



Our Reference: 66869_Approved

8th December 2020

Dr Alex Tan
Radiation Oncology
Townsville Hospital and Health Service
Alex.Tan@health.qld.gov.au

Dear Dr Tan,

HREC Reference number: HREC/QTHS/66869
Project title: Feasibility of Magnetic Resonance Imaging guided stereotactic reirradiation of locally recurrent prostate tumours

Thank you for re-submitting the above project for ethical and scientific review. The response was considered by Townsville Hospital and Health Service Human Research Ethics Committee (HREC) Chairperson on 02/12/2020.

The Townsville Hospital and Health Service HREC is constituted according to the National Health and Medical Research Council's '*National Statement on Ethical Conduct in Human Research*' (NHMRC, 2007, updated 2018). The Townsville Hospital and Health Service HREC operates in accordance with the '*Ethical conduct in research with Aboriginal and Torres Strait Islander Peoples and communities: Guidelines for researchers and stakeholders*' (NHMRC, 2018); and the '*National Statement on Ethical Conduct in Human Research*' (NHMRC, 2007, updated 2018). Attached is the HREC composition with membership category and affiliation with the Hospital (Attachment I).

The HREC has granted approval of this research project. The research proposal meets the requirements of the '*National Statement on Ethical Conduct in Human Research*' (NHMRC, 2007, updated 2018).

Documents reviewed and approved:	Version	Date
Response		27.11.2020
Participant Information Sheet	1.1	01.12.2020
Participant Consent Form	1.1	01.12.2020
Study Protocol	1.1	01.12.2020
Documents noted:	Version	Date
Human Research Ethics Application Form	Ver 2	27.11.2020
Radiation Safety Report		29.09.2020
EQ-5D-5L Questionnaire	Ver 1.2	2009
EORTC QLQ - PR25		1999
EORTC QLQ-C30	Ver 3	1995
Curriculum Vitae – A.Tan		
Curriculum Vitae – A-M.Nguyen		26.10.2020
Curriculum Vitae – B.Shirley		23.10.2020
Curriculum Vitae – C.Fisher		
Curriculum Vitae – C.Rumley		23.10.2020
Curriculum Vitae – M.Y.Tan		23.10.2020

The research project has ethical approval for the following sites:
Townsville Hospital and Health Service

Please note the following key dates for this study:

HREC approval expiry: 04/12/2025
Annual report due: 30/04/2021

You are reminded that this letter constitutes ethical approval only. You must not commence this research project at a site until separate authorisation from the CEO or Delegate of that site has been obtained.

If conducting this study in a Health Service, a copy of this approval must be submitted to the Health Service Research Governance Officer/Delegated Personnel with a completed Site Specific Assessment (SSA) Form and supporting study documents for authorisation from the CEO or Delegate to conduct this research at the approved sites. Refer to the local THHS website for further information on Site Specific Assessment:

<https://www.health.qld.gov.au/townsville/tresa/index>

Please note the following conditions of approval:

1. The Principal Investigator or study sponsor will report anything to the Committee which might warrant review of ethical approval of the project, including:
 - a) Within 72 hours of becoming aware of the event:
 - i. All significant safety issues and urgent safety measures taken as a response to the significant safety issue.
 - b) Within 15 calendar days of a sponsor's decision:
 - i. Notification of temporary halt of a study for safety reasons,
 - ii. Early termination of a study for safety reasons.
 - c) Within 15 calendar days of becoming aware of the event or report:
 - i. Any unforeseen events that might affect the ethical acceptability of the project,
 - ii. Any protocol violations and deviations from the study protocol that implicate participant consent, participant safety or data integrity,
 - iii. If the project is discontinued at a site before the expected date of completion,
 - iv. Where applicable, all industry safety monitoring and or Data and Safety Monitoring Board (DSMB) reports.

Do not submit individual line listings or individual adverse event reports to the HREC.
2. The Principal Investigator will provide an **annual progress report** and a final report at the completion of the study in the specified format to the HREC. The final report should include a copy of the results and/or publication, if not available at the time of reporting these must be provided in a timely manner. For clinical trials the annual report must include a safety report including a clear summary of the evolving safety profile of the trial.
3. Proposed amendments to the research protocol or conduct of the research which may affect the ongoing ethical acceptability of the project must be submitted first to the HREC for review, then to the relevant Research Governance Offices (RGO).

Proposed amendments to the research protocol or conduct of the research which only affects the ongoing site acceptability of the project are to be submitted to the relevant RGOs only. Further advice on submitting amendments is available from

<https://www.health.qld.gov.au/townsville/tresa/index/human-research-ethics-committee-hrec/amendments>

4. The HREC may undertake active monitoring of this research at any time. This may include random inspections of research sites, data, or consent documentation; and or interviews with research participants or other forms of feedback from them.

The HREC Terms of Reference, Standard Operating Procedures, membership and standard forms are available from <https://www.health.qld.gov.au/townsville/tresa/index/human-research-ethics-committee-hrec>. The HREC wishes you every success in your research.

Kind regards,



Dr Hudson Birden
Chairperson
Human Research Ethics Committee
Townsville Hospital and Health Service