This panel is composed of 22 members

VOTING MEMBERS

Eleven members affiliated with Stanford University

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

FAUSTMAN, William (Ph.D.)  
Clinical Professor, Affiliated  
Psychiatry and Behavioral Sciences (VA)

ARIAGNO, Ronald L. (M.D.)  
Professor, Emeritus  
Pediatrics

GILL, Manjit (CIP)  
Nonscientific Member

COHANE, Carol A. (R.N., B.S.N.)  
Clinical Research Coordinator  
Department of Anesthesiology, Pain and Perioperative Medicine

LEMMENS, Hendrikus J.M. ((M.D., Ph.D.)  
Professor  
Department of Anesthesiology, Pain and Perioperative Medicine

DOHERTY, Anastasia (CIP)  
Nonscientific Member

PERKASH, Inder (M.D.)  
Professor, Emeritus  
Urology

DRESCHER, Kent D. (Ph.D.)  
Health Science Specialist (Retired)  
Psychiatry (VA)

RASMUSSEN, Lucille E. (Sc.D.)  
Senior Research Scientist (retired)

ECKERT, Penelope D. (Ph.D.)  
Professor  
Linguistics
Four Outside Nonscientific Members Otherwise Unaffiliated with Stanford

ODA, Robert
Director (retired)
Camp Glenwood
(Prisoner Representative)

ROE, Richard R. (M.Th.)
Executive Director
Council of Churches of Santa Clara County

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

STOVEL, John E. (Jack) (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

MCCLELLAND, 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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