**AUBURN UNIVERSITY HUMAN RESEARCH PROTECTION PROGRAM (HRPP)**

**Developmental Approval Request Form**

For assistance, contact: **The Office of Research Compliance (ORC)**Phone: **334-844-5966** E-Mail: [IRBAdmin@auburn.edu](mailto:IRBAdmin@auburn.edu) Web Address: <http://www.auburn.edu/research/vpr/ohs>

Research Title (should match the title of your grant application): \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_­­­­­­­­­­­­­­­\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Source of Proposed Funding: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

AU OSP Proposal Number: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Proposed Study Dates: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

* Anticipated start date for funding: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_
* Anticipated start date of research activities involving human subjects: \_\_\_\_\_\_\_\_\_\_\_\_

Principal Investigator name \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_­­­­­­

* PI title and degree(s) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_
* AU Department: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_
* Campus Address: \_\_\_\_\_\_\_\_\_\_\_\_\_­\_\_\_\_\_\_\_\_\_\_\_\_
* Preferred telephone: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_
* AU E-mail address: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_
* Signature of PI: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Principal Investigator Assurance Statement: I understand that as Principal Investigator, I have ultimate responsibility for the protection of the rights and welfare of human subjects and the ethical conduct related to this research study. I have carefully read all instructions and information included on the AU Office of Human Research (IRB) website. I further understand no human subjects research activities can occur without AU IRB approval.

PI initials \_\_\_

1. Describe the purpose of the research study.

1. Describe the role of the Auburn University research team.
2. Describe the research activities involving human subjects. Specify whether existing data or specimens will be used and, if YES, specify the source of the data or specimens.
3. Provide the anticipated timeline for human subjects components to be developed and implemented and anticipated timeline when the completed IRB application will be submitted for IRB review.
4. Attach a copy of the proposal submitted to the sponsor or at a minimum, provide the scope of work.

Submit a copy of this completed and signed form and other requested documents to [irbsubmit@auburn.edu](mailto:irbsubmit@auburn.edu).