**Adverse Event Report Form**

Research Ethics Board



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**Principle Investigator Protocol Number Date**

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**Project Title**

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**Email Phone Number**

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Adverse events can be defined as unexpected potential harms to participants, breaches of confidentiality, newly arising conflicts of interests, or any other event that could compromise the ethical integrity of a research project.

Provide the following information about the adverse event you are reporting:

1. The details of the adverse event.
2. Details on how the research team managed the adverse event or how they propose to manage it.
3. Please email this form to reb@rrc.ca

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Recommendation of the Principal Investigator:

1. The study should continue without change to the protocol Yes [ ]  No [ ]
2. The study should continue without change to the consent form Yes [ ]  No [ ]

\*If no to either question, please submit a study amendment form with this adverse events report form.