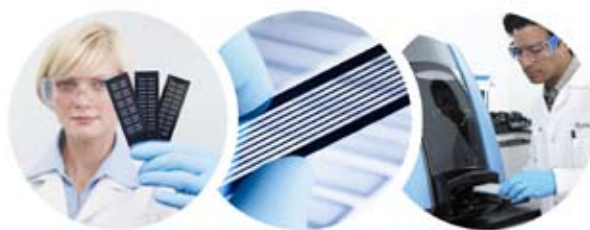


Illumina Seminar Series



From Whole-Genome to Whole-Solution, Disease Analysis Tools for the Next Generation

In the study of diseases, their causes and the development of therapies to treat them, flexible tools that look at RNA- and DNA-based changes across the genome are mission critical. Illumina is committed to providing such tools, from genome-wide analysis to low-multiplex target validation.

This seminar series will bring you compelling updates on new tools for genetic analysis from Illumina and presentations from scientists using Illumina technology to power their next generation studies.

MONDAY, MAY 4, 2009

UNIVERSITY OF CALGARY
HEALTH SCIENCES CENTRE
3330 HOSPITAL DRIVE NW, CALGARY
THEATRE 3

*Register early to secure a seat as space is limited.

SEMINAR SCHEDULE

9:00	REGISTRATION	
9:30	Introduction	
9:40	<i>An Overview of Illumina's Sequencing, Expression and Regulation Portfolio</i>	Chris Streck, Illumina
10:10	<i>ChIP-Sequencing: Centromeres and Circadian Rhythms</i>	Michael Freitag, Ph.D., Oregon State University
10:40	<i>Sequencing Cancer Genomes and Transcriptomes</i>	Richard Moore, Ph.D., BC Cancer Agency Genome Sciences Centre
11:10	<i>An Overview of Illumina's DNA Analysis Products – From Genotyping to CNV</i>	David Dailey, Ph.D., Illumina
11:40	<i>Pharmacogenomics of Severe Adverse Drug Reactions</i>	Colin Ross, Ph.D., University of British Columbia
12:10	CONCLUSION	
12:15	LUNCH AND NETWORKING	

Register now at:
www.illumina.com/seminars

