



Government of South Australia

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Research Ethics Committee

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21 December 2006

**Dr Marianne Chapman
Intensive Care Unit
ROYAL ADELAIDE HOSPITAL**

Dear Dr Chapman,

**Re: "Glucose feedback inhibition of gastro-duodenal motility in critical illness."
Volunteer Information Sheet & Consent Form, Version 1. Relative Information Sheet &
Consent Form, Version 1.
RAH PROTOCOL NO: 061208.**

I am pleased to advise that Research Ethics Committee approval has been given to the above project. Please quote the RAH Protocol Number allocated to your study on all future correspondence.

Research Ethics Committee deliberations are guided by the NHMRC National Statement on Ethical Conduct in Research Involving Humans.

The general conditions of approval follow:

- Adequate record-keeping is important. If the project involves signed consent, you should retain the completed consent forms which relate to this project and a list of all those participating in the project, to enable contact with them in the future if necessary. The duration of record retention for all research data is 15 years.
- You must notify the Research Ethics Committee of any events which might warrant review of the approval or which warrant new information being presented to research participants, including:
 - (a) serious or unexpected adverse events which warrant protocol change or notification to research participants,
 - (b) changes to the protocol,
 - (c) premature termination of the study,
 - (d) completion of the study with a study completion summary.
- The Committee must be notified within 72 hours of any serious adverse event occurring at this site.
- Approval is ongoing, subject to satisfactory annual review.

Yours sincerely,

**Dr M James
CHAIRMAN
RESEARCH ETHICS COMMITTEE**