

INSTITUTIONAL REVIEW BOARD FOR THE PROTECTION OF HUMAN SUBJECTS

Move your cursor over the fields below, then click or	TAP TO ENTER TEXT IN THE	HIGHLIGHTED AREA.
PROJECT TITLE:	ANTICIPATED START DATE:	ANTICIPATED END DATE:
PRINCIPAL INVESTIGATOR:	TELEPHONE:	E-MAIL:
PRINCIPAL INVESTIGATOR DESIGNATION:		
□FACULTY □ UNDERGRADUATE STUDENT □GRADUATE STUDENT	г	
CO-INVESTIGATOR(S):	TELEPHONE:	E-MAIL:
DEPARTMENT:		
DETERMINATION OF RISK/	REVIEW STATUS	
PROTECTED POPULATIONS AND OTHER FULL BOARD DETE	RMINANTS	
\square Minors \square Illegal Behavior \square Sensitive Con	tent 🗆 Cognitively In	npaired/Mentally Ill
□ Pregnant Women □ Prisoners □ Fetuses	□ Veterans/Milit	tary Personnel
☐ FULL BOARD REVIEW ☐ EXPEDITED REVIEW	□ Ехемрт	
PRINCIPAL INVESTIGATOR CERTIFICATION I certify that this research conforms to campus and federal regular justified by sound research design; will adhere to ethical principle and/or objectives of my department/unit and college/division.		
Electronic Signature:		
Co-Investigator(s) Certification (If Applicable):		
I certify that I am familiar with campus policies and procedures ruphold high ethical principles in all research using human subject		of human subjects and v
Electronic Signature:	Date:	

Describe project goal(s):
Describe project participants and how you plan to recruit participants:
Does your project pose any known risk to participants? If so, please describe:
boes your project pose any known risk to participants: It so, please describe.

Does your project involve deception? If so, please describe:
Project Procedures:
 Please describe procedures for obtaining informed consent from participants (Note: At the end of this document please attach the specific informed consent document your participants will view and/or sign).
Please describe specific procedures you plan to use in your project (i.e. Describe what participants will experience).

Please describe how you will safeguard the identity of participants:		
	Please note any additional informatio	
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Use the space at the end of this document to provide the following attachments. These attachments must be complete to receive IRB approval.

- Informed consent document
- Demographic questions
- All information participants will read, hear, or view during the study.
 - o All instruments used in the study should be attached for IRB review.
- Description of participant debriefing procedures

Human Subjects - Institutional Review Board Action

THIS SECTION FOR IRB USE ONLY

PLEASE SAVE THE COMPLETED DOCUMENT IN PDF FORMAT AND EMAIL IT TO THE PRINCIPAL INVESTIGATOR AND THE VPAA OFFICE: RDODSON@STERLING.EDU. FORMS WILL BE KEPT ON FILE IN THE VPAA OFFICE.

Request Denied □	Reason request was denied
More Information Required □	Description of additional information
Request Approved	
Reviewer's Electronic Signature:	Date:

Please use the space below to provide the attachments requested above.				