

ETHICS APPROVAL & GOVERNANCE AUTHORISATION



8 November 2019

Professor Kim Cornish
School of Psychological Sciences
Monash University

Dear Professor Cornish,

Project Title: Training Attention in Children with Acquired Brain Injury: A Randomised Control Trial of the TALI Attention Training Program

HREC Reference Number: 38132

I am pleased to advise that the below amendment has received **ethical approval** from The Royal Children's Hospital Melbourne Human Research Ethics Committee (HREC).

The HREC confirms that your proposal meets the requirements of the National Statement on Ethical Conduct in Human Research (2007). This HREC is organised and operates in accordance with the National Health and Medical Research Council's (NHRMC) National Statement on Ethical Conduct in Human Research (2007), and all subsequent updates, and in accordance with the Note for Guidance on Good Clinical Practice (CPMP/ICH/135/95), the Health Privacy Principles described in the Health Records Act 2001 (Vic) and Section 95A of the Privacy Act 1988 (and subsequent Guidelines).

The amendment has also received **governance authorisation at the Melbourne Children's Campus** (incorporating The Royal Children's Hospital, Murdoch Children's Research Institute and the University Of Melbourne Department Of Paediatrics).

HREC Approval Date: 8 November 2019

Approved Documents:

The following documents have been reviewed and approved:

Document	Version	Date
Protocol	7.0	18 September 2019
Parent/Guardian ICF	7.0	24 October 2019
Tracing Letter	5.0	24 October 2019
Clinician Information Sheet	3.0	24 October 2019
Participant Flyer	2.0	24 October 2019

Conditions of Ethics Approval:

- You are required to submit to the HREC:
 - An Annual Progress Report (that covers all sites listed on approval) for the duration of the project. This report is due on the anniversary of HREC approval. Continuation of ethics approval is contingent on submission of an annual report, due within one month of the approval anniversary. Failure to comply with this requirement may result in suspension of the project by the HREC.
 - A comprehensive Final Report upon completion of the project.

- Submit to the reviewing HREC for approval any proposed amendments to the project including any proposed changes to the Protocol, Participant Information and Consent Form/s and the Investigator Brochure.
- Notify the reviewing HREC of any adverse events that have a material impact on the conduct of the research in accordance with the NHMRC Position Statement: *Safety monitoring and reporting in clinical trials involving therapeutic goods November 2016*.
- Notify the reviewing HREC of your inability to continue as Coordinating Principal Investigator.
- Notify the reviewing HREC of the failure to commence the study within 12 months of the HREC approval date or if a decision is taken to end the study at any of the sites prior to the expected date of completion.
- Notify the reviewing HREC of any matters which may impact the conduct of the project.
- If your project involves exposure of persons to ionising radiation, you must ensure that your research is carried out in accordance with the Australian Radiation Protection and Nuclear Safety Agency (ARPANSA) Code of Practice for the 'Exposure of Humans to Ionizing Radiation for Research Purposes (2005)' (Radiation Protection series Publication No.8)
- The HREC, authorising institution and/or their delegate/s may conduct an audit of the project at any time.

Yours sincerely



Karen Sosa

Research Ethics and Governance Officer (HREC)

Research Ethics and Governance

The Royal Children's Hospital Melbourne

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