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 | 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.

<table>
<thead>
<tr>
<th></th>
<th>Principal Investigator(s) (PI) or Faculty Advisor(s)</th>
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<tbody>
<tr>
<td>Full Name(s)</td>
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<td>Department(s)</td>
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<td>Email Address(es)</td>
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<td>Institution (if other than CCSU)</td>
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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.)