



Human Research Protection
Office of Research Protections

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STATEMENT OF COMPLIANCE

The University of Pittsburgh IRB is duly constituted, has written procedures for initial and continuing review of research studies; prepares written minutes of convened meetings, and retains records pertaining to the review and approval process; all 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 Food and Drug Administration. The University of Pittsburgh has an active FWA (FWA00006790) and is registered with OHRP/FDA. In addition, the University of Pittsburgh HRP has full accreditation by the Association for Accreditation of Human Research Protection Programs, Inc. (AAHRPP).

A handwritten signature in black ink, appearing to read "M. Hsieh".

Margaret Hsieh, MD
IRB Chair

December 2, 2025