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09 March 2018

Dr Dominic Lomiwes The New Zealand Institute of Plant and Food Research Ltd. Private Bag 11600 Palmerston North 4442

Dear Dr Lomiwes

Re:	Ethics ref:	17/STH/250		
	Study title:	The effect of long term storage on the acute bioefficacy of blackcurrant juice in reducing exercise-induced oxidative stress following sustained medium-intensity cycle bout		

I am pleased to advise that this application has been <u>approved</u> by the Southern Health and Disability Ethics Committee. This decision was made through the HDEC-Expedited Review pathway.

Conditions of HDEC approval

HDEC approval for this study is subject to the following conditions being met prior to the commencement of the study in New Zealand. It is your responsibility, and that of the study's sponsor, to ensure that these conditions are met. No further review by the Southern Health and Disability Ethics Committee is required.

Standard conditions:

- 1. Before the study commences at *any* locality in New Zealand, all relevant regulatory approvals must be obtained.
- Before the study commences at *any* locality in New Zealand, it must be registered in a clinical trials registry. This should be a WHO-approved (such as the Australia New Zealand Clinical Trials Registry, <u>www.anzctr.org.au</u>). However <u>https://clinicaltrials.gov/</u> is acceptable provided registration occurs prior to the study commencing at *any* locality in New Zealand.
- 3. Before the study commences at *a given* locality in New Zealand, it must be authorised by that locality in Online Forms. Locality authorisation confirms that the locality is suitable for the safe and effective conduct of the study, and that local research governance issues have been addressed.

After HDEC review

Please refer to the *Standard Operating Procedures for Health and Disability Ethics Committees* (available on www.ethics.health.govt.nz) for HDEC requirements relating to amendments and other post-approval processes.

Your next progress report is due by 09 March 2019.

Participant access to ACC

The Southern Health and Disability Ethics Committee is satisfied that your study is not a clinical trial that is to be conducted principally for the benefit of the manufacturer or distributor of the medicine or item being trialled. Participants injured as a result of treatment received as part of your study may therefore be eligible for publicly-funded compensation through the Accident Compensation Corporation (ACC).

Please don't hesitate to contact the HDEC secretariat for further information. We wish you all the best for your study.

Yours sincerely,

Ms Raewyn Idoine Chairperson Southern Health and Disability Ethics Committee

Encl: appendix A: documents submitted appendix B: statement of compliance and list of members

Document	Version	Date
Covering Letter: Cover letter	1	13 December 2017
CV for CI: MBIE formatted CV for HDEC application	1	19 December 2017
Evidence of scientific review: Scientific peer review - P Gopal	1	06 December 2017
Investigator's Brochure: Study 1 Poster	1	13 December 2017
Investigator's Brochure: Study 2 Poster	1	13 December 2017
List of foods to avoid during trial days	1	19 October 2017
PIS/CF: Study 1 Consent Form	3	16 February 2018
PIS/CF: Study 2 Consent form	3	20 February 2018
PIS/CF: Study 1 Information sheet	4	16 February 2018
PIS/CF: Study 2 Information sheet	4	16 February 2018
Protocol: Study protocol	4	13 December 2017
Survey/questionnaire: Rating of perceived exertion scale	1	19 December 2017
Survey/questionnaire: Study 1 Health Screening Form	1	19 October 2017
Survey/questionnaire: Study 2 Health screening form	1	19 October 2017
Survey/questionnaire: Study 2 Baecke Questionnaire	2	13 December 2017
Application	1	-
Covering Letter: Cover letter - response to committee feedback	2	27 February 2018
Response to Request for Further Information	1	-

Appendix B Statement of compliance and list of members

Statement of compliance

The Southern Health and Disability Ethics Committee:

- is constituted in accordance with its Terms of Reference
- operates in accordance with the Standard Operating Procedures for Health and Disability Ethics Committees, and with the principles of international good clinical practice (GCP)
- is approved by the Health Research Council of New Zealand's Ethics Committee for the purposes of section 25(1)(c) of the Health Research Council Act 1990
- is registered (number 00008713) with the US Department of Health and Human Services' Office for Human Research Protection (OHRP).

List of members

Name	Category	Appointed	Term Expires
Ms Raewyn Idoine	Lay (consumer/community perspectives)	27/10/2015	27/10/2018
Dr Sarah Gunningham	Non-lay (intervention studies)	27/10/2015	27/10/2018
Assc Prof Mira Harrison-Woolrych	Non-lay (intervention studies)	27/10/2015	27/10/2018
Dr Fiona McCrimmon	Lay (the law)	27/10/2015	27/10/2018
Dr Anna Paris	Lay (other)	24/08/2017	24/08/2020
Dr Nicola Swain	Non-lay (observational studies)	27/10/2015	27/10/2018
Dr Devonie Waaka	Non-lay (intervention studies)	13/05/2016	13/05/2019

Unless members resign, vacate or are removed from their office, every member of HDEC shall continue in office until their successor comes into office (HDEC Terms of Reference)

http://www.ethics.health.govt.nz