THE UNIVERSITY OF IOWA

Human Subjects Office/
Institutional Review Board (IRB)
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STATEMENT OF COMPLIANCE

The University of Iowa has filed an federalwide assurance (FWA) of compliance with the Office of Human Research Protections under FWA000003007. The University of Iowa has assured activities related to human subjects research, regardless of the source of support, will be guided by the Belmont Report as its statement of principles governing the institution for protecting the rights and welfare of human subjects.

The University of Iowa FWA includes three registered internal Institutional Review Boards (IRB) under IORG# IORG0000070.

IRB-01(Biomedical) IRB00000099
IRB-02(Social Science\Behavioral) IRB00000100
IRB-03(Biomedical\VA Review only) IRB00006022

The University of Iowa Institutional Review Boards are duly constituted with written procedures for initial and continuing review of human subjects research; written minutes of convened IRB meetings, an electronic research application system to adhere to records retention requirements of the review and approval process. All documented processes are in compliance with requirements of FDA regulations 21 CFR Parts 50 and 56, HHS regulations 45 CFR 46, and International Conference on Harmonization (ICH) E6, Good Clinical Practice (GCP) as adopted by the FDA, as applicable.

The University of Iowa also reviews in compliance with other applicable federal and state laws and regulations governing IRBs and human subjects research. A full copy of the IRB standard operating procedures can be found on the Human Subjects website at www.hso.research.uiowa.edu.

The University of Iowa has been fully accredited by the Association for the Accreditation of Human Research Protection Programs (AAHRPP) since 2003.