STANFORD UNIVERSITY

Administrative Panel on
Human Subjects in Medical Research

2014-2015
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

SLAMOWITZ, Debra (R.N.)
Nurse
Medicine/Infectious Diseases

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

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

CLARK-WORLEY, Sarah (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

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

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

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

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

BANHART, 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).

Last updated: July 2015