## A signed copy of this page must be included in all protocols submitted for IRB review

## PRINCIPAL INVESTIGATOR ASSURANCE

NAME:	DEPARTMENT:	
PROJECT TITLE:		
project according to participating in all Wide Assurance where Protections (OHRP) Services (DHHS) as (HUIRB) policies as forms will be kept as Howard University notified of any adversal measures employed apprised of any chan prior to their initiation of the IRB which with recruited for this proconsent forms; (3) Assurance with the project of the IRB which with the I	Investigator, give my assurance the rules and regulations govern projects as stipulated in Howard is on file with the Office of the United States Department and Howard University Instituted procedures. Properly exemples part of the records of this project Institutional Review Board (Institutional Report will be sufficient in this protocol and Board at the contain the following: (1) The piect; (2) The number and location and adverse reactions or events the to correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (4) Any adverse reactions or events the correct them; and (5) Any adverse reactions or events the correct them; and (6) Any adverse reactions or events the correct them; and (6) Any adverse reactions or events the correct them; and (6) Any adverse reactions or events the correct them; and (6) Any adverse reactions or events the correct them; and (6) Any adverse reactions the correct them; and (6) Any adverse reactions are correct them;	ning the rights of humans and University's Federal ce for Human Research ent of Health and Human attutional Review Board ecuted informed consent ect. The Chairman of the RB) will be immediately to may occur and of the man of the IRB will be approval will be obtained abmitted to the Chairman ne number of participants ion of executed informed a that may have occurred
Signature of Principal	Investigator	Date