in superficial areas of the brain. Recent years have seen a growth in the use of TMS to investigate modulation of neural networks involved in substance use disorders.

Method: This study randomised participants to active TMS and Sham (placebo) arms. Participants received Theta burst Stimulation (TBS) to the left dorsolateral prefrontal cortex and orbitofrontal cortex (12 sessions over 4 weeks). Participants were offered weekly counselling and an opt-in neuroimaging sub-study. Feasibility measures including recruitment, treatment adherence, patient experience, as well as outcomes such as reported substance use and cognitive testing were examined.

Results: 72 referrals were received during the 44 week study period. 19 progressed to consent and 14 started and completed study treatments. All completed at least the minimum 6 of 12 eligible TMS sessions. TMS was well tolerated with 14 adverse events reported, the majority being minor (e.g. headache) or unrelated to the study intervention.

Discussions: The use of TMS in MA use disorder in a public outpatient setting was feasible and acceptable for patients. Many referrals were lost before screening or had TMS exclusion criteria (e.g. history of seizures, bipolar disorder, psychosis). Those that started, completed most treatments.

Implications for Practice: We demonstrate that TMS for MA use disorder is feasible, if resource intensive, in a public outpatient D&A setting.

Disclosure of Interest Statement: PF has received equipment for research from MagVenture A/S, Nexstim, Neuronetics and Brainsway Ltd and funding for research from Neuronetics. He is a founder of Monarch Mental Health Group and Resonance Therapeutics.