Northwestern | RESEARCH

Northwestern University
Institutional Review Board
Biomedical IRB
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Social & Behavorial Sciences IRB
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**Statement of Compliance with FDA 21 CFR Part 11**

Date: April 18, 2023

Northwestern University's Researchers conducting human participant research, are required to use an electronic system (eIRB+) to submit applications to the Institutional Review Boards (IRBs). Northwestern University's IRB Office and IRBs also use eIRB+ to conduct and record the review and approval process of human participant research, and applicable compliance processes. eIRB+ is a closed system.

This is to certify that the Northwestern University IRBs indicated below are in compliance with the U.S. Food and Drug Administration federal regulations found at 21 CFR Part 11 pertaining to Electronic Records: Electronic Signature Standards.

DocuSigned by:
Nathalia Henry-Whitely
9C51B42440934AD...

**Nathalia Henry Whitely, MS, CIP, CHRC**
Executive Director, IRB Office

Name of the IRB: Northwestern University IRB Panels A, B, C, D, E and Q

Address:

C/O IRB Office:

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