Central Connecticut State University
Institutional Review Board

For IRB use only:
___ Full committee
___ Expedited

Continuing Review and Renewal Form

Along with this form you must submit:

- Any currently used consent and/or assent form(s)
- A separate modification form if any new changes are proposed
- Any publications based on data from the study

Complete and submit this form to irb@ccsu.edu at least 30 days before the expiration date of your approved research.

Investigator Information	**FACULTY MEMBERS:** ***check one*** ☐ Principal Investigator(s) (PI) or Faculty Advisor(s)	**STUDENTS:** *If a student was the lead investigator (LI) for this project, please complete entire section below. If there were multiple students who were lead investigators, please list each LI.* ***NOTE**: Faculty advisors are to review and forward all submissions to the IRB on their advisee's behalf.*
Full Name(s)		
Department(s)		
Email Address(es)		
Institution *(if other than CCSU)*		

IRB approval number:

Study title:

Expected end date:

Current protocol status (please check the appropriate status):

☐	Open to recruitment of new participants.
☐	Closed to recruitment but data collection with existing participants is on going.
☐	Closed to recruitment and data collection is completed; data analysis is on going or to be conducted.
☐	Closed to recruitment, and data collection and data analysis is complete; study is closed.

1. How many participants have enrolled in the study?

2. How many (if any) participants began the study but discontinued participation prior to completion? Provide a summary of any withdrawal of participants from the research and the reasons for withdrawal, if known.

3. Note the nature of any change(s) since last review: *Check all that apply.*

- ☐ *Change in Principal Investigator (PI)*
- ☐ *Addition of subjects to be recruited and/or change to recruitment process.*
- ☐ *Change/addition to the data gathering process-- procedures, interventions or instruments used.*
- ☐ *Change to process for insuring privacy and confidentiality.*
- ☐ *Change to consenting process -- form/script content.*
- ☐ *Other*
- ☐ *None*

4. Provide a brief summary of any modifications to the research approved by the IRB as noted above and the dates these modifications were approved.

5. Provide a summary of any unanticipated problems and/or adverse events. *(In many cases, such a summary could be a brief statement that there have been no unanticipated problems or adverse events.)*

6. Provide a summary of any complaints about the research from participants or others since the last IRB review. *(In many cases, such a summary could be a brief statement that there have been no complaints.)*