Institutional Review Board
2500 West North Ave.,
Baltimore, MD 21216
Phone: 410.951.3510

Application to Use Human Subjects in Research

Date

Application Type ◯ New Application ◯ Resubmission

Principal Investigator

E-mail Phone

Principal Investigator

E-mail Phone

Principal Investigator

E-mail Phone

University

Mailing Address

Project Title

Type of Project

◯ Faculty or Staff Research Project ◯ Undergraduate or Graduate Student Thesis or Research Project ◯ Doctoral Student Dissertation, Thesis, or Research Project

External Funding Source

Anticipated Project Start Date (Allow 4 weeks from submission)

Type of Review ◯ Exempt from Full Board Review ◯ Full Board Review ◯ Expedited Review

Students are not considered P.I. for purposes of the IRB Application. The P.I. is the faculty advisor or dissertation chair.

Student Name

Student Mailing Address

Address Line Two

City State Zip

Student E-mail Address

Student Telephone Number

Brief Description. A brief description (one paragraph) of the significance of this project in lay terms.

Methods and Procedures. Describe the methods and procedures to be used during the research project. Outline the sequence of events involving human subjects.

Benefits. Describe the benefits (if any) to the subjects involved in the research. (See Human Subjects Handbook).

Risks. Describe the risks (if any) to the subjects involved in the research. (See Human Subjects Handbook).

Study Participants. Describe the study participants, including number, characteristics, and method of participant selections. If a random sample is to be drawn, specify the specific random technique to be used. Justification is required if study participants is restricted to one gender or ethnic group.

Sample Size. A 10% sample frame is recommended for statistical analysis.
In each independently drawn sample, the number of cases should not be lower than 30 cases.
Justification is required if the study utilizes a smaller sample.

Informed Consent. A description of what the Principle Investigator will do to insure the study participants will be informed of all details of the study and consent to participate in the study.

Confidentiality and/or Anonymity. A description of how confidentiality and/or anonymity will be maintained.