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APPROVAL OF SUBMISSION

Date: January 16, 2019

From: Philip Frum, IRB Analyst

To: Latoya Haynes-Thoby

Type of Submission:	Initial Study
Title of Study:	Factors That Influence Resilience for Transition-Aged Black Mothers with Histories of Trauma
Principal Investigator:	Latoya Haynes-Thoby
Study ID:	STUDY00011139
Submission ID:	STUDY00011139
Funding:	Not Applicable
IND,IDE, or HDE:	Not Applicable
Documents Approved:	• Connor-Davidson Resilience Scale (not to be printed in entirity within dissertation) (0.01), Category: Data Collection Instrument • Demographic Survey.docx (0.01), Category: Data Collection Instrument • Eligibility Questions.docx (0.01), Category: Other • Follow-Up Questions.docx (0.01), Category: Data Collection Instrument • HRP-589 - ORP Consent Form (Waiver of Written Documentation of Consent)-2.pdf (0.02), Category: Consent Form • HRP-591 - Protocol for Human Subject Research-3-1.pdf (0.03), Category: IRB Protocol • Research Study Flyer-2.docx (0.01), Category: Recruitment Materials • Spirituality Scale.pdf (0.01), Category: Data Collection Instrument • Trauma History Questionnaire (0.01), Category: Data Collection Instrument
Review Level:	Expedited

On 1/16/2019, the IRB approved the above-referenced Initial Study. This approval is effective through 1/15/2020 inclusive. You must submit a continuing review form with all required explanations for this study at least 45 days before the study's approval end date. You can submit a continuing review by navigating to the active study and clicking 'Create Modification / CR'.

If continuing review approval is not granted before 1/15/2020, approval of this study expires on that date.
Attached are stamped approved consent documents. Use copies of these documents to document consent.
In conducting this study, 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 CATS IRB (http://irb.psu.edu). These requirements include, but are not limited to:

- Documenting consent
- Requesting modification(s)
- Requesting continuing review
- Closing a study
- Reporting new information about a study
- Registering an applicable clinical trial
- Maintaining research records

This correspondence should be maintained with your records.