



Cancer Genomics Consortium, 16th Annual Meeting

August 3-6, 2025

The Royal Sonesta Houston Galleria, Houston, Texas

All session times are in US Central Daylight Time.

SATURDAY, AUGUST 2, 2025

In-Person Unconference

Unconference participation incurs an additional registration fee.

9:00 AM – 5:00 PM **In-Person Variant Interpretation Unconference**
Presenters TBA

6:30 – 8:30 PM **Evening Social for Unconference Attendees**
Location TBA

SUNDAY, AUGUST 3, 2025

Pre-Meeting Workshops

Workshop participation incurs an additional registration fee.

8:30 – 10:00 AM **Navigating Academic vs. Non-academic Clinical Pathways and Achieving Success/Happiness in Academic Medicine**
Presenters:
Rachel Burnside, *University of Florida College of Medicine, FL*
Marilyn Bui, *Moffitt Cancer Center and Research Institute, FL*

10:00 – 10:15 AM **Coffee Break for Workshop Attendees**

10:15 – 11:45 AM **Bioinformatics Workshop on Recurrent Biomarkers in Cancer Genomes**
Presenters:
JianJiong Gao, *Memorial Sloan Kettering Cancer Center, NY*
Olena Vaske, *University of California Santa Cruz, CA*
Jinghui Zhang, *St. Jude Children's Research Hospital, TN*

Opening of the 16th Annual Cancer Genomics Consortium

1:00 – 1:05 PM **Welcome**

Fady Mikhail, *University of Alabama at Birmingham, AL*

1:05 - 1:15 PM

Presidential Address

Jane Houldsworth, *Icahn School of Medicine at Mount Sinai, NY*

1:15 – 2:15 PM

Keynote Presentation

Clinical application of genomics for childhood cancer patients - an update on two decades of precision oncology

Will Parsons, *Baylor College of Medicine and Texas Children's Hospital, TX*

2:15 – 3:15 PM

Platform Session 1: Solid Tumors

Rare oncogenic structural variations in *FGFR* genes in childhood brain tumors provide potential therapeutic targets

Mary Clay Bailey, *Baylor College of Medicine, TX*

Chromothripsis in meningiomas is associated with more aggressive behavior

Ha Nguyen, *Northwestern Medicine, IL*

Clinicopathologic features and outcomes for colorectal carcinomas with *KRAS* codon 146 mutations

Bryan Iorgulescu, *MD Anderson Cancer Center, TX*

Complex copy number variation in atypical melanocytic neoplasms and early melanoma: A focused study of 60 cases

Katherine Geiersbach, *Mayo Clinic, MN*

3:15 – 4:00 PM

Coffee Break

Science Café Presentations

4:00 – 4:30 PM

Diamond Exhibitor Showcase

(No CME or CEUs available)

4:30 – 4:45 PM

CGC Annual Award

4:45 – 5:15 PM

CGC 2025 Business Meeting

5:15 – 6:15 PM

Platform Session 2: Variant Curation, Interpretation, and Standardization

Advancing gene curation in hereditary cancer: A comprehensive re-curation of breast/ovarian and colon cancer genes

Jasmine Baker, *Baylor College of Medicine, TX*

The landscape of germline and somatic cancer variants in tumor suppressor genes

Suhasini Lulla, *Baylor College of Medicine, TX*

Application of expert panel-derived oncogenicity guidelines in *BCR::ABL1*-like B-lymphoblastic leukemia/lymphoma

Mark Evans, *Caris Life Sciences, UT*

Systematic curation of defining and supportive diagnostic variants to enhance genomic diagnostics in pediatric oncology

Morteza Seifi, *ClinGen, WI*

6:15 – 8:15 PM

Welcome Reception

CGC Exhibit Hall, Legends Ballroom

MONDAY, AUGUST 4, 2025

8:30 – 9:30 AM

Platform Session 3: Emerging Technologies in Clinical Genomics and Technical Lab Topics

Clinical utility of optical genome mapping (OGM) for B-cell acute lymphoblastic leukemia (B-ALL)

Angela Lager, *University of Chicago, IL*

Optical genome mapping on pediatric leukemia samples: A single hospital experience

John Herriges, *Children's Mercy Hospital, Kansas City, MO*

Impact of long-term plasma storage on cell free DNA biomarker studies

Zejuan Li, *Houston Methodist Hospital, Weill Cornell Medical College, TX*

Analytical validation and cross-platform analysis of the oncomine TCR pan-clonality assay for T-Cell receptor profiling

Mohamed Maher, *University of Texas MD Anderson Cancer Center, TX*

9:30 – 10:30 AM

Invited Session 1

Federal regulations affecting molecular diagnostic testing and gene patenting

Donald Karcher, *George Washington University Medical Center*

Roger Klein, *Arizona State University Law School*

10:30 – 11:15 AM

Coffee Break with Exhibitors

Science Café Presentations

11:15 – 11:30 AM

Platinum Exhibitor Showcase

(No CME or CEUs available)

11:30 AM – 12:00 PM

CGC Updates

12:00 – 1:00 PM

Platform Session 4: Hematologic Malignancies

Diagnostic utility of FISH testing for Ph-like B-ALL: A 5-year institutional experience

Fady Mikhail, *University of Alabama at Birmingham, AL*

Genomic characterization of *CRLF2*-rearranged pediatric B-ALL using transcriptome sequencing (RNA-Seq)

Gordana Raca, *Children's Hospital Los Angeles, CA*

Genomic Proximity Mapping (GPM): Evaluation of a next generation cytogenomic assay for acute myeloid leukemia

Min Fang, *Fred Hutchinson Cancer Center, WA*

Comparative analysis of targeted RNA-Seq and optical genome mapping for detecting clinically significant gene rearrangements

Gokce Toruner, *University of Texas MD Anderson Cancer Center, TX*

1:00 – 2:00 PM

Buffet Lunch and Networking

Exhibit Hall and Foyer

2:00 – 3:00 PM

Invited Session 2

Genome mapping and sequencing technologies for characterizing structural variants

Lisa Lansdon, *Children's Mercy Hospital, Kansas City and University of Missouri-Kansas City School of Medicine, MO*

Fritz Sedlazeck, *Baylor College of Medicine, TX*

3:00 – 4:00 PM

Keynote Presentation

Molecular-genomic findings in leukemias: Diagnostic, prognostic, and therapeutic implications

Hagop Kantarjian, *University of Texas MD Anderson Cancer Center, TX*

4:00 – 4:45 PM

Coffee Break with Exhibitors

Science Café Presentations

4:45 – 5:30 PM

Speed Abstract Session 1: Solid Tumors, Bioinformatics, Artificial Intelligence, Emerging Technologies, and Other Advances in Clinical Genomics

Tumor mutation burden (TMB) as a biomarker in advanced and metastatic breast cancers - An analysis of clinical NGS data in a 74 cases series

Liu Liu, *Brown University Health, RI*

Multi-modal validation and pan-cancer analysis of 9p21 loss: Diagnostic and prognostic implications

Holly Hill, *University of Texas MD Anderson Cancer Center, TX*

Expanding MAVE data maps for use in human genomics applications

Jeremy Arbesfeld, *The Ohio State University, OH*

New Q-Band artificial intelligence-based chromosome analysis and karyotyping - A pilot evaluation

Cristina Steele, *Applied Spectral Imaging for Laboratorio Clinico Colcan, Columbia*

5-hydroxymethylcytosine as a liquid biopsy biomarker in colorectal cancer

Christopher Maher, *Washington University School of Medicine, MO*

Accurate and reliable detection of clonal hematopoiesis in plasma cell-free DNA

Alyssa Parker, *Vanderbilt University, TN*

5:30 – 6:30 PM

Poster Session (Odd Numbered Posters), Cash Bar

CGC Exhibit Hall

7:00 PM

Early Career Social

For attendees in training or recently out of training

TUESDAY, AUGUST 5, 2025

8:30 – 9:30 AM

Platform Session 5: Bioinformatics, Artificial Intelligence, and Machine-Learning

Prediction of BRAF V600 mutation status in cutaneous melanoma using an explainable deep learning model

Vibha Rao, *Dartmouth Health, NH*

Prediction of gene expression in NSCLC tumors and their microenvironment using an explainable machine learning model

Shrey Sukhadia, *Dartmouth Hitchcock Medical Center, NH*

Improving fusion detection sensitivity in the TruSight Oncology 500 Panel through DNA-based structural variant analysis

Qiliang Ding, *Mayo Clinic, MN*

Improved copy number estimation in central nervous system (CNS) tumors using methylation array

Drew Duckett, *Northwestern University, IL*

9:30 – 10:30 AM

Keynote Presentation

Discovering and defining the telomere biology disorders

Sharon Savage, *National Cancer Institute, National Institutes of Health, MD*

10:30 – 11:15 AM

Coffee Break with Exhibitors

Science Café Presentations

11:15 – 11:30 AM

Platinum Exhibitor Showcase

(No CME or CEUs available)

11:30 – 11:45 AM

Platinum Exhibitor Showcase

(No CME or CEUs available)

11:45 AM – 12:45 PM

Invited Session 3

Cancer genome bioinformatics in a clinical setting

Mark Cowley, *Children's Cancer Institute, Australia*

Sheryl Elkin, *QIAGEN Digital Insights, MA*

12:45 – 2:00 PM

In-Person Round Table Discussions with Lunch

Table topic selection is during conference registration.

2:00 – 2:45 PM

Speed Abstract Session 2: Variant Curation, Interpretation and Standardization, and Equity/Access in Genomic Medicine

Development of standards for the assessment of gene fusion oncogenicity

Karen Tsuchiya, *Nationwide Children's Hospital, OH*

The actionable transcriptome: A framework for incorporating transcriptional profiling into precision oncology

Amber Johnson, *MD Anderson Cancer Center, TX*

Introduction of an online portal for cancer-specific gene list curation
Beth Pitel, *Mayo Clinic, MN*

Enhancing data standardization and structure to improve clinical variant classification

Wesley Goar, *Nationwide Children's Hospital, OH*

Somatic variant classifications in ClinVar: An update after one year

Melissa Landrum, *NCBI/NLM/NIH, MD*

Clinical impact of in-house molecular testing of solid tumors in resource-limited settings: A pilot study in Pakistan

Romena Qazi, *Shaukat Khanum Memorial Cancer Hospital and Research Centre, Pakistan*

2:45 – 3:45 PM

Poster Session (Even Numbered Posters)

CGC Exhibit Hall

3:45 – 4:30 PM

Coffee Break with Exhibitors

Science Café Presentations

4:30 – 4:45 PM

Platinum Exhibitor Showcase

(No CME or CEUs available)

4:45 – 5:00 PM

Platinum Exhibitor Showcase

(No CME or CEUs available)

5:00 – 6:00 PM

Panel Discussion on the Integration of Cytogenetics and Molecular Genetics Laboratories

Panelists:

Yasmine Akkari, *Nationwide Children's Hospital, OH*

Shashikant Kulkarni, *MD Anderson Cancer Center, TX*

Gordana Raca, *Children's Hospital Los Angeles, CA*

Dayna Wolff, *Medical University of South Carolina, SC*

7:00 – 8:30 PM

Optional After Hours Social Activities

Register for activities during meeting registration. Participants can pick up tickets at the CGC 2025 Registration Desk.

Ice at the Galleria

Museum of Illusion

WEDNESDAY, AUGUST 6, 2025

9:00 – 10:00 AM

Platform Session 6: Germline/Constitutional Genomics, Mosaicism, and Equity/Access in Genomic Medicine

Assessing the efficacy of additional cell analysis in detecting low-level constitutional mosaic karyotypes

Aiko Otsubo, *University of Michigan, MI*

Best practices for testing low-level mosaic variants: Recommendations from CGC somatic overgrowth & vascular anomalies working group
Avinash Dharmadhikari, *Children's Hospital Los Angeles, CA*

Genetic regulation of inflammatory proteomes: Bridging genomic disparities between African and European populations
Caroline Amour, *Kilimanjaro Christian Medical University College, Tanzania*

Current state of molecular cancer testing in resource-limited and underserved communities
Casey Brewer, *Cincinnati Children's Hospital Medical Center, OH*

10:00 – 11:00 AM

Invited Session 4

Clinical variant interpretation knowledgebases in cancer genomics

Beth Pitel, *Mayo Clinic and Variant Interpretation for Cancer Consortium (VICC), MN*

Alex Wagner, *Nationwide Children's Hospital and Variant Interpretation for Cancer Consortium (VICC), OH*

11:00 – 11:30 AM

Coffee Break with Exhibitors

11:30 AM – 12:30 PM

Platform Session 7: Diagnostic Challenge Case Studies

Detection of *IGH::MYC* and a cryptic *IGH::BCL6* rearrangement refines diagnosis in high-grade B-Cell lymphoma

Will Coward IV, *University of North Carolina Healthcare Cytogenetics Laboratory, NC*

Decoding the genetic complexity of B-ALL through long-read and RNA sequencing methods

Jiadi Wen, *Yale School of Medicine, CT*

Rare *TBL1XR1::JAK2* fusion in a patient with pediatric T-ALL identified by optical genome mapping and long-read sequencing

Aravindh Nagarajan, *Children's Mercy Hospital, Kansas City, MO*

A rare recurrent 11;19 rearrangement resulting in *KMT2A::MYO1F* fusion in pediatric AML

Manisha Sutariya, *University of Minnesota, MN*