RESEARCH ETHICS BOARD

REB REVIEWER CONSENT FORM CHECKLIST

Consent Information for prospective participants	TCPS-2 Reference	Included	Not included
Explanation of the responsibilities of the participant.	3.2(b)	☐	☐
Description of the potential benefits of the study, both to participants and to others (in plain language)	3.2 (c)	☐	☐
Description of all the foreseeable risks, side effects and discomforts (in plain language)	3.2 (c)	☐	☐
Statement of withdrawal to include:			
• That potential participants are under no obligation to participate	3.2 (d)	☐	☐
• That participants are free to withdraw at any time	3.2 (d)	☐	☐
• An explanation of the rights of the participant to request the withdrawal of data, and any limitations on the feasibility of that withdrawal	3.2 (d)	☐	☐
Where appropriate, information concerning the possibility of the commercialization of findings and/or real or perceived conflicts of interest	3.2 (e)	☐	☐
Explanation of the dissemination of the results (e.g. will participants be identified directly or indirectly)	3.2 (f)	☐	☐
Identify the name and contact information of a qualified representative who can explain scientific or scholarly aspects of the research to participants	3.2 (g)	☐	☐
Statement informing participants of the names and telephone numbers of the REB that he/she can contact if there are questions about his/her rights as a participant in the study or to report possible ethical issues in the research	3.2 (h)	☐	☐
Description of what information will be collected about participants and for what purposes	3.2 (i)	☐	☐
Description of who will have access to information collected about the identity of the participant	3.2 (i)	☐	☐
Description of how confidentiality and anonymity will be protected	3.2 (i) & 5.2	☐	☐
Description of the anticipated uses of data	3.2 (i)	☐	☐
Where appropriate, information indicating who may have a duty to disclose information collected (and to whom such disclosures could be made)	3.2 (i)	☐	☐
Information about any payments, incentives, reimbursements, and/or compensations for participants	3.2 (j)	☐	☐

Statement to the effect that, by consenting, participants have not waived any rights to legal recourse in the event of research related harm	3.2 (k)	☐	☐
Consent shall be documented in either a signed consent form or in documentation by the researcher of another appropriate means of consent	3.12	☐	☐