The Panel is composed of 25 members

VOTING MEMBERS

Fourteen members affiliated with Stanford University

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

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

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

DOHERTY, Anastasia (CIP)
Nonscientific Member

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

ECKERT, Penelope D. (Ph.D.)
Professor
Linguistics

HOLMAN, Halsted (Hal)(M.D.)
Professor, Emeritus
Medicine/Immunology and Rheumatology

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

MOLVIN, Celia (CIP)
Nonscientific Member

MURPHY, Kevin (CIP)
Nonscientific Member

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

RASMUSSEN, Lucille E. (Sc.D.)
Senior Research Scientist (retired)
FAUSTMAN, William (Ph.D.)
Clinical Associate Professor, Affiliated
Psychiatry and Behavioral Sciences (VA)

GILL, Manjit (CIP)
Non-scientific Member

Four Outside Non-scientific Members Otherwise Unaffiliated with Stanford

ODA, Robert
Director (retired)
Camp Glenwood

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

STOVEL, John E. (Jack) (M.A.)
Teacher (retired)

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

NON-VOTING MEMBERS

Six 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

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

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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