

THURSDAY, APRIL 18, 2024 PRE-CONFERENCE PRECISION MEDICINE IN PRACTICE PRIMER AND CASE-BASED WORKSHOP

APRIL 19-21, 2024 | FAIRMONT CHICAGO MILLENNIUM PARK, CHICAGO, IL

ACCELERATING PROGRESS IN PRECISION MEDICINE FOR CANCER: THE TIME IS NOW





THIS ACTIVITY IS BROUGHT TO YOU IN PARTNERSHIP WITH CITY OF HOPE

INFORMATION

COURSE DESCRIPTION

Rapid advances in genomic technologies have revolutionized our approach to risk-adapted cancer treatments and screening for early detection or prevention of cancer, heralding the era of precision health care across the continuum of cancer care. Knowledge of the genomic landscape and immune microenvironment of common cancers has fueled the need to identify cancer patients and healthy individuals at risk of the most lethal but treatable types of cancers where more intensive surveillance and personalized management plan can be implemented to improve overall survival. A classic example is identifying BRCA1 gene mutation carriers to preempt or prevent the development of lethal basal-like breast cancers and treating BRCA-Associated cancer with Parp Inhibitors. Identifying Lynch Syndrome patients led to the development of curative neoadjuvant immunotherapy approaches for Rectal Cancer.

Accelerating Progress in Precision Medicine for Cancer: The Time is Now will provide an interdisciplinary educational forum in which to update state-of-the-art science in cancer genomics, including Artificial Intelligence, discuss real-world cases, curate interesting pedigrees, and prepare case summaries to disseminate new knowledge and evidence-based management of high-risk patients and their at-risk family members. The goal of this educational activity is to equip healthcare providers with practical tools and knowledge to accelerate progress in precision medicine for cancer.

TARGET AUDIENCE

This interdisciplinary and inter-professional educational activity has been designed for health care providers including advanced nurse practitioners, genetic counselors, oncology nurses, pharmacists, physicians, physician assistants, and other healthcare professionals involved in providing cancer genomics risk assessment services or initiating such services in their health care system. Special efforts will be made to meet the needs of clinicians practicing in non-academic centers and geographically and/or socioeconomically underserved communities.

LEARNING OBJECTIVES

At the conclusion of this activity, participants will be able to:

Review advances in genetics, genomics, and "Omics" technologies transforming precision medicine for cancer;

- Gain a better understanding of controversies in germline genomic testing for risk-adapted screening and integration of polygenic risk score for cancer prevention;
- Recognize advances in clinical research with special emphasis on patientcentric and biomarker informed clinical trials in high risk populations;
- Identify molecular determinants of response to therapy in hematologic malignancies and solid tumors;
- Appreciate patient advocacy and partnerships to accelerate progress in development of novel therapies and vaccines for cancer control;
- Learn how Artificial Intelligence is transform health care to promote health equity in the US and across the world.

CONFERENCE LOCATION

Fairmont Chicago Millennium Park 200 N. Columbus Drive Chicago, IL 60611 United States

ACCREDITATION AND CREDIT DESIGNATION

PHYSICIAN CREDIT

The University of Chicago Pritzker School of Medicine is accredited by the Accreditation Council for Continuing Medical Education (ACCME) to provide continuing medical education for physicians.

The University of Chicago Pritzker School of Medicine designates this live activity for a maximum of 21.50 *AMA PRA Category 1 Credits*[™]. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

INFORMATION

ACCREDITATION AND CREDIT DESIGNATION CONTINUED

GENETIC COUNSELOR CREDIT

The National Society of Genetic Counselors (NSGC) has authorized the University of Chicago Pritzker School of Medicine to offer up to 2.150 CEUs or 21.50 Category 1 contact hours for the activity 12th International Clinical Cancer Genomics Conference – Accelerating Progress in Precision Medicine for Cancer: The Time is Now! The American Board of Genetic Counseling (ABGC) will accept CEUs earned at this program for the purposes of genetic counselor certification and recertification.

AMERICAN BOARD OF INTERNAL MEDICINE MOC PART II CREDIT

Successful completion of this CME activity, which includes participation in the evaluation component, enables the participant to earn up to 21.50 MOC points in the American Board of Internal Medicine's (ABIM) Maintenance of Certification (MOC) program. Participants will earn MOC points equivalent to the amount of CME credits claimed for the activity. It is the CME activity provider's responsibility to submit participant completion information to ACCME for the purpose of granting ABIM MOC credit.

NURSING CREDIT

University of Chicago Medicine is accredited as a provider of continuing nursing professional development by the American Nurses Credentialing Center's Commission on Accreditation.

Participants who successfully complete the entire activity and complete an evaluation form will earn 21.50 contact hours.

OTHER HEALTHCARE PROFESSIONS CREDIT

Other healthcare professionals will receive a Certificate of Participation. For information on the applicability and acceptance of Certificates of Participation for educational activities certified for AMA PRA Category 1 Credit[™] from organizations accredited by the ACCME, please consult your professional licensing board.

CME CERTIFICATES/ COURSE EVALUATION

The evaluation will be done when the participant comes to claim credit. Go to www.cancergeneticsconference.com/ to get information on how to claim CME.

PLANNED BY





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NON PROFIT EXHIBITORS:

- Clinical Cancer Genomics Community of Practice/ City of Hope
- My Faulty Gene
- National Consortium of Breast Centers
- WISDOM/CAPS

FACULTY

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CONTINUED...

CLINICAL CANCER GENOMICS CONFERENCE

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AGENDA

THURSDAY, APRIL 18, 2024

PRE-CONFERENCE: Precision Medicine in Practice Primer and Case-Based Workshop ROUGE ROOM (1ST FLOOR)

1:00 pm	WELCOME Kathleen Blazer, EdD, MS, CGC
1:10 pm	The Two Genomes of Cancer in Precision Medicine: Overview of the Tumor and Germline Genomes- Key Concepts and Terms in Precision Medicine Padma Sheila Rajagopal, MD, MPH, MSc, NCI
1:30 pm	Technical Considerations in Tumor and Germline Testing: What Test Tells Us What Colin Pritchard, MD, PhD
1:50 pm	Fundamental Concepts in Targeted Cancer Therapeutics Katherine Roth, MD
2:10 pm	Germline Signatures Indicating Candidacy for Targeted Treatments and Clinical Trials and Somatic Signatures Indicating Candidacy for Germline Testing Joanne Jeter, MD
2:40 pm	Q&A
2:50 pm	BREAK
3:10 pm	BREAKOUT SESSIONS ROUGE ROOM (1 st FLOOR) Putting It All Together: Team-Based Learning (TBL)
Moderators:	Kathleen Blazer, EdD, MS, CGC Sandra Dreike, MS, CGC Rachelle Manookian, MS, CGC Bita Nehoray, MS CGC Christina Rybak, MS, CGC Susan Shehayeb, MS, CGC Elise Sobotka, MS, MPH, CGC Ilana Solomon MA, ScM, CGC Elyssa Zukin, MS, CGC
3:50 pm	Team-Based Learning Case Presentations with Panel Discussion
Moderators:	Listed above
Faculty Panel:	Joanne Jeter, MD, MS Colin Pritchard, MD, PhD Padma Sheila Rajagopal, MD, MPH, MSc, NCI Kathleen Roth, MD
5:00 pm	ADJOURN

FRIDAY, APRIL 19, 2024

7:00 am REGISTRATION, BREAKFAST AND EXHIBITS INT'L. BALLROOM FOYER

SESSION 1: Survivorship in Patients at Risk for Multiple Cancers INTERNATIONAL BALLROOM

Moderators:	Lorraine Canham, MD Sharon Savage, MD	
8:00am	WELCOME & ANNOUNCEMENTS Iris Romero MD, MS	
8:10am	Celebrating 50 Years of Excellence in Cancer Research: University of Chicago Cancer Center and the CCGOP Kathleen Blazer, EdD, MS, CGC Olufunmilayo Olopade, MD	
8:30 am	The Complex Needs of Families at a High Genetic Risk of Cancer – Learning from Telomere Biology Disorders Sharon Savage, MD	
8:55 am	KEYNOTE ADDRESS I: Li-Fraumeni Syndrome as a Case Study of the Effect of Early Diagnosis on Outcomes Judy Garber, MD, MPH	
9:20 am	Genomic Testing in Childhood: Lessons Learned About Family Dynamics Melody Perpich, MS, CGC	
9:35 am	Q&A	
9:55 am	BREAK/EXHIBITS	
SESSION 2: Gastro-intestinal Diseases INTERNATIONAL BALLROOM		
Moderators:	Sonia Kupfer, MD Jeremy Segal, MD, PhD	
10:30 am	KEYNOTE ADDRESS II: Toward Precision Prevention: Vaccines in Lynch Syndrome Matthias Kloor, MD	
11:00 am	New Mechanisms of Carcinogenesis in Lynch Syndrome & Clinical Implications Aysal Ahadvoa, PhD	
11:20 am	The Rising Tide of Colorectal Cancer in Younger Adults: Why Is This Happening & What Do We Do About It? Cathy Eng, MD	
11:40 am	Shifting Paradigms in CDH1 Management Ophir Gilad, MD	
12:00 pm	Breakthroughs in Molecular Diagnosis Colin Pritchard, MD, PhD	

12:20 pm **Q&A**

12:40 pm **GENERAL LUNCHEON SEATING** STATE ROOM

12:50 pm OPTIONAL NON CME PRODUCT THEATER GOLD ROOM

Presented by Cancer IQ Expert Roundtable on the Value of Genetic Testing for Cancer Risk Assessment

SESSION 3: Concurrent Case Presentations with Interprofessional Panel Discussions

with inter	rprofessional Panel Discussions
1:45 pm	1. Molecular Tumor Boards – How Are Suspected Germline Variants Typically Handled at Your Institution's Molecular Tumor Board: Is There a Set-up Process for Germline Referral/Testing? ROUGE ROOM (1 ST FLOOR)
Moderators:	Cara Essling, MMSPA Bita Nehoray, MS, CGC
Faculty Panel:	Kara Maxwell, MD, PhD Colin Pritchard, MD Padma Sheila Rajagopal, MD, MPH, MSc, NCI Ari Rosenberg, MD Jeremy Segal, MD, PhD Rajat Thawani, MBBS
	2. Breast, Ovarian, GI, Endometrial Cancers High and Moderate Risk, & Risk Management INTERNATIONAL BALLROOM
Moderators:	Christine Drogan, MS, CGC Emma Keel, MS, CGC
Faculty Panel:	Cathy Eng, MD Judy Garber, MD, MPH Matthias Kloor, MD Sonia Kupfer, MD Jeffrey Weitzel, MD
	3. Hematologic, Pediatric and New Syndromes CUVEE (1ST FLOOR)
Moderators:	Rachelle Manookian, MS, CGC Melody Perpich, MS, CGC
Faculty Panel:	Lorraine Canham, MD Jane Churpek, MD, MS Michael Drazer, MD, PhD William Foulkes, MBBS, PhD Sharon Savage, MD
2:45 pm	BREAK/EXHIBITS
Early Diag	4: New Approaches to gnosis and Treatment ed Cancers IONAL BALLROOM
Moderators:	Jane Churpek, MD, MS Padma Sheila Rajagopal, MD, MPH, MSc, NCI
3:00 pm	KEYNOTE ADDRESS III: The Promise of Multi-Cancer Detection Assays Peter Kuhn, MD
3:30 pm	MicroRNA Biogenesis Proteins, Ontogenesis to Oncogenesis William Foulkes, MBBS,PhD
	CONTINUED

PROGRAM AGENDA AND SPEAKER SELECTION SUBJECT TO CHANGE.

3:50 pm **Predicting Response and Toxicities** to Immunotherapy Peter O'Donnell, MD 4:30 pm Q&A

4:50 pm ADJOURN

5:00 pm **POSTER RECEPTION**

GOLD ROOM Food and beverage to be served

See back pages for poster numbers and presenters

SATURDAY, APRIL 20, 2024

- **REGISTRATION, BREAKFAST** 7:30 am AND EXHIBITS INT'L. BALLROOM FOYER
- ANNOUNCEMENTS 8.10am Kathleen Blazer, EdD, MS, CGC Olufunmilayo Olopade, MD

SESSION 5: Research Highlights INTERNATIONAL BALLROOM

- Sonia Kupfer, MD Moderator: Dengzeng Huo, PhD
- Advanced Personalized Risk 8:15 am Assessment: Making Meaning out of PRS and Model Output Lisa Madlensky, PhD, CGC
- Update from International 8:35 am **Consortia Studies** Katherine Nathanson, MD
- **KEYNOTE ADDRESS IV:** 8:55 am Update from All of Us Study Karriem Watson, DHSc, MS, MPH
- No One Left Behind: Enhancing 9:25 am Equity in Genomic Testing and Screening - Highlights from the WISDOM Study Yiwey Shieh, MD

9:45 am Q&A

10:00 am BREAK/EXHIBITS

SESSION 6: Updates from the Global Community of Practice INTERNATIONAL BALLROOM

10:30 am	RESEARCH PRESENTATIONS
Moderators:	Raja Pramanik, MBBS, MD, DM, BA Jeffrey Weitzel, MD
Faculty Panel:	Oral Abstract Presenters
	Please reference the oral abstract presentations page for more info
11:30 am	Patient Centered Genetic Counseling – What are We Learning from Our Patients?
Moderators:	Grace-Ann Fasaye, ScM, CGC Feighanne Hathaway, MS, CGC
Faculty Panel:	Michelle Bruszer

Wenora Johnson Helen Palmguist Parul Somani

- 12:30 pm **GENERAL LUNCH SEATING** STATE ROOM
- **OPTIONAL NON CME** 12:50 pm PRODUCT THEATER GOLD ROOM Presented by Astra Zeneca **Precision Medicine**

Detecting PIK3CA, AKT1, and PTEN alterations using NGS Justin Grahl, PharmD Precision Medicine and Clinical **Oncology Pharmacist at Froedtert** Cancer Center and the Medical College of Wisconsin

SESSION 7: Using Genetics for Precision Prevention

Speaker:

INTERNATIONAL BALLROOM Iris Romero, MD, MS Moderators: Lucy Godley, MD, PhD 1:30 pm Introduction to Harvey Golomb **Distinguished Lecture Series** Olufunmilayo Olopade, MD **Distinguished Lecturer Lessons** 2:00 pm Learned over 30 years as a Clinical **Cancer Geneticist** Stephen Gruber, MD, PhD, MPH 2:30 pm Q&A **BREAK/EXHIBITS** 2:45 pm **SESSION 8:** Town Hall Case Discussions with Interprofessional Panel 3:00 pm 1. Solid Tumor Breakout INTERNATIONAL BALLROOM Olufunmilayo Olopade, MD Moderator: Rodrigo Santa Cruz Guindalini, MD, PhD Faculty Panel: Stacy Gray, MD Nora Jaskowiak, MD Kara Maxwell, MD, PhD Jeremy Segal, MD, PhD David Vanderweele, MD, PhD 2. Hematologic Malignancies GOLD ROOM Moderators: Michael Drazer, MD, PhD Feighanne Hathaway, MS, CGC Faculty Panel: Daniel Arber, MD Jane Churpek, MD, MS Afaf Osman, MD Caner Saygin, MD Wendy Stock, MD 5:00 pm **ADJOURN** 6:30-**EVENING GALA** 9:30pm ROUGE ROOM (1st FLOOR) Sponsored by Astra Zeneca **Precision Medicine**

SUNDAY, APRIL 21, 2024

SUNDA	AY, APRIL 21, 2024	
7:30 am	REGISTRATION, BREAKFAST AND EXHIBITS INT'L. BALLROOM FOYER	
8:10am	ANNOUNCEMENTS Kathleen Blazer, EdD, MS, CGC Olufunmilayo Olopade, MD	
SESSION 9: At the Frontier of Breast Cancer Research INTERNATIONAL BALLROOM		
Moderators:	Frederick Howard, MD Loren Saulsberry, PhD	
8:15 am	Al for Breast Screening Adam Yala, PhD	
8:35 am	Targeting DNA Repair Pathways and Immune Modulation in TNBC Rita Nanda, MD	
9:00 am	KEYNOTE ADDRESS V: Developing Vaccines for Cancer Interception Susan Domchek, MD	
9:30 am	MRI Surveillance and Mortality in BRCA1 and BRCA2 Mutation Carriers Steven Narod, MD	
10:00 am	Breast Cancer in Young Women – Results from Breast Cancer Family Registry Mary Beth Terry, PhD	
10:30 am	Q&A	
10:45 am	BREAK/EXHIBITS	
SESSION 10: At the Frontier of Ovarian Cancer Research INTERNATIONAL BALLROOM		
Moderators:	Pamela Ganschow, MD Kathryn Mills, MD	

	Kathryn Mills, MD
11:00 am	Clinical Classification of Inherited BRCA2 Variants Fergus Couch, PhD
11:30 am	KEYNOTE ADDRESS VI: Impact of Bilateral Salpingo- Ooporectomy on Cancer Incidence and Mortality in Women with a BRCA1 or BRCA2 Pathogenic Varian Kelly Metcalfe, RN, PhD
12:00 pm	Curing Ovarian Cancer: Are We There Yet From a Surgical Perspective? John Moroney, MD
12:30 pm	Is Advanced Ovarian Cancer Curable; Update on Integration of PARP Inhibitors? Gini Fleming, MD
1:00 pm	Q&A
1:15 pm	CONCLUDING REMARKS Kathleen Blazer, EdD, MS, CGC Tuya Pal, MD Olufunmilayo Olopade, MD

1:30 pm

ORAL ABSTRACT PRESENTATIONS

SESSION 6: RESEARCH PRESENTATIONS

Kenn Chi Ndi, MD Setting Up Cancer Genetics Counseling and Testing Services in Cameroon

Carolin Solomi, V MBBS, DGO, DNB, MRCOG

Genetic Threads of Hope: Unraveling Germline Mutations in Ovarian Cancer, their Impact on Clinical Outcomes and Empowering Ovarian Cancer Defense with Cascade Genetic Testing — an Indian Perspective

Marina Candido Visontai Cormedi, MD

Developing a Polygenic Risk Score for Precision Prevention of HR+/HER2- Breast Cancer in Brazilian Women

Linda Ann Smith, MD

Following Circulating Tumor DNA in Breast Cancer Patients to Enhance Treatment Options

Daniel G. Luba, MD

Hereditary Cancer Screening in a GI Practice in Monterey County, California

Lauren Banaszak, MD

Implementation and Practicality of Guideline-Driven Germline Genetic Evaluation Criteria in the Care of Patients with Myelodysplastic Syndrome and Acute Myeloid Leukemia

Ta-Chueh Melody Rosenberg, MSN, AGACNP-BC, OCN Reducing A Specific Barrier to Cancer Genetic Testing for Veterans: Service Connection Discrimination Concern





POSTERS



POSTER 1

- C

Morgan Devlin, MS, LCGC Ascension St. John Patient Insurance Literacy and Awareness of Financial Resources: A Quality Improvement Project in a Detroit Community Cancer Genetics Clinic

POSTER 2

Lauren Banaszak, MD University of Wisconsin-Madison Implementation and Practicality of Guideline-Driven Germline Genetic Evaluation Criteria in the Care of Patients With Myelodysplastic Syndrome and Acute Myeloid Leukemia

POSTER 3

Vera Cherepakho, MS, LGC Kaiser Permanente Division of Research Validation of Patient Facing Online Risk Assessment Tool for Population Screening of Hereditary Cancer

POSTER 4

Sarah (Nielsen) Young, MS, CGC Invitae Corp. Real-World Breast Surgery Utilization among Breast Cancer Patients with Germline Variants of Uncertain Significance

POSTER 5

Lee Ann Daly, MS, CGC NYU Langone Health Biallelic BRCA2 Mutations and Negative Chromosome Breakage Analysis in a Patient With Adult Onset Breast and Colon Cancers

POSTER 6

Ashley Lynn Hendershot, DNP, FNP-C, ACGN University of Rochester Medical Center Evaluation of a Collaborative Institutional Relationship to Increase Regional Hereditary Cancer Genetic Testing in Individuals with Cancer

POSTER 7

Guillermo Alberto, MD Instituto Alexander Fleming Co-Occurrence of Germline Pathogenic Variants In BRCA1 or BRCA2 and Lynch Syndrome Genes: First Report in South American Population of Three Unrelated Argentinian Families

POSTER 8

Ophir Gilad, MD University of Chicago Birth Cohort Effect on Age of Colorectal Cancer Onset in Lynch Syndrome

POSTER 9

Kenn Chi Ndi, MD Yaounde General Hospital The University of Yaounde 1, Cameroon Setting Up Cancer Genetics Counseling and Testing Services in Cameroon

POSTER 10

Heather Hannon, DNP APRN-NP ACGN Bon Secours Cancer Institute Promoting Adherence to Germline Genetic Testing for Pancreatic Adenocarcinoma

POSTER 11

Britt Johnson, PhD, FACMG Invitae, Corp Tracking Uncertainty in Germline Genetic Testing for Hereditary Cancer Syndromes: Sources, Attributes, and Resolution of Variants of Uncertain Significance in Over 1 Million Individuals

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Ed Esplin, MD, PhD, FACMG, CGAF, FACP Invitae One of These is Not Like The Others:

A Descriptive Study of the Attenuated Phenotype of PMS2

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Jennifer Diaz, AOCNP, CGRA, APRN Medical University of South Carolina **Experience with a Pathogenic Variant Clinic**

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Kristie Bobolis, MD Sutter Medical Group Pure Lobular Carcinoma In Situ Presenting as a Palpable Mass in an Individual with a Germline Pathogenic Variant in CDH1

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Thomas (T.J.) Slavin, MD HALO Research Institute Cancer Risks Associated with Germline Pathogenic Variants in MLH1, MSH2, MSH6, PMS2, and EPCAM genes

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Jennifer Thalita Targino dos Santos, MS University of São Paulo Using Design Thinking to Improve the Identification of Patients At Risk For Hereditary Neoplastic Syndromes

POSTER 17

Alexandra Capasso, MS, CGC City of Hope Genetic Counselors' Facilitation of Informed Consent and the Utility of Supplemental Education Tools for Alternate Approaches to Pre-test Counseling for Hereditary Cancer Predisposition

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Carolin Solomi. V, MBBS, DGO, MS, DNB, MRCOG All India Institute of Medical Sciences, New Delhi, India

Genetic Threads of Hope: Unraveling Germline Mutations in Ovarian Cancer, Their Impact On Clinical Outcomes and Empowering Ovarian Cancer Defense With Cascade Genetic Testing – An Indian Perspective

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Daniel G Luba, MD Monterey Bay GI Consultants Hereditary Cancer Screening in a GI Practice In Monterey County, California

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Owen Mitchell, B.S. University of Chicago Germline Genetic Testing Hesitancy in Patients With Mesothelioma

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Achille Van Christ Manirakiza, MD University of Chicago Germline Sequence Variation in Rwandan Breast and Prostate Cancer Cases

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Stephane Maria Diaz Montes, MD Instituto Nacional de Enfermedades Neoplásicas — Li-Fraumeni Syndrome: A Case Report

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Analí Pamela Mora Alferez, MD Instituto Nacional de Enfermedades Neoplásicas Constitutional Mismatch Repair Deficiency (CMMRD) With An Incidental Finding in ATM Gene in a Patient With a Malignant Brain Tumor

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Cathryn Koptiuch, MS, CGC Department of Veterans Affairs Implementation of a National Cancer Genetics Service for the Veterans Health Administration

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Tessa Bachinski, MS, CGC Advocate Health Cancer Diagnoses in Individuals Identified to Have a CHEK2 Pathogenic or Likely Pathogenic Variant: A Multi-Center, Multi-State Retrospective Cohort Study

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Rachel Epstein, MS, CGC Endeavor Health (formerly NorthShore University HealthSystem) The Breast Health Assessment: A family Health

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Lindsay Fosler, MS, CGC Invitae

The Estimated Healthcare Spend to Identify 1 Hereditary Cancer Syndrome Patient with a Deep Intronic Pathogenic Variant By RNA Sequencing is Nearly 500x More Costly Than Identifying 1 Patient With Multi-Gene Panel DNA Testing

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Lindsay Fosler, MS, CGC Invitae Utility of RNA Analysis in Genetic Testing for Hereditary Cancer is Highly Similar Between Large Cohorts From Two Independent Laboratories

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Marina Candido Visontai Cormedi, MD University of Sao Paulo Developing A Polygenic Risk Score For Precision Prevention of HR+/HER2- Breast Cancer in Brazilian Women

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Michelle Weaver Knowles, FNP, CBCN, ONNCG, ACGN, CGRA Ava Health **Ava Health the Voice for Others**

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Angelina Izguerra, Research Associate, BPH University of Illinois Cancer Center Universal Hereditary Cancer Risk Assessment (uHCRA) with Navigation in an FQHC Primary Care Setting

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Heather Hannon, DNP APRN-NP ACGN Bon Secours Cancer Institute Beyond the Obvious Involving the Genetic Expert is Key to Ensuring Comprehensive, Guideline-Based Care

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Linda Ann Smith, MD X-Ray Associates of New Mexico Following Circulating Tumor DNA in Breast Cancer Patients to Enhance Treatment Options

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Najeb Ullah Khan, PhD Institute of Biotechnology and Genetic Engineering Genetic Susceptibility and Epidemiological Trends of Breast Cancer in the Pashtun Population of Khyber Pakhtunkhwa, Pakistan: Implications for Precision Medicine

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Brenda R. Grimes, PhD Eli Lilly and Company Genomic and transcriptomic profiling of primary tumors from patients with HR+, HER2-, node-positive, high-risk early breast cancer in the monarchE trial

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Susan Shehayeb, MS, CGC City of Hope Prevalence and Penetrance of LZTR1 Pathogenic Variants: To Screen or Not to Screen?

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Elyssa Zukin, MS, CGC City of Hope Uptake of a Chatbot for Disclosure of Germline Genetic Test Results through a Universal Genetic Testing Program

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Bita Nehoray, MS, CGC City of Hope Feasibility of Whole Body MRI and Multicancer Early Detection Testing in People at High Risk for Cancer Development

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Bita Nehoray, MS, CGC City of Hope Variants at Variance: An Assessment of Discordant Classifications and Phenotypes among TP53 Variants from the International LiFT UP Study

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Rachelle Manookian, MS, CGC City of Hope Have We Considered Mosaicism? A Nearly Missed Molecular Diagnosis of von Hippel-Lindau

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Elise Sobotka, MS, MPH, CGC City of Hope **Prevalence of CDKN2A Pathogenic Variants** in a Large Institutional Cohort: Scrutinizing the Common p.Ile49Thr Variant

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Anuja Chitre, MS, CGC City of Hope Prevalence and Penetrance of SPINK1 Pathogenic Variants: A Burden to Patients and Providers

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Anuja Chitre, MS, CGC City of Hope Healthcare Utilization Among Individuals Diagnosed with Lynch Syndrome Through a Universal Germline Genetic Testing Program

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Aleck Cervantes City of Hope Bridging Data For LiFT UP: Successes and Challenges in International Pedigree Data Collaboration

POSTER 45

Rossella Graffeo MD Oncology Institute of Southern Switzerland Breast Cancer Patients with Pathogenic Variants in High Penetrance Genes: Risk Reducing Contralateral Mastectomy Decision in a Breast Centre of Southern Switzerland

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Kathleen R. Blazer, EdD, MS receives research support from Pfizer and Astra Zeneca.

Michelle Bruszer is employeed by Sonova Hearing Aid Manufacturer.

Marina Candido Visontai Cormedi, MD serves as an sub-investigator for Astra Zeneca, as a speaker for Fleury Genomica, and receives institutional research funding from Novartis.

Fergus J. Couch, PhD receives research support from Grail.

Susan M. Domchek, MD served as a consultant for AstraZeneca and GSK.

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Gini F. Fleming, MD has received institutional research funding from Roche, Iovance, Sermonix, Computer, Celldex, Correct, Astra Zeneca, Molecular Templates, Astellas, K Group Beta, and Pfizer. Dr. Fleming plans to discuss investigational/unapproved use of withdrawn PARPI indications and ongoing studies, as well as trials attempting to overcome PARPI resistance, ATR inhibitors, and weel inhibitors.

Feighanne Hathaway, MS, CGC serves as a consultant for AstraZeneca.

Frederick M. Howard, MD serves as a consultant for Novartis AG.

Joanne M. Jeter, MD, MS served as a private investigator for research study for BioAtla. Dr. Jeter plans to discuss investigational/unapproved use of vaccination for Lynch syndrome, PARP inhibitors for non-BRCA mutations in HRD tumors, and tumor type-independent matching of mutations and targeted therapies.

Wenora Johnson serves as a speaker for GSK.

Matthias Kloor, MD serves as a speaker for Roche.

Peter Kuhn, PhD serves as an advisor to Quest Diagnostics and as a consultant to Epic Sciences and Cansera.

Kelly Metcalfe, RN, PhD serves as a consultant for AstraZeneca.

Kathryn Mills, MD serves as a speaker and consultant for Stryker.

Rita Nanda, MD serves on the advisory board for AstraZeneca, Beyond Spring Pharma, Daiichi Sankyo, Fujii Film/GE, Gilead, Infinity Pharmaceuticals, iTeos Therapeutics, MacroGenics, Merck, Novartis, OBI Pharma, OncoSec, Pfizer, Sanofi, Seagen, and Stemline Therapeutics. Dr. Nanda has received grant funding to her institution from Arvinas, AstraZeneca, Celgene, Corcept Therapeutics, Genentech/Roche, Gilead/ Immunomedics, Merck, Novartis, OBI Pharma, OncoSec, Pfizer, Relay Therapeutics, Seattle Genetics, Sun Pharma, and Taiho Oncology.

Katherine Nathanson, MD serves on the advisory board for Merck.

Peter O'Donnell, MD receives honoraria from Adept Field Solutions, Advarra, Merck, Astellas, AbbVie, Pfizer, Custom Learning Designs, Axiom Healthcare Strategies, EMD Serono, IntrinsiQ, ISMIE, NAMCP, Seagen, Curio Science, FirstWord, MedLearning Group, Research to Practice, Great Debates and Updates, MJH Life Sciences, Peerview, Vaniam Group, Institute for Enquiring Minds, Pharmavision UK, PRIME Education LLC, Amerisource Bergen, Health Advances, Parexel Intl Corp, Vida Ventures LLC, Curio Science, Astellas, Seagen, Janssen, Nektar, DragonflyTherapeutics, and G1 Therapeutics.

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Ari J. Rosenberg, MD serves as a consultant for Astellas, EMD Serono, Novartis, Galectin, Privo, Elsai, Vaccitech, and Regeneron, and receives honorarium for speaking from Coherus. Dr. Rosenberg plans to discuss investigational/unapproved clinical trials under IND.

Christina C. Rybak, MS owns stock in Natera.

Rodrigo Santa Cruz Guindalini, MD, PhD serves as a consultant and on the advisory board for AstraZeneca and on the Speakers' Bureau for AstraZeneca, Roche, Gilead, GSK, MSD, BMS, Pfizer, and Daiichi Sankyo.

Jeremy Segal, MD, PhD serves as a consultant for Abbvie, Amgen, and as a scientific advisor for OrisDx.

Parul Somani, MBA serves on the speaker's bureau for Bristol Meyers Squibb, WebMD/Medscape, and Stanford Health Care.

 $\ensuremath{\mathsf{Wendy}}\xspace$ Servier, MD serves as a consultant for Newave, Servier, Jazz, and Amgen.

David J. VanderWeele, MD, PhD serves on the advisory committee for Bayer and on the speaker bureau for Astra Zeneca and Astellas. Dr. VanderWeele plans to discuss investigational/unapproved use of biomarker-driven therapies approved for the same biomarker but in a different disease.

Jeffrey N. Weitzel serves as a for Natera, MyOme, Cancer IQ, and as a speaker for Astra Zeneca.

Adam Yala, PhD serves as an advisor for Janssen R&D, Dewpoint Therapeutics, Merck, HuroneAI, and EarlyByrd.

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