



***Examining Our Present to Illuminate Our Future***  
**August 20, 2021**

**PROGRAM**

*All times listed are in Pacific Daylight Time (PDT)*

- 8:00-8:15**                    **Welcome and Opening Remarks**  
Amy Poole, MS, CGC and Jamie Beavers, MS, CGC (AEC Co-Chairs)  
Michelle Fox, MS, CGC (President, SCGC)
- 8:20-9:05**                    **Plenary 1: Giving Back to Get Ahead: Mentoring the Next Generation**  
Deanna Erwin, MS, CGC; Elise Sobotka, MS, MPH, CGC; Lauren Gima,  
MS, CGC
- 9:10-10:10**                    **Educational Breakout Session**
- Track A: How Small DNA Fragments Changed Prenatal Diagnosis*  
*Deborah Krakow, MD*
- Track B: Molecular Landscape of Colorectal Cancer: Practicing at*  
*the Intersection of Somatic and Germline Genomics*  
Heinz-Joseph Lenz, MD; Jacob Comeaux, LCGC; Charite Ricker, MS,  
CGC; Julie Culver, MS, LCGC, CCRP; Daisy Hernandez, MS, CGC
- 10:10-10:25**                    **BREAK**
- 10:25-11:10**                    **Plenary 2: Federal Bill Access to Genetic Counselor Services Update**  
Sandra Brown, MS, LCGC
- 11:15-12:00**                    **Platform research presentations**
- Gender-Expansive GC Practice, Kayla Lam-Little, MSGC
  - Protestant Perspectives on Prenatal Genetic Services:  
Congruence Between Pastors and Laity, Ellen Brudi, MS
  - Ethnic Differences in the Frequency of Cancer Reported from  
Family Pedigrees in the Prenatal Genetic Counseling Setting, Alex  
Palacios, MS

**12:00-1:00**

**LUNCH AND SPONSORED TALKS**

- **Current Testing Landscape and the Future of Next-Generation Cyto-genomics** presented by Abby Stevens, MS, CGC with Lineagen and Bionano
- **Introduction to PGx with Fulgent Genetics** presented by Gregory Kellogg, MS, CGC
- **No Resting on Laurels: Continual Improvement in Variant Interpretation** presented by Robert Nussbaum, MD with Invitae
- **Single-gene NIPT for Recessive Conditions: A New Carrier Screening Workflow** presented by Jennifer Hoskovec, MS, CGC with BillionToOne

**1:00-1:45**

**POSTER SESSION WITH AUTHORS**

**1:50-2:50**

**Plenary 3:**

**Fostering Inclusive Excellence in Genetic Counseling Graduate Program Admissions**

Christina Palmer, PhD, MS

**Genetic Counselors in the DEI Space: Courage in Practice**

Aarin Ables Williams, MS, CGC

**2:50-3:00**

**BREAK**

**3:00-3:45**

**Plenary 4: The Importance of a Clinical Validity Database in the Age of Rapid Gene Discovery**

Bess Wayburn, PhD, LCGC

**3:45-4:00**

**Closing Remarks**

Amy Poole, MS, CGC and Jamie Beavers, MS, CGC (AEC Co-Chairs)

**4:00-5:00**

**ENGAGEMENT HOUR: SPONSORED TALKS**

- **Veronica: A Fabry Patient Story** presented by Amicus Therapeutics
- **Tools for Managing Urea Cycle Disorder Patients** presented by Horizon Therapeutics
- **Mosaicism ratio: Application to singleton and multifetal gestations** presented by LabCorp
- **Single-gene NIPT for Noonan Syndrome** presented by Natera
- **Stay in the Know: What's New With Whole Genome Sequencing?** presented by PerkinElmer Genomics
- **An IDEA(L) test: How comprehensive genetic testing is improving care for patients with ASD and ID** presented by Prevention Genetics
- **The Exome Blindspot: Variantyx Whole Genome Sequencing Finds what Exomes Miss** presented by Variantyx

Total CEUs available: 0.575 (5.75 contact hours)

**POSTERS AND AUTHORS**  
**(listed alphabetically by title)**

“Area of Practice Change in Genetic Counselor: Perspectives, Motivation, and Helpful Skills”

Primary author: Cynthia Haag

“Attitudes and Perspectives on Genetic Counseling and Genetic Testing for Breast Cancer Risk Amongst Native Hawaiian Breast Cancer Survivors”

Primary author: Nicole Choy

“Genetic Counselor Attitudes on the Reporting of Variants of Uncertain Significance for Multigene Cancer Panels”

Primary author: Emmeline Chang

“Genetic Counselors' Knowledge of State Abortion Laws”

Primary author: Josephine Gao

““LFS and Living My Best!” Pilot Study”

Primary author: Isabella Brink

“The Current Practice of Genetic Counseling in China: Preparing for the Challenges of Genomic Medicine”

Primary author: Lindsey Walker

“The value of mitochondrial genome analysis and approaches for difficult-to-sequence regions in hereditary hearing loss panel testing”

Primary author: Alicia Scocchia