

# Advances in Genome Science.



## Using New Technologies to Study the Genetics of Disease

**Thursday 12th September, 2.00 pm**

**Aula Riunioni Dipartimento, Palazzo Bellini  
Università del Piemonte Orientale  
via Solaroli 17, 28100 Novara**

Discover how new technologies are being used to identify genetic variations associated with human disease.

In this seminar, you'll learn how high-throughput genetics is expanding the field of human disease research; from traditional genome-wide association studies (GWAS) to targeted resequencing approaches that can identify both common and exceedingly rare causal variations.

Hear from scientists who use Illumina technologies to address their research questions.

*Il seminario si terrà in Italiano. La partecipazione al seminario è gratuita. .*

### Agenda

<b>2:00 pm</b>	<b>Registration</b>
<b>2:30 pm</b>	<b>Illumina's Mission</b> - Luca Beretta, Illumina
<b>2:45 pm</b>	<b>Genetic disease introduction</b> - Nicola Cirenei, Illumina
<b>3:15 pm</b>	<b>Illumina technology in Genetic disease</b> - Nicola Cirenei, Illumina
<b>3:45 pm</b>	<b>Coffee Break</b>
<b>4:05 pm</b>	<b>Molecular Diagnosis of Genetic Diseases: from 1 Gene to 1000s</b> - Dr.ssa Laura Pezzoli, Genetica Medica AO Papa Giovanni XXIII Bergamo
<b>4:45 pm</b>	<b>My samples, My studies, MiSeq: From custom amplicon to exome sequencing on a benchtop instrument</b> - Luca Beretta, Illumina

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