

MEETING AGENDA

The Future of Investigational Medicine: Utilizing Science to Optimize the Early Phase Oncology Clinical Trial Effort

March 23-26, 2023 The Banbury Center, Lloyd Harbor, New York

This meeting was organized by

Tobias Janowitz, Cold Spring Harbor Laboratory

Lillian Siu, Princess Margaret Cancer Centre

and generously funded by the Cold Spring Harbor Laboratory – Northwell Health Affiliation

THURSDAY, MARCH 23

Afternoon Arrivals and check-in at Robertson House

6:00 pm Reception and dinner

FRIDAY, MARCH 24

7:15 am Breakfast

8:30 am Welcome; Overview of Meeting, Goals

Rebecca Leshan, The Banbury Center

Tobias Janowitz, Cold Spring Harbor Laboratory Lillian Siu, Princess Margaret Cancer Centre

Robert Maki, Memorial Sloan Kettering Cancer Center

9:15 am SESSION 1A: Trial Operations: Collaborations Across Disciplines, Sectors, and

Institutions to Enable Early Phase Clinical Trials

Sumithra Mandrekar, Mayo Clinic

S. Michael Rothenberg, Pfizer Inc.

Daniel Tan, National Cancer Centre Singapore



10:55 am SESSION 1B: Trial Operations: Regulations and Cost-Effective Delivery of

Early Phase Clinical Trials

(v) Andrew Lo, Massachusetts Institute of Technology

Naranjargal Dashdorj, Onom Foundation

Ke Liu, Marengo Therapeutics

12:20 pm Luncheon

2:05 pm SESSION 2: Incorporating New Technologies and Biomarkers into Clinical Trials

Minetta Liu, Natera, Inc.

Lillian Siu, Princess Margaret Cancer Centre Christina Yap, Institute of Cancer Research

3:45 pm SESSION 3: What Are the Best Trial Endpoints, Especially if Remote Trial Technology

Is Utilized?

Shaalan Beg, Science 37

Neal Meropol, Flatiron Health

Christopher Hartshorn, National Institutes of Health (NIH)

5:00 pm Day One Wrap-up

6:00 pm Reception and dinner

SATURDAY, MARCH 25

7:45 am Breakfast

9:00 am SESSION 4: Patient Inclusion and Trial Access

Deborah Collyar, Patient Advocates In Research (PAIR)

Karen Mustian, Wilmot Cancer Institute

Elizabeth Fox, St. Jude Children's Research Hospital

Nyasha Chambwe, Feinstein Institutes for Medical Research

11:05 am SESSION 5: Combinatorial Strategies in Drug Development – how to Nominate the

Most Relevant and Truly Synergistic Combinations for Clinical Testing

Helen Chen, National Cancer Institute Cancer Treatment Evaluation Program (CTEP)

Marcus Goncalves, Weill Cornell Medicine

Daniel King, Feinstein Institutes for Medical Research

12:30 pm Luncheon





2:20 pm SESSION 6: Biological Validation in Early Phase Trials: Approaches and

Implementation Strategies

Keith T. Flaherty, Massachusetts General Hospital

Kurt A. Schalper, Yale University

Christophe Le Tourneau, Institut Curie

Yu Shyr, Vanderbilt University Medical Center Tobias Janowitz, Cold Spring Harbor Laboratory

5:15 pm Day Two Wrap-up

6:00 pm Reception and dinner

SUNDAY, MARCH 26

7:45 am Breakfast

9:00 am SESSION 7A: Summarizing & Consensus-Building

Moderated Discussion: key concepts, recommendations

11:00 am SESSION 7B: Meeting Output, Next Steps

Moderated Discussion: planning output(s), responsibilities, next steps and timelines

12:00 pm Luncheon

1:30 pm Participant Departures

PARTICIPANTS

Victoria Aranda, Nature

Shaalan Beg, Science 37

Nyasha Chambwe, Feinstein Institutes for Medical

Research

Helen Chen, National Cancer Institute Cancer

Treatment Evaluation Program (CTEP)

Deborah Collyar, Patient Advocates In Research (PAIR)

Naranjargal Dashdorj, Onom Foundation

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Kurt A. Schalper, Yale University

Yu Shyr, Vanderbilt University Medical Center

Lillian Siu, Princess Margaret Cancer Centre

Daniel Tan, National Cancer Centre Singapore

Andrew Whiteley, Cold Spring Harbor Laboratory

Christina Yap, Institute of Cancer Research