



**NEPEAN BLUE MOUNTAINS LOCAL HEALTH DISTRICT HUMAN RESEARCH ETHICS COMMITTEE
CERTIFICATE OF APPROVAL**

Tuesday 17th April 2018

A/Professor Virginia Skinner
Centre for Nursing and Practice Development
Nepean Hospital

Dear A/Professor Skinner,

HREC study reference: Study - HREC/18/Nepean/30
Study title: Does using a peanut ball during labour with an epidural affect birth outcomes? A pilot Study.

Principal Investigator: A/Professor Virginia Skinner.

Your request to undertake the above protocol was considered by the NBMLHD Human Research Ethics Committee (HREC) at its meeting held on the 27th February 2018.

*On receipt of your responses dated 9/3/2018, 20/3/2018 and 17/4/2018 to the concerns of the Committee dated 7/3/2018, 20/3/2018 and 17/4/2018, we are satisfied that your protocol meets the requirements of the National Statement on Ethical Conduct in Human Research (2007) and was **APPROVED** on 17th April 2018 to be conducted as a Multisite study.*

It is the Principal Investigator's responsibility to ensure that all researchers associated with this project are aware of the conditions of approval and which documents have been approved.

The Principle Researcher is required to note the following conditions of approval:

- The coordinating investigator will immediately report anything which might warrant review of ethical approval of the project in the specified format, including unforeseen events that might affect continued ethical acceptability of the project.
- The coordinating investigator will immediately report any protocol deviation / violation, together with details of the procedure put in place to ensure the deviation / violation does not recur.
- The Coordinating Investigator will provide to the HREC in the specified format proposed amendments to the protocol or conduct of the research which may affect the ethical acceptability of the project. Copies of all proposed changes when approved by the HREC must also be provided to the research governance officer.
- The HREC must be notified, giving reasons, if the project is discontinued at a site before the expected date of completion.

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HREC/18/NEPEAN/30 – Cont'd

- The Coordinating Chief Investigator must provide an annual report to the HREC and a final report at completion of the study, in the specified format. HREC approval is granted for a period of 5 years and ongoing approval is contingent upon annual report submission. Annual Reports for all studies should be submitted on the anniversary of the approval date of project. They will be processed and presented to the HREC at the next scheduled meeting. A copy of the Annual / Final Research Report Form can be obtained electronically from the Research Office on request.
- It should be noted that compliance with the ethical guidelines is entirely the responsibility of the investigators.
- The HREC has the discretion to adopt other appropriate mechanism for monitoring depending on the complexity, design and risk perceived, including:
 - Discussion of relevant aspects of the project with investigators at any time.
 - Random inspection of research sites, data or consent
 - Interview with research participants or other formats of feedback from them.
 - Request and review reports from independent agencies such as Data and Safety Monitoring Board.
- For clinical trials using implantable medical devices, the coordinating investigator will confirm to the HREC that a process has been established for tracking participants with consent for the lifetime of the device and will immediately report any device incidents to the Therapeutic Goods Administration (TGA).
- If your research project is an interventional trial please ensure you register your trial onto one of the clinical trial registries, for example. <http://www.anzctr.org.au/>

The NBMLHD HREC has been accredited by the NSW Ministry of Health as a lead HREC to provide the single ethical and scientific review of proposals to conduct research within the NSW public health system. This lead HREC is constituted and operates in accordance with the National Health and Medical Research Council's *National Statement on Ethical Conduct in Human Research* and the *CPMP/ICH Note for Guidance on Good Clinical Practice*

SPECIAL CONDITIONS:

- **If you are using a flyer please ensure to obtain approval from the Facility Manager, Peter Hinrichsen- Corporate Services. Please email Lana Hul- Lana.Hul@health.nsw.gov.au to obtain this approval prior to placing flyers in NBMLHD for study recruitment.**

Approved Documents:

Documents reviewed and approved at the meeting were:

Document	Version	Date
NEAF Submission code AU/1/C4E339	-	-
Protocol	18	17/4/2018
Participant Information Sheet and Consent Form - BMDAMH	6	20/3/2018

Participant Information Sheet and Consent Form – Lithgow Hospital	6	20/3/2018
Peanut Ball Education Package	2	20/3/2018
Flyer	6	17/4/2018
Peanut Ball Positioning Chart	-	-
EQ-5D-5L Health Questionnaire	2	2009
Data Collection Sheet	1	16/2/2018
Online Survey for Women who use Peanut Ball	2	8/3/2018
Master Study Participant List	1	16/2/2018

Approved Sites:

Approval is given for this research project to be conducted at the following sites and campuses:

Insert Site / Sites

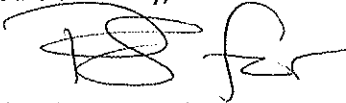
- **Blue Mountains District Anzac Memorial Hospital**
- **Lithgow Hospital**

RESEARCH GOVERNANCE - Site-Specific Assessment (SSA):

SSA authorisation is required at all sites participating in the study. SSA must be authorised at a site before the research project can commence via the Chief Executive or their delegate – the Research Governance Officer.

The completed Site-Specific Assessment Form as well as a copy of this ethics approval letter and all approved documents must be submitted to the Research Governance Officer for authorisation by the Chief Executive or delegate. This applies to each site participating in the research.

Yours Sincerely,



A/Prof Ian Seppelt
Chair, Human Research Ethics Committee
Nepean Blue Mountains LHD

Cc: Gina Oliver Research Governance

Please quote project number and title in all correspondence

