



8:00 AM - 8:15 AM	Breakfast & Session 1 Registration
8:15 AM – 8:20 AM	Intro: The New Agilent - The Premier Laboratory Partner for a Better World Heidi L. Kijenski <i>Global Marketing Director – Clinical Genomics, Agilent Technologies, Inc.</i>
8:20 AM – 8:40 AM	Intro: Integrated Pathology Solutions for Cancer Research Tyler J. Hendershot PhD <i>Technical Business Manager, Agilent Technologies, Inc.</i>
8:40 AM – 9:10 AM	Workflow Challenges and Solutions in FISH Testing – a CGI Experience V. Subhadra Nandula, PhD, FACMG <i>Associate Director Cytogenetics, Cancer Genetics, Inc.</i>
9:10 AM – 9:20 AM	Break and Session 2 Registration
9:20 AM – 9:30 AM	Targeting Large and Small - OneSeq: One Assay for Genome-Wide CNVs, LOH and Targeted Mutations Corinna Nunn <i>Global Senior Product Manager, OneSeq, Agilent Technologies, Inc.</i>
9:30 AM – 9:50 AM	Agilent Genomic Solutions: Paving the Way to Making ACTG Spell A-N-S-W-E-R-S Maria Celeste M. Ramirez, PhD <i>Global Product Manager, NGS-SureSelect, Agilent Technologies, Inc.</i>
9:50 AM -10:15 AM	A Pilot Study to Evaluate the Use of Isolated Circulating Fetal Nucleated Cells (CFNCs) for Aneuploidy Screening by aCGH Lian Liu, MD <i>CEO, PacGenomics</i>
10:15 AM – 10:40 AM	Optimizing Exome Sequencing by Increasing Coverage for Genes Associated with Disease Madhuri Hegde, PhD, FACMG <i>Professor of Human Genetics, Executive Director, Emory Genetics Lab, Emory University</i>
10:40 AM – 11:05 AM	Utility of an Automated SureSelect NGS Panel for Understanding Marfan and Marfan-like Syndromes Featuring Aortopathy Whitney Wooderchak-Donahue, PhD <i>Adjunct Assistant Professor, Department of Pathology, University of Utah, Research and Development Scientist, ARUP Laboratories</i>
11:05 AM – 11:30 AM	Assessing Copy Number from Exome Sequencing and Exome Array CGH Based on CNV Spectrum in a Large Clinical Cohort Jane Juusola, PhD, FACMG <i>Director, Whole Exome Sequencing, GeneDx</i>
11:30 AM – Noon	Panel Q&A and Wrap Up