Enquiries to: Deborah Wainwright Telephone: (61 7) 4616 6696

Our Ref: HREA/2020/QTDD/62491

Dr Catriona Duncan Urology Registrar Toowoomba Hospital Government

Allied Health Services

Darling Downs Hospital and Health Service

Dear Dr Duncan

HREC Reference number: HREA/2020/QTDD/62491 Project title: MMaLeS: Markers in Male Lichen Sclerosis.

Thank you for submitting the above project for ethical and scientific review. This project was considered by the Darling Downs Health Human Research Ethics Committee (HREC).

This HREC is constituted and operates in accordance with the National Health and Medical Research Council's (NHMRC) National Statement on Ethical Conduct in Human Research 2007 (updated 2018), NHMRC and Universities Australia Australian Code for the Responsible Conduct of Research (2018) and the CPMP/ICH Note for Guidance on Good Clinical Practice.

I am pleased to advise that the Human Research Ethics Committee has granted approval of this research project at the following sites:

- Toowoomba Hospital
- St Vincent's Private Hospital Toowoomba

Please note: This letter constitutes ethical approval only. A copy of this approval must be submitted to the HHS Research Governance Officer (RGO) along with a completed Site Specific Assessment (SSA) Application and applicable documents for authorisation from the CE or delegate to conduct this research within the HHS.

The documents reviewed and approved include:

Document	Version	Date
HREA application		8 May 2020
Protocol	V 3	4 May 2020
Ethically Defensible Plan		
Participant Information and Consent Form	V 3	22 May 2020
Participant Screening Questionnaire		
CV – D Desai		
CV – C Duncan		
CV – E Whiteside		
CV – S Ellem		
CV – L Windus		

Office of the Executive Director Allied Health Services Cnr Hogg & Tor Streets PO Box 405 Toowoomba Queensland 4350 Australia Telephone +61 7 4699 8060 Facsimile +61 7 4699 8211 www.health.qld.gov.au/darlingdowns Please note the following conditions of approval:

- 1. The Principal Investigator will immediately report anything which might warrant review of ethical approval of the project in the specified format, including:
 - a. Unforeseen events that might affect continued ethical acceptability of the project.
 - b. Serious Adverse Events must be notified to the Committee as soon as possible. In addition, the Investigator must provide a summary of the adverse events, in the specified format, including a comment as to suspected causality and whether changes are required to the Patient Information and Consent Form. In the case of Serious Adverse Events occurring at the local site, a full report is required from the Principal Investigator, including duration of treatment and outcome of event.
- 2. Amendments to the research project which may affect the ongoing ethical acceptability of a project must be submitted via Ethical Review Manager to the HREC for review. Amendments to the research project which only affect the ongoing site acceptability of the project are not required to be submitted to the HREC for review. These amendment requests should be submitted directly to the RGO via Ethical Review Manager (by-passing the HREC).
- 3. Proposed amendments to the research project which may affect both the ethical acceptability and site suitability of the project must be submitted firstly to HREC for review and, once HREC approval has been granted, then submitted to the RGO.
- 4. Amendments which do not affect either the ethical acceptability or site acceptability of the project (e.g. typographical errors) should be submitted via Ethical Review Manager to the HREC coordinator. These should include a cover letter from the principal investigator providing a brief description of the changes and the rationale for the changes and accompanied by all relevant updated documents with tracked changes.
- 5. The HREC will be notified, giving reasons, if the project is discontinued at a site before the expected date of completion.
- 6. The Principal Investigator will provide an annual report to the HREC and at completion of the study in the specified format.
- 7. The District administration and the HREC may inquire into the conduct of any research or purported research, whether approved or not and regardless of the source of funding, being conducted on hospital premises or claiming any association with the Hospital; or which the Committee has approved if conducted outside Darling Downs Health.

HREC approval is valid for 3 years from the date of this letter.

Should you have any queries about the HREC's consideration of your project please contact the Chair of the Darling Downs Health Human Resource Ethics Committee. The HREC terms of Reference, Standard Operating Procedures, membership and standard forms are available from http://www.health.gld.gov.au/ohmr/html/regu/regu home.asp

The HREC wishes you every success in your research.

Yours sincerely

Angela O'Shea

Chair

Human Research Ethics Committee

Darling Downs Health

B.M.O'

25 / 05 / 2020

Page 2 of 2