STANFORD UNIVERSITY

Administrative Panel on
Human Subjects in Medical Research

2018-2019
IRB #6: Roster

Palo Alto, CA  94306
Assurance #FWA00000935

This panel is composed of 23 members.

VOTING MEMBERS

Fourteen Members affiliated with Stanford University

OAKES, David D. (M.D.) (CHAIR)
Professor, Emeritus
Surgery

LANE, Alfred (M.D.)
Professor
Dermatology and Pediatrics

AMYLON, Michael D. (M.D.)
Professor, Emeritus
Pediatrics

MEYER, Timothy W. (M.D.)
Professor
Medicine/Nephrology (VA)

DOAN, Samantha (CIP)
Nonscientific Member

MUNDY, David C.H. (M.Div.)
Reverend
Nonscientific Member

DOHERTY, Anastasia (CIP)
Nonscientific Member

RECHT, Lawrence D. (M.D.)
Professor
Neurology and Neurological Sciences

GILL, Manjit (CIP)
Nonscientific Member

SCANDLING, John D. (M.D.)
Professor
Medicine/Nephrology

GOEL, Ashima (M.S., CIP)
Nonscientific Member

STOCKDALE, Frank E. (M.D.)
Professor, Emeritus
Medicine/Oncology

HOLZER, Alison (Ph.D.)
Clinical Research Manager
Pediatrics

WILSON, Darrell M. (M.D.)
Professor
Pediatrics/Endocrinology
Two Outside Nonscientific Members Otherwise Unaffiliated with Stanford

EIGENBROD, Richard A. (M.B.A.)
Business Consultant

PETERHANS, Laura (M.A.)
Teacher (Retired)

NON-VOTING MEMBERS

Seven Ex Officio Members

BANGHART, Dawn (C.H.P.)
Sr. Health Physicist
Environmental Health and Safety (EH&S)
(Alternate for Lance Phillips)

SEGAL, Ellyn D. (Ph.D.)
Biosafety Officer
Environmental Health and Safety (EH&S)

CAPLUN, Elizabeth
Deputy Director
Office of the Vice Provost and Dean of Research
(Alternate for Kathy McClelland)

STAYN, Susan (J.D.)
Senior University Counsel
Office of the General Counsel

MCULELLAND, Kathy
Research Compliance Director
Office of the Vice Provost and Dean of Research

THOMPSON, Kathleen
Director, Research Management Group
Office of the Dean of the School of Medicine

PHILLIPS, Lance (M.S., C.H.P., C.S.P.)
Radiation Safety Officer
Environmental Health and Safety (EH&S)

ICH/GCP: Stanford University Administrative Panels on Human Subjects in Medical Research (IRB) are in compliance with Good Clinical Practices as consistent with U.S. Food and Drug Administration Code of Federal Regulations (21 CFR 50 and 56) and DHHS (45 CFR Part 46).

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