

CGC 14th Annual Meeting

August 13 - 16, 2023 St. Louis, MO



Cancer Genomics Consortium, 14th Annual Meeting

August 13-16, 2023 Hyatt Regency at the Arch, St. Louis, Missouri

SUNDAY, AUGUST 13, 2023

Pre-Meeting Workshops

9:00 – 10:30 AM Bioinformatics Workshop: Exploring the Clinical Interpretation Resource

Landscape

Beth Pitel, Mayo Clinic

Brendan Reardon. Dana Farber Cancer Center

Morial Singer-Berk, Broad Institute

10:30 – 10:45 AM Coffee Break for Workshop Attendees

10:45 AM – 12:15 PM Case-Based Workshop: Assessment of Genomic Changes in Clinical

Cases: Things to Think About and How We Approached Them

Lisa Brailey, Mt. Sinai School of Medicine

Celeste Eno, Cedars-Sinai

Jimbo Fan, *University of Virginia*

Keela Scott, University of Missouri-Columbia

Teresa Smolarek, Cincinnati Children's Hospital Medical Center

Opening of the 14th Annual Cancer Genomics Consortium

1:00 – 1:05 PM Welcome

Rashmi Kanagal-Shamanna, MD Anderson Cancer Center

1:05 - 1:15 PM Presidential Address

Obi Griffith, Washington University School of Medicine

1:15 – 2:15 PM Keynote Presentation

John DiPersio, Washington University School of Medicine

2:15 – 2:30 PM Session 1: Applications of Emerging Technologies in Clinical Genomics 1

Clinical Validation of Plasma Whole Genome Sequencing for Detection of Minimal Residual Disease from Solid Tumours

Felix Beaudry, Ontario Institute for Cancer Research

Personalized sequencing assays for cerebrospinal fluid liquid biopsies in children with brain tumors

Katherine Miller, Nationwide Children's Hospital

Application of optical genome mapping to identify samples with homologous recombination deficiency

Alex Hastie, Bionano Laboratories

Comprehensive next generation cytogenomics improves risk stratification of Acute Myeloid Leukemia

Stephen Eacker, Phase Genomics/Fred Hutchinson Cancer Center

4:00 – 4:30 PM Invited Speaker Presentation

Robert Hasserjian, Mass General Brigham

4:30 – 5:30 PM Session 2: Bioinformatics, Artificial Intelligence, Machine-Learning 1

Overcoming challenges in semantic alignment of therapeutics knowledge using Therapy

James Stevenson, Nationwide Children's Hospital

Technologist Award

Tracking Immunotherapy Response with Single Cell T Cell Receptor Profiling in Canine Models of Cancer

Obi Griffith, Washington University School of Medicine

Al-Based Algorithms for Neoplastic Metaphase Cells Boost Efficiencies in the Cytogenetics Laboratory

Bo Hong, ARUP Laboratories

Mapping variants from multiplex assays of variant effect (MAVEs) to human reference sequences

Jeremy Arbesfeld, The Ohio State University

Trainee Award

5:30 – 6:15 PM CGC Scientific and Program Updates

Implementation survey of the ACMG/CGC standards for interpretation of acquired CNAs and CN-LOH in neoplastic disorders

Fady Mikhail, *University of Alabama at Birmingham*

Implementation survey of the ACMG/CGC standards for interpretation of acquired CNAs and CN-LOH in neoplastic disorders

Kathleen Schieffer, Nationwide Children's Hospital

Early Career / CCGA / Webinar Overview

Evaluation of Hi-C versus Optical Genome Mapping for Diagnosing Constitutional Genomic Structural Variants

He Fang, University of Washington

Endothelial cells are a key target of IFN-g during response to combined PD-1/CTLA-4 ICB treatment in bladder cancer

Sharon Freshour, Washington University School of Medicine

HPV forms chimeric virus-human transcripts that affect host gene expression in cervical tumors

Kay Jayachandran, Washington University School of Medicine

Concurrent Systemic Mastocytosis and T-Lymphoblastic Lymphoma Unified by a Novel Cryptic JAKMIP2::PDGFRB Rearrangement Kevin Shopsowitz, *University of British Columbia*

Loss of MSH2 and MSH6 is frequently observed in prostate neoplasms with mismatch repair deficiency

Gokce Toruner, The University of Texas MD Anderson Cancer Center

6:45 – 7:00 PM Platinum Vendor Showcase: Pacific BioSciences

(No CME or CEUs available)

7:00 – 7:15 PM Platinum Vendor Showcase

(No CME or CEUs available)

7:15 - 9:00 PM Welcome Reception

Exhibit Hall

MONDAY, AUGUST 14, 2023

8:00 – 9:00 AM Session 3: Genomic resources for variant curation and standardization 1

ClinGen Cancer Variant Interpretation (CVI) Committee: Pilot Guidance for Somatic Cancer Variant Curation Expert Panels

Deborah Ritter, Baylor College of Medicine

Djerba: A Modular System to Generate Clinical Genome Interpretation Reports for Cancer

lain Bancarz, Ontario Institute for Cancer Research

Investigation of pathogenic and truncated variants of RUNX1 and DDX41 in All of Us

Huan Mo, National Human Genome Research Institute

Developing a generalized model for variants in CIViC

Arpad Danos, Washington University in St. Louis

9:00 – 10:00 AM Keynote Presentation

Olufunmilayo Olopade, University of Chicago

10:45 – 11:45 AM Session 4: Solid Tumors 1

Comparative analysis of RNA expression identifies druggable targets in difficult-to-treat pediatric solid tumors

Yvonne Vasquez, University of California, Santa Cruz

Comprehensive 'Omic' Profiling Reveals 'Atypical Oligodendrogliomas' which Challenge CNS Diagnostic Classification

Adrian Dubuc, Brigham and Women's Hospital

Cell-free DNA genomic and epigenomic analysis to predict survival in mCRPC patients treated with AR-directed therapy

Predeep Chauhan, Washington University School of Medicine

Optical Genome Mapping Reveals New Insights into ZFTA Fusion in Supratentorial Ependymomas

Jianling Ji, Children's Hospital Los Angeles, USC

11:45 – 12:15 PM Speed Abstracts Session II

Clinical Utility of Optical Genome Mapping: Comparison with Standard Cytogenomics Work-up for Hematological Malignancies

Gokce Toruner, The University of Texas MD Anderson Cancer Center

High-risk genetic variants underlie unfavorable prognosis of Blymphoblastic leukemia patients of Hispanic ethnicity

Wengyn Maximilian, Children's Hospital Los Angeles

Improving interoperability of therapeutics and their targets for clinical and precision medicine applications

Matthew Cannon, Nationwide Children's Hospital

Five-year Experience of Evaluating Individuals At-risk for Underlying Genetic Predisposition to Hematologic Malignancy

Min Fang, Fred Hutchinson Cancer Center

Analytical validation of an optical genome mapping assay for structural variant detection in hematologic malignancies

Trilochan Sahoo, Bionano Laboratories

12:15 – 12:30 PM Platinum Vendor Showcase: Thermo Fisher Scientific

Using OncoScan to address challenges calling genomic instability in solid tumor samples.

(No CME or CEUs available)

12:30 – 1:00 PM Diamond Vendor Showcase: Bionano Genomics

Panel Discussion

Moderator: Alka Chaubey, Bionano Genomics

(No CME or CEUs available)

1:00 – 2:00 PM Buffet Lunch

(Exhibit Hall)

2:00 – 3:00 PM Session 5: Hematological Malignancies 1

Real-World Analysis of Cytopenic Patients for Identification of Clonal

Cytopenia(s) of Undetermined Significance (CCUS)

Anwar Iqbal, University of Rochester Medical Center

Chromosomal Microarray Analysis Work-up for Hypocellular MDS Patients with Inconclusive Cytogenetics

Ha Nguyen, Northwestern Medicine

Technologist Award

Prognostic significance of copy number gain of MYC detected by FISH analysis in large B-cell lymphoma

Madina Sukhanova, Northwestern Medicine Feinberg School of Medicine

Prognostic significance of copy number gain of MYC detected by FISH analysis in large B-cell lymphoma

Victória Tomaz, Hospital Israelita Albert Einstein

Trainee Award

3:00 – 3:30 PM Invited Speaker Presentation

Francesc Sole, Josep Carreras Leukaemia Research Institute

3:30 – 4:00 PM Session 6: Bioinformatics and Genomic Resources

Optical Genome Mapping identifies additional cytogenetic abnormalities in patients with hematologic malignancies

Sachin Jadhav, Healthcare Global

Comprehensive Genomic Characterization of Infantile Cancers Reveals High Yield of Therapeutically Targetable Alterations

Mariam Mathew, Nationwide Children's Hospital

4:45 – 5:00 PM Platinum Vendor Showcase

(No CME or CEUs available)

5:00 – 5:30 PM Speed Abstracts Session III

Evolution of a variant curation procedures in the open-access cancer variant interpretation knowledgebase CIViC

Kilannin Krysiak, Washington University School of Medicine

Gene Normalizer: a tool to resolve genetic ambiguity through data harmonization

Anastasia Smith, The Ohio State University

Cell-type-specific genetic-to-epigenetic relationships in the human breast Axel Hauduc, *University of British Columbia*

Assessment of TRG and TRB Clonality by NGS of Dermatologic Specimens is Impacted by Biopsy Type, DNA and Amplicon Sizes

Jane Houldsworth, Icahn School of Medicine at Mount Sinai

Genomic microarray analysis reveals heterogeneity in high hyperdiploid B-cell acute lymphoblastic leukemia

Julie Feusier, ARUP Laboratories - Phoenix Children's

5:30 – 6:00 PM Spotlight Symposium: ISCN Nomenclature

Rosalind Hastings, ISCN Standing Committee Chair & GenQA Consultant

Oxford University Hospitals NHS Foundation Trust

6:00 – 7:00 PM Poster Viewing Session

Exhibit Hall

7:00 – 8:00 PM Early Career Social

For attendees in training or recently out of training

TUESDAY, AUGUST 15, 2023

8:00 – 9:00 AM Session 7: Bioinformatics, Artificial Intelligence, Machine Learning 2

Cell-free DNA fragmentation profiling as a method for tumor fraction assessment and treatment monitoring in NSCLC

Zachary Skidmore, Delfi Diagnostics

Resolving Ambiguities in Copy Number Variation Representation Kori Kuzma, *Nationwide Children's Hospital*

Automated Deep Aberration Detection from Chromosome Karyotype Images

Min Fang, Fred Hutchinson Cancer Research Center

Fusion Curation Interface: an educational tool to explore a unified framework for representing & curating gene fusions

Kathryn Stahl, Nationwide Children's Hospital

Technologist Award

9:00 – 10:00 AM Keynote Presentation

Guillermo Garcia-Manero, MD Anderson Cancer Center

10:45 – 11:45 AM Session 8: Hematological Malignancies 2

Frequency and Etiology of Cytogenetically Cryptic Oncogenic Fusions in Pediatric AML

Gordana Raca, Children's Hospital Los Angeles

Clonal Hematopoiesis in Childhood Cancer Survivors

Irenaeus Chan, Washington University in St. Louis - School of Medicine

Integrative Cytogenetic and Molecular Studies Unmasks 'Chromosomal Mimicry' in Hematologic Malignancies

Adrian Dubuc, Brigham and Women's Hospital

Optical genome mapping in hematological malignancy: Clinical outcomes

in a 2-year follow-up retrospective study

Nikhil Sahajpal, *Greenwood Genetic Center*

11:45 AM – 12:15 PM Spotlight Symposium: WHO/IARC Overview Dilani Lokuhetty, International Agency for Research on Cancer (IARC) 12:15 - 12:30 PM Platinum Vendor Showcase: Imagia Canexia Health (No CME or CEUs available) 12:30 - 12:45 PM **Platinum Vendor Showcase: Qiagen** (No CME or CEUs available) **Round Table Discussions with Lunch** 12:45 - 2:00 PM Please sign up for table topics during conference registration. 2:00 - 3:00 PM Session 9: Applications of Emerging Technologies in Clinical Genomics 2 Targeted RNA-Seq on fresh frozen and methanol/acetic acid fixed cells in diagnostic workup of hematologic malignancies Xiaoyu Qu, Fred Hutchinson Cancer Center A novel method for detection of loss of heterozygosity using B-allele frequency from optical genome mapping data Aliz Raksi, Bionano Laboratories - Tesa Consulting Analysis of Gene Rearrangements in Neoplasms with Hi-C Sequencing **Using Fresh-Frozen and FFPE Specimens** Yajuan Liu, University of Washington-Seattle Rare SRY-positive derivative X chromosome in female fetus with apparently normal development Casey Brewer, Cincinnati Children's Hospital Medical Center 3:00 - 3:30 PM **Invited Speaker Presentation** Joseph Khoury, University of Nebraska Medical Center 3:30 - 4:15 PM **Coffee Break** 4:15 - 4:30 PM Platinum Vendor Showcase: SOPHiA Genetics Liquid Biopsy – From individual markers to tumor signal (No CME or CEUs available) 4:30 - 5:00 PM **Speed Abstracts Session IV**

ClinGen Pediatric Cancer Taskforce initiatives to advance pediatric clinical interpretations through expert curation

Jason Saliba, Washington University School of Medicine

Uveal Melanoma - The New Zealand Perspective

Amanda Dixon-McIver, IGENZ

VMD4Kids: A highly sensitive NGS panel to detect low-level mosaic variants in vascular anomalies & overgrowth disorders.

Avinash Dharmadhikari, *Children's Hospital Los Angeles*

Clinical validation and Implementation of exome, transcriptome and whole genome sequencing for pediatric cancers

Alexandre Rouette, CHU Sainte-Justine - Molecular Diagnostic Lab

Whole-exome sequencing identifies somatic mutations penile squamous cell carcinoma

Kelly Duarte, University of Sao Paulo

5:00 – 5:30 PM Spotlight Symposium: GOAL Consortium

Dara Aisner, *University of Colorado* Jeremy Segal, *University of Chicago*

6:00 – 9:00 PM Optional After Hours Social Activities

WEDNESDAY, AUGUST 16, 2023

8:30 – 9:30 AM Session 10: Genomic resources for variant curation and standardization

Curating Variants of Established Clinical Significance

Mariam Khanfar, Washington University School of Medicine

Implementing the ClinGen/CGC/VICC Oncogenicity Guidelines in a Pediatric Variant Classification Workflow

Wesley Goar, Nationwide Children's Hospital

Oncogenic assessment of FLT3 Variants by the ClinGen FLT3 Somatic Cancer Variant Curation Expert Panel

Jason Saliba, Washinton University School of Medicine

Feasibility, Accuracy and usability analysis of MapAML, a first-in-class app for integrated diagnosis in AML

Thais Moyen, Hospital Israelita Albert Einstein

9:30 – 10:30 AM Keynote Speaker

David Beck, New York University School of Medicine

10:30 - 11:15 AM Coffee Break

11:15 – 12:15 PM Session 11: Solid Tumors 2

A cell-free DNA 5-hydroxymethylcytosine marker predicts immunotherapy response in lung cancer

Zejuan Li, Houston Methodist

Profiling PIK3CA Variants - a highlight of C2 domain variants in Disorders of Somatic Mosaicism

Yang Cao, Washington University in St. Louis

A female-specific chimeric RNA with differential expression in COVID patients

Xinrui Shi, University of Virginia

Trainee Award

Utilizing Rapid Molecular Testing to Reduce Disparites in Pediatric Cancer in Sub-Saharan Africa

Julie Gastier-Foster, Baylor College of Medicine/Texas Children's Hospital