

MELBOURNE HEALTH HUMAN RESEARCH ETHICS COMMITTEE

ETHICAL APPROVAL

Dr Kimberley Haines
Western Health
Level 3 CHRE,
176 Furlong Rd
St Albans, VIC 3021

30 January 2018

Dear Dr Kimberley Haines,

HREC Reference Number: HREC/17/MH/432

Melbourne Health Site Reference Number: 2017.371

Project Title: icuRESOLVE (Intensive Care Unit REcovery Solutions cO-Led through surVivor Engagement) Co-designing Peer Support With ICU Survivors: Phase 2

I am pleased to advise that the above project has **received ethical approval** from the Melbourne Health 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).

HREC Approval Date: 30 January 2018

Ethical approval for this project applies at the following sites:

Site
Western Health

Approved Documents:

The following documents have been reviewed and approved:

Document	Version	Date
Protocol	2	08 January 2018
Participant Information Sheet/Consent Form	2	19 December 2017
Outcomes Measure	1	21 November 2017
Consent Script	1	21 November 2017
Case Report Form - Patient	1	21 November 2017
Case Report Form – Carer or Relative	1	21 November 2017
Screening Log	1	21 November 2017
Attendance Adherence Log	1	21 November 2017

Letter of Invitation for Peer Support Session	1	21 November 2017
Reminder Letter for Subsequent Peer Support Session	1	21 November 2017
Letter of Invitation to Participate	1	21 November 2017

Governance Authorisation:

Governance Authorisation is required at each site participating in the study before the research project can commence at that site.

You are required to provide a copy of this HREC approval letter to the principal investigator for each site covered by this ethics approval for inclusion in the site specific assessment application.

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's Safety Monitoring and Reporting in Clinical Trials Involving Therapeutic Goods (2016) guideline.
- 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 radiation, you are legally obliged to conduct your research in accordance with the Australian Radiation Protection and Nuclear Safety Agency Code of Practice 'Exposure of Humans to Ionizing Radiation for Research Purposes' Radiation Protection series Publication No.8 (May 2005)(ARPANSA Code).

Please note: Template forms for reporting Amendments, safety reporting, Annual/Final reports, etc. can be accessed from: <https://www2.health.vic.gov.au/about/clinical-trials-and-research/clinical-trial-research/monitoring-reporting>

The HREC may conduct an audit of the project at any time.

Yours sincerely,



Prof Peter Colman
Chair – Melbourne Health Human Research Ethics Committee (HREC)