

**Application for Initial Review
Institutional Review Board (IRB)
Chicago State University
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Investigator Agreement

Please read and initial each of the following items in the space provided:

- I agree to conduct the study in accordance with the relevant, current protocol and will only make changes in a protocol after notifying the sponsor and the CSU IRB, except when necessary to protect the safety, rights, or welfare of the research participants.
- I agree to personally conduct or supervise the described investigation.
- I agree to ensure that all of the requirements relating to the recruitment and consent process are met.
- I agree to report to the sponsor and the CSU IRB any adverse experiences and/or events that occur during the course of the experiment.
- I agree to ensure that all associates, colleagues, and employees assisting in the conduct of the study are informed about their obligations consistent with and in meeting the above commitments.
- I agree to maintain adequate and accurate records in accordance with IRB regulations and to make those records available for inspection in accordance with those regulations.
- I ensure that I will submit a request for initial and continuing review and approval to the CSU IRB within the appropriate period of review.
- I agree to report promptly to the IRB any and all changes in the research activity and all unanticipated problems involving risk to the participants and/or others.
- I agree to comply with all other requirements regarding the ethical and legal obligations of clinical investigators and all other pertinent requirements found in the IRB regulations.
- I agree to submit a copy of the final report of the results and a summary of those results upon completion of the study.
- I agree to attach a *Certificate of Completion* from the NIH "Human Participant Protection" website dated during the previous two years to the completed application materials. A link to this website is accessible at the CSU IRB webpage.

The CSU IRB reserves the right to audit any/all IRB approved protocols to inquire about the progress of the study, inspect accrued consent documents, inspect accrued data, and/or observe the consent and recruitment process utilized. The Principal Investigator must cooperate fully with the IRB staff in making such visits.

Signature of Principal Investigator

Signature of Supervising Faculty

Print Name

Print Name

Date

Date