

2500 W. North Avenue Baltimore, Maryland 21216 Office of the Institutional Review Board (IRB)

IRB Chair: Dr. Michelle Pointer

Dear Researcher:

We are pleased by your plan to conduct research at Coppin State University. The Institutional Review Board (IRB) and entire Coppin family are committed to the development and perpetuation of research that is grounded in conventional and accepted standards and practices. As such, we believe it is important to provide a "user friendly" method by which you can seek approval to conduct research using human subjects at Coppin State.

This letter and attachments will provide you with the information required by Coppin and the Federal Government (U.S. Department of Health & Human Services, DHHS/Office of Human Research Protections/OHRP) to insure that your research does not present a risk to human subjects. To insure that we comply with federal regulations, please provide all requested information.

The attached "Application to Use Human Subjects in Research" and appropriate forms, should be completed and returned to the IRB office. You will receive a response within 30 days from the date of receipt of the application packet. Should you have questions, please do not hesitate to contact the IRB Chair Ft00 kej gng'Rqkpvgt at <u>o rqkpvgt@coppin.edu</u>

Sincerely, Dr. Michelle Pointer

Application to Use Human Subjects in Research

Cover Sheet

The Coppin State University Institutional Review Board (IRB) for the Protection of Human Subjects is charged with the responsibility of reviewing, prior to its initiation, all research involving human subjects. The IRB is concerned with justifying the participation of subjects in research and protecting the welfare, rights, and privacy of subjects.

All material, including this cover sheet, should be submitted to the Chair, IRB, Coppin State University, Dr. Michelle Pointer, in the number of copies required for the type of review. (See below). Incomplete applications will be returned. Call 410-951-3516, if you have questions.

New Application Resubmission	
	/
Principal Investigator (Faculty must serve as Principal Investig	gator for students) Email
Department	Telephone
Project Title	
Study Participants (e.g., students, faculty, parents) External F applicable	Funding Source (present or proposed), if
Student name	/ Telephone
	/
Student mailing address	E-mail
Type of Project:	□ Student Project, Thesis or Dissertation
Anticipated Project Start Date (Collection of data from	human subjects)
Please select the type of review you believe the applie □ Exempt from Full Board Review (Submit two copies) □ Full Board Review (Submit sixteen copies)	cation should receive.
The IRB will not review applications for projects that are alrunderway, research should be immediately suspended until the Return complete application Chair: Dr. Michell (410) 951-3516	he application has been reviewed.

E-mail: mpointer@coppin.edu

APPLICATION FORMAT AND DOCUMENTS

The following information must be attached to the Cover Sheet (see above). Use the headings specified below and in the order presented below, with each item identified and addressed separately, otherwise the application will be returned without review. Center the research topic, PI name, phone and email address at the top of the page.

- 1. **Brief Description** A brief description (one paragraph) of the significance of this project in lay terms.
- 2. **Methods and Procedures** Describe the methods and procedures to be used during the research project. Outline the sequence of events involving human subjects.
- 3. **Benefits** Describe the benefits (if any) to the subjects involved in the research. (See page 27 of Human Subjects Handbook)
- 4. **Risks** Describe the risks (if any) to the subjects involved in the research. (See page 27 of Human Subjects Handbook)
- 5. **Study Participants** Describe the study participants, including number, characteristics, and method of participant selection. 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.
- 6. **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.
- 7. **Informed Consent** A description of what the Principal Investigator will do to insure that study participants will be informed of all details of the study and consented to participation in the study.
- 8. **Confidentiality and/or Anonymity** A description of how confidentiality and/or anonymity will be maintained.

Note: Make sure that the entire application is typed. Handwritten applications will be returned without approval.

Note: The narrative descriptions should be double-spaced.

See "Important Attachments" sheet, below.

IMPORTANT ATTACHMENTS

Applications must include each of the following items, if appropriate to the proposed research

- Informed Consent Document The informed consent document must include the pertinent items from the "Basic Elements of Informed Consent" (See Human Subjects Handbook or Sample on the K drive).
- Questionnaire, Survey, Testing Instrument A copy of any questionnaire, survey, or testing instrument (if any) to be used in this project must be attached. There must be separate validation of instruments that are not established, not vetted, or not in the public domain.
- > Institutional Review Board Authorization Form
- > Advertisements or Posters A copy of any advertising that will be used to recruit subjects.
- Telephone Scripts or Other Recruitment Scripts A copy of any telephone scripts, or other recruitment scripts that will be used.
- ▶ **Debriefing Materials.** Any written or orally presented information indicating that study participants will have the opportunity to contact the Principal Investigator.
- Letters of Approval Letters of approval from each cooperating school, hospital organization, club, or similar type of group (If subjects are obtained through this type of group or organization, a written letter of approval, from an individual authorized to approve such activities, is required). Projects that utilize Coppin's data (i.e. student records, names, etc.) must have the authorization of the appropriate person authorized to release such data.



Institutional Review Board Authorization Form

Name of Studer	nt:	Date:	-
Title of Study			-
Name of Instru	ment(s)		_

I ______, a student at _____College/University having recognized my responsibility to obtain written permission to use the above stated tests/instruments in my research, have rightly done so. Therefore, appropriate documentation and a copy of the instrument are attached for the Principal Investigator to review and for the Principal Investigator to submit to the IRB. The above document(s) has __ has not __ been obtained via public domain usage.

Student Signature: Date:	Date:
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Principal Investigator _____ Date: _____

Institutional Review Board (IRB) Coppin State University

Application to Use Human Subjects in Research

Cover Sheet

The Coppin State University Institutional Review Board (IRB) is charged with the responsibility of reviewing all research involving human subjects, prior to the initiation of such research. The IRB is concerned with protecting the welfare, rights, and privacy of subjects. This application cover sheet, and supporting documents, should be submitted to Dr. Michelle Pointer, Chair, IRB (Health and Human Services Building, 353A). If you have any questions, please call or e-mail Dr. Michelle Pointer at: mpointer@coppin.edu.

Please Check One:	New application	Resubmission of ea	arlier application		
Principal Investigator		E-r	E-mail		
(F	aculty must serve as Principal Investi	gator for student research projects)			
Department	Telephone				
Project Title					
		Will Be Collected (Briefly			
External Funding Sour	rce (if applicable)				
Type Of Project:	_ Faculty/staff research j	project Student r	esearch project or thesis		
Student name (if applicab	le)	E-n	nail		
Student Address		Telephone _			
	art Date (for collection of data fr	om human subjects)			
Type Of Review Requ	iested:Full Bo	oard Review (five copies)			
	Expedit	ted Review (three copies)	Exempt Review		
Please return this appl consent form, to:	ication form, and suppor	ting documents such as co	opy of the informed		
,	Dr. Michelle Poi	nter, Chair, IRB			
	<u>mpointer@coppi</u>	<u>n.edu</u>			