15th Annual AACI CRI Meeting

JUNE 26-28 2023

Loews Chicago O'Hare Hotel



PO Box 7317 Pittsburgh, PA 15213

www.aaci-cancer.org



CANCER CLINICAL RESEARCH FROM ABSTRACT TO REALITY



AACI CRI 2023 Steering Committee

Chair: Tara L. Lin, MD The University of Kansas Cancer Center

Chair-elect: Thomas J. George, Jr., MD, FACP University of Florida Health Cancer Center

Frances Brogan, MSN, RN, OCN, CCRP

Herbert Irving Comprehensive Cancer Center Columbia University Irving Medical Center

Arcadia Cruz, PhD UC San Diego Moores Cancer Center

Angela Fritsche, MPA Mayo Clinic Comprehensive Cancer Center

Margaret Kasner, MD, MSCE Sidney Kimmel Cancer Center at Jefferson Health

Matthew Kovak, MS, CCRP UAMS Winthrop P. Rockefeller Cancer Institute Wendy Law, PhD Fred Hutchinson Cancer Center

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Anne Schnatterly, MBA, BSN, RN, CCRP WVU Cancer Institute

Susanna Sellmann, MRT, BSc, CCRP Princess Margaret Cancer Centre University Health Network

Rachna Shroff, MD, MS The University of Arizona Cancer Center

Meeting App and Social Media

The agenda, presentations, exhibitor information, and attendees list are available on the meeting app.

To access the app:

- Download "Cvent Events" from your mobile app store.
- Within the Cvent Events app, search for "15th Annual AACI Clinical Research Innovation Meeting" and download the meeting.
- Enter your name and email used during registration.
- You will receive a 6-digit verification code. Enter your code and log in.

The meeting app can also be accessed on your desktop at: https://cvent.me/80OeXR

Twitter: @AACI_Cancer Facebook: AACICancer LinkedIn: https://linkedin.com/company/aaci-cancer Hashtag: #CRI2023 Wireless Network: AACI CRI Wi-Fi Password: 2023cri

AACI CRI Welcome

he Association of American Cancer Institutes (AACI) Clinical Research Innovation (CRI) program serves as a network for research leaders to develop and share best practices for the efficient operation of clinical trials offices (CTO) at AACI cancer centers.

The programming of the 15th Annual AACI CRI Meeting, "Taking Innovations in Cancer Clinical Research From Abstract to Reality," aims to stimulate cancer center interactions and maximize resources by creating opportunities for peer-to-peer networking and collaboration.

CRI Strategic Plan Goals and Projects

- Increase AACI cancer center member participation in CRI and integrate
 CRI into AACI programs and initiatives
- Collect and disseminate benchmarking data to develop and support cancer center clinical trials
- Encourage cancer centers to share emerging practices for improving community outreach and engagement in clinical trials
- Promote diversity, equity, and inclusion for training and career development
- Develop clinical research education, resources, and networks for professional development to continue fostering communication and mentoring opportunities
- Increase engagement with industry and other stakeholders to support CRI
- Develop financially sustainable models for facilitating efficient and rapid clinical trials office operations

AACI CRI Meeting Objectives

- Discuss ways to create and maintain a positive environment in the clinical trials office (CTO) to ensure staff retention and top performance
- Share strategies for developing a community network for cancer clinical trials by reassessing long-standing operational practices
- Learn from patient advocates' unique perspectives on recruitment challenges and solutions
- Share successful and innovative strategies to broaden the outreach and recruitment of diverse patients in clinical trials
- Understand potential disconnections between clinical care and research by examining relationships between departments and aligning reporting structures to share data as a collective effort
- Review the differences and common operational challenges between pragmatic trials and decentralized trials to understand how to overcome them
- Review tips for preparing a Cancer Center Support Grant (CCSG) application, with a focus on the Clinical Protocol and Data Management (CPDM) and Protocol Review and Monitoring System (PRMS) components
- Network with like-minded professionals to help build relationships and support
 efficiency in all AACI cancer center member CTOs

Who Attends This Meeting?

- Individuals from AACI member cancer centers, including:
 - CTO administrative directors, medical directors, managers, and supervisors
 - Deputy/associate directors of clinical research administration
 - Cancer center administrators
 - Research regulatory management and staff
 - Clinical research finance directors, managers, and supervisors
 - Biostatisticians and informatics specialists
- Employees of U.S. Department of Health and Human Services agencies and offices, including the National Cancer Institute and the U.S. Food and Drug Administration
- AACI sustaining members
- AACI Corporate Roundtable members
- Representatives from industry, including drug development companies, clinical research organizations, and consultants
- Employees of information technology companies that support cancer center clinical research management
- Representatives of like-minded organizations promoting patient access to clinical trials

MEETING PROGRAM

15th Annual AACI Clinical Research Innovation (CRI) Meeting *Taking Innovations in Cancer Clinical Research From Abstract to Reality Monday, June 26 – Wednesday, June 28*

All sessions are in central time. Sessions are held in Cassatt unless otherwise noted.

Monday, June 26

11:00 AM Registration Opens Artist Foyer

11:00 AM Exhibits Open Artist Foyer

1:00 PM Welcome

Tara L. Lin, MD The University of Kansas Cancer Center

1:30 PM Exploring Unique and Innovative Strategies for Clinical Trial Recruitment

This panel discussion will focus on unique approaches to trial recruitment, incorporating contemporary innovations like social media campaigns and mobile outreach. Experts from AACI cancer centers will share insights into various strategies to optimize trial recruitment with a particular focus on groups under-represented in clinical research.

Moderator: Thomas J. George, Jr., MD, FACP

University of Florida Health Cancer Center

Lisa Carter-Bawa, PhD, APRN, ANP-C, FAAN Center for Discovery and Innovation at Hackensack Meridian Health

Kearston Ingraham, MPH Duke Cancer Institute, Duke University Medical Center

Mel Mann, MBA, MEd Patient Advocate

2:30 PM Networking Break Artist Foyer

3:00 PM Manage, Review, Monitor: Preparing the CPDM and PRMS Components of the CCSG

National Cancer Institute (NCI) officials will provide an overview of recent and proposed changes to NCI's Cancer Center Support Grant (CCSG), including new guidelines for reporting data for pragmatic trials. Panelists will also review the practical implications of preparing a CCSG competing renewal, with a focus on the Clinical Trials Reporting Program, revisions to the Funding Opportunity Announcement— including funding of investigator-initiated trials and catchment area considerations—new diversity and inclusion reporting requirements, and prospects for continued virtual site visits.

Moderator: Wendy Law, PhD

Fred Hutchinson Cancer Center

Min He, PhD National Cancer Institute

Gisele Sarosy, MD National Cancer Institute

4:00 PM Poster Session Avedon

5:30 PM Welcome Reception Artist Foyer Supported by Actalent and Florence

Tuesday, June 27

7:00 AM General Breakfast Guggenheim – Museum Wing

- 7:00 AM CTO Administrative Directors' Breakfast (invitation only) Prado Room
- 7:00 AM CTO Medical Directors' Breakfast (invitation only) Metropolitan Room
- 8:00 AM Exhibits Open Artist Foyer

8:15 AM Welcome

Tara L. Lin, MD The University of Kansas Cancer Center

8:30 AM Keynote Presentation: The Power of Positivity: Building a Stronger, More Resilient Clinical Cancer Research Team

Dr. Ted A. James will offer valuable insights and actionable tactics for promoting a positive workplace culture, raising morale and retention, and improving outcomes. Other discussion topics include the significance of positivity in clinical cancer research and how it can be leveraged to enhance the institution's mission and achieve greater professional satisfaction. Participants will leave with tangible strategies they can adopt to improve their performance and transform their work environment.

Moderator: Tara L. Lin, MD

The University of Kansas Cancer Center

Ted A. James, MD, MHCM, FACS

Beth Israel Deaconess Medical Center, Harvard Medical School

9:30 AM Modernizing Clinical Trial Conduct: Operational Readiness for Pragmatic and Decentralized Trials

Managing pragmatic versus decentralized trials first requires a deep understanding of the differences between them. This session will aim to provide clear definitions of the two types of trials, along with a review of both their unique and shared challenges, including activation, staffing, and training.

Moderator: Angela Fritsche, MPA Mayo Clinic Comprehensive Cancer Center

Mark Agulnik, MD City of Hope Comprehensive Cancer Center

Shaalan Beg, MD Science 37

Ana Patricia Ortiz, MPH, PhD University of Puerto Rico Comprehensive Cancer Center

Harpreet Singh, MD U.S. Food and Drug Administration

10:45 AM Breakout Sessions

Community Outreach and Engagement & Diversity, Equity, and Inclusion Louvre 1

Co-moderator: Patricia Hurley, MSc *American Society of Clinical Oncology*

Co-moderator: Grzegorz Nowakowski, MD Mayo Clinic Comprehensive Cancer Center

Quality Assurance & Remote Monitoring and Auditing *Warhol B*

Co-moderators: Adrian Granobles, CCRP, and Karima Yataghene, MD Memorial Sloan Kettering Cancer Center

Regulatory Pollock A

Co-moderators: J. Kaitlin Morrison, PhD, and Shaw Scott UNC Lineberger Comprehensive Cancer Center University of North Carolina at Chapel Hill

Resource Management and Finance *Warhol A*

Co-moderators: Phillip Eggleton, BBA, and Ronni Geary, MBA, CPC UAMS Winthrop P. Rockefeller Cancer Institute

Training, Career Development, and Staff Retention *Louvre 2*

Moderator: Anna Kukulka, RN, BSN, MEd, CCRC University of Florida Health Cancer Center

Trial Recruitment and Study Conduct (IITs) Pollock B

Co-moderators: Lillian Neal, MSc, CCRP, and Katherine Vosburgh, BSN, RN, BMTCN Hollings Cancer Center, Medical University of South Carolina

Trial Start-up, Activation, and Protocol Development *Louvre 3*

Co-moderators: Amanda Spratt, CCRP, and Lauren Wall, MS The University of Chicago Medicine Comprehensive Cancer Center

11:45 AM Lunch Guggenheim – Museum Wing

1:15 PM 2023 CRI Abstract Presentations

Abstracts from AACI cancer center members focus on oncology research that illuminates clinical research challenges and solutions, accelerating cancer drug development. The CRI steering committee has selected three abstracts for presentation at this year's meeting. The abstract presentations will be followed by a Q&A session.

Moderator: Tara L. Lin, MD The University of Kansas Cancer Center

Jennifer Bollmer, PhD Medical College of Wisconsin Cancer Center

Christy Spalink, DNP, MSN, RN Laura and Isaac Perlmutter Cancer Center at NYU Langone

Christina Wiess, CCRP Yale Cancer Center, Yale School of Medicine

2:15 PM Breakout Sessions

Community Outreach and Engagement & Diversity, Equity, and Inclusion Louvre 1

Co-moderator: Jose Gomez, MSW Cedars-Sinai Cancer

Co-moderator: Nedra Johnson, BSN, RN, CCRP Medical College of Wisconsin Cancer Center

Quality Assurance & Remote Monitoring and Auditing *Warhol B*

Co-moderators: Susan Puleio and Jacqueline Simpronio *Memorial Sloan Kettering Cancer Center*

Regulatory Pollock A

Moderator: Stacy Mercado, MS, CCRC, CHRC Abramson Cancer Center of the University of Pennsylvania

Resource Management and Finance Warhol A

Moderator: Joshua Plassmeyer, MS, CCRP UPMC Hillman Cancer Center

Training, Career Development, and Staff Retention *Louvre 2*

Moderator: Carrie Lee, MD, MPH UNC Lineberger Comprehensive Cancer Center University of North Carolina at Chapel Hill

Trial Recruitment and Study Conduct (IITs) *Pollock B*

Co-moderators: Christina Crabtree-Ide, PhD, MPH, and Rachel Frascati, PhD

Roswell Park Comprehensive Cancer Center

Trial Start-up, Activation, and Protocol Development *Louvre 3*

Co-moderators: Philip Arlen, PhD, Lindsey Mooney and Kelly Williams

Sylvester Comprehensive Cancer Center, University of Miami Health System

3:15 PM Networking and Dessert Break Artist Foyer

3:45 PM "We Are (Work) Family": Building a Positive CTO Team Culture

Along with the daily tasks that keep a clinical trials office running smoothly, creating a positive environment for staff helps to ensure efficiency and top performance. In this session, panelists discuss defining the "work family," building values and culture, hiring practices, remote work vs. in-person policies, communication, and maintaining a positive culture at community sites.

Moderator: Matthew Kovak, MS, CCRP

UAMS Winthrop P. Rockefeller Cancer Institute

Zeno Ashai, MBBS, MPH USC Norris Comprehensive Cancer Center

Kristin Herman, MBA

Sidney Kimmel Cancer Center at Jefferson Health

Michelle Liendo, MSHI, CCRP

VCU Massey Cancer Center

4:45 PM Vendor Presentation: Triomics

Activating Clinical Data for Research

Clinical data is the linchpin for advancing medical research. However, the lack of standardization and interoperability poses challenges to the automation and scaling of these processes. As a result, researchers are stuck with laborious manual procedures such as patient chart reviews and transferring data from EHR to EDCs.

Triomics is at the forefront of solving these challenges by harnessing the power of data standards such as FHIR, mCode, and AI systems, including sophisticated language models that can extract relevant information from complex medical data.

In this interactive session, we'll explore new developments in health care data interoperability, AI, and Triomics' innovative technology. This session is ideal for cancer center administrators, clinical trialists, health care researchers, and data scientists.

8:00 PM Hospitality Ice Bar, Lobby Level

Wednesday, June 28

7:00 AM	Breakfast Guggenheim – Museum Wing
8:00 AM	Exhibits Open Artist Foyer
8:00 AM	Welcome
	Tara L. Lin, MD The University of Kansas Cancer Center

8:15 AM It Takes a Community: Overcoming the Barriers to Decentralized Trials

Developing a community network for cancer clinical trials is a multistep and often complex process. Challenges can include finding the right space, thinking outside the box on budgeting, and adapting sometimes long-standing administrative practices. Panelists for this session will discuss how to start from scratch when setting up a decentralized trial infrastructure.

Moderator: Arcadia Cruz, PhD

UC San Diego Moores Cancer Center

Amishi Dhadwal The Tisch Cancer Institute at Mount Sinai

Jesse Nodora, DrPH UC San Diego Moores Cancer Center

Grzegorz Nowakowski, MD Mayo Clinic Comprehensive Cancer Center

Triomics

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9:15 AM Part 1: Bridging the Gap Between Clinical Care and Research Operations

Cancer centers routinely wrestle with disconnections between clinical care and research that can sometimes compromise optimal clinical trial operations. Panelists will look at ways to foster collaboration across the campus, including building relationships between the cancer center and investigational drug services, infusion center, radiology, ophthalmology, and other departments.

Moderator: Frances Brogan, MSN, RN, OCN, CCRP

Herbert Irving Comprehensive Cancer Center Columbia University Irving Medical Center

Heidi Finnes, PharmD, BCOP, FHOPA Mayo Clinic Comprehensive Cancer Center

Shahnaz Singh-Kandah, MSN

Herbert Irving Comprehensive Cancer Center Columbia University Irving Medical Center

Christy Spalink, DNP, MSN, ACNP-BC, ACHPN, OCN, RN

Laura and Isaac Perlmutter Cancer Center at NYU Langone

10:15 AM Break

10:30 AM Part 2: Bridging the Gap Between Clinical Care and Research Operations

A collaborative work culture is key to forming strong bonds between clinical care and research operations. Aligning reporting structures and sharing data like catchment area parameters for use in the Cancer Center Support Grant application are among the aspects of collective effort that will be examined in this session.

Moderator: Anne Schnatterly, MBA, BSN, RN, CCRP

WVU Cancer Institute

Todd Burus, MAS UK Markey Cancer Center

Christopher Loertscher, MA USC Norris Comprehensive Cancer Center

Dhaval Mehta, MD UPMC Hillman Cancer Center

11:30 AM Closing Remarks

11:45 AM Adjourn

AACI CRI Meeting 2023 Abstracts

FIRST PLACE:

Hybrid Decentralization of Early Phase Cancer Clinical Trials to Enhance Study Recruitment of Underrepresented Minorities C. Wiess, A. Rodrigues, I. Palma, D. Wall, P. LoRusso Yale Cancer Center, Yale School of Medicine

SECOND PLACE:

Strategies to Improve Clinical Research Staff Engagement, Retention, Career Development, and Performance at an NCI-Designated Cancer Center

C. Spalink, A. Joshi, A. Husni, P. Patel, A. Haegler, E. Waalkes, N. Chowdhury, B. Pothuri, J. Mehnert

Laura and Isaac Perlmutter Cancer Center at NYU Langone

THIRD PLACE:

Protocol Prioritization Scores: Are They Predictive? J. Bollmer, J. Thomas, B. George, M. Larson, K. Schroeder, S. Zindars, R. Kurzrock Medical College of Wisconsin Cancer Center

Abstracts are organized by category and completion status, then in alphabetical order by cancer center.

CATEGORIES:

Community Outreach and Engagement & Diversity, Equity, and Inclusion

Quality Assurance & Remote Monitoring and Auditing

Regulatory

Resource Management and Finance

Training, Career Development, and Staff Retention

Trial Recruitment and Study Conduct (IITs)

Trial Start-up, Activation, and Protocol Development

Community Outreach and Engagement & Diversity, Equity, and Inclusion

- Partnering and Building Opportunities Within North Carolina: A Qualitative Analysis of the Lineberger Comprehensive Cancer Center Clinical and Research Internship for Black, Indigenous, and People of Color (BIPOC) Undergraduate Students
 A. Daye, S. Godfrey, A. Walens, V. Carlisle, B. Austin, C. Lee, A. Leak Bryant UNC Lineberger Comprehensive Cancer Center, University of North Carolina at Chapel Hill
- 2. A Multimodal Approach to Increasing Participation of Underrepresented Communities in Investigator-Initiated Cancer Clinical Trials

J. Gomez, G. Gresham, E. Hautamaki, M. Malikowski, K. Reckamp, B. Rimel Cedars-Sinai Cancer

- 3. A Multichannel Approach to Reducing the Health Equity Gap in the Black Community J. Gomez, A. Levi, A. Hendifar Cedars-Sinai Cancer
- 4. Development of a Process to Share Plain Language Summaries of Clinical Research Results With Participants at Princess Margaret Cancer Centre K. Zeman, H. Cole, S. Sellmann Princess Margaret Cancer Centre, University Health Network
- 5. Evaluating Clinical Trial Participation Across the Catchment Area: A Data-Driven Approach K. Sinclair, D. Forsyth, K. Hamade, C. McNair Sidney Kimmel Cancer Center at Jefferson Health
- 6. Increasing Clinical Trial Accrual of Minority Patients by **Expanding Clinical Operations at Satellite Sites** A. Dhadwal, D. Catmaero, A. Lieberman-Cribbin, C. Rodriguez, J. Richter, S. Jagannath The Tisch Cancer Institute at Mount Sinai



Quality Assurance & Remote Monitoring and Auditing

- 8. Development of a Digital Audit Tracking Tool for FDA Audit Readiness K. MacLennan, B. Koch Abramson Cancer Center of the University of Pennsylvania
- 9. Improving Quality: First and Third Patient Review A. Fritsche, K. Croghan, J. Zbacnik, A. Youssef, L. Winkowski, A. Holland, G. Nowakowski

Mayo Clinic Comprehensive Cancer Center

- 10. Improving Quality: Audit Readiness Team L. Winkowski, K. Croghan, K. Severson, H. Kogut, A. Jurrens, A. Fritsche, G. Nowakowski, A. Mansfield Mayo Clinic Comprehensive Cancer Center
- 11. Too Many Studies to Audit and Monitor? Let the Protocol Risk **Assessment Tool System Help You Prioritize** A. Granobles, K. Yataghene Memorial Sloan Kettering Cancer Center

- 12. Getting Monitoring Deficiencies Resolved A. Granobles, K. Mantha-Thaler, K. Yataghene Memorial Sloan Kettering Cancer Center
- 13. Saved by Automation! A Continuation of the Story of How Technology and Innovative Thinking Significantly Increased **Productivity Surrounding CAPA Completion** J. Simpronio, S. Puleio, M. Ayerov, H. Daggumati, K. Yataghene Memorial Sloan Kettering Cancer Center
- 14. How to Conduct a Regulatory Review to Ensure a Quality FDA Inspection G. Grimaldi, M. Reynolds, P. Chadha, S. Kling, V. Michel, C. Luk, F. Yeh, D. De Blasi, K. Yataghene, C. Houston, A. Drilon, M. Gounder Memorial Sloan Kettering Cancer Center
- 15. Innovative Approaches to Clinical Research Monitoring: The Power of Ingenuity at Memorial Sloan Kettering Cancer Center S. Sanchez-Molero Perez, A. Granobles, K. Mantha-Thaler, L. Bello-Matricaria, K. Yataghene Memorial Sloan Kettering Cancer Center
- 16. The Impact of Having a "Quality" Quality Assurance System on Audit Findings from 2020-2022 J. Brown, M. Martinez, N. Surana, P. Seo, E. Dawkins Sylvester Comprehensive Cancer Center, University of Miami Health System
- 17. Creating a Robust Quality Assurance Program to Ensure **Compliance in Research** S. Achberger, K. McCaffrey, M. Kilbane Cleveland Clinic Cancer Center
- 18. Reduced Research Patient Wait Times Using Automated Dispensing Cabinet (ADC) Technology for Oral Investigational Drug at an NCI-Designated Comprehensive Cancer Center E. Waalkes, C. Spalink, J. Scagliola, A. Joshi, B. Pothuri, J. Mehnert, D. Ayoubi Laura and Isaac Perlmutter Cancer Center at NYU Langone
- 19. Keeping an Eye on RNI: Frequent Monitoring to Eliminate Preventable Reportable New Information E. O'Donovan, P. Patel, E. Yepes, A. Joshi, C. Spalink, A. Goutzinopoulos, B. Pothuri Laura and Isaac Perlmutter Cancer Center at NYU Langone
- 20. Meeting a National Need: Implementing an NCTN Quality **Assurance Program** R. Selle, C. Gill, S. Zindars, K. Schroeder, B. George Medical College of Wisconsin Cancer Center

- 21. Path to Improved Trial Management and FDA Inspection Readiness G. Grimaldi, M. Reynolds, P. Chada, C. Luk, F. Yeh, K. Yataghene, C. Houston, A. Drilon, M. Gounder Memorial Sloan Kettering Cancer Center
- 22. Implementation of an Audit Assessment Category Guidance System to Define Audit Deficiencies as Critical, Major, or Minor M. Storms, K. Bogaard

The University of Texas MD Anderson Cancer Center

23. A Formal Dose Escalation/Safety Lead-In Request and Approval Process M. Gawliu

UCSF Helen Diller Family Comprehensive Cancer Center

24. Taking Data Validation to the Next Level: Automating Data Validation Using CDASH-Standardized Global eCRFs S. Rachuri, M. O'Dwyer, K. Douglas, L. Logan, J. Tewell, S. Balu, C. Lee, J.K. Morrison, E. Crecelius

UNC Lineberger Comprehensive Cancer Center, University of North Carolina at Chapel Hill

Regulatory

25. Implementation of a Dashboard to Improve Protocol Oversight and Data and Safety Monitoring Committee (DSMC) Reviews C. Kolenut, K. Napolitano, X. Lekperic, C. Zamore, A. Bijwe, D. Caron, S. Hanley, J. Chaft, S. Slovin

Memorial Sloan Kettering Cancer Center

26. Creation of the Performance Monitoring Committee: Optimizing Review of the MSK Clinical Research Portfolio X. Lekperic, K. Napolitano, C. Kolenut, S. Hanley, A. Rodavitch, C. Houston, D. Rathkopf

Memorial Sloan Kettering Cancer Center

- 27. A Review and Recommendations for Implementing eRegulatory Investigator Site File Systems (eBinder, eISF) M. Blair, C. Trani, L. McHugh, K. Tang, V. Chan Abramson Cancer Center of the University of Pennsylvania
- 28. Development and Implementation of a Research Study Regulatory Complexity Assessment Tool M. Blair, C. Trani, L. McHugh Abramson Cancer Center of the University of Pennsylvania
- 29. Eliminating Unnecessary Review of Offsite Adverse Event (Expedited IND Safety) Reports: Departmental Collaboration Leading to Institutional Position M. Blair, S. Mercado, M. Hendricks, D.T. Vogl Abramson Cancer Center of the University of Pennsylvania

30. Closing Time: Protocol Scoring & Remote Closeout for Portfolio Optimization M. Ismailzadah, C. Rivera, S. Mistretta, D. Agrinsoni, T. Negri, R. Shelton, J. Jurcic, A. Lassman

Herbert Irving Comprehensive Cancer Center, Columbia University Irving Medical Center

- 31. Using HL7-FHIR to Automate Mandatory Reporting of Bone Marrow Transplant Data Decreases Staff Effort and Improves Data Quality C. Thomas, R. Panchal, J. Konecny, T. Casali, M. Buckley, E. Klein Memorial Sloan Kettering Cancer Center
- 32. Regulatory Burden of IRB Submissions: Commercial vs. Internal IRBs
 E. Sibilsky Enselman, J. Humfleet, D. Bashllari
 University of Michigan Rogel Cancer Center

Resource Management and Finance

- **33. Clinical Research Scorecard Performance Metrics M. Hendricks** *Abramson Cancer Center of the University of Pennsylvania*
- 34. Creation of a Budget Workload Score for Analysis B. Search, H. Hampton, K. Kaufman, E. Lascu, B. Zakrzewski Memorial Sloan Kettering Cancer Center
- 35. Automation of Clinical Research Administrative Fees for Internal Recovery
 B. Search, J. Chen, K. Kaufman, L. Lupkin, J. Yan Memorial Sloan Kettering Cancer Center
- 36. Fostering Portfolio Stewardship Through a Trial Portfolio Balancing Framework J. Lebsack, H. Soliman Moffitt Cancer Center
- Development of an Enhanced Clinical Trial Workload Assessment Tool – The BC Clinical Trial Complexity Tool M. Sadiq, S. Sundquist, D. Kato, R. Xu, D. Curman, P. Pollock, K. Sit, K. Halvorsen, J. Clark, M. Abacan, C. Kollmannsberger, B. Eigl BC Cancer
- Will They Pay? Let's Find Out First!: Saving Time and Money in Industry-Trial Activation
 E. Lebleu, S. Ford, L. Hayes, J. Moehle, H. Soares
 Huntsman Cancer Institute, University of Utah
- 39. Leveraging Automation to Increase Time Savings for Processing Research Non-Billables (RNBs)
 S. Siamwalla, R. Panchal, M. Buckley, J. Lengfellner
 Memorial Sloan Kettering Cancer Center

- 40. Development of a Clinical Research Coordinator Capacity Model J. Johnson, M. Ugrenovic-Petrovic, E. Royster, B. Jones-Lombard, B. Mack, A. Patel, I. Krupitsky, K. Hulse, J. Lebsack Moffitt Cancer Center
- 41. Improving PRMC Accrual Monitoring Procedures: Making it Count S. Osipowicz, R. Dampman Weiss, J. Curry, J. Johnson, M. Kasner Sidney Kimmel Cancer Center at Jefferson Health
- 42. Clinical Research Following the Money, Phase 4 R. Geary, P. Eggleton, M. Kovak, M. Birrer, A. Smith, Z. Feng, N. Pruss UAMS Winthrop P. Rockefeller Cancer Institute
- 43. Developing a Scoring Tool to Calculate Protocol Acuity for **Clinical Research Nurse Workload** C. Jones¹, M. McAdoo¹, K. Mack¹, A. Hanlyn² ¹UAMS Winthrop P. Rockefeller Cancer Institute ²UAMS IT Research Systems

Training, Career Development, and Staff Retention

- 44. #ResearchOnResearch A Research Training Initiative for Clinical **Research Professionals** T. Waite Abramson Cancer Center of the University of Pennsylvania
- 45. CROSS to CRES: The Evolution of a Clinical Research Operations Supplemental Series to an Accredited Clinical Research Education Series

T. Waite, C. Redlinger-Tabery, E. Dahlmeier Abramson Cancer Center of the University of Pennsylvania

46. New Employee Orientation – Joining the 21st Century F. Kerr, S. Asche, C. Bucks

Indiana University Melvin and Bren Simon Comprehensive Cancer Center

47. Strategies to Improve Clinical Research Staff Engagement, Retention, Career Development, and Performance at an NCI-**Designated Cancer Center** C. Spalink, A. Joshi, A. Husni, P. Patel, A. Haegler, E. Waalkes, N. Chowdhury, B. Pothuri, J. Mehnert

Laura and Isaac Perlmutter Cancer Center at NYU Langone

48. The Dynamic Duo: Dyad Mentorship of Cancer Clinical Trials **Office (CCTO) Leadership** A. Fritsche¹, G. Nowakowski¹, J. Moehle², T. Werner²

¹Mayo Clinic Comprehensive Cancer Center ²Huntsman Cancer Institute, University of Utah

- 49. Retaining Staff Through Surveys: 6-Month, Stay and Exit J. Ludescher, G. Nowakowski, T. Halfdanarson, A. Fritsche, C. Griffin, R. Platou, M. Wrenn, K. Croghan Mayo Clinic Comprehensive Cancer Center
- 50. A Cancer Clinical Trials Office (CCTO) Orientation Course Reduces **Insufficiencies Among Study Coordinators** K. Croghan, G. Boe, J. Zbacnik, A. Youssef, A. Holland, G. Nowakowski, A. Fritsche Mayo Clinic Comprehensive Cancer Center
- 51. Going From an In-Person to Remote Training Program: How to **Ensure Engagement** V. Tomaselli, M. Nicola Memorial Sloan Kettering Cancer Center
- 52. The Great Rebound: Successful Clinical Trials Office Staffing **Recovery Strategies** K. Morrison, S. Ladd, J. Huamani-Bundy, C. Hilliard, L. Schreiner, N. Whitman, M. Roxas, S. Scott, B. Adams, E. Moore, J. Maccarone, E. Kelly, J. Mayfield, E. Riley, M. Laffan, C. Tew, P. Derebail, S. Rego, L. Kiefer, T. Conrad, B. Marini, G. Harrison, W. Sarratt, V. Bae-Jump, L. Carey, C. Lee

UNC Lineberger Comprehensive Cancer Center, University of North Carolina at Chapel Hill

- 53. Innovative and Inclusive Approach to Clinical Research Pharmacist Development K. Pavlik, R. Sawant, L. Coleman, S. Brogan, T. Ferencz, P. Patel Yale Cancer Center, Yale School of Medicine
- 54. Training for Excellence in Clinical Research: 5 Years of Growth M. Guy, M. Kilbane, K. McCaffrey Cleveland Clinic, Taussig Cancer Institute
- 55. Investing in the Future: Protocol Review Mentorship Program for Oncology Fellows at the Duke Cancer Institute C. Riggan, A. Bender, A. Armstrong Duke Cancer Institute, Duke University Medical Center
- 56. Narrowing the Gap The Synergistic Effect of the Clinical Trials Nurse Liaison L. Lujan, S. Sharry, J. Moehle, R. Doering, A. Emett, J. Jones, C. Kotobalavu, M. Dolim, T. Werner, H. Soares Huntsman Cancer Institute, University of Utah
- 57. Adding to the Career Ladder of Clinical Research Staff at IUSCCC L. Haney, F. Kerr, J. Spittler, L. Rohn, J. Corman, L. Sego, C. Nelson, S. Bailey, M. Contreraz, T. Lautenschlaeger Indiana University Melvin and Bren Simon Comprehensive Cancer Center

58. Development and Implementation of Micro-Trainings as Part of **Continued Education for Clinical Research** K. Croghan, G. Boe, J. Zbacnik, A. Youssef, A. Holland, G. Nowakowski, A. Fritsche Mayo Clinic Comprehensive Cancer Center

59. Managing Flexible Work Schedules Within a Disease-Specific Team

C. Dwight, S. Zindars, H. Heaviland Medical College of Wisconsin Cancer Center

- 60. Peer Support for Second Victim Syndrome S. Eberhardt, A. Pilarski, T. Klatt, S. Babe, H. Nestle, K. Schroeder, M. Lingongo Medical College of Wisconsin Cancer Center
- 61. Integrating Technology to Support Data Management Abstraction of Adverse Events (AE) and Concomitant Medications (ConMed) From the Electronic Health Record (EHR) to Sponsor Electronic Data Capture (EDC) Systems Using Design Thinking Methodology to Increase Efficiency and Help Reduce Staff Turnover

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