UCONN HEALTH

Statement of Compliance

The University of Connecticut (UConn) Health Center Institutional Review Boards (UConn Health IRBs) are organized and operate in accordance with applicable laws, regulations, and guidelines in the United States including, but not limited to, U.S. Food and Drug Administration (FDA) 21 CFR Parts 50 and 56, U.S. Department of Health and Human Services regulations 45 CFR Part 46, International Conference on Harmonisation (ICH) Good Clinical Practice (GCP), and the Belmont Report. Where appropriate, UConn Health IRBs comply with additional regulations and guidelines as required in specific research jurisdictions.

UConn Health IRBs are registered with FDA and OHRP and UConn Health Federalwide Assurance (FWA) is approved by OHRP:

- UConn Health IRB Panel 1 Registration #: IRB00000451
- UConn Health IRB Panel 2 Registration #: IRB00000452
- IRB Organization (IORG) #: IORG0000266
- FWA#: 00006064

The UConn Health Human Subjects Protection Program has been fully accredited by the Association for the Accreditation of Human Research Protection Programs (AAHRPP) since 2006.

The primary responsibility of UConn Health IRBs is to ensure that the rights and welfare of the human subjects who participate in research studies are protected.

Version dated 7/16/2024

Human Subjects Protection Program
Institutional Review Board
263 FARMINGTON AVENUE
FARMINGTON, CT 06030-1511
PHONE 860.679.3054
FAX 860.679.1005
research.uchc.edu/rcs/hspp