



Final Review of Human Subjects Research Investigation

Required for All Human Subjects Research Studies

Study Title:

Principal Investigator:

IRB Protocol Number:

1) Please confirm the status of your project?

☐ All research-related activities are complete and I want to close the study.

2) Provide the following information about subject enrollment:

- Number of subjects approved for enrollment by the IRB
- Number of subjects who provided informed consent to participate in the study since initial approval
- Number of subjects terminated from study (lost to follow-up, terminated by PI, withdrew consent, etc.)
- Number of subjects who provided informed consent to participate in the study during the most recent approval period
- Number of subjects that have completed all study procedures and follow-up visits
- Number of subjects currently participating in study-related activities *(If this number is greater than 0, you must submit a continuing review or annual progress report to keep the study open).*

3) Please include a statement indicating whether any complaints were received about the research study (if none, enter NONE):



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- 4) Provide a list of all study-related adverse events and summarize any study-related adverse events that resulted in changes to the protocol or informed consent documents.

- 5) Describe any other unanticipated problems involving risks to subjects, withdrawal of subjects from the research due to non-compliance, or any issues with recruitment, enrollment and/or retention of subjects.



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- 6) Please provide a succinct narrative (no more than 2 paragraphs) about the study's results and findings.



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- 7) Has any data from this study been disseminated by the investigator in a peer-reviewed forum (e.g., conference, abstract, publication, etc.)?

☐ No.

☐ Yes. If so, please provide an appropriate citation of the work.

Name of PI:

Signature:

Date: