

Preliminary Program

This program is subjected to changes. Also, we will select additional talks based on incoming abstracts.

Wednesday, May 31

10–11.00 Registration, setting up posters, and welcoming coffee.

SESSION 1

Clinical status and perspectives for hereditary breast and ovarian cancer (HBOC) risk prediction.

Fergus Couch

The Mayo Clinic Cancer Center and the Center for Individualized Medicine, USA

Diana Eccles

University of Southampton, UK

Rita Schmutzler

University Hospital Cologne, Germany

Sabine Linn

The Netherlands Cancer Institute, The Netherlands

+Talks selected from incoming abstracts

DISCUSSION

The unmet needs in the clinic for genetic counselling of HBOC.

Key Questions:

- At what point are we safe to decide whether a VUS is pathogenic or not?
- How do different countries deal with VUS with respect to the use of prediction programs, conjoint databases, the offer of intensified screening and a recall system?
- How to fill in the gap between research and clinic?
- What are the important measurements for risk prediction?
- Infrastructure and databases for germline and tumour material.

Chair: Diana Eccles

Panel: Rita Schmutzler, Sabine Linn, Jan Schellens, Marjanka Schmidt, and Fergus Couch

SESSION 2

Identification of novel HBOC genes and variants

Claus Storgaard Sørensen

Biotech Research and Innovation Centre, University of Copenhagen, Denmark

“New mechanisms guarding genome integrity”

Bjarni Jóhann Vilhjálmsson

Center for Bioinformatics (BiRC), University of Aarhus, Denmark

“Joint re-analysis of GWAS summary statistics identifies new variants associated with human traits and diseases”

Stefano Annunziato

The Netherlands Cancer Institute, The Netherlands

“Somatic engineering of the mammary gland for the development of novel mouse models of breast cancer.”

+Talk selected from incoming abstracts

Thursday, June 1

SESSION 3

Analysis of the effects of HBOC gene variants on treatment response.

Lisa Wiesmüller

University Hospital Ulm, Germany

“Error-prone DNA repair signature for detection of breast and ovarian cancer risk”

Violeta Serra

Vall d’Hebron Institute of Oncology, Spain

“Functional homologous recombination DNA repair as a major in vivo mechanism of PARP inhibitor resistance in germline BRCA1/2-mutated breast tumors.”

Oded Kopper

Hubrecht Institute, The Netherlands.

“Organoids representing human ovarian cancer”

+Talks selected from incoming abstracts

SESSION 4

Model systems and assays for the functional classification of HBOC gene variants.

Bas Vroling

Bio-Product, The Netherlands

Joey Riepsaame

Sir William Dunn School of Pathology, University of Oxford, UK

Michel Cannieux

Integrated DNA Technologies, USA

“Strategies for optimization of CRISPR editing and HDR using Cas9 and Cpf1 ribonucleoproteins (RNPs)”

Peter Bouwman

The Netherlands Cancer Institute, The Netherlands

“Functional analysis of BRCA1 variants and domains using genetically engineered mouse model systems”

Maaïke Vreeswijk

Leiden University Medical Center, The Netherlands

“Functional characterization of variants of uncertain significance in BRCA2”

Haico van Attikum

Leiden University Medical Center, The Netherlands

Sean Tavtigian

Huntsman Cancer Institute, University of Utah, USA

Alvaro NA Monteiro

Moffitt Cancer Center, USA

“Incorporating Functional Data into Clinical Annotation.”

+Talks selected from incoming abstracts

Friday, June 2

SESSION 4 (CONT.)

DISCUSSION

How to push forward the gene variant classification from functional assay into the clinic?

Key Questions:

- What are the requirements for gene models and functional assays to have clinical impact on diagnostic and treatment decision?

- Consequences for assay design and biological controls.
- Current status and the road forward for genetic variant risk prediction.
- Cancer risk associated with hypomorphic variants.
- How to incorporate functional data into multifactorial likelihood models.

Chair: Maaïke Vreeswijk

Panel: Lisa Wiesmüller Peter Bouwman, Fergus Couch, and Alvaro NA Monteiro

SESSION 5

Novel insights in HBOC suppression and consequences for functional assays.

Shyam Sharan

Center for Advanced Preclinical Research, National Cancer Institute, USA

“Synthetic viability: how cells overcome BRCA2-loss induced cell-lethality”

Ralph Scully

Beth Israel Deaconess Medical Center, Harvard University, USA

Arnab Ray Chaudhuri

National Cancer Institute, USA

“Remodelling DNA replication forks – From synthetic viability to chemoresistance”

+Talks selected from incoming abstracts

14.30–15.00 Concluding remarks by Jos Jonkers.

