Cardiovascular Toxicity Assessment In Oncology Trials Workshop September 22, 2016

Organized by FDA with support from: AACR, ACC, AHA & ASCO

Speakers and Panelists

Darrell Abernethy, MD, PhD

Associate Director for Drug Safety Office of Clinical Pharmacology (OCP) Office of Translational Science (OTS) CDER/FDA Silver Spring, MD

Laleh Amiri-Kordestani, MD

Clinical Team Leader Breast/Gynecology Team Division of Oncology Product 1 Office of Hematology and Oncology Products CDER/FDA Silver Spring, MD

Ana Barac, MD, PhD, FACC

Director, Cardio-oncology Program Medstar Heart Institute Medical Director Cardiac Rehabilitation Program Medstar Washington Hospital Center Associate Professor of Medicine Georgetown University Washington, DC

Paul W. Burridge, PhD

Northwestern University Feinberg School of Medicine Chicago, IL

Robert Califf, MD, MACC Commissioner

U.S. Food & Drug Administration Silver Spring, MD

Kevin J. Croce, MD, PhD Director, Complex Coronary Artery Intervention Program Director, Translational Discovery Program Interventional Cardiology Brigham and Women's Hospital Harvard Medical School Boston, MA

Chau T. Dang, MD

Chief, Westchester Medical Oncology Service Breast Medicine Service Memorial Sloan Kettering Cancer Center New York, NY

R. Angelo de Claro, MD

Clinical Team Leader Hematology/Benign Hematology Team 4 Office of Hematology and Oncology Products CDER/FDA Silver Spring, MD

Michael S. Ewer, MD, MPH, JD

Special Assistant to the Vice President of Medical Affairs Professor MD Anderson Cancer Center University of Texas Houston, TX

Gary Gintant, PhD

Research Fellow Dept. Integrative Pharmacology Integrated Sciences & Technology AbbVie North Chicago IL

Juan Carlos Plana Gomez, MD

Chief of Clinical Operations, Cardiology Department of Medicine, Baylor College of Medicine Director of the Cardio-Oncology Center and co-Director, Center for Advanced Cardiac Imaging Baylor St. Luke's Medical Center Houston, TX

Nicole Gormley, MD

Clinical Team Leader (Acting) Hematology/Benign Hematology Team 2 Office of Hematology and Oncology Products CDER/FDA Silver Spring, MD

Cardiovascular Toxicity Assessment In Oncology Trials Workshop September 22, 2016

Organized by FDA with support from: AACR, ACC, AHA & ASCO

Speakers and Panelists

Benjamin D. Humphreys MD, PhD

Joseph Friedman Associate Professor & Chief Division of Nephrology Washington University in St. Louis School of Medicine St. Louis, MO

W. Gregory Hundley, MD, FACC, FAHA

Professor Department of Internal Medicine & Radiology Wake Forest Baptist Medical Center Winston-Salem, NC

Bonnie Ky, MD, MSCE

Univ. of Pennsylvania School of Medicine Asst. Professor of Medicine & Epidemiology Director, Penn Center for Quantitative Echocardiography Chair, ECOG Cardiotoxicity Working Group Philadelphia, PA

Daniel Lenihan, MD

Director, Clinical Research Program, Professor of Medicine Vanderbilt Health Nashville, TN

Steven E. Lipshultz, MD

Department of Pediatrics Wayne State University School of Medicine Children's Hospital of Michigan Detroit, MI

Debra Madden

Advocate/Patient Representative, FDA Patient Representative, ECOG/ACRIN Cancer Research Group Patient Representative PCORI Inaugural Advisory Panel on the Assessment of Prevention, Diagnosis, and Treatment Options, National Breast Cancer Coalition

Michael Mauro, MD

Hematologist Leader, Myeloproliferative Neoplasms Program Memorial Sloan Kettering New York, NY

Myrtle D. Millin, DVM, PhD, Fellow, ATS

Chief, Toxicology and Pharmacology Branch Developmental Therapeutics Program Div. of Cancer Treatment and Diagnosis National Cancer Institute, NIH Bethesda, MD

Lori Minasian, MD

Deputy Director NCI/ Division of Cancer Prevention Bethesda, MD

Javid J. Moslehi, MD

Director, Cardio-Oncology Program Assistant Professor of Medicine Vanderbilt School of Medicine Nashville, Tenn

Babak Navi, MD, MS

AHA Representative Director, Stroke Center Assistant Professor of Neurology Weill Cornell Medical College New York, NY

Todd Palmby, PhD

Pharmacology/Toxicology Supervisor Div. of Hematology, Oncology, Toxicology Office of Hematology and Oncology Products CDER/FDA Silver Spring, MD

Thomas Papoian, PhD

Division of Cardiovascular & Renal Products Office of Drug Evaluation I (ODEI) CDER/FDA Silver Spring, MD

Cardiovascular Toxicity Assessment In Oncology Trials Workshop September 22, 2016

Organized by FDA

with support from: AACR, ACC, AHA & ASCO

Speakers and Panelists

Richard Pazdur, MD

Acting Director Oncology Center of Excellence (OCE), FDA Director Office of Hematology & Oncology Products CDER, FDA

Sandra M. Swain, MD, FACP, FASCO

Associate Dean for Research Development Professor of Medicine Georgetown University Medical Center Washington, DC

Shari Targum, MD

Clinical Team Leader Division of Cardiovascular & Renal Products Office of Drug Evaluation I (ODEI) CDER/FDA Silver Spring, MD

Hugo M. Vargas, PhD, DSP

Integrated Discovery & Safety Pharmacology Comparative Biology and Safety Sciences Amgen, Inc. Thousand Oaks, CA

Suparna B. Wedam, MD

Breast Oncology Group Breast Cancer Scientific Liaison Division of Oncology Products 1 Office of Hematology & Oncology Products CDER/FDA Silver Spring, MD

Neil J. Weissman, MD, FACC, FASE

Director, Cardiovascular Core Laboratories President, MedStar Health Research Institute Professor of Medicine, Georgetown University Washington, DC