University of CINCINNATI
Institutional Review Board
FWA #: 00003152

APPROVAL

February 18, 2021

Kimberly McCormick
CECH Education

Dear Kimberly McCormick,

Type of Submission:	Initial Study
Title:	Teaching Practices for the Gifted During COVID 19
Investigator:	Kimberly McCormick
IRB ID:	2020-1248
Funding:	None
Study Risk Level:	No greater than minimal risk
Documents Reviewed:	• Parent Permission-COVIDStudy-Final-21Jan21.doc, Category: Consent Form; • TeacherSurvey-COVIDStudy-FINAL-07Dec2020.docx, Category: Data Collection Tools; • HRP-503-McCormickGuilbaultProtocol-21Jan2021-FINAL, Category: IRB Protocol; • FocusGroupProtocol-Teaching Practices for the Gifted During COVID-FINAL-21Jan2021, Category: Data Collection Tools; • Child Assent Information sheet-COVIDStudy-FINAL-21Jan21.doc, Category: Consent Form; • StudentSurvey-COVIDStudy-FINAL-07Dec2020.docx, Category: Data Collection Tools; • HRP-502I-Information Sheet-COVIDStudy-Final-21Jan21.docx, Category: Consent Form; • Parent_Student-Recruitment-21Jan2021.docx, Category: Recruitment Materials; • Teacher-Recruitment-21Jan2021.docx, Category: Recruitment Materials;

On **1/28/2021**, the IRB reviewed the above submission using an EXPEDITED review procedure in accordance with 45 CFR 46.110(b)(1) which was given approval pending the response to modifications required. Response to the modifications required provided to the IRB were reviewed and approved on **2/17/2021** under the following category(ies):

- (7)(a) Behavioral research
- (7)(b) Social science methods

The IRB approved the protocol from **1/28/2021** to **1/27/2024**. Thirty days before **1/27/2024**, you are to submit a completed continuing review and required attachments to request continuing approval or closure. You can submit a continuing review by navigation to the active study and clicking Create Modification/CR. If continuing review approval is not granted before the expiration date of **1/27/2024**, approval of this study expires on that date.

THE IRB HAS DETERMINED THE FOLLOWING CONSENTING REQUIREMENTS:

- **Per 45 CFR 46.117 (21 CFR 56.109)** the IRB has waived the requirement to obtain DOCUMENTATION of informed consent for all adult participants.
- **Per 45 CFR 46.117 (21 CFR 56.109)** the IRB has waived the requirement to obtain DOCUMENTATION of parental permission for all child participants.
- **Per 45 CFR 46.408 (21 CFR 50.55)** the IRB has waived the requirement to obtain DOCUMENTATION of assent for all child participants.
- The Board determined that this study meets the criteria for enrollment of minors as described in UC Human Research Protection Program Policy V.01 and Procedure 332.

PI NOTIFICATIONS:

This approval is through the IRB only. You may be responsible for reporting to other regulatory officials. Please check with your institution and department to ensure you have met all reporting requirements.

INTERNATIONAL CONFERENCE ON HARMONIZATION AND GOOD CLINICAL PRACTICES STATEMENT:

The Institutional Review Board is duly constituted (fulfilling FDA requirements for diversity), has written procedures for initial and continuing review of clinical trials: prepares written minutes of convened meetings and retains records pertaining to the review and approval process all in compliance with requirements defined in 21 CFR Parts 50, 56 and 312 Code of Federal Regulations. This institution is in compliance with the ICH GCP as adopted by FDA/DHHS.

Thank you for your cooperation during the review process.