EXECUTIVE SUMMARY
Symptom Management and Quality of Life Steering Committee
Clinical Trials Planning Meeting
Building Bridges: the Identification of Core Symptoms and Health-Related Quality of Life Domains for use in Cancer Research
September 22-23, 2011

Meeting Co-chairs: Deborah Watkins-Bruner, Ph.D., R.N. and Bryce Reeve, Ph.D.
SxQOL SC Co-chairs: Michael Fisch, M.D., M.P.H. and Deborah Watkins-Bruner, Ph.D., R.N.

Introduction
The National Cancer Institute (NCI) Symptom Management & Quality of Life Steering Committee (SxQOL SC) convened a Clinical Trials Planning Meeting for identifying core symptoms and domains for use in cancer clinical trials on September 22-23, 2011 in Washington, DC.

The objectives and goals of the meeting were to:

1) Identify a standard core set of patient-reported symptoms and/or health-related quality of life (HRQOL) domains to be assessed in clinical trials with cancer patients. Selected symptom and HRQOL domains should be ones that are commonly experienced across cancers and are helpful to inform clinical research findings and policy decisions.

2) Identify a core set of symptoms and/or HRQOL domains that should be assessed in clinical trials that include patients with either a head and neck cancer, prostate cancer, or gynecological cancer.

3) Identify patient-reported questionnaires that are appropriate to capture relevant symptoms and HRQOL domains selected in objectives 1 and 2.

Meeting attendees included SxQOL SC members, clinicians, clinical trials experts, biostatisticians, translational scientists, health-related quality of life scientists, patient advocates, and NCI staff.

Background
In 2001, the NCI created the Cancer Outcomes Measurement Working Group (COMWG) consisting of 35 experts convened to examine the state of the science and identify future priorities for outcomes assessment in cancer research. After an extensive review of the cancer outcomes research field over the previous two decades, the COMWG found that assessing health-related quality of life (HRQOL) and symptom burden is feasible using questionnaires that meet established criteria for reliability and validity.
Building on the COMWG findings, the NCI sponsored an international conference in 2006 entitled, “Patient-Reported Outcomes Assessment in Cancer Trials (PROACT): Evaluating and Enhancing the Payoff to Decision Making.” The meeting resulted in a 2007 JCO monograph\(^3\) which identified significant issues and challenges for the incorporation of patient-reported outcomes (PROs: includes HRQOL and symptom burden) in cooperative group trials. Among them was the recognition of the potential for patient-reported symptoms to enhance adverse event monitoring in cancer trials.

Both the COMWG and PROACT reported that a key impediment to move the field forward was a lack of universally recognized standard set of PRO domains to routinely be collected in cancer trials. Clinical trial investigators struggle with the task of knowing what domains to measure in their study that would inform the understanding of the safety and efficacy of the intervention under investigation. The literature is vast on this topic, yet there lacks one source where consensus has been reached on the key PRO domains. As a result, investigators may drop consideration of a PRO endpoint or spend significant time up front to review the literature, consult with co-investigators, and agree on measured endpoints. Further, there lacks consistency from one study to the next on what PRO endpoints are measured which reduces our ability to compare or combine results across trials. Thus, identification of a core set of PRO domains has multiple advantages:

1) Enables clinical trial investigators to come to one source to know what HRQOL and symptom domains to include in their study. This source document will list which domains need to be assessed by cancer type and/or treatment mode and associated questionnaires that measure the domain.

2) Allows researchers and funding agencies to identify domains that lack good quality measures or identify existing questionnaires that require more validation evidence for use in clinical trials. This will lead to a research agenda for measures development and validation in NCI-sponsored trials.

3) Identifies a core set of data elements that will facilitate comparison and combination of data across research studies around the world. This is in line with bio-informatics database initiatives to identify common data elements for meta-analysis types of studies.

**Consensus and Recommendations**

There was general enthusiasm for identifying a core set of symptoms to be assessed in all studies along with other disease/site specific symptoms. However, there were a number of caveats identified related to the purpose, screening, efficacy, and timing of assessments. Researchers currently do not have a good way to look across various studies, and compare and follow patients over time. The assessment of these core patient-reported symptoms would not be mandated nor
replace any other hypothesis driven assessment. Limitations acknowledged were that there remains a need for a thorough, systematic review on functional status and quality of life before specific recommendations could be made on these patient reported outcomes.

**Proposed core set of symptoms to assess in all cancer trials:**

Due to concern for patient burden, core symptoms have been separated into first and second tier symptoms. First tier symptoms are those highly recommended to be assessed in all cancer patients regardless of disease site, stage, or treatment modality. The second tier symptoms are recommended for assessment if other hypothesis-driven, site, stage or modality specific patient reported outcomes would not overwhelm subjects or be likely to increase lack of compliance.

**Proposed core set of first tier symptoms.** Insomnia/ Sleep Disturbance, Pain (general), Fatigue, Nausea, Depressed mood/ sad feelings, Anorexia/ decreased appetite, Anxiety, Concentration Problems

**Proposed core set of second tier symptoms:** Dyspnea/ shortness of breath, Vomiting, Neuropathy/ Numbness or tingling in hands or feet, Diarrhea

Issues to consider with use of the proposed core set of symptoms for all cancers include:

- The need to consider spectrum (local, advanced, metastatic) of the disease and treatment intention (curative, palliative, adjunctive) as the core set of symptoms may differ
- The need for the symptoms to be benchmarked in context of population norms, when in the course of treatment/ follow-up these symptoms are assessed, and the question of why symptoms should be monitored if it will not change treatment.
- Consideration of latent variables not identified
- How the tier may change based upon patient attribution

**Proposed core set of symptoms to assess in three site specific cancer trials:** In addition to the core set of symptoms recommended above, it is recognized that there are common disease site-specific symptoms that should be assessed across site-specific trials. The conference had time and resources to make recommendations for patient self-reported symptoms in three common disease sites: prostate, head and neck, and ovarian cancers.

**Recommendations for PROs to collect in prostate cancer trials**

Localized prostate cancer: urinary incontinence (how much control, how often leak, how much leak), urinary obstruction/irritation (burning, slow flow, night-time urination, frequency), bowel/rectal symptoms (diarrhea, urgency, blood, cramping, pain), sexual dysfunction (erectons, orgasm, desire), hormonal symptoms (hot flashes, breast tenderness, weight change)

Advanced prostate cancer: pain (intensity, interfering with daily activities), energy/vitality (fatigue), emotional well-being (feeling depressed, nervous, trouble sleeping), physical capacity (ability for work, physical activities, ADLs)

Other important measures: regret, satisfaction with care, anxiety about disease
Recommendations for PROs to collect in head and neck cancer trials
Swallowing, pain/oral, skin changes, dry mouth, dental health, opening mouth/trismus, taste, excess/thick mucus/saliva, shoulder disability/motion, social functioning, and functional status.

Recommendations for PROs to collect in ovarian cancer trials
Abdominal core (abdominal pain, appetite loss, bloating, constipation, cramping, indigestion, nausea, vomiting, weight gain or loss), Neuropathy, Fear of recurrence or death, Sexual function, Overall quality of life

This Executive Summary presents the consensus arising from the CTPM. These recommendations are not meant to address all clinical contexts, but rather represent priorities for publicly funded clinical research.

Anticipated Action(s)
- Publish white papers (4) related to recommendations for all cancers, prostate cancer, head and neck cancer, and ovarian cancer
- Present findings at national meetings
- Disseminate recommendations to treatment trialists (i.e., NCI Disease-Specific Steering Committees, Task Forces and NCI Cancer Clinical Trial Cooperative Groups)
- Work with the Cooperative Groups to determine feasibility of incorporating core sets of symptoms

References
AGENDA

NCI Symptom Management & Quality of Life Steering Committee
Clinical Trials Planning Meeting on
Building Bridges: the Identification of Core Symptoms and Health-Related Quality of Life Domains for use in Cancer Research
September 22-23, 2011

Hyatt Regency Washington on Capitol Hill
Capitol Room
400 New Jersey Avenue, NW
Washington, DC 20001

Meeting Goals:
1) Identify a standard core set of patient-reported symptoms and/or health-related quality of life (HRQOL) domains to be assessed in clinical trials with cancer patients. Selected symptom and HRQOL domains should be ones that are commonly experienced across cancers and are helpful to inform clinical research findings and policy decisions.

2) Identify a core set of symptoms and/or HRQOL domains that should be assessed in clinical trials that include patients with either a head and neck cancer, prostate cancer, or gynecological cancer.

3) Identify patient-reported questionnaires that are appropriate to capture relevant symptoms and HRQOL domains selected in goals 1 and 2.

<table>
<thead>
<tr>
<th>Session Topic</th>
<th>Presenter</th>
<th>Time</th>
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<tbody>
<tr>
<td>Sept. 22, 2011 (Thursday)</td>
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<tr>
<td>Registration and Dinner</td>
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<td>5:30 pm</td>
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<tr>
<td>Welcome and Introductions</td>
<td>Deborah Bruner, Bryce Reeve</td>
<td>6:00 pm</td>
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<tr>
<td>Charge from cancer survivor</td>
<td>Mary Lou Smith</td>
<td>6:10 pm</td>
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<tr>
<td>NCI: Perspective on Patient Reported Outcomes in Cancer Clinical Trials</td>
<td>Lori Minasian</td>
<td>6:20 pm</td>
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<tr>
<td>Goals and Limitations of Selection of a Core Set of PROs for Cancer Clinical Trials</td>
<td>Deborah Bruner</td>
<td>6:30 pm</td>
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Agenda as of 09-20-11
<table>
<thead>
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<th>Time</th>
<th>Event</th>
<th>Presenter(s)</th>
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<tbody>
<tr>
<td>6:40 pm</td>
<td>Meaningful Use of Patient-Reported Data in Cancer Treatment Trials: Balancing Decision Making Utility, Measurement Standards, and Response Burden</td>
<td>Bryce Reeve</td>
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<tr>
<td>6:50 pm</td>
<td>All Cancers: Identifying Core Symptoms and HRQOL Domains</td>
<td>Amylou Dueck</td>
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<tr>
<td>7:10 pm</td>
<td>Patient-Reported Outcomes for Adverse Event Reporting</td>
<td>Ethan Basch</td>
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<tr>
<td>7:25 pm</td>
<td>From Screening to Efficacy Endpoints</td>
<td>David Cella</td>
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<tr>
<td>7:40 pm</td>
<td>All Cancers Core Set Recommendations</td>
<td>Carolyn Reilly</td>
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<td>7:50 pm</td>
<td>Panel Members Introduction and Charge for Panel Discussion</td>
<td>Deborah Bruner</td>
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<td>7:55 pm</td>
<td>Panel Discussion</td>
<td>Ethan Basch, Deborah Bruner, David Cella, Corneel Coens, Amylou Dueck, Lori Minasian, Sandra Mitchell, Carolyn Reilly, Mary Lou Smith</td>
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<tr>
<td>8:55 pm</td>
<td>Panel Wrap-up</td>
<td>Deborah Bruner</td>
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<td>9:00 pm</td>
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**Sept 23, 2011 (Friday)**

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<tbody>
<tr>
<td>8:00 am</td>
<td>Welcome, Summary of Day 1, Overview of Day 2</td>
<td>Deborah Bruner, Bryce Reeve</td>
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<tr>
<td>8:10 am</td>
<td>Prostate Cancer Survivor Perspective</td>
<td>Richard J. Vetter</td>
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<td>8:20 am</td>
<td>Prostate Cancer Use of Patient Reported Outcomes in a Clinical Trial</td>
<td>Himu Lukka</td>
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<td>8:40 am</td>
<td>Prostate Cancer PRO Expert Perspective</td>
<td>Ronald Chen</td>
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<td>9:00 am</td>
<td>Prostate Cancer Core Set Recommendations</td>
<td>Howard Sandler</td>
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<td>Panel Members Introduction and Charge for Panel Discussion</td>
<td>Howard Sandler</td>
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<td>Panel Discussion</td>
<td>Neil Aaronson, Peter Chang, Ronald Chen</td>
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Agenda as of 09-20-11
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<tr>
<td>10:15 am</td>
<td>Panel Wrap-up</td>
<td>Howard Sandler</td>
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<tr>
<td>10:20 am</td>
<td>Break</td>
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<tr>
<td>10:30 am</td>
<td>Head and Neck Cancer Survivor Perspective</td>
<td>Pat Gavin</td>
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<tr>
<td>10:40 am</td>
<td>Head and Neck Cancer - Treatment and Toxicity</td>
<td>Drew Ridge</td>
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<tr>
<td>11:00 am</td>
<td>Patient Reported Outcomes: Clinical Trials Issues in Head and Neck Cancer</td>
<td>Barbara Murphy</td>
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<tr>
<td>11:20 am</td>
<td>What can PROs Teach Us?</td>
<td>Avi Eisbruch</td>
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<tr>
<td>11:40 am</td>
<td>Head and Neck Cancer Core Set Recommendations</td>
<td>Bhishamjit Chera</td>
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<td>11:50 am</td>
<td>Panel Members Introduction and Charge for Panel Discussion</td>
<td>Ben Movsas</td>
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<tr>
<td>11:35 am</td>
<td>Panel Discussion</td>
<td>Bhishamjit Chera, Avi Eisbruch, Pat Gavin, Ben Movsas, Barbara Murphy, Drew Ridge</td>
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<td>12:35 pm</td>
<td>Panel Wrap-up</td>
<td>Ben Movsas</td>
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<tr>
<td>12:40 pm</td>
<td>Working Lunch</td>
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<tr>
<td>1:10 pm</td>
<td>Measuring Miracles: Reflections of an Ovarian Cancer Survivor</td>
<td>Meg Gaines</td>
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<tr>
<td>1:20 pm</td>
<td>How and Why PROs inform GOG 252: A Phase III Trial of Bevacizumab with Intravenous versus Intraperitoneal Chemotherapy</td>
<td>Lari Wenzel, Steve Plaxe</td>
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<tr>
<td>1:40 pm</td>
<td>Priority Symptoms and QOL Domains for Women with Recurrent Ovarian Cancer based on data from GOG 259: The WRITE Symptoms Study</td>
<td>Heidi Donovan</td>
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<tr>
<td>2:00 pm</td>
<td>Ovarian Cancer Core Set Recommendations</td>
<td>Lari Wenzel</td>
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<tr>
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<td>Panel Members Introduction and Charge for Panel Discussion</td>
<td>Lari Wenzel</td>
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<tr>
<td>2:15 pm</td>
<td>Panel Discussion</td>
<td>David Cella, Heidi Donovan, Kristine Donovan, Meg Gaines, Richard Penson</td>
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<tr>
<td>3:15 pm</td>
<td>Panel Wrap-up</td>
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<tr>
<td>3:20 pm</td>
<td>Meeting Wrap-up</td>
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<td>4:00 pm</td>
<td>Adjourn</td>
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Agenda as of 09-20-11
# NCI Symptom Management & Quality of Life Steering Committee

**Clinical Trials Planning Meeting on**

*Building Bridges: the Identification of Core Symptoms and Health-Related Quality of Life Domains for use in Cancer Research*

*September 22-23, 2011*

## Participant List

<table>
<thead>
<tr>
<th>Name</th>
<th>Affiliation</th>
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<tbody>
<tr>
<td><strong>Bryce Reeve, Ph.D.</strong></td>
<td>University of North Carolina at Chapel Hill</td>
</tr>
<tr>
<td><strong>Kathleen Castro, R.N.</strong></td>
<td>National Cancer Institute</td>
</tr>
<tr>
<td><strong>Lori Frank, Ph.D.</strong></td>
<td>Medimmune, LLC</td>
</tr>
<tr>
<td><strong>Patricia Ganz, M.D.</strong></td>
<td>Jonsson Comprehensive Cancer Center</td>
</tr>
<tr>
<td><strong>Martha Gaines, J.D.</strong></td>
<td>University of Wisconsin - Madison</td>
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<tr>
<td><strong>Heidi Donovan</strong></td>
<td>University of Pittsburgh School of Nursing</td>
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<tr>
<td><strong>Carol Ferrans, Ph.D.</strong></td>
<td>University of Illinois at Chicago</td>
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<tr>
<td><strong>Jane Meza, Ph.D.</strong></td>
<td>University of Nebraska Medical Center</td>
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<tr>
<td><strong>Patrick Gavin</strong></td>
<td>Grand Rapids Clinical Oncology Program</td>
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<tr>
<td><strong>Ashley Smith, Ph.D., M.P.H.</strong></td>
<td>National Cancer Institute</td>
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<tr>
<td><strong>Richard Vetter, Ph.D.</strong></td>
<td>Mayo Clinic</td>
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<tr>
<td><strong>Leorey Saligan, Ph.D.</strong></td>
<td>National Institute of Nursing Research</td>
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<tr>
<td><strong>Jeffrey Krischer, Ph.D.</strong></td>
<td>University of South Florida</td>
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<tr>
<td><strong>Andrea Barsevick, Ph.D.</strong></td>
<td>Fox Chase Cancer Center</td>
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<tr>
<td><strong>Barbara Murphy, M.D.</strong></td>
<td>Vanderbilt Ingram Cancer Center</td>
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<tr>
<td><strong>Joseph Kelaghan, M.D.</strong></td>
<td>National Cancer Institute</td>
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<tr>
<td><strong>Ann O'Mara, Ph.D.</strong></td>
<td>National Cancer Institute</td>
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<tr>
<td><strong>Clement Gwede, Ph.D.</strong></td>
<td>H. Lee Moffitt Cancer Center and Research Institute</td>
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<tr>
<td><strong>Jennifer Hayes, Ph.D.</strong></td>
<td>National Cancer Institute</td>
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<tr>
<td><strong>Bhupinder Mann</strong></td>
<td>National Cancer Institute</td>
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<tr>
<td><strong>Michael Fisch, M.D.</strong></td>
<td>The University of Texas M. D. Anderson Cancer Center</td>
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<tr>
<td><strong>Joanna Brell, M.D.</strong></td>
<td>National Cancer Institute</td>
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</table>
Ethan Basch, M.D.
Memorial Sloan-Kettering Cancer Center
New York, NY

Jay Pearson, Ph.D.
Merck Research Laboratories
North Wales, PA

Amylou Dueck, Ph.D.
Mayo Clinic
Scottsdale, AZ

Carolyn Reilly, Ph.D.
Emory University
Atlanta, GA

Steven Plaxe, M.D.
University of California, San Diego
La Jolla, CA

Edward Trimble
National Cancer Institute
Bethesda, MD

Mary Lou Smith, J.D.
Research Advocacy Network
Naperville, IL

Charles Cleeland, Ph.D.
The University of Texas M. D. Anderson Cancer Center
Houston, TX

Julia Rowland, Ph.D.
National Cancer Institute
Bethesda, MD

Lisa Kachnic, M.D.
Boston Medical Center
Boston, MA

Jennifer Squires
National Cancer Institute
Bethesda, MD

Andrea Denicoff, R.N.
National Cancer Institute
Bethesda, MD

Claire Snyder, Ph.D.
Johns Hopkins School of Medicine
Baltimore, MD

Karen Mustian, Ph.D.
University of Rochester Medical Center
Rochester, NY

Vivian von Gruenigen, M.D.
Summa Health System
Akron, OH

Lori Minasian, M.D.
National Cancer Institute
Bethesda, MD

Debra Barton, Ph.D.
Mayo Clinic
Rochester, MN

Charles Shapiro, M.D.
The Ohio State University Medical Center
Columbus, OH

Susan Rossi, Ph.D., M.P.H.
National Cancer Institute
Bethesda, MD

Deborah Watkins Bruner, R.N., Ph.D.
University of Pennsylvania
Philadelphia, PA

David Cella, Ph.D.
Northwestern University
Chicago, IL

Bhisham Chera, M.D..
University of North Carolina
Chapel Hill, NC

Ronald Chen, M.D.
University of North Carolina
Chapel Hill, NC

Howard Sandler
Cedars-Sinai Medical Center
Los Angeles, CA

Carol Moinpour, Ph.D.
Fred Hutchinson Cancer Research Center
Seattle, WA
Himu Lukka  
Juravinski Cancer Centre  
Hamilton, Ontario  
Canada

Diane St. Germain, R.N.  
National Cancer Institute  
Bethesda, MD

Eileen Dimond  
National Cancer Institute  
Bethesda, MD

Donald Manning, M.D., Ph.D.  
University of Virginia  
Bloomsbury, NJ

Charles Loprinzi, M.D.  
Mayo Clinic  
Rochester, MN

Neil Aaronson, Ph.D.  
The Netherlands Cancer Institute  
Amsterdam, The Netherlands

Benjamin Movsas, M.D.  
Henry Ford Health System  
Detroit, MI

Donna Berry, Ph.D.  
Dana-Farber Cancer Institute  
Chestnut Hill, MA

Stephen Joel Coons, Ph.D.  
Critical Path Institute  
Tucson, AZ

Michael Brundage, M.D.  
Queen's University  
Kingston, Ontario  
Canada

Marge Good, R.N.  
National Cancer Institute  
Bethesda, MD

John Ridge, M.D., Ph.D.  
Fox Chase Cancer Center  
Philadelphia, PA

Electra Paskett, Ph.D.  
Ohio State University  
Columbus, OH

Jin-Shel Lai, Ph.D.  
Northwestern University  
Chicago, IL

Kristine Donovan, Ph.D.  
Moffitt Cancer Center and H. Lee Research Institute  
Tampa, FL

LeeAnn Jensen, Ph.D.  
National Cancer Institute  
Bethesda, MD

Jeff Sloan, Ph.D.  
Mayo Clinic  
Rochester, MN

Amy Guo, Ph.D.  
Novartis Oncology  
East Hanover, NJ

Antonia Bennett, Ph.D.  
Memorial Sloan-Kettering Cancer Center  
New York, NY

Alice Chen, M.D.  
National Cancer Institute  
Bethesda, MD

Steven Clauser, Ph.D.  
National Cancer Institute  
Bethesda, MD

Russell Glasgow, Ph.D.  
National Cancer Institute  
Bethesda, MD

Chao-Pin Hsiao, Ph.D.  
National Institute of Nursing Research  
Bethesda, MD

Virginia Kwitkowski  
U.S. Food and Drug Administration  
Silver Spring, MD

Denny Kwock  
Daily Wellness Company  
Honolulu, HI

William Lenderking, Ph.D.  
United BioSource Corporation  
Lexington, MA
Wolf Lindwasser, Ph.D.
National Cancer Institute
Bethesda, MD

Sue Marden
National Institute of Nursing Research
Bethesda, MD

Andrew Miller, M.D.
Emory University School of Medicine
Atlanta, GA

Sandra Mitchell, Ph.D.
National Cancer Institute
Bethesda, MD

Rick Moser, Ph.D.
National Cancer Institute
Bethesda, MD

Jean Paty, Ph.D.
invivodata, inc.
Pittsburgh, PA

Stephen Raymond, Ph.D.
PHT Corporation
Boston, MA

Stephanie Shook, Ph.D.
American College of Radiology
Philadelphia, PA

Shaia Sutherland, Ph.D.
Afexa Life Sciences
Edmonton, Alberta

Bhadrasain Vikram
National Cancer Institute
Bethesda, MD

Avraham Eisbruch, M.D.
University of Michigan
Ann Arbor, MI

Richard Penson, M.D.
Harvard Medical School
Boston, MA

Alexandra Hanlon, Ph.D.
University of Pennsylvania
Philadelphia, PA

Worta McCaskill-Stevens
National Cancer Institute

Yvette Ortiz
The EMMES Corporation
Rockville, MD

Lynne Wagner, Ph.D.
Northwestern University
Chicago, IL

Lari Wenzel, Ph.D.
University of California, Irvine
Irvine, CA

John Wilson, Ph.D.
National Surgical Adjuvant Breast and Bowel Project Biostatistic Center
Pittsburgh, PA

Maria Katapodi, Ph.D.
University of Michigan
Ann Arbor, MI

Teresa Deshields, Ph.D.
Siteman Cancer Center
St. Louis, MO

Benjamin Greer, M.D.
University of Washington
Seattle, WA

Jamie Von Roenn, M.D., Ph.D.
Northwestern Memorial Hospital
Chicago, IL

Cornel Coens
European Organisation for Research and Treatment of Cancer
Brussels, Belgium

Peter Chang, M.D.
Beth Israel Deaconess Medical Center
Brookline, MA

Jonathan Cho, M.D.
National Cancer Institute
Bethesda, MD

Paul Kluetz, M.D.
U.S. Food and Drug Administration
Silver Spring, MD