# Supplemental Tables

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Supplemental Table 1: Post-Workshop Ratings of Workshop Objectives by the Students - Classes 2001 to 2014															
Includes of Biostatisticians and/or Cor	Percentages of participants at Workshops held from 2001 to 2014 indicating "Agree" or "Strongly Agree" with the listed statement											tement			
Class	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	Mean
N total	109 <sup>L</sup>	105 <sup>L</sup>	107 <sup>L</sup>	106 <sup>L</sup>	104 <sup>L</sup>	101	102 <sup>L</sup>	102 <sup>L</sup>	98*	101 <sup>L</sup>	100 <sup>L</sup>	100	97	99	102
Total Jr. Faculty or Fellow	100	100	102	100	101	101	101	100	99	100	99	100	97	99	100
Total Other Participant	9	5	5	6	3	0	1	2	0	1	1	0	0	0	2
Rate of return (%)	87%	87%	88%	89%	75%	98%	98%	98%	96%	96%	99%	94%	86%	97%	97%
Mean percentage ratings								-							
Workshop objectives								-							
In this workshop I was introduced to the principles of good clinical trial design	100	100	100	100	100	100	100	100	100	100	100	100	99	99	99.9
I was given the tools needed to conduct clinical trials that will yield clear results which investigators can use to proceed to the next level research	100	100	100	98	100	99	99	100	100	100	100	100	98	99	99.5
The workshop exposed me to the full spectrum of challenges in clinical research, from conventional antineoplastics to gene therapy	97	95	96	96	98	99	97	95	97	96	97	96	96	98	96.6
I was given adequate information in the protocol development sessions to develop a clinical trial protocol	99	98	99	98	99	98	98	99	100	99	98	98	96	99	98.4
My interest in conducting cancer clinical trials increased as a result of attending the workshop	99	97	98	96	97	96	99	99	100	99	97	99	100	96	98.0
The workshop increased my understanding of cancer and enabled me to understand new concept	95	94	96	97	91	100	95	97	95	95	97	97	96	95	95.7
I have made contacts with faculty members that will be useful in my future research	99	99	99	99	97	96	96	96	99	95	100	100	96	98	97.8
I have made contacts or networked with other participants	97	96	95	93	96	97	96	100	100	96	100	98	99	96	97.1

that will be useful in my future								
research								

# Supplemental Table 2: Supplemental Table 2: Mean Post-Workshop Ratings by the Students - Classes 2001 to 2014 <sup>L</sup>Includes Jr Biostatisticians and/or Corporate participants in analysis \*A total of 99 students attended in 2009, but one student failed to receive an evaluation

	2001	2002	2002	2004	2005	2007	2007	2000	2000	2010	2011	2012	2012	2014	M
	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	Mean
N total	109 <sup>L</sup>	105 <sup>L</sup>	107 <sup>L</sup>	106 <sup>L</sup>	104 <sup>L</sup>	101	102 <sup>L</sup>	102 <sup>L</sup>	98*	101 <sup>L</sup>	100	100	97	99	102
Total Jr. Faculty or Fellow	100	100	102	100	101	101	101	100	99	100	99	100	97	99	100
Total Other Participant Type	9	5	5	6	3	0	1	2	0	1	1	0	0	0	2
Rate of return (%)	87%	87%	88%	89%	75%	98%	98%	102 <sup>L</sup>	96%	96%	99%	97%	86%	97%	92%
Didactic lectures				-											
Overall, the lecture addressed the stated instructional objectives	3.6	3.7	3.7	3.7	3.7	3.7	3.7	3.7	3.7	3.7	3.8	3.8	3.8	3.7	3.7
Overall, I learned a great deal from this lecture	3.4	3.5	3.6	3.6	3.5	3.6	3.5	3.5	3.5	3.6	3.7	3.7	3.6	3.6	3.6
The lecture will be helpful to me in designing/conducting clinical trials	3.2	3.4	3.5	3.6	3.5	3.6	3.5	3.4	3.5	3.6	3.6	3.7	3.6	3.6	3.6
The lecture was delivered in an effective manner	3.5	3.6	3.6	3.7	3.6	3.7	3.6	3.6	3.6	3.6	3.7	3.8	3.7	3.7	3.7
Protocol Development Groups															
In the protocol development group session, I received constructive feedback	3.8	3.8	3.8	3.8	3.8	3.9	3.8	3.8	3.9	3.8	3.8	3.8	3.9	3.9	3.8
I am leaving the workshop with a completed protocol	3.5	3.5	3.6	3.6	3.7	3.6	3.6	3.8	3.7	3.7	3.7	3.7	3.8	3.7	3.7
I expect to implement the trial described in my protocol within the next year	3.5	3.6	3.5	3.6	3.6	3.7	3.7	3.7	3.6	3.7	3.7	3.7	3.8	3.7	3.7
Faculty in Protocol Development Groups	3.8	3.8	3.8	3.8	3.9	3.8	3.9	3.7	3.9	3.8	3.9	3.8	3.9	3.9	3.8
Small Group Discussions															1
Overall I learned a great deal from the discussion session	3.3	3.1	3.4	3.6	3.4	3.5	3.5	3.3	3.4	3.4	3.5	3.6	3.6	3.3	3.5
The discussion will be helpful to me in designing/conducting research protocols	3,2	3.3	3.5	3.6	3.4	3.5	3.5	3.3	3.4	3.4	3.6	3.6	3.6	3.3	3.5
The discussion session was conducted in an effective manner	3.4	3.5	3,2	3.7	3.5	3.6	3.5	3.3	3.5	3.5	3.5	3.6	3.6	3.4	3.5

	1-2 years	3-4 years	$\geq$ 5 years
	surveys	surveys	surveys
	Class of	Class of	Class of
	2001 to 2007	1998 to 2005	1996 to 2003
N return / N total	488 / 839	287 / 736	266 / 835
Rate of return (%)	58	38	32
Completion of medical training	73	96	99
Current position in:			
Academics/University	88	85	79
Activity main focus:			
1. Patient-oriented clinical research	49	53	60
2. Translational research	31	31	19
3. Basic research	3	2	5
4. Clinical practice	14	13	11
5. Other	2	1	5
Time spent on patient-oriented clinical research:			
1. Substantial	61	68	73
2. Some	31	29	21
3. Little	7	3	6
4. None	1	0	1
My participation in the workshop has been valuable to me in			
conducting clinical trials.			
If yes, why?	99	100	99
- Oriented me to principles of good clinical trial design	94	90	93
- Increased my understanding of the full spectrum of challenges in	86	88	86
clinical cancer research			
- Familiarized me with the process of approval/ implementation of	74	71	76
clinical trials			
- Raised my awareness of the importance of good clinical trial design	80	61	81
for the benefit of cancer patients			
- Other	7	8	11
My participation to the workshop has been valuable in advancing			
my career.			
If yes, why?	93	89	91
- Enhanced my clinical research productivity	63	61	68
- Stimulated me to continue my career in clinical cancer research	79	76	77
- Provided opportunities for networking	64	56	60
- Other	5	5	5
Have you maintained contact with any of the workshop faculty			
and/or fellow participants?			
If yes, why?	63	57	48
- Provided faculty mentoring	37	40	21
- Offered new prospects for research collaborations	41	48	44
- Opened up new job opportunities in clinical research	17	22	10
- Other	20	11	10
Status of workshop protocol/clinical trial			
- Submission to IRB	54	60	69
- Approval by IRB *	48	57	68
- Funding*	43	44	52
How many additional protocols have been submitted or approved			
by an IRB?			

# Supplemental Table 3 - Mean Percentage Outcome Measures Obtained from Students at 1 to 2Years, 2 to 3 Years, and $\geq$ 5 Years after the Workshop

Ye	ears, 2 to 3 Years, and $\geq$ 5 Years after the Workshop			
		1-2 years	3-4 years	$\geq$ 5 years
		surveys	surveys	surveys
		Class of	Class of	Class of
		2001 to 2007	1998 to 2005	1996 to 2003
1.	0	22	6	2
2.	1	26	10	2
3.	2	23	16	6
4.	3	16	17	14
5.	4	15	13	2
6.	≥ 5	8	39	49
Ho	w many have been implemented?			
1.	0	28	7	2
2.	1	29	15	5
3.	2	23	17	11
4.	3	11	16	7
5.	4	4	12	1
6.	$\geq$ 5	6	33	50

# Supplemental Table 3 - Mean Percentage Outcome Measures Obtained from Students at 1 to 2

## **Supplemental Material**

## S1. Teaching Schedule, Teaching Methods, and Evaluation Methods

The schedule was quite intense, with some participants referring to it as a "boot camp." (Refer to Supplemental Material S2 for a recent definitive example of the program.)

To impart the most knowledge during the workshop and have the students depart from the venue with an actionable protocol, the following teaching and evaluation methods were used.

Teaching methods

- 1. Didactic lectures A total of 20 lectures including discussion was scheduled throughout the course.
- 2. Protocol Development Groups (PDG) This was a particularly unique aspect of the workshop. Each student was assigned to a PDG for the week. Most PDGs comprised 8-10 students led by 2-3 clinical investigator faculty members, a biostatistician, and a patient advocate. The PDGs met at least 8 times throughout the workshop beginning on the first night with each student presenting his/her concept followed by initial feedback and questions from the faculty and fellow students. Throughout the workshop there were deadlines for deliverables, such as: (a) a formal concept sheet; (b) a draft protocol; (c) a patient informed consent form; and (d) a completed protocol suitable for submission to an IRB. Each document was reviewed and returned with in-person and written feedback from PDG mentors. The PDG sessions were intense, constructive, and highly interactive. Students often asked questions of each other and offered suggestions, and the faculty, by working together on the students' projects, had the opportunity to model the collaborative behavior needed for team science.
- 3. Other teaching methods These included interactive Small Group Discussion Sessions covering a wide range of special topics such as biomarkers; selection of appropriate clinical trial endpoints; special populations; role play of an informed consent (along with a patient); Meet-the-Professor Sessions; career counseling sessions including advice on scientific writing, oral presentations, and grantsmanship; Special Interest Groups (e.g., immunology, radiation oncology, etc.); and mentoring, consultations, one-on-one meetings, and multiple other informal interactions at nearly every meal or break.

Evaluation methods

- 1. A pre-workshop test was given upon arrival on the first day (Saturday afternoon), consisting of 50-60 multiple-choice questions. This provided a baseline to determine how much didactic material had been mastered when compared to the same questions answered in a post-workshop exit test. These pre- and post-tests were mandatory for students and optional for junior biostatisticians and corporate attendees.
- 2. Daily and overall workshop evaluation questionnaires in which the participants (mandatory for students and optional for junior biostatisticians and corporate attendees) anonymously rated, on a scale from 1 (Strongly Disagree) to 4 (Strongly Agree), their level of satisfaction with the program, learning activities, and faculty.

- 3. Faculty ratings of students' achievement and progress in developing and completing their protocols during the Protocol Development Groups.
- 4. Intermediate and long-term surveys administered, respectively, 1 to 4 and 5 or more years after the workshop, to follow up on the students' academic position, level of activity in clinical research, and networking with workshop faculty and student colleagues. The very first survey was administered in 2002, and since then a total of seven surveys and eight workshops have been included in this review. Of the seven surveys, the first two were postal surveys, while the last five were administered online.
- 5. Comparisons by Thomson Reuters, through their data-mining method, of the numbers of publications, grants received, collaborations, and clinical trials implemented between 2002 and 2010 by those applicants who were selected versus those not selected for the workshop. The methodology for this study is outlined in S7.

# S2. Typical Schedule for Workshop (2014 provided as an example)

# Saturday, July 25

- Registration
- Administration of Pretest
- Welcome and Overview
- Protocol Development Group Session 1 / Dinner

# Sunday, July 26

- Continental Breakfast
- Lecture Session 1
  - Statistical Considerations in Protocol Development: From Hypothesis to Analysis
  - Design and Analysis of Phase I Clinical Trials in Cancer Therapy
  - Novel Statistical Designs in Phase I Studies
- Lecture Session 2
  - Principles of Clinical Pharmacology
  - Phase II Trials
  - Novel Phase II Designs
- Protocol Development Group Session 2
- Lunch
- Office Hours 1: Protocol Related Questions
- Independent Study Time
- ASSIGNMENT DUE: PROTOCOL SYNOPSIS SHEETS
- Reception / Dinner

# Monday, July 27

- Continental Breakfast
- Lecture Session 3
  - Incorporation of Biological Correlative Studies into Early Clinical Trials

- Feasibility and Reproducibility in Biomarker Studies
- Statistical Aspects of Correlative Studies
- "Omics" Platform Considerations in Clinical Trials
- Lecture Session 4
  - Imaging in the Era of Targeted Therapy
  - Taking Your Product to Clinic
  - Barriers to Accrual and How to Overcome Them
- Protocol Development Group Session 3
- Lunch
- Small Group Discussion Sessions 1-2
  - Choosing Appropriate Clinical Endpoints
  - Imaging Endpoints in Clinical Trials
- Small Group Discussion Sessions 3-4
  - Inflammatory and Immunologic Biomarkers
  - Laboratory Correlates
- Independent Study Time / Dinner on Own
- ASSIGNMENT DUE: STATISTICAL SECTION

### Tuesday, July 28

- Continental Breakfast
- Lecture Session 5
  - Navigating the Clinical Trials System and Clinical Trials in the Cooperative Groups
  - Quality of Life and Survivorship
- Break / Group Photo
- Protocol Development Group Session 4
- Lunch
- Small Group Discussion Sessions 5-7
  - First in Human Trials: Understanding the Regulatory Process
  - Written and Oral Research Presentations That Audiences Remember
  - Work-Life Balance (for Both Men and Women)
- Office Hours 2: Protocol Related Questions
- Independent Study Time / Dinner on Own
- ASSIGNMENT DUE: PROTOCOL DRAFT

#### Wednesday, July 29

- Continental Breakfast
- Lecture Session 6
  - Everything You Need to Know about Clinical Trial Ethics and Informed Consent: A Case History
  - You're in Charge: Investigator Responsibilities
- Lecture Session 7

- Panel Discussion: Special Populations in Clinical Trials
- Protocol Development Group Session 5
- Lunch
- Special Interest Group and Career Sessions
  - Surgery and Surgery-related Specialties
  - Radiation Oncology
  - Pediatric Oncology
  - Becoming a Phase I Investigator
- Office Hours 3: Any Questions or Topics, including Career Related
- Independent Study Time / Dinner on Own
- Optional Session
  - Women in Academia
- ASSIGNMENT DUE: INFORMED CONSENT DOCUMENTS

### Thursday, July 30

- Continental Breakfast
- Lecture Session 8
  - Career Development Opportunities: The Government Can Help
  - Successful Grantsmanship
- Protocol Development Group Session 6
- Administration of Postworkshop test
- Lunch
- Independent Study Time
- ASSIGNMENT DUE: FINAL PROTOCOL, INFORMED CONSENT DOCUMENTS, WORKSHOP EVALUATIONS
- Reception / Banquet / Dance

#### Friday, July 31

• Continental Breakfast

Departure

# S3: Student Affiliations (1996-2014)

Abbott Biotherapeutics
Abramson Cancer Center of the University of
Pennsylvania
Abraxis BioScience, Inc.
Aichi Cancer Center Research Institute
Alaska Native Tribal Health Consortium
Albany Medical College
Albert Einstein College of Medicine
Amgen, Inc.
Arizona Oncology
AstraZeneca
Atlantic Health Carol G. Simon Cancer Center
Aton Pharma, Inc.
Avera Cancer Institute
Barbara Ann Karmanos Cancer Institute
Barts & The London School of Med. & Dent.
Baylor College of Medicine
BC Cancer Agency
Belfast City Hospital
Beth Israel Deaconess Medical Center
Billings Clinic
Brigham & Women's Hospital
Bristol-Myers Squibb Co.
Brooke Army Medical Center
Brown University Medicine School
Cabrini Center for Cancer Care
California Pacific Medical Center
Cancer Center of North Carolina
Cancer Center of the Rockies
Cancer Research Center of Hawaii
Cancer Therapy & Research Center
Case Comprehensive Cancer Center
Case Western Reserve University
Cedars-Sinai Medical Center
Celgene Corporation
Centro Nacional de Investigaciones Oncológicas
(CNIO)
Children's Hospital & Research Medical Center
Children's Hospital Colorado
Children's Hospital Los Angeles
Children's Hospital of Philadelphia
Children's National Medical Center
Chinese University of Hong Kong

Christian Medical College, Vellore
Christie Hospital
Cincinnati Children's Hospital Medical Center
City of Hope National Medical Center
Colorado Integrative Cancer Care
Columbia University Medical Center
Creighton University Medical Center
Cross Cancer Institute
Crozer-Keystone Health System
Daiichi-Sankyo, Inc.
Dana-Farber Cancer Institute
Dartmouth-Hitchcock Medical Center
Duke University Medical Center
East Tennessee State University
Eli Lilly and Company
Emory University Winship Cancer Institute
Erasmus Medical Center
Everett Clinic
Every
FDA-CDER
Flinders Medical Center
Fox Chase Cancer Center
Franklin Square Hospital Center
Fred Hutchinson Cancer Research Center
FUNDALEU
Garvan Institute of Medical Research
Genentech, Inc.
Georgetown Lombardi Comp. Cancer Center
Georgia Health Sciences University
German Cancer Research Center (DKFZ)
Good Samaritan Medical Center
H. Lee Moffitt Cancer Center & Res. Institute
Harbor-UCLA/City of Hope
Harvard Medical School
Harvard Medical School/Massachusetts General
Hospital
Henry Ford Health System
Hillel Yaffe Medical Center
Hospital Geral de Fortaleza
Imclone Syst Int Gmbh
Indiana University
Indiana University Simon Cancer Center
Indiana University-Purdue University Indianapolis
Instituto Nacional de Cancerología
IU Health Goshen Center for Cancer Care
Jacobi Medical Center

Jamia Millia Islamia	New York University Langone Medical Center
Jennerex Biotherapeutics	New York University School of Medicine
John Wayne Cancer Institute	Northwestern University Lurie Comp. Cancer Center
Johns Hopkins Kimmel Comp. Cancer Center	Norton Radiation Onc.
Johnson & Johnson Pharmaceutical R&D	Novartis
Kaiser Permanente	NRG Oncology
King Abdulaziz Medical City	Ochsner Medical Center
Levine Cancer Institute, Carolinas HealthCare	Ohio State University Comprehensive Cancer Center
System	OHSU Knight Cancer Institute
Liverpool Hospital	Oncology and Therapeutic Development
Loma Linda University Medical Center	Oncology Center Antwerp (OCA)
Louisiana State University Health Sciences Center	Oregon State University
Loyola University Medical Center	Ottawa Hospital Regional Cancer Center
LSU Feist-Weiller Cancer Center	Penn State University Hershey Medical Center
Lucile Packard Children's Hospital	Pfizer Oncology
Magee-Womens Hospital	Pfizer, Inc.
Maidstone General Hospital	PharmaMar S.A.
Marshall University School of Medicine	PharmaNet
Massachusetts General Hospital Cancer Center	PMK Consulting
Mayo Clinic	PRA International
Mayo Clinic Arizona	Providence Cancer Center
Mayo Clinic Florida	Queens Long Island Medical Group
McGill University	R&D Systems, Inc.
McMaster University Juravinski Cancer Center	Rainbow Babies & Children Hospital
Medical College of Wisconsin	Rockefeller University
Medical Oncology and Hematology Assoc.	Roswell Park Cancer Institute
Medical University of Gdansk	Royal Brisbane Hospital
Medical University of SC Hollings Cancer Center	Rush University Medical Center
Medical University of South Carolina	Rutgers R.W. Johnson Medical School
Melbourne Blood Specialists	Saint Louis University
Memorial Sloan Kettering Cancer Center	San Diego Healthcare System
Merck Research Laboratories	Sanofi-Aventis Oncology
Michigan State University	Sarah Cannon Research Institute
Montefiore Medical Center	Scottsdale Healthcare/TGEN
Mount Auburn Hospital	Searle
Mount Sinai Medical Center	Seattle Genetics, Inc.
Mundipharma International Ltd	Sinai Hospital
National Cancer Center Hospital (Japan)	Southeast Gynecologic Oncology Assoc.
National Cancer Center Singapore	Spring Bioscience/Ventana
National Cancer Institute U.S.	St. George Hospital
National Cancer Institute Canada	St. Jude Children's Research Hospital
National Cancer Institute Naples	St. Luke's Hospital & Health Network
Nemrock, Cairo University School of Medicine	St. Mary's Hospital
Neurohealth Center	St. Michael's Hospital
New England Medical Center	Stanford Group Company
New York Medical College	Stanford University School of Medicine
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START
State University of New York
Stony Brook University
SUNY University Hospital & Medical Center
Sydney Cancer Center
Tata Cancer Hospital
TetraLogic Pharmaceuticals
Texas Children's Cancer Center
Texas Oncology, PA
Texas Tech University Health Sciences Center
TGen/Virginia G Piper Cancer Center
The Christie NHS Foundation Trust
The Cleveland Clinic Taussig Cancer Institute
The Rockwood Clinic
The Royal Marsden Hospital
Therakos Johnson & Johnson
Thomas Jefferson University Kimmel Cancer Center
Toledo Radiation Oncology, Inc.
Tulane University Medical Center
U.S. Army Medical Res. & Materiel Command
U.S. Oncology
UC Davis Cancer Center
UCLA David Geffen School of Medicine
UCLA Jonsson Comprehensive Cancer Center
UCSD
UCSD Moores Cancer Center
UCSF Comprehensive Cancer Center
UMDNJ-The Cancer Institute of New Jersey
UNC Lineberger Comp. Cancer Center
University Health Network Ontario Cancer Institute
University Health Network Princess Margaret
Hospital
University Hospitals of Cleveland
University Hospitals/ Seidman Cancer Center
University Medical Center Groningen
University of Alabama Comp. Cancer Center
University of Arizona Cancer Center
University of British Columbia
University of California Irvine
University of California, Los Angeles
University of Chicago Medical Center
University of Cincinnati
University of Colorado Denver
University of Colorado/Children's Hospital Colorado
University of Florida
University of Illinois at Chicago

University of Iowa Holden Comp. Cancer Center
University of Kansas Medical Center
University of Kentucky Markey Cancer Center
University of Louisville Brown Cancer Center
University of Maryland Greenebaum Cancer Center
University of Miami
University of Michigan Comp. Cancer Center
University of Minnesota Masonic Cancer Center
University of Nebraska Medical Center
University of New Mexico - Albuquerque
University of North Carolina Chapel Hill
University of Nottingham
University of Oklahoma Health Sciences Center
University of Pennsylvania
University of Rochester Cancer Center
University of Rochester Medical Center
University of South Alabama Mitchell Cancer
Institute
University of Toronto Hospital for Sick Children
University of Utah Huntsman Cancer Institute
University of Vermont College of Medicine
University of Virginia
University of Washington
University of Washington Medical Center
University of Western Ontario
University of Wisconsin Comprehensive Cancer
University Surgical Associates
LIPMC Cancer Center
USC Keck School of Medicine
USC Children's Hospital Los Angeles
USC Norris Comprehensive Cancer Center
UT Health Science Center at Houston
UT Health Science Center at Nousion
UT MD Anderson Conson Conter
UT Modicel Prench
UT Southwastern Madical Contar
VA Madical Cantan Albana
V A Medical Center Albany
Vall d'Hebron University Hospital
vanderollt-Ingram Cancer Center
vermont Cancer Center
Virginia Commonwealth University
VU University Medical Center
Wake Forest University Baptist Medical Center
Walter Reed Army Medical Center
Washington Cancer Institute

Washington Hospital Center
Washington University Siteman Cancer Center
Waterford Regional Hospital
Weill Cornell Medical College of Cornell University
West Virginia University Randolph Cancer Center
Westmead Hospital
Weston Park Hospital

Wilford Hall Medical Center

William Beaumont Hospital
Women & Infants Hospital
Yale Cancer Center
Yale Oncology Fellowship Program
Yale-New Haven Hospital
Yonsei University Cancer Center

#### S4: Faculty History (1996-2014)

(Affiliations listed were current during the years of service)

James L. Abbruzzese, MD, FACP (1999-2006) The University of Texas MD Anderson Cancer Center Specialty Preferences: Clinical study and treatment of pancreatic cancer

Alex A. Adjei, MD, PhD (2004-2008) Roswell Park Cancer Institute State University of New York at Buffalo Specialty Preferences: Phase I clinical trials; biomarkers; lung cancer; pharmacogenetics; clinical trials in special populations

David S. Alberts, MD (1996-1998) University of Arizona Cancer Center Specialty Preferences: Clinical trials in cancer prevention, biomarkers, chemoprevention

James R. Anderson, PhD. (1998, 2000) University of Nebraska Medical Center Specialty Preferences: Study design, biostatistics

Kenneth C. Anderson, MD (2007) Dana-Farber Cancer Institute Harvard Medical School Specialty Preferences: Multiple myeloma; drug development; translational research

Karen H. Antman, MD (2003) Boston University, School of Medicine Specialty Preferences: Breast cancer; mesotheliomas; sarcomas

Jean-Pierre Armand, MD (1997-1998) Institut de Cancérologie Gustave Roussy Specialty Preferences: Early drug development; Phase I and II clinical trials; novel anticancer agents

Carlos L. Arteaga, MD (2005) Vanderbilt-Ingram Cancer Center Breast Cancer Program Vanderbilt University School of Medicine Vanderbilt-Ingram Comprehensive Cancer Center Specialty Preferences: Oncogenes and breast tumor initiation and progression; targeted therapies and biomarkers of drug action and resistance; clinical trials in breast cancer

Steven D. Averbuch, MD (2002- 2003)S.D. Averbuch Consulting, LLCSpecialty Preferences: Biochemical modulators of therapy and toxicity; biomarkers; clinical trials in solid tumors; diagnostic techniques

Rochelle Bagatell, MD (2004-2008) University of Arizona Specialty Preferences: Neuroblastoma; early phase clinical trials in pediatrics; pediatric cooperative group studies Joseph S. Bailes, MD (2006) US Oncology Specialty Preferences: Clinical practice; legislation, public policy, and advocacy

Dean F. Bajorin, MD (2006) Memorial Sloan Kettering Cancer Center Specialty Preferences: Bladder cancer; testicular cancer; renal cell cancer; prostate cancer

Charles M. Balch, MD (2000-2005) American Society of Clinical Oncology Johns Hopkins Medical Institutions Specialty Preferences: Melanoma and breast cancer research; tumor immunology and human T lymphocyte differentiation

Lodovico Balducci, MD (2001-2003) Moffitt Cancer Center Specialty Preferences: Geriatric oncology; clinical trials in solid tumors; combined modalities of therapy

Gary J. Becker, MD (2005) National Cancer Institute Specialty Preferences: Vascular and interventional radiology

Stacey L. Berg, MD (2010-2014) Texas Children's Cancer Center Specialty Preferences: Pediatric oncology; biomedical ethics; palliative care

Monica M. Bertagnolli, MD (2006-2007) Harvard University Medical School Dana-Farber Cancer Institute Brigham and Women's Hospital Specialty Preferences: Surgical oncology; colorectal cancer pathogenesis; familial colon cancer; colorectal cancer prevention trials

Michael R. Bishop, MD (2011-2013) University of Chicago, Department of Medicine Specialty Preferences: Clinical transplantation therapy; novel clinical trials in allogeneic and autologous stem cell transplantation; B cell malignancies, T cell reconstitution

A. William Blackstock, MD, (2007-2012) Wake Forest University School of Medicine Specialty Preferences: Phase I/II chemoradiation trial design; preclinical chemoradiation trials; chemoradiation translational studies

Douglas W. Blayney, MD (2008) Stanford Cancer Center Specialty Preferences: Oncology; quality of life; informatics; hematological malignancies W. Archie Bleyer, MD (1998, 2000)
The University of Texas MD Anderson Cancer Center
Specialty Preferences: Clinical trials in leukemias and lymphomas; clinical trials in solid tumors; combined modalities of therapy; drug delivery systems; drug evaluation methodology

James M. Boyett, PhD (1998, 2000, 2002) St. Jude Children's Research Hospital Specialty Preferences: Biostatistics; pediatric brain tumors

Thomas M. Braun, PhD, MS, BBA (2013-2014) University of Michigan School of Public Health Specialty Preferences: Adaptive phase I trial designs in oncology; multivariate longitudinal modeling of periodontal outcomes

Otis W. Brawley, MD (1996-1998) National Cancer Institute Specialty Preferences: Clinical trials; breast cancer; epidemiology

Dean E. Brenner, MD (2000) University of Michigan Specialty Preferences: Pharmacodynamics of anticancer drugs; chemoprevention

Diane A. Bronzert , MD (1998) National Cancer Institute Specialty Preferences: Brain and breast cancers; drug and multidrug resistance; phase I-II studies

Paul A. Bunn, MD (2002) University of Colorado Specialty Preferences: Lung and Thoracic Cancers

Gregory Burke (1996, 1998) Novartis Specialty Preferences: Drug development

Michael A. Caligiuri, MD (2003-2007) The Ohio State University Specialty Preferences: Immunology; leukemia and lymphoma; Epstein-Barr virus

Rick J. Chappell, PhD (2007-2014) University of Wisconsin Specialty Preferences: Statistics; goals and aims (primary/secondary); power/sample size calculations; analysis plans; noninferiority trials; radiation oncology; phase I clinical trial design

Michaele C. Christian, MD (2003) National Institutes of Health Specialty Preferences: NCI Clinical Trials Monitoring Branch

Edward Chu, MD (2014) University of Pittsburgh School of Medicine Specialty Preferences: Clinical trials/biostatistics/epidemiology; gastrointestinal cancer; general oncology; cancer education Gary M. Clark, PhD (1996-2001, 2010-2011) Array BioPharma, Inc. Specialty Preferences: Identification and evaluation of prognostic and predictive biomarkers for breast cancer, lung cancer, and other solid tumors; design and analysis of phase I, II, and III clinical trials of targeted therapies

Neil J. Clendeninn, MD, PhD (2004-2008) CANAID, Inc. Specialty Preferences: Clinical drug development; clinical pharmacology; regulatory issues; industrybiotech/pharma

Laura Cleveland (2008, 2010-2013) The Ohio State University Specialty Preferences: Patient advocacy; trial design; patient accrual issues; regulatory mandates and informed consent

Jerry M. Collins, PhD (1996-1997) National Cancer Institute Specialty Preferences: Clinical pharmacology

Deborah Collyar (2002) Patient Advocates in Research (PAIR) Specialty Preferences: Patient advocacy

Charles A. Coltman, MD (1996-1998, 2000-2001) The University of Texas Health Science Center. at San Antonio Specialty Preferences: Breast cancer; combined modalities of therapy; drug evaluation

John J. Crowley, PhD (1996-1998, 2000) Cancer Research and Biostatistics Specialty Preferences: Biostatistics

Ramzi N. Dagher, MD (2003) Pfizer, Inc. Specialty Preferences: Sarcomas and soft tissue cancers; clinical trials in solid tumors; combined modalities of therapy; study design and methods

Otilia Dalesio, M.Sc (1998, 2000, 2003) Netherlands Cancer Institute Specialty Preferences: Multicenter trials; lung and colorectal cancers

Mary B. Daly, MD, PhD, FACP (2002) Fox Chase Cancer Center Specialty Preferences: Clinical genetics; breast cancer; risk assessment; chemoprevention research; family risk assessment and genetic testing; epidemiology; gynecologic cancers

Gwen Darien (2003-2005) American Association for Cancer Research Specialty Preferences: Patient advocacy and engagement; community engagement in genomics Christopher K. Daugherty, MD (1998, 2000) University of Chicago Specialty Preferences: Ethical issues in cancer care and clinical research; stem cell transplantation; leukemia

Angela M. Davies, MD, FRCP(C) (2009-2010) OSI Pharmaceuticals Specialty Preferences: Medical oncology; small molecule therapeutics; lung cancer

Thomas Davis, MD (1997) Celldex Therapeutics Specialty Preferences: Drug development; immune-oncology

Angelo M. De Marzo, MD, PhD (2009-2014) Johns Hopkins University School of Medicine Specialty Preferences: Pathology; prostate cancer; biomarkers

Patricia Delaney (2001) U.S. Food and Drug Administration Specialty Preferences: Public education about and inclusion of cancer survivors; cancer in drug development

Peggy Devine (2007) University of California San Francisco Specialty Preferences: Writing a quality consent form; retention strategies; accrual strategies; community/minority outreach

Mary L. (Nora) Disis, MD (2001-2004, 2007-2012) Washington School of Medicine Fred Hutchinson Cancer Research Center Specialty Preferences: Breast and ovarian cancer; tumor immunology; translational oncology; biomarkers

Chaitanya R. Divgi, MD (2011-2012) Columbia University Specialty Preferences: Renal cancer; ovarian cancer; antibodies in cancer imaging and therapy; PET; radiochemistry

Everett D. Dodson (2013-2014) Howard University Cancer Center Specialty Preferences: Colon cancer; prostate cancer; patient and research subject advocacy

Afshin Dowlati, MD (2013-2014) Case Western Reserve University Specialty Preferences: Experimental therapeutics; hematology/oncology; lung cancer; phase I clinical trials; thoracic cancer

Brian J. Druker, MD (2004) OHSU Knight Cancer Institute Specialty Preferences: Leukemia; precision medicine; targeted therapies John R. Durant, MD (1997-1998) American Society of Clinical Oncology Specialty Preferences: Translational and clinical research; combination chemotherapy

Janice J. Dutcher, MD (1996-1998, 2000-2002) St. Luke's-Roosevelt Hospital Center. Specialty Preferences: Clinical trials; renal cell cancer; immunotherapy

S. Gail Eckhardt, MD (2000-2003, 2006-2014) University of Colorado Cancer Center Specialty Preferences: Early clinical trials; colorectal cancer research; preclinical and clinical development of molecularly targeted compounds; melanoma; predictive biomarkers

Merrill J. Egorin, MD, FACP (1996-2004, 2007-2009) University of Pittsburgh School of Medicine Specialty Preferences: Clinical pharmacology; phase I trials; pharmacokinetics; cancer in the elderly

Lawrence Einhorn, MD (2000) Indiana University School of Medicine Specialty Preferences: clinical trials and supportive care for lung and testicular cancers

Lee M. Ellis, MD (2007-2012) The University of Texas MD Anderson Cancer Center Specialty Preferences: Biomarkers; targeted therapies; angiogenesis; surgical oncology: GI cancer

Ezekiel J. Emanuel, MD (2001-2004) National Institutes of Health Specialty Preferences: Medical ethics; breast oncology

Charis Eng, MD, PhD (2001) The Cleveland Clinic Specialty Preferences: Genomic medicine

Elihu H. Estey, MD (2002) Fred Hutchinson Cancer Research Center Specialty Preferences: Trial design that can monitor multiple outcomes; acute myeloid leukemia

Martine Extermann, MD, PhD (2006) University of South Florida H. Lee Moffitt Cancer Center Specialty Preferences: Geriatric oncology; comprehensive assessment/multivariate analyses/outcome; breast cancer; AML in the elderly (clinical); comorbidity; decision analysis

John M. Falleta, MD (2003) Duke University Medical Center Specialty Preferences: Pediatric oncology

Ann T. Farrell, MD (2004-2005)U. S. Food and Drug AdministrationSpecialty Preferences: FDA regulations; endpoints; hematologic malignancies; trial design

David W. Feigal, MD (1996-1997) U. S. Food and Drug Administration Specialty Preferences: Drug development

Ellen G. Feigal, MD 1997) California Institute for Regenerative Medicine Specialty Preferences: Clinical study design; drug development

Scott Z. Fields, MD (2004-2005) EISAI Columbia University College of Physicians and Surgeons NY Presbyterian Hospital Specialty Preferences: Hematology and oncology disorders

Robert A. Figlin, MD (1998, 2000) Cedars-Sinai Medical Center Specialty Preferences: Kidney cancers; translational research; clinical trials

Lawrence H. Fong, MD (2010-2014) University of California San Francisco Specialty Preferences: Prostate cancer; cancer immunotherapy; kidney cancer; melanoma; immunologic biomarkers

Margaret Foti, PhD, MD (hc) (1998-2014) American Association for Cancer Research (AACR) Specialty Preferences: Nonprofit association management; scholarly publishing; scientific communications; science policy and government affairs; foundation support; research administration

Wilbur A. Franklin, MD (2006-2008) University of Colorado Denver Health Sciences Center Specialty Preferences: Pathobiology; biomarkers; lung cancer

Patricia A. Ganz, MD (2002, 2005) UCLA Schools of Medicine and Public Health Jonsson Comprehensive Cancer Center Specialty Preferences: Cancer survivorship; late effects of cancer treatment; care quality of cancer patients; cancers in the elderly

Elizabeth Garrett-Mayer, PhD (2002-2004, 2006-2014) Medical University of South Carolina Hollings Cancer Center Specialty Preferences: Biostatistics; predictive models; phase I and phase II study designs; power and sample size

Robert A. Gatenby, MD (2006-2007) H. Lee Moffitt Cancer Center & Research Institute Specialty Preferences: Evolutionary dynamics in cancer and cancer drug resistance

Constantine A. Gatsonis, PhD (2005) Brown University Specialty Preferences: Development of statistical methods for diagnostic tests and biomarker evaluation; design and analysis of clinical trials of diagnostic and screening modalities Stephen George, PhD (1997-1998, 2000, 2003) Duke University Medical Center Specialty Preferences: Statistical issues related to the design, conduct, and analysis of clinical trials

Stanton L. Gerson, MD (2003) Case Comprehensive Cancer Center Specialty Preferences: DNA repair and stem cell therapy

Jorge Gomez, MD, PhD (2003) University of Arizona Health Sciences Center Specialty Preferences: Community-based research; cancer health disparities.

Mithat Gönen, PhD (2007-2012) Memorial Sloan Kettering Cancer Center Specialty Preferences: Clinical trials with imaging endpoints; surgical oncology clinical trials; staging and prognosis; general biostatistics

Steven Goodman, MD, PhD (1998) Johns Hopkins University Specialty Preferences: Reproducibility and efficiency of biomedical research

David J. Gordon, MD (2000) University of Iowa Specialty Preferences: Ewing sarcoma; pediatric oncology

Gary B. Gordon, MD, PhD (2000) Ayala Pharmaceuticals Specialty Preferences: Drug development

Lester S. Gorelic, PhD (1998, 2001-2009) National Cancer Institute Specialty Preferences: Support for cancer research training and career development

Sylvan B. Green, MD (1999-2005) University of Arizona Arizona Cancer Center Specialty Preferences: Design and analysis of clinical trials; prevention trials, epidemiologic studies; biostatistical and computer methodology application

Charles K. Grieshaber (1996) Oncology Research & Regulatory Consulting Associates (ORRCA), Inc. Specialty Preferences: Pharmacokinetics; drug evaluation

Louise B. Grochow, MD (2000-2002) AstraZeneca Specialty Preferences: Drug development

Susan G. Groshen, PhD (2001-2004, 2006) University of Southern California Keck School of Medicine USC/Norris Comprehensive Cancer Center Specialty Preferences: Biostatistics; urology/bladder cancer; phase I trials

Stuart A. Grossman, MD (2008-2014) Johns Hopkins University Specialty Preferences: Primary brain tumors; CNS metastases; cancer pain; clinical trial design and conduct

Stephen J. Gwyther, MD (2003) East Surrey Hospital Specialty Preferences: Clinical radiology, imaging

Stephen M. Hahn, MD (2004-2008) University of Pennsylvania Hospital Specialty Preferences: Radiation oncology; non-small cell lung cancer; combined chemoradiotherapy; prostate cancer

Lyndsay N. Harris, MD (2013-2014) Case Western Reserve University Specialty Preferences: Breast cancer; cancer prevention; psychosocial issues/psychiatry; tumor biology/molecular biology/growth factors/cytokines

Ernest T. Hawk, MD, MPH (2008-2009) The University of Texas MD Anderson Cancer Center Specialty Preferences: Cancer prevention; GI oncology; cancer screening

Daniel F. Hayes, MD (2010) University of Michigan Comprehensive Cancer Center Specialty Preferences: Breast cancer; clinical trials; new therapies and tumor markers; translational and clinical studies of breast

Daniel F. Heitjan, PhD (2004-2007) Abramson Cancer Center University of Pennsylvania Specialty Preferences: Biostatistics; quality of life; clinical trials; health economics

Glenn Heller, PhD (2002- 2003) Memorial Sloan Kettering Cancer Center Specialty Preferences: Memorial Sloan Kettering Cancer Center

Roy S. Herbst, MD, PhD (1998-2000, 2007-2008) The University of Texas MD Anderson Cancer Center Specialty Preferences: Lung cancer; phase I trials

Kenneth R. Hess, PhD (2001-2005) The University of Texas MD Anderson Cancer Center Rice University Specialty Preferences: Biostatistics Manuel Hidalgo, MD, PhD (2005-2008) CEU Universidad San Pablo Centro Integral de Oncologia "Clara Campal" Centro Nacional de Investigaciones Oncológicas Specialty Preferences: Phase I trials; targeted agents; pharmacodynamics

Bruce J. Hillman, MD (2004-2007) University of Virginia Specialty Preferences: Diagnostic imaging; screening; multicenter trials

Bruce E. Hillner, MD Virginia Commonwealth University Specialty Preferences: Outcomes research in oncology; breast cancer

Susan G. Hilsenbeck, PhD (1998-2007) Baylor College of Medicine Specialty Preferences: Biostatistics; breast cancer; clinical trials; genomics and expression arrays

Waun Ki Hong, MD The University of Texas MD Anderson Cancer Center Specialty Preferences: Head and neck cancers; targeted therapy; chemoprevention

Ivan D. Horak, MD, FACP (2004-2007) Enzon Pharmaceuticals, Inc. Specialty Preferences: Antisense, siRNA, and micro RNA; monoclonal antibodies; role of biomarkers in drug development; signal transduction pathways as a target for therapy

Sandra J. Horning, MD (2005, 2010-2013) Genentech, Inc. Specialty Preferences: Drug development

Susan Band Horwtiz, PhD (2002) Albert Einstein College of Medicine Specialty Preferences: Development of drugs derived from natural products; drug resistance

S. Percy Ivy, MD (2013-2014) National Cancer Institute Specialty Preferences: Early drug development; clinical trial design; regulatory affairs including IND, IDE, and adverse event reporting; molecular characterization and biomarker development; experimental therapeutics: antiangiogenesis; embryonic signaling; targeted kinases; protein processing

Mary Jackson Scroggins (2012-2014) In My Sister's Care Specialty Preferences: Patient advocacy; gynecologic cancer

C. Carl Jaffe, MD, FACC (2004-2005) Yale University School of Medicine Specialty Preferences: Diagnostic radiology; nuclear medicine

Elizabeth M. Jaffee, MD (2005, 2008-2009) Sidney Kimmel Comprehensive Cancer Center Johns Hopkins University Specialty Preferences: Immunotherapy; pancreatic cancer therapy; biomarker clinical development

Steven Joffe, MD, MPH (2005-2009) Harvard Medical School Dana-Farber Cancer Institute Children's Hospital Specialty Preferences: Pediatric oncology; research ethics; clinical trial design; hematopoietic stem cell transplantation

Bruce E. Johnson, MD (2008) Dana-Farber Cancer Institute Specialty Preferences: Targeted treatment of thoracic malignancies

David H. Johnson MD (2004) UT Southwestern Medical Center Specialty Preferences: Developmental therapeutics; lung cancer therapeutics

Roy B. Jones PhD, MD (2014) MD Anderson Cancer Center Specialty Preferences: Bone marrow transplantation (autologous/allogeneic/stem cell)

Robert L. Justice, MD U. S. Food and Drug Administration Specialty Preferences: Oncology drug products; drug evaluation and research

Judith E. Karp, MD (2008-2012) Sidney Kimmel Comprehensive Cancer Center Johns Hopkins University Specialty Preferences: Acute leukemias; signal transduction; cell kinetics

Kay F. Kays (2014) Pancreatic Cancer Action Network Specialty Preferences: GI/Pancreatic cancer; survivorship; patient advocacy; quality of life; patient accrual

M. Margaret Kemeny, MD, FACS (1996-1998, 2006)
Queens Cancer Center of Queens Hospital
Mt. Sinai School of Medicine
Specialty Preferences: GI cancers; hepatobiliary tumors; surgical oncology; cancer in the elderly

Larry G. Kessler, ScD (2010) University of Washington School of Public Health Specialty Preferences: Cost effectiveness and diagnostic value of medical technology in screening for cancer and other diseases

KyungMann Kim, PhD (2004-2006, 2008) University of Wisconsin Comprehensive Cancer Center Specialty Preferences: Developmental therapeutics; lung cancer; chemoprevention; biostatistics Mark D. Krailo, PhD (2006-2014) University of Southern California Specialty Preferences: Statistical design; data collection; interim monitoring

Larry E. Kun, MD (1997, 2006-2008) St. Jude Children's Research Hospital The University of Tennessee College of Medicine Specialty Preferences: Radiation oncology; neuro-oncology; pediatric oncology; late effects/pediatric cancer

Brenda F. Kurland, PhD (2012-2014) University of Pittsburgh Specialty Preferences: Quantitative imaging biomarkers; analysis of correlated data; early-phase cancer clinical trials

Daniel H. Laheru, MD (2010-2014) Sidney Kimmel Comprehensive Cancer Center Johns Hopkins University Specialty Preferences: Gastrointestinal oncology, specifically pancreatic and colon cancers

Kathleen R. Lamborn, PhD (2001-2003) University of California San Francisco Specialty Preferences: Biostatistics; glioma

Quynh-Thu Le, MD (2008) Stanford Cancer Center Specialty Preferences: Radiation oncology; head and neck cancers

J. Jack Lee, PhD (1998, 2000) The University of Texas MD Anderson Cancer Ctr. Specialty Preferences: Biostatistics; multiple biomarkers and adaptive designs to develop more efficient and ethical clinical trials

Ming Lei, PhD (2010-2011) Center for Cancer Training National Cancer Institute Specialty Preferences: National management of grant-supported research training; DNA replication and chromatin assembly; training and career development of cancer scientists

Steven J. Lemery, MD (2008)U. S. Food and Drug AdministrationSpecialty Preferences: Review products for gastrointestinal malignancies, melanoma, and other advanced skin cancers and sarcomas

Barbara K. LeStage, MHP (2011-2014) Specialty Preferences: Patient advocacy; breast cancer; clinical imaging

Allen S. Lichter, MD (2007) American Society of Clinical Oncology Specialty Preferences: Radiation oncology Stuart M. Lichtman, MD, FACP (2004-2005, 2007-2009) Memorial Sloan-Kettering Cancer Center Specialty Preferences: Gynecologic oncology; geriatric oncology

Michael P. Link, MD (2001-2003) Stanford University School of Medicine Specialty Preferences: Pediatric oncology; stem cell transplant

David A. Lipschitz, MD (2001-2003) University of Arkansas for Medical Sciences Specialty Preferences: Gerontology

Scott M. Lippman, MD (1996-2005) The University of Texas MD Anderson Cancer Center Specialty Preferences: Clinical and translational research: head, neck, and lung cancers; genetic drivers of cancer; predictive molecular signatures; biomarkers for clinical response in solid tumors; design of trials using molecular targets and markers for cancer prevention and therapy

Alan F. List, MD (2000-2003, 2006)H. Lee Moffitt Cancer Center and Research InstituteUniversity of South FloridaSpecialty Preferences: Clinical studies; leukemia; myelodysplastic syndromes

John H. Littlefield, MD (2000) The University of Texas Health Science Center at San Antonio Specialty Preferences: Health profession education, evaluation

Albert F. Lobuglio, MD (1997) UAB Comprehensive Cancer Center Specialty Preferences: Immunological therapeutics

Wendy B. London, PhD (2011-2014) Harvard University Specialty Preferences: Neuroblastoma; prognostic stratification; clinical trial design

Patricia M. LoRusso, DO (2004-2005, 2009-2012) Wayne State University Karmanos Cancer Institute Specialty Preferences: Phase I; breast cancer

Michael T. Lotze, MD (2004-2008, 2013-2014) University of Pittsburgh Specialty Preferences: Biologic therapy/immunobiology; gastrointestinal cancer; liver and pancreas tumors; melanoma/skin cancer

H. Kim Lyerly, MD (1997-1998) Duke University School of Medicine Specialty Preferences: Applied therapeutics; immunology; surgery

James C. Lynch, PhD (2001-2002, 2004-2006) Eppley Cancer Center University of Nebraska Medical Center Specialty Preferences: Biostatistics; childhood cancer; clinical trials; cancer survivorship; non-Hodgkin lymphoma; stem cell transplantation

David A. Mankoff, MD, PhD (2007) University of Washington Specialty Preferences: Cancer imaging; breast cancer; nuclear medicine/PET; cancer biomarkers

Guido G. Marcucci, MD (2009-2010) The Ohio State University Comprehensive Cancer Center Specialty Preferences: Molecular biology and cancer genetics; experimental therapeutics; acute myeloid leukemia; chronic lymphocytic leukemia

Lynn M. Matrisian, PhD (2004) Pancreatic Cancer Action Network Specialty Preferences: Pancreatic cancer, basic science

Mary S. McCabe, RN, MA (2004-2006) Memorial Sloan Kettering Cancer Center Specialty Preferences: Informed consent; specimen collection; quality of life research; patient-centered endpoints

W. Gillies McKenna, MD, PhD (1998) University of Oxford Specialty Preferences: Radiation oncology; treatment resistance

Neal J. Meropol, MD (2007-2014) Fox Chase Cancer Center (2007-2008) Case Western Reserve University (2009-2014) Specialty Preferences: Developmental therapeutics; gastrointestinal cancer; liver and pancreas tumors; medical oncology; physician-patient communication

Rosemarie Mick, MS (1997-1998, 2000, 2003) University of Pennsylvania Perelman School of Medicine Specialty Preferences: Biostatistics; clinical trial methodology, optimal designs for early-stage cancer trials, and the development of immunotherapies

Carol M. Moinpour, PhD (1996, 2006) Fred Hutchinson Cancer Research Center University of Washington Specialty Preferences: Quality of life assessment in clinical trials; analysis issues for quality of life outcomes; clinically meaningful differences in quality of life scores; outcomes assessment in symptom management trials

Jeffrey F. Moley, MD (1996) Washington University Siteman Cancer Center Specialty Preferences: Endocrine and oncologic surgery; multiple endocrine neoplasia

Wayne L. Monsky, MD, PhD (1997) University of California Davis Cancer Center Specialty Preferences: Diagnostic and interventional radiology; minimally invasive therapies for cancer Vicki A. Morrison, MD (2010-2014) University of Minnesota Specialty Preferences: Lymphoma; non-Hodgkin lymphoma; bone marrow transplantation; infectious complications in oncology patients

Heidi Nelson, MD (2001-2003) Mayo Clinic Specialty Preferences: Colon and rectal cancer surgery

Donna S. Neuberg, ScD (2004-2006) Harvard School of Public Health Dana-Farber Cancer Institute Specialty Preferences: Statistical design and analysis; pediatrics; vaccines/immunotherapy; leukemia and lymphoma

Donna Niedzwiecki, PhD (2001) Duke University Medical Center Specialty Preferences: Biostatistics, clinical trials design, GI cancer, lymphoma, melanoma

Daniel P. Normolle, PhD (2008-2010) University of Pittsburgh Cancer Institute Specialty Preferences: Translational research design; phase I clinical trial design; biomarker analysis

Larry Norton, MD (2001) Mayo Clinic Specialty Preferences: Breast cancer prevention, diagnosis, and treatment

Allen Oliff, MD (1997) GlaxoSmithKline Specialty Preferences: Drug discovery; molecular pathogenesis: RNA tumor viruses, oncogenes, and tumor suppressor genes

Katherine S. Panageas, DrPH (2011-2014) Memorial Sloan Kettering Cancer Center Specialty Preferences: Clinical trial design, melanoma; immunotherapy; neuro-oncology

William Pao, MD (2012) Vanderbilt University Ingram Cancer Center Specialty Preferences: Translational research; lung cancer; mechanisms of sensitivity and resistance to targeted agents in solid tumors

David R. Parkinson, MD (2000-2003) ESSA Pharma, Inc. Specialty Preferences: Oncology drug development

Electra D. Paskett, PhD (2004) The Ohio State University Comprehensive Cancer Center Specialty Preferences: Epidemiology and cancer health disparities; breast cancer; colorectal cancer; uterine, cervical, and endometrial cancers Jyoti D. Patel, MD (2013-2014) Northwestern University Specialty Preferences: Lung cancer

Richard Pazdur, MD (2001-2002) U. S. Food and Drug Administration Specialty Preferences: Oncology product regulation and clinical review

Edith A. Perez, MD (2003-2006, 2009-2014) Mayo Clinic Specialty Preferences: Breast cancer; targeted therapies; translational clinical trials

Jane Perlmutter, PhD (2006-2010) Gemini Group Specialty Preferences: Patient advocates in research; innovative trial design–Bayesian approaches and adaptive trials; improving patient communication and decision making, especially the informed consent process; biomarkers in breast cancer

Gina R. Petroni, PhD (1996-2003, 2005) University of Virginia Cancer Center Specialty Preferences: Design and analysis of clinical trials; data and safety monitoring; vaccine trials

Steven Piantadosi, MD, PhD (1996,1998) Cedars-Sinai Medical Center Specialty Preferences: Biostatistics; multi-center and international trials; innovative clinical trial methods

Tatiana M. Prowell, MD (2014)U. S. Food and Drug AdministrationJohns Hopkins University Cancer CenterSpecialty Preferences: Breast cancer; general oncology

Alfred W. Rademaker, PhD (2006-2014) Northwestern University Specialty Preferences: Early-phase cancer clinical trial designs; sample size determination; statistical methods for diagnostic testing; cluster randomized clinical trials

Gregory H. Reaman, MD (2009-2010) Children's Oncology Group Specialty Preferences: Pediatric phase I and II trials; clinical trials in childhood leukemia; correlative biomarkers in pediatric cancer clinical trials

Michael P. Redden, JD, (2008-2009) Duke Clinical Research Institute Specialty Preferences: Patient advocacy; quality of life issues; informed consent language

Meredith M. Regan, ScD (2008-2014) Dana-Farber Cancer Institute Specialty Preferences: Biostatistics, particularly as applied in breast and genitourinary cancers

Brian I. Rini, MD, FACP (2011-2014) Case Western Reserve University Specialty Preferences: Genitourinary malignancies; renal cell carcinoma; antiangiogenic therapy; immunotherapy

Paula K. Roberson, PhD (2001) University of Arkansas for Medical Sciences Specialty Preferences: Design and analysis of clinical trials; variable selection; nonparametric methods

Lorna Rodrìguez-Rodrìguez, MD, PhD (2001-2006) The Cancer Institute of New Jersey University of Medicine and Dentistry of New Jersey Robert Wood Johnson Medical School Specialty Preferences: Gynecologic oncology; laboratory correlates; multimodality trials

Denise Roe, PhD (2000-2003) University of Arizona Cancer Center Specialty Preferences: Biostatistics and epidemiology; developing and evaluating statistical methods useful in clinical trials; prevention studies; pharmacokinetics; longitudinal studies

Stan Rosenfeld (2000) University of California San Francisco Specialty Preferences: Prostate cancer patient advocacy

David Rosenthal, MD (2000) The University of Texas MD Anderson Cancer Center Specialty Preferences: Radiation oncology; head and neck cancers

Mace L. Rothenberg, MD (2000- 2004) Pfizer, Inc. Specialty Preferences: Oncology drug safety and regulation

Erkki Ruoslahti, MD, PhD (1998) Sanford-Burnham Medical Research Institute Specialty Preferences: Targeted therapy; breast cancer and glioblastoma

John H. Sampson, MD (1998) Duke University Medical Center Specialty Preferences: Brain cancer; neurosurgery

Howard M. Sandler, MD (2008-2009) Samuel Oschin Comprehensive Cancer Institute Cedars-Sinai Medical Center Specialty Preferences: Prostate cancer; brain tumors; bladder cancer; radiation oncology

Richard L. Schilsky, MD (2002-2006) The University of Chicago Pritzker School of Medicine Specialty Preferences: GI cancer; clinical trial endpoints/drug approval process; phase I trials; pharmacology; cooperative groups

Lowell E. Schnipper, MD (2007-2009) Harvard Medical School Beth Israel Deaconess Medical Center Specialty Preferences: Breast cancer clinical trials; non-Hodgkin lymphoma; experimental therapeutics; drug resistance

Gary K. Schwartz, MD (2010) Memorial Sloan Kettering Cancer Center Weill Cornell Medical College Specialty Preferences: Sarcoma; melanoma; new drug development

Karl L. Schwartz, MFA (2011-2014) Patients Against Lymphoma Specialty Preferences: Clinical trial accrual challenges – how patients see trials; eligibility criteria as they relate to enrollment and generalizability of trial results; simplifying the informed consent document; improving readability; biospecimen samples anticipating obstacles to consent

Mary J. Scroggins, MA (2006-2007, 2009-2014) In My Sister's Care Specialty Preferences: Health disparities; community partnerships; vulnerable populations; informed consent; quality of life; international health

Lalitha K. Shankar, MD, PhD (2005) Cancer Imaging Program National Cancer Institute Specialty Preferences: Functional and molecular imaging-diagnosis and treatment of cancer; clinical trial establishment and monitoring to evaluate imaging tracers and techniques to improve prevention, diagnosis, and treatment of cancer

Westley Sholes, MPA (2008-2012) California Prostate Cancer Coalition Specialty Preferences: Patient advocacy issues; health care management; prostate cancer

Yu Shyr, PhD (2004-2014) Vanderbilt University Specialty Preferences: Biostatistics; bioinformatics; clinical trial design

Rache M. Simmons, MD, FACS (2004-2005, 2010) Joan and Sanford Weill Medical College of Cornell University New York Presbyterian Hospital Specialty Preferences: Oncoplastic surgery; breast cancer surgery

Lillian L. Siu, MD, FRPCPC (1998-1999, 2003-2007) Princess Margaret Hospital University of Toronto Specialty Preferences: Head and neck cancer; targeted agents; GI cancer

George W. Sledge, Jr., MD (2011-2014) Stanford University Cancer Center Specialty Preferences: Breast cancer; novel biologic therapies; angiogenesis and antiangiogenic therapy

Franklin O. Smith, III, MD (2011-2014) University of Cincinnati Specialty Preferences: Hematology/oncology; pediatric oncology; hematopoietic stem cell transplantation; acute myeloid leukemia; Fanconi anemia

Anil K. Sood, MD (2009-2014) The University of Texas MD Anderson Cancer Center Specialty Preferences: Gynecologic oncology; novel therapeutics; angiogenesis

Barry E. Storer, PhD (2001-2005) University of Washington Fred Hutchinson Cancer Research Center Specialty Preferences: Clinical trial design; statistical method development to identify the impact of the disease incidence and therapy outcomes

Ellen L. Stovall (2002, 2004-2005) National Coalition for Cancer Survivorship Specialty Preferences: Patient advocacy

Steven Stratton, MD (2005-1009, 2011-2013) University of Arizona Cancer Center Specialty Preferences: Skin cancers; early detection; cancer interception; chemoprevention

Daniel C. Sullivan, MD (2003-2004) Duke University Comprehensive Cancer Center Specialty Preferences: Radiation oncology; clinical evaluation and validation of imaging biomarkers

Vera J. Suman, PhD (2004-2010) Mayo Foundation Specialty Preferences: Biostatistics; clinical trial design; immunologic endpoints; data safety monitoring plans/boards; surrogate endpoints; biomarkers—evaluation of prognostic value

Sandra M. Swain, MD (2012) Washington Cancer Institute Specialty Preferences: Targeted therapeutics for metastatic and inflammatory breast cancer

G. Marie Swanson, PhD (1996-1997)Indiana Univ. School of MedicineSpecialty Preferences: Epidemiology; population diversity in cancer risk; breast cancer

Alphonse G. Taghian, MD, PhD (2010) Harvard Medical School Massachusetts General Hospital Specialty Preferences: Radiation biology; breast radiation oncology; lymphedema; cardiac exposure to radiation

Chris H. Takimoto, MD, PhD (2014) Janssen Research and Development Specialty Preferences: Pharmaceutical/drug development; gastrointestinal cancer

Ian F. Tannock, MD, PhD (1996-1998, 1999-2000, 2002-2003) University of Toronto University Health Network Specialty Preferences: Breast, head and neck, kidney, prostate, and urinary bladder cancers; microenvironmental causes of resistance to drugs

Lisa L. Taylor, MBA (2011-2014) NRG Oncology Specialty Preferences: Uterine cancer; cervical cancer; ovarian cancer; prostate cancer

Margaret Tempero, MD (1998) University of California San Francisco Pancreas Center Specialty Preferences: Investigational therapeutics; antibody-based therapies; pancreatic cancer

Joel E. Tepper, MD (2001-2006) University of North Carolina School of Medicine Specialty Preferences: GI cancers; soft tissue sarcomas; radiation oncology; combined modality therapies

Charles Richard Thomas, Jr., MD (2009-2012) The Oregon Health & Science University Specialty Preferences: Radiation oncology with a thoracic focus; radiation oncology with a GI focus

Andrea B. Troxel, ScD (2013-2014) University of Pennsylvania Specialty Preferences: Longitudinal studies; missing data; quality of life; oncology

Donald L. Trump, MD (2004-2005) Roswell Park Cancer Institute State University of New York Specialty Preferences: Genitourinary cancer; prostate and bladder cancer; drug discovery; biomarker development; clinical genomics and clinical cancer trials

Miguel A. Villalona-Calero, MD (2008-2011) The Ohio State University College of Medicine and Public Health Arthur James Cancer Hospital Specialty Preferences: Phase I trials; EGFR pathway; lung cancer; DNA repair

Daniel D. Von Hoff, MD (1996-1999, 2003-2004, 2005-2007, 2009, 2013-2014) Translational Genomics Institute (TGen) Specialty Preferences: Clinical trials; biostatistics; epidemiology; liver and pancreas tumors; pharmaceutical/drug development

Jamie H. Von Roenn, MD (2010-2014) American Society of Clinical Oncology Specialty Preferences: Palliative care; chemotherapy; symptom management

Julie M. Vose, MD, MBA (1997) University of Nebraska Medical Center Specialty Preferences: Clinical trials in leukemias and lymphomas

Nu Viet Vu, PhD (2001-2003, 2005-2014) University of Geneva Specialty Preferences: Educational testing and evaluation; development of clinical knowledge; assessment of clinical reasoning and competence; program evaluation Richard L. Wahl, MD (2008-2010) Johns Hopkins Medical Institutions Specialty Preferences: Radiology; PET; imaging

Vivian K. Weinberg, PhD (2007-2014) University of California San Francisco Specialty Preferences: Biostatistics; clinical trials; prostate cancer; other GU cancers; survival analysis

Louis M. Weiner, MD (1998-2000, 2003-2009) Fox Chase Cancer Center Temple University School of Medicine Specialty Preferences: Immunotherapy; career development; gastrointestinal cancer

Heidi L. Weiss, PhD (2007-2012) Baylor College of Medicine Specialty Preferences: Biostatistics; design and analysis of cancer immunotherapy trials; design and analysis of cancer chemoprevention trials

H. Samuel Wieand, PhD (2006) NSABP Biostatistical Center Specialty Preferences: Biostatistics; breast and bowel cancers

Jonathan S. Wiest, PhD (2012-2013) National Cancer Institute Specialty Preferences: Lung cancer; cancer genetics

George T. Wilding, MD (1997-1998) University of Wisconsin Carbone Cancer Center Specialty Preferences: Genitourinary cancers; clinical trials

Christopher G. Willett, MD (2013-2014) Duke University Cancer Center Specialty Preferences: Radiation oncology; multimodality management of GI cancer; clinical trials in GI cancer

James E. Williams (2007) Intercultural Cancer Council Specialty Preferences: Patient advocacy; prostate cancer; cancer disparities research; prevention research

Terence Z. Wong, MD, PhD (2013-2014) University of North Carolina Specialty Preferences: Oncologic imaging; PET/CT; image processing; radiation therapy treatment planning; radionuclide therapy

Donn C. Young, PhD (2004-2009) The Ohio University Comprehensive Cancer Center Biostatistics Core Specialty Preferences: Biostatistics; clinical endpoints/quality of life; phase I/II clinical trials **S5: Thomas Reuters Report** (Authors refer to provided attached PDF).