

The University of Montana
Institutional Review Board (IRB)
Continuation Report for Human Subject Research

This report must be completed if data collection is still in progress one year past last IRB approval.

The Institutional Review Board (IRB) is required by Title 21, Code of Federal Regulations (Part 56.109) and Title 45, Code of Federal Regulations (Part 46.109) to conduct continuing review of ongoing projects not less than once per year. Assistance in meeting these federal requirements is appreciated.

Project Director: _____ Dept.: _____

Signature: _____

Phone Number: _____ E-mail Address: _____

Faculty Supervisor: _____ Dept.: _____

Signature : _____ Phone Number: _____

Name of project: _____

Date of last approval: _____

1. Approximately how many subjects have you tested? _____

2. Describe any adverse effect or unanticipated problems involving risks to subjects: _____

3. Describe the circumstances surrounding the withdrawal of any subjects from this research: _____

4. Describe any complaints received from subjects about the research: _____

5. Summarize below any recent findings or publications regarding risks or adverse effects associated with research like yours:

6. If there are any changes you wish to make to your current consent form, attach a copy of the amended form to this document.

If your project originally required action by the full Board, rather than being exempt or administratively approved, it will again be reviewed by the full Board upon receipt of this document.

IRB Determination

For IRB Use Only

____ Exempt from Review

____ Approved by Administrative Review

____ Full IRB Determination:

____ Approved

____ Conditional Approval (see attached memo)

____ Resubmit Proposal (see attached memo)

____ Disapproved (see attached memo)

Signature / IRB Chair _____ Date: _____