Office of Research Compliance
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NOTICE OF APPROVAL OF EXPEDITED HUMAN SUBJECTS RESEARCH

DATE:	September 08, 2020
TO:	Gunty, Mark, PhD, Office of Strategic Planning and Institutional Res
FROM:	Hall, Matthew, PhD, Expedited Review
PROTOCOL TITLE:	Mission Identity New Students Survey
FUNDING SOURCE:	None
PROTOCOL NUMBER:	20-07-6140
APPROVAL PERIOD:	Approval Date: September 08, 2020 Expiration Date: September 07, 2025

The University of Notre Dame Institutional Review Board (IRB) has reviewed the research protocol described above. The research has been approved as described in the protocol. In compliance with (as applicable) 21 C.F.R. 56.109(e) and 46 C.F.R. 46.109(d), this notice serves as written notification of the IRB#s determination. This approval is based on your agreement to abide by the policies and procedures of the University of Notre Dame Human Research Protection Program and does not supercede any other approvals required for you to conduct your human subjects research. Relevant policies and procedures can be found at:
research.nd.edu/our-services/compliance/human-research

The research procedures described in this protocol have been approved under the following Expedited categories:
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If applicable, renewal may be required for the human subjects research activities to continue past any expiration date indicated. IRB approval is also required prior to implementing any changes or amendments to the protocol, regardless of how minor, except to eliminate immediate hazards to subjects. No changes to the informed consent document may be made without prior IRB approval.

The initial approval period is noted above. Continued approval may be contingent upon the submission of a renewal application to the Human Subjects Office in a timely fashion. Failure to submit a renewal notice in a timely fashion may result in the expiration and subsequent suspension of all protocol activities. Failure to receive notification from the Notre Dame Research Compliance Office will not relieve you of your responsibility to ensure compliance with Federal Regulations regarding annual review [as applicable, 21 C.F.R. 56.109(f) and 45 C.F.R. 46.109(e)]

Unanticipated problems involving risks to subjects and others, incidents of non-compliance, protocol violations, and sponsor or investigator initiated suspensions of research must be reported to the IRB following IRB SOPs.

You should retain a copy of this letter and all associated approved study documents for your records. Please refer to the assigned study number and study title in future correspondence with our office.

This approval is issued under University of Notre Dame's Federal Wide Assurance (FWA00002462) with the Office for Human Research Protections (OHRP). If you have any questions regarding your obligations under Committee's Assurance, please do not hesitate to contact us.

Hall, Matthew, PhD

Chair, Notre Dame Research Compliance