Navigating the Genomic Regulatory Landscape

September 12 & 13, 2017 | San Diego, CA

Join us for this in-depth, two-day workshop and explore current regulatory issues and challenges related to genomics. Lecturers from academia, industry, and the legal profession will discuss the impact of regulations on genomics in the context of medicine, ethics and patient privacy, legal strategies, the use of big data, and the commercial landscape. The workshop will consist of a combination of lectures, regulatory case studies, an onsite tour at the biotechnology company Illumina and opportunities for networking.

Why Attend?

- Gain an in-depth understanding of the current challenges and opportunities faced by the genomics regulatory environment
- Apply critical skills in regulatory science and how it applies to an examination of translational, clinical, ethical, legal and data challenges for genomic topic
- Evaluate the impact of laws, policies and regulations on the genomic industry

Speakers:

- **Paul Rejto**
  Head of Oncology Translational Research, Pfizer, Inc.
- **Ruth Waterman**
  Assistant Clinical Professor, Anesthesiology, UC San Diego
- **Dale Hunt**
  Partner, Hahn Loeser & Parks
- **Ashley Van Zeeland**
  Chief Technology Officer, Human Longevity, Inc.
- **Camille Nebeker**
  Director, Research Ethics, Scripps Translational Science Institute
- **Michael Kalichman**
  Director, Research Ethics Program, UC San Diego
- **Hannah Carter**
  Assistant Professor, School of Medicine, UC San Diego

Tuesday, September 12,
7:30 a.m. - 6:15 p.m

Wednesday, September 13,
8:30 a.m. - 12:45 p.m

Location:
UC San Diego Extension
6256 Greenwich Dr., San Diego, CA

Price:
$299

What's included:

- World-class speakers & instruction
- Tour and lecture at Illumina, Inc.
- Breakfast and lunch included
- Reception at Rock Bottom Restaurant and Brewery in La Jolla, CA
- 1.5 units of credit

ENROLL NOW!

SPACE IS LIMITED!

FOR MORE INFORMATION:
858-534-9151 | drhammon@ucsd.edu

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extension.ucsd.edu/genomicworkshop