



Kwantlen
UNIVERSITY COLLEGE

RESEARCH ETHICS BOARD

APPLICATION FOR ETHICS REVIEW

Cover Page

Application # _____
(For REB use only)

(Please submit your application via e-mail to research@kwantlen.ca)

Title of project

Principal Investigator

Position

Department

Contact telephone numbers

Email

Other contact methods (optional)

Fax

Pager

Preferred address to receive correspondence and/or approval if other than email

Co-investigator(s) (Name, Position, Department)

Additional assistants and their roles

4 key words

Proposed Start Date: _____ Proposed End Date: _____
(Expect a minimum of 6 – 8 weeks for a response from the Research Ethics Board.)

Signature:

Principal Investigator: _____

Date: _____

Full title of research

1. Describe the purpose of the research.
2. Describe the research question or hypothesis to be tested if known.
3. Describe the methodology of the research study/project.
4. Describe the method(s) of recruiting participants.
5. Describe the participant groups in this study.
6. Will the study involve any potential risks to the participants? If so, please describe the risks.
7. Describe your informed consent procedures where applicable. Where it is not applicable, explain why it is not, e.g. where one is studying the public activities of politicians who have agreed to publicity, consent is already given. (Be sure to consider all of the elements of the Requirement for Free and Informed Consent in the guidelines.)
8. Where deception is used, please include your rationale and debriefing procedures.
9. What provisions are made for informing participants, for follow-up with participants?
10. How do you plan to handle the requirement of confidentiality and/or anonymity where applicable?
11. Describe any potential conflict of interest of anyone involved in the research.
12. Describe any provisions for compensation of participants if applicable.

This form must be resubmitted after approval if there are major changes to your study. See part C of the Kwantlen University College policy on post approval monitoring. Major changes include changes in protocol, consent, risks, participant groups, recruitment, compensation, deception, confidentiality, anonymity, researchers involved, or other ethically sensitive matters. Please highlight changes on the resubmitted form.

Date: _____

Signature: _____