**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](mailto: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.