Subject: E19135 PI: Andrew Heiss IRB Determination: APPROVAL

From: Human Subjects Committee - To: andrew_heiss@byu.edu - Cc: - Date: May 9, 2019 at 17:44, Attachments: image003.jpg
Heiss E19135 consent.pdf

INSTITUTIONAL REVIEW BOARD
FOR HUMAN SUBJECTS

Memorandum

To: Professor Andrew Heiss
Department: RIPM
College: MSB
From: Sandee Aina, MPA, IRB Administrator
Bob Ridge, PhD, IRB Chair
Date: May 9, 2019
IRB#: E19135
Title: *"Why Donors Donate: Disentangling Organizational and Structural Heuristics for International Philanthropy"*

Brigham Young University's IRB has approved the research study referenced in the subject heading as exempt level, category 2. This category does not require an annual continuing review. Each year near the anniversary of the approval date, you will receive an email reminding you of your obligations as a researcher and to check on the status of the study. You will receive this email each year until you close the study.

The study is approved as of **May 9, 2019**. Please reference your assigned IRB identification number in any correspondence with the IRB.

Continued approval is conditional upon your compliance with the following requirements:

1. A copy of the informed consent statement is attached. No other consent statement should be used. Each research subject must be provided with a copy or a way to access the consent statement.
2. Any modifications to the approved protocol must be submitted, reviewed, and approved by the IRB before modifications are incorporated in the study.
3. All recruiting tools must be submitted and approved by the IRB prior to use.
4. In addition, serious adverse events must be reported to the IRB immediately, with a written report by the PI within 24 hours of the PI's becoming aware of the event. Serious adverse events are (1) death of a research participant; or (2) serious injury to a research participant.
5. All other non-serious unanticipated problems should be reported to the IRB within 2 weeks of the first awareness of the problem by the PI. Prompt reporting is important, as unanticipated problems often require some modification of study procedures, protocols, and/or informed consent processes. Such modifications require the review and approval of the IRB. Please refer to the IRB website for more information.

Rachel Teki
IRB Secretary
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Brigham Young University
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