

SHORT COURSE – WEDNESDAY OCTOBER 21

Thinking outside the box: Alternative service delivery models

9:00 – 9:10 a.m.	Opening Remarks: SPC
9:10 – 10:55 a.m.	<p>Plenary Session #1: INTRO AND PEOPLE DOING THE WORK</p> <p>9:10 – 10:00 a.m. Alternative Service Delivery Models: 5 Years of Experience at the Hereditary Cancer Program in British Columbia Zoe Lohn</p> <p>10:00 – 10:25 a.m. The Mainstreaming Process: Oncologist-Mediated Genetic Testing for Hereditary Cancer Stephanie Desmarais, McKenzie Mitchell</p> <p>10:25 – 10:55 a.m. Undergraduate co-Operative Students in the Genetics Clinic Brittney Johnstone, Karen Ott</p>
10:55 – 11:25 a.m.	Break
11:25 – 12:45 p.m.	<p>Plenary Session #2: PHONE COUNSELING, AND CHANGING MODELS OF GENETIC COUNSELLING IN A PANDEMIC</p> <p>11:25 – 12:05 p.m. Telephone: a means to scale & increase access, with high patient and genetic counselor satisfaction Colleen Caleshu</p> <p>12:05 – 12:45 p.m. A Complete 180 – How a community hospital changed to phone and virtual counselling due to the COVID-19 pandemic Ingrid Ambus, Islay Fitzgerald, Kristen Miller</p>
12:45 – 1:15 p.m.	Break
1:15 – 2:45 p.m.	<p>Plenary Session #3: USING THE INTERNET IN NEW WAYS</p> <p>1:15 – 1:45 p.m. Genetic counselling delivery by webinar: The Alberta experience with hypertrophic cardiomyopathy Raechel Ferrier</p> <p>1:45 – 2:15 p.m. Implementation of Online Learning Module for Hereditary Breast and Ovarian Cancer Kelly Anderson</p> <p>2:15 – 2:45 p.m. Effectiveness of the Genomics ADvISER, a decision aid for the selection of incidental genome sequencing results</p>

	Salma Shickh
2:45 – 3:15 p.m.	Break
	Plenary Session #4: UNIVERSAL GENETIC SCREENING
	3:15 – 4:00 p.m. Genetic testing for all breast and ovarian cancer patients at the time of diagnosis: Can (and should) it be done? Jeanna McCuaig, Kelly Metcalfe
	4:00 – 4:30 p.m. Genetic Screening in ALS – A National Approach Kristiana Salmon
3:15 – 5:00 p.m.	4:30 – 5:00 p.m. Universal testing of endometrial cancer: Reaching an under-served population (the UTERUS project) Candice Jackel-Cram
ANNUAL EDUCATION CONFERENCE (AEC) – OCTOBER 22-23	
THURSDAY OCTOBER 22	
ALL TIMES ARE CENTRAL TIME (WINNIPEG TIME)	
9:00 – 9:10 a.m.	Opening Remarks: AEC Committee Co-Chairs
9:10 – 9:45 a.m.	Melanie Care: CAGC Presidential Address
9:45 – 10:45 a.m.	Keynote: Twisted Science: Are Public Representations of Genetics Doing Harm? Timothy Caulfield
10:45 – 11:15 a.m.	Break
	Abstract Session
	11:15 – 11:30 a.m. An internship in psychiatric genetic counseling: Impact on genetic counseling graduates' practice and career choices Brianna Van Den Adel
	11:30 – 11:45 a.m. Proactive genetic screening addresses limitations of family history and access to genetic testing Jessica Gu
11:15 – 12:30 p.m.	11:45 – 12:00 p.m. Exploring perceptions and attitudes towards integrating a genetic counsellor into a multidisciplinary primary care practice team. A qualitative GenCOUNSEL study

	<p>Caitlin Slomp</p> <p>12:00 – 12:15 p.m. A nationally agreed cross-professional competency framework to facilitate genomic testing Amanda Pichini</p> <p>12:15 – 12:30 p.m. The costs associated with conventional diagnostic testing compared to genomic sequencing in patients with adult-onset neurological conditions Yun Amber Zhu</p>
12:30 – 1:00 p.m.	<p><u>CAGC Research Grant Presentation – The Whole Exome Sequencing Experience in Canada: Current Practices and Anticipated Challenges to Widespread Implementation (Not Available OnDemand)</u> Salma Shickh</p>
1:00 – 1:30 p.m.	Break
1:30 – 2:15 p.m.	<p><u>General Public Inquiries – How Far should you go?</u> Tina Babineau Sturk</p>
2:15 – 3:00 p.m.	<p>Posters</p> <p>2:15 – 2:20 p.m. Solving a Cold Case: A Lesson on Making Assumptions Natasha Osawa</p> <p>2:20 – 2:25 p.m. A woman-centered, constructivist grounded theory of decision making regarding antidepressants in pregnancy (Not Available OnDemand) Catriona Hippman</p> <p>2:25 – 2:30 p.m. A novel model for the management of clinically relevant incidental findings Rita Kodida</p> <p>2:30 – 2:35 p.m. Talking about Mental Health and 22q11DS Corey Filiaggi</p> <p>2:35 – 2:40 p.m. Far and Wide: Uptake and acceptability of remote service provision for genome-wide sequencing in Canada: a GenCOUNSEL substudy Emily Enns</p> <p>2:40 – 2:45 p.m. Do genetic counselling students use self-help online resources for mental health support? A survey looking at usage, perception, and drawbacks Kelsey Kalbfleisch</p> <p>2:45 – 2:50 p.m. Performance of an eHealth decision support tool (MIPOGG) for identifying children with Li-Fraumeni syndrome, Gorlin syndrome, DICER1 syndrome, and constitutional mismatch repair deficiency</p>

	<p><i>Robyn Hebert</i></p> <p>2:50 – 2:55 p.m. The expansion of prognostic tumor genetic testing and emerging roles for genetic counsellor <i>Jaime Jessen</i></p>
2:55 – 3:30 p.m.	Break
3:30 – 4:30 p.m.	<p><u>The Cost of Free Testing: Sponsored Genetic Testing in Canadian Health Care</u> <i>Kirsten Bartels, Jennifer Nuk</i></p>
4:30 – 5:30 p.m.	<p><u>Supporting a Lost Dream</u> <i>Danielle Holm, Jennifer Karpes, Patti Walker</i></p>
FRIDAY OCTOBER 23	
ALL TIMES ARE CENTRAL TIME (WINNIPEG TIME)	
9:00 – 9:15 a.m.	Opening Remarks, CAGC Award Presentations
9:15 – 10:00 a.m.	<p><u>Improving the Experience for LGBTQ2S+ People</u> <i>Marni Panas</i></p>
10:00 – 10:45 a.m.	<p><u>Genetic Counselling for Intersex Conditions: A Community Approach to Depathologization</u> <i>Katie Saulnier</i></p>
10:45 – 11:15 a.m.	Break
	<p>Posters</p> <p>11:15 – 11:20 a.m. The utility of rapid whole exome sequencing for recurrence risk counselling and subsequent pregnancy management: a case report <i>Islay Fitzgerald</i></p> <p>11:20 – 11:25 a.m. A method to improve genetic diagnostic yield among patients suspected to have primary immunodeficiency <i>Christine Davies</i></p> <p>11:25 – 11:30 a.m. Frequency and characteristics of copy number variants identified by a whole-exome sequencing platform across medical specialities <i>Allison Sluyters</i></p> <p>11:30 – 11:35 a.m. A new clinical exome assay developed for the high-throughput diagnosis of genetic conditions <i>Julie Hathaway</i></p>
11:15 – 12:00 p.m.	11:35 – 11:40 a.m.

	<p>Congenital disorder of fucosylation caused by homozygous mutations in FUT8 undetectable with traditional transferrin screening <i>Cassandra McDonald</i></p> <p>11:40 – 11:45 a.m. Challenges to the detection and interpretation of single nucleotide variants in close proximity to copy number variants: a CHEO Genetics Diagnostic Laboratory experience <i>Patricia Harper</i></p> <p>11:45 – 11:50 a.m. The Value of Whole Exome Sequencing in Prenatal Diagnosis: The Experience of a Canadian Tertiary Hospital <i>Anna Pan</i></p> <p>11:50 – 11:55 a.m. Exploring the impact of a decision aid: Engagement and personalization during pre-test genetic counselling for secondary finding selection from genomic sequencing among adults affected by cancer <i>Sara Rafferty</i></p>
12:00 – 1:00 p.m.	<p>Introduction to Preimplantation Genetic Testing for the Clinical Counsellor (Not Available OnDemand) <i>Charlotte Emmerson, Erica Pai, Jaspreet Sekhon-Warren</i></p>
1:00 – 1:30 p.m.	Break
1:30 – 3:15 p.m.	<p>Abstract Session</p> <p>1:30 – 1:45 p.m. Retrospective analysis of >25,000 rare disease patients confirms biallelic variants in the NRAP gene are a significant cause of dilated cardiomyopathy <i>Kim Gall</i></p> <p>1:45 – 2:00 p.m. The diagnostic yield of multi-gene intellectual disability panels in 140 patients with neurodevelopmental disorders <i>Mireille Cloutier</i></p> <p>2:00 – 2:15 p.m. Should we be offering pharmacogenetic testing for CYP2D6 and CYP2C19 to pregnant women taking SSRIs? (Not Available OnDemand) <i>Catriona Hippman</i></p> <p>2:15 – 2:30 p.m. Implementing Genomics in the Neonatal Period: An Assessment of Parental Decision-Making and Anxiety <i>Tasha Wainstein</i></p> <p>2:30 – 2:45 p.m. It meant a lot and it helped a lot”: A retrospective study of parental experiences with rapid genome-wide sequencing in a neonatal intensive care unit <i>Caitlin Aldridge</i></p> <p>2:45 – 3:00 p.m. “What doesn’t kill you makes you stronger”: Parents’ perspectives after the return of variants of uncertain significance from multigene panel testing for hearing loss</p>

	Rosettia Ho 3:00 – 3:15 p.m. Sickle cell trait newborn screen results: disclosure and management Margaret Lilley
3:15 – 3:45 p.m.	Break
3:45 – 5:00 p.m.	Beyond the Bubble: SCID newborn screening and management in Alberta Dr. Stacey Hume, Dr. Ross Ridsdale, Dr. Sneha Suresh, and Cowie family
5:00 – 5:10 p.m.	Closing Remarks Charlotte Fung, Danna Hull

Please note: Speakers were selected to present based on their area of expertise; however, the opinions of the speakers are not necessarily reflective of the opinions of the CAGC.