

# IRB Statement of Assurance

Investigator: \_\_\_\_\_ Department: \_\_\_\_\_

Co-Investigator \_\_\_\_\_ Telephone Number \_\_\_\_\_

Title of Project \_\_\_\_\_

Investigators must sign and submit electronically the following Statement of Assurance:

The proposed investigation involves the use of human subjects. I am submitting this form with a description of my project, prepared in accordance with the “Assurance of Compliance of HHS Regulations for Protection of Human Research Subjects.”

I agree:

- 1) to obtain informed consent of subjects who are to participate in this project;
- 2) to report to the Institutional Research Board any unanticipated effects on subjects which become apparent during the course or as a result of experimentation and the actions taken as a result;
- 3) to cooperate with members of the Institutional Research Board with the continuing review of this project;
- 4) to obtain prior approval from the Institutional Research Board before amending or altering the scope of the project or implementing changes in the approved consent form;
- 5) to maintain documentation of consent forms and progress reports as required by the “Assurance” cited above.
- 6) to complete the required training and submit documentation of this training to the IRB chairperson. Please contact the IRB to obtain information on the required training (email: [IRB@presby.edu](mailto:IRB@presby.edu)).

Signature of Primary Investigator \_\_\_\_\_ Date \_\_\_\_\_

Signature of Co-Investigator (if applicable): \_\_\_\_\_ Date \_\_\_\_\_