UNIVERSITY OF CENTRAL FLORIDA

Institutional Review Board
FWA00000351
IRB00001138, IRB00012110
Office of Research
12201 Research Parkway
Orlando, FL 32826-3246

APPROVAL

June 18, 2024

Dear Christina Torres:

On 6/18/2024, the IRB reviewed the following submission:

Type of Review:	Modification Update, Category 6,7a
Title:	COLORS Repository
Investigator:	Christina Torres
IRB ID:	MOD00005560
Funding:	None
IND, IDE, or HDE:	None
Documents Reviewed:	• HRP-258 - FORM Memo Cultural Appropriateness(1)_Firmado.pdf, Category: International; •2024Fall_TESOL_Services_Enrollment_Request3.docx, Category: Recruitment Materials; • Consent Document, learners, COLORS Repository5.pdf, Category: Consent Form; • Consent Document, teacher training, COLORS Repository5.pdf, Category: Consent Form; • MOD 5560 HRP-503, COLORS Repository Protocol7.docx, Category: IRB Protocol;

The IRB approved the protocol on 6/18/2024.

In conducting this protocol, you are required to follow the requirements listed in the Investigator Manual (HRP-103), which can be found by navigating to the IRB Library within the IRB system. Guidance on submitting Modifications and a Continuing Review or Administrative Check-in is detailed in the manual. If continuing review is required and approval is not granted before the expiration date, approval of this protocol expires on that date.

If this protocol includes a consent process, use of the time-stamped version of the consent form is required. You can find the time-stamped version of the consent form in the "**Documents**" tab under the "**Final**" column.

When you have completed your research, please submit a Study Closure request so that IRB records will be accurate.

If you have any questions, please contact the UCF IRB at 407-823-2901 or irb@ucf.edu. Please include your project title and IRB number in all correspondence with this office.

UCF IRB approval is not a substitute for international requirements to conduct research. Investigators are responsible for obtaining documentation of local IRB, Ethics Board or equivalent body or committee review of the research or documentation that local ethics review is not required.
Please visit the 2019 Edition of the International Compilation of Human Research Standards
at: https://www.hhs.gov/ohrp/sites/default/files/2019-International-Compilation-of-HumanResearch-Standards.pdf for laws, regulations and guidelines that govern Human Subject Research in 104 countries.

Sincerely,

Kristin Badillo
Designated Reviewer