Institutional Review Board **Human Subjects Office** 612 Boyd GSRC Athens, GA 30602-7411



Phone: 706-542-3199 Fax: 706-542-3360

Email: irb@uga.edu

IRB CONTINUING REVIEW/AMENDMENT FORM

Principal Investigator (PI): Andrea Knapp									
Co-Principal Investigator (Required, if co-PI is a student): Andrea Knapp									
Project #: 2009-10070-0 Title of Study: Student Understanding and Sustainability: Challenge Involvement Programs in Mathematics									
PLEASE ANSWER ALL QUESTIONS (Use the text boxes for explanation/additional information or attach a separate cover letter.)									
Have you started data collection for this research project?			\boxtimes						
2	How many total participants have been accrued since the beginning of the research project? (Note: This corresponds to the number of individuals who gave consent; this number should include withdrawals but actual number of withdrawals is reported in #7 below.)			~300					
3	Do you plan to continue to <u>recruit</u> participants for this research project? (If you answered YES , please skip to Question #6.)			\boxtimes					
4	If you answered NO to question #3, do you plan to continue to <u>collect</u> data with previously recruited participants?			\boxtimes					
5	If you answered NO to questions #3 and #4 above, do you plan to continue to <u>analyze</u> previously collected data that is individually-identifiable?								
6	Have there been any complaints about the research since the protocol was approved by the IRB? If YES , please provide complete information on the complaints made.			\boxtimes					
7	Have any participants withdrawn, dropped out, or were lost to follow-up from participation since the protocol was last approved by the IRB? If YES , please indicate the number and provide detailed information/reason(s).								
8	Have there been any adverse events or unanticipated problems involving risks to the participants or others since the protocol was last approved by the IRB? If YES , please contact the IRB office immediately to request an adverse event/incident report form.			\boxtimes					
9	Have there been any chang	ges to the study population? If YES , please explain changes.		\boxtimes					
10	Have the <u>procedures</u> changed in any way since the protocol was last approved by the IRB? If YES , please explain.			\boxtimes					
11	Have any <u>materials or instruments</u> changed in any way since the protocol was last approved by the IRB? If YES , please explain.			\boxtimes					
12	_	fic literature, or interim experience with this or related studies, f potential risks or benefits to study participants? If YES , please evant literature.		\boxtimes					
13	·	s changed in any way since the protocol was last approved by the and attach copy of the revised document(s).		\boxtimes					
14	A <u>clean</u> copy of the current version of the consent document(s) <u>must</u> be submitted with the request to continue if you plan to recruit new participants, or if a revised consent document is necessary as a result of an amendment. Have you attached a clean copy of your current consent document(s)?			\boxtimes					
15	· · · · · · · · · · · · · · · · · · ·	ges to the members of the research team (e.g., change in PI; estigators)? If YES , please describe personnel change(s). Note: plete the CITI training. Racheal Landers , a master's student,							
Princ	Date: 7	19-10							
For electronic submission, a check in this box is acceptable as a signature:									

interventions, <u>and</u> the research will remain active only for lo individually-identifiable private information.	ng-term follow-up of subject	rs; or if the remaining research	activities are limited to analysis o	of