Informed Consent for Human Participants
CHECKLIST FOR RESEARCHERS

This checklist is intended to assist researchers with information that may be required in developing written Informed Consent documentation. It must accompany the Human Participants Research form.

☐ Yes ☐ No Have you provided contact information for yourself as the researcher (your name, telephone number, email address, status — BES/MES/PhD student)

☐ Yes ☐ No Have you included a brief description of the purpose/rationale of the study? Explain to participants why the research is being conducted and what the researcher is looking to achieve.

☐ Yes ☐ No Have you included a brief description of the study design? What are participants expected to do and what is the expected time commitment.

☐ Yes ☐ No Have you included a brief description of the potential risks and discomforts to the participants and applicable mitigation methods?

☐ Yes ☐ No ☐ N/A Have you included any benefits of the research and benefits to participants, if there are any? This may contribute to why people may choose to participate in your research.

☐ Yes ☐ No Have you indicated whether and what incentives, if any, are offered to participants and why?

☐ Yes ☐ No Have you included statements of the following (as applicable):
  i) Participation in the study is completely voluntary and participants have the right to withdraw at any time.
  ii) Should a participant withdraw from the study, all data generated by their participation shall be destroyed.
  iii) Participants have the right to answer questions.
  iv) Indicating how the research will be presented and reported? (e.g., “This research is part of my MES Major Paper”. Please also indicate if your research may be used beyond your current academic work.

☐ Yes ☐ No Have you described the methods by which confidentiality and anonymity will be attained and maintained?

Indicate if the interviewing or recording of the participant will be associated with identifying information.

☐ Yes ☐ No Have you described the storage method, length of retention and disposal method of all data gathered during the study? Researchers must keep consent forms for 2 years following the completion of the study.

☐ Yes ☐ No Have you included a statement indicating that the research has been reviewed and approve by the FES Human Participants Research Committee on behalf of York University?

☐ Yes ☐ No Have you provided contact information for participants should they have any questions for the Senior Manager & Policy Advisor for the Office of Research Ethics, 5th floor, York Research Tower, York University, 416-736-5914 or ore@yorku.ca?

☐ Yes ☐ No Have you included a signature line and a date line for participants if a written consent form is being used?

☐ Yes ☐ No Have you included a signature line and a date line for yourself as researcher if either a written consent form or letter is being used?

☐ Yes ☐ No ☐ N/A If the research involves a written questionnaire or a survey, have you attached the survey to the application?

☐ Yes ☐ No ☐ N/A If the study involves the use of a minor, have you included:
  i) A separate information letter to the parents of the minor?
  ii) A separate parental permission letter to be attached to the minor’s letter of consent.
  iii) A signature line for the parent/guardian of the minor?
  iv) A line for the parent/guardian to indicate their relationship to the minor?