STANFORD UNIVERSITY

Administrative Panel on Human Subjects in Medical Research

2015-2016
IRB #6: Roster

Palo Alto, CA 94306
Assurance #FWA00000935

The Panel is composed of 23 members

VOTING MEMBERS

Thirteen 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

JACOB, Theodore (Ph.D.)
Career Research Scientist
Psychology Service (VA)

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

ZARCONE, Vincent P. (M.D.)
Professor, Emeritus
Psychiatry and Behavioral Sciences

MOLVIN, Celia (CIP)
Nonscientific Member

Three Outside Nonscientific Members Otherwise Unaffiliated with Stanford

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

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

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

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)

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

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

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

JAMES, Ann (Ph.D., 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

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

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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