

## **Application for Continuing Review of Human Participant Research**

**Instructions**: Complete this form by checking all appropriate boxes, answering questions completely, attaching required documents and signing the Certification Statement.

Submit application form electronically to irb@westga.edu. (Incomplete applications will be returned unreviewed.)

## **Section I: Study Status** 1. Study Title: 2. IRB Number: 3. IRB Expiration Date: 4. Principal Investigator:\_\_\_\_\_ Responsible faculty member if student is the PI: Department(s): 5. Indicate all that apply: \_\_\_\_\_ Active (study is ongoing with no changes) \_\_\_\_\_ Active with approved changes (study is ongoing with approved modifications) Active with changes being requested with this application (changes are highlighted in attached documents-modification form attached) \_\_\_\_\_ Inactive (study was initiated but data collection and analysis are inactive) \_\_\_\_\_ Inactive - study was not started. Anticipated start date is: 6. Since the last review, are there any new relationships between the researcher(s) and agencies (e.g., schools, hospitals, homes) involved in the research? No Yes If yes, explain and attach a current statement of approval (e.g., letter of agreement) from any agencies that will be involved with the research: 7. Since the last review, have any new or known or potential conflicts of interest related to this research been identified? Conflict of interest relates to situations in which financial or other personal considerations may compromise or involve the potential/have the appearance for compromising an employee's objectivity in meeting University responsibilities including research activities. \_\_No \_\_\_\_Yes If yes, describe the known or potential conflicts of interest and explain how participants will be protected from the influence of competing interests:

## **Section II: Study Modifications/Amendments**

1. Si	nce the last review:		
	There have been no changes in research personnel or their roles and responsibilities.		
	There have been changes in research personnel and/or their roles and responsibilities as indicated		
	below:		
	The following research personnel are no longer associated with the study:		
	The following research personnel are new or have changed roles with the study:		
	New team members or team member with changed roles must complete the required CITI training if the will: 1) access participants' private identifiable information, 2) obtain informed consent, or 3) interact with participants.)		
2. Si	nce the last review. Select one of the following.		
	There have been <u>no</u> approved modifications or addendums to the research study or consent.		
	There have been approved modifications or addendums to the research study or consent.		
	There are requested modifications or addendums to the research study and/or consent form. <i>An IRB</i>		
	Study Modification Form is being submitted with this renewal form and revised application materials reflecting these changes are attached.		
	rejiecting these changes are attached.		
3. Se	lect one of the following.		
	_ This is not a funded study		
	_ There have been no amendments or modifications to the grant since the last review		
	_ There have been amendments or modification to the grant since the last review. A copy of the updated		
	grant materials with changes outlined or highlighted is attached.		
	Section III: Participant Activity/Complaints/Adverse Events		
1. N	umber of participants approved by the IRB:		
2. N	umber of participants accrued to date:		
	escribe any difficulties in participant enrollment is enrollment goals have not been met. How will this impac se study? (attach additional pages if necessary)		
	you have exceeded the sample size initially proposed, explain why (attach additional pages if necessary ibmit a revised application reflecting change in the number of participants):		
 5. N	umber of participants withdrawn prior to completing the study:		
	f applicable, provide specific details about any participant withdrawals from the study, whether voluntary or initiated by the investigator: (attach additional pages if necessary)		
5. No. 15	umber of participants withdrawn prior to completing the study:  f applicable, provide specific details about any participant withdrawals from the study, whether voluntary		

6. Were there any complaints regarding the research	?
No Yes  If yes, please provide a detailed explanation and in	clude copies of the complaints.
Attach as separate sheet.	
7. Were there any adverse events or unanticipated porNo Yes	roblems involving risks to the subjects or others?
If yes, please provide a detailed explanation: <i>Attac</i>	h as separate sheet.
Certification:	
responsibility for ensuring that all members of the • Complete the required CITI training and any o	(and responsible faculty member if PI is a student) accepts a research team: other necessary training to fulfill their study responsibilities. In the IRB approved application letter and comply with the
University of West Georgia's Guidelines for th communication.	e Review of Research Involving Human Subjects and all IRB
<ul> <li>To uphold the rights and welfare of all study p</li> </ul>	participants.
	rand responsible faculty member if PI is a student) have ctronic means, and this application is signed electronically le faculty member if a student is the PI.
with the intent to sign this application, and	titute the symbol and/or process I have adopted my name and email address, set out below, thus signature to this application.
PI Name	PI Email address
Application should be completed by student and sent research proposed, the responsible faculty member stapplication to the UWG IRB at irb@westga.edu.	
scientific merit, rational, and significance. I furth	t I have reviewed and approved the protocol for her acknowledge that I approve the ethical basis for study.
Responsible Faculty Name if PI is a student	Responsible Faculty Email address
Please send an electronic attachment of this application	on and any accompanying materials to irb@westga.edu.
Questions or comments, please contact Charla Campb	