STANFORD UNIVERSITY

Administrative Panel on Human Subjects in Medical Research

2017-2018
IRB #6: Roster

Palo Alto, CA 94306
Assurance #FWA00000935

This panel is composed of 24 members

VOTING MEMBERS

Fourteen Members affiliated with Stanford University

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

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

DOHERTY, Anastasia (CIP)
Nonscientific Member

GILL, Manjit (CIP)
Nonscientific Member

GOEL, Ashima (M.S.)
Nonscientific Member

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

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

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

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

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

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

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

WILSON, Darrell M. (M.D.)
Professor
Pediatrics/Endocrinology

WITWER, Chris, CIP
Nonscientific Member

Three Outside Nonscientific Members Otherwise Unaffiliated with Stanford

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

PARKER, George W. (M.B.A.)
Controller (retired)

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)

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

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

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

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

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

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

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).

Last updated: April, 2018