From TCPS2 Article 3.2

Consent forms should include the following information. Frequently overlooked items are in bold.

- (a) information that the individual is being invited to participate in a research project;
- (b) the research purpose in plain language, the identity of the researcher, the **identity of the funder or sponsor**, the **expected duration** and nature of participation, and an explanation of the responsibilities of the participant;
- (c) all reasonably foreseeable risks and potential benefits, both to the participants and in general, that may arise from research participation;
- (d) an assurance that prospective participants:
 - are under no obligation to participate; are free to withdraw at any time without prejudice to pre-existing entitlements;
 - will be given, in a timely manner, information that is relevant to their decision to continue or withdraw from participation; and
 - will be given information on their right to withdraw data or human biological materials, including any limitations on the feasibility of that withdrawal. Include a date or deadline after which participants may no longer withdraw their data.
- (e) any possibility of commercialization of research findings or any potential conflicts of interest on the part of the researchers, their institutions or the research sponsors;
- (f) how research results will be disseminated and whether participants will be identified directly or indirectly;
- (g) the identity and contact information of a qualified person who can explain scientific or scholarly aspects of the research to participants;
- (h) the identity and contact information of the Acadia REB Chair provided at http://reb.acadiau.ca.
- (i) what information will be collected about participants and for what purposes; who will have access to information collected about the identity of participants; how confidentiality will be protected; anticipated uses of data; any relevant duty to disclose information collected, and to whom such disclosures could be made;
- (j) any payments, including incentives for participants, reimbursement for participation-related expenses and compensation for injury; and
- (k) a statement to the effect that, by consenting, participants have not waived any rights to legal recourse in the event of research-related harm.

The Acadia Research Ethics Board requires these additional warnings in all consent forms:

- 1. Employers may have legal access to any information transmitted on employer-owned equipment. Participants may therefore wish to use their own equipment if available.
- 2. Any information sent or stored online may be legally accessed by domestic or foreign authorities.
- 3. Participants' data may be accessed by designated employees of Acadia University as needed to fulfill their duties in managing and maintaining Acadia's software and hardware systems.
- 4. Compensated participants will be identified to Financial Services at Acadia University (if compensated through Acadia University).