

## Program

### Tuesday, December 11, 2012

#### Satellite Symposia - Krehl Klinik Auditorium, INF 410

12:00 - 2:00 pm **Satellite Symposium:** Small RNAs (Org. J. Haas and J. Brosius)  
Bryan Cullen - Sebastián Pfeffer - Gunter Meister -Richard Reinhardt

02:00 - 2:30 pm **Coffee Break**

02:30 - 4:00 pm **Satellite Symposium:** Next-Generation Sequencing (Org. S. Wiemann and B. Timmermann)  
Stephan Wolf - Wolfgang Huber - Benedikt Brors - Saskia Biskup

04:00 - 4:30 pm **Coffee Break** (DKFZ Communication Center)

#### Welcome - DKFZ Communication Center, Main Auditorium

4:30 - 5:00 pm **Wolfgang Wurst**, Helmholtz Zentrum Munich, Germany,  
Speaker Project Committee of NGFN-Plus / NGFN-Transfer in the Program of Medical Genome Research

**N.N.**, Federal Ministry of Education and Research, Germany

**Christof von Kalle**, National Center for Tumor Diseases (NCT)/  
German Cancer Research Center (DKFZ), Heidelberg, Germany

#### Symposium I: International Projects in the Program of Medical Genome Research

5:00 - 5:30 pm *International Cancer Genome Consortium (ICGC)*  
**Michael Taylor**, The Hospital for Sick Children, Toronto, Canada

5:30 - 6:00 pm *International Cancer Genome Consortium (ICGC)*  
**Peter Lichter**, DKFZ, Heidelberg, Germany  
*Novel findings in pediatric brain tumors revealed by high resolution genome, methylome and transcriptome analyses*

6:00 - 6:30 pm *International Cancer Genome Consortium (ICGC)*  
**Matthew Meyerson**, Dana Farber Cancer Institute, Boston, USA  
*Genome Alterations in Human Lung Cancers*

6:30 - 7:00 pm *1000 Genomes Project*  
**Hans Lehrach**, MPI for Molecular Genetics, Berlin, Germany  
*1000 Genomes Project - An integrated map of genetic variation*

7:00 pm **Supper**

### Wednesday, December 12, 2012

#### Symposium II: International Projects in Genome Research

9:00 - 9:30 am **John Blangero**, Texas Biomedical Research Institute, San Antonio, USA  
*Identification of Complex Disease Genes Using Whole Genome Sequencing in Large Pedigrees*

9:30 - 10:00 am **Joris A. Veltman**, Radboud University Nijmegen Medical Centre, Nijmegen, The Netherlands  
*De novo diagnostics in patients with intellectual disability*

10:00 - 10:30 am *Alzheimer's Disease Genetics Consortium (ADGC)*  
**Gerard D. Schellenberg**, University of Pennsylvania School of Medicine, Philadelphia, USA  
*Neurodegenerative disease genetics; GWAS, exomes and beyond*

10:30 - 11:00 am **Coffee Break**

11:00 - 11:30 am *International Human Epigenome Consortium (IHEC)*  
**Peter Jones**, USC Norris Comprehensive Cancer Center, Los Angeles, USA  
*The Cancer Epigenome*

11:30 - 12:00 pm *International Human Epigenome Consortium (IHEC)*  
**Jörn Walter**, Saarland University, Germany  
*Control of DNA-methylation in development*

12:00 - 2:00 pm **Lunch Break and Poster Session I**  
12:00 – 1:00 pm odd numbers  
01:00 – 2:00 pm even numbers

#### Company Satellite Session

*Affymetrix - Caliper a Perkin Elmer Company - Illumina - Bio-Rad Laboratories*

2:00 - 2:30 pm *International Knockout Mouse Consortium / EUCOMM*  
**Wolfgang Wurst**, Helmholtz Zentrum Munich, Germany  
*International Mouse Knock-out Consortium: Resource to functionally validate all genes*

2:30 - 3:00 pm *International Mouse Phenotyping Consortium / EUMODIC*  
**Martin Hrabě de Angelis**, Helmholtz Zentrum Munich, Germany  
*Creating a comprehensive encyclopedia of mammalian gene function*

3:00 - 3:30 pm **Matthias Mann**, Max Planck Institute for Biochemistry, Munich, Germany  
*High resolution, quantitative mass spectrometry combines proteomics and genomics*

3:30 - 4:00 pm **Coffee Break**

#### Symposium III Genomics of Common Disease I

4:00 - 4:15 pm **André Reis**, University Erlangen-Nuremberg, Germany  
*Range of genetic mutations associated with severe non-syndromic sporadic intellectual disability: An exome sequencing study*

4:15 - 4:30 pm **Vanessa Nieratschker**, Central Institute for Mental Health, Mannheim, Germany  
*Pre-, peri- and postnatal stress in human and non-human off-spring: A convergent approach to study genetic and epigenetic impact on depression*

4:30 - 4:45 pm **Andre Franke**, University Hospital Schleswig-Holstein, Kiel, Germany  
*Host-microbe interactions shape genetic risk for inflammatory bowel disease*

4:45 - 5:00 pm **Robert Häslér**, Christian Albrechts University of Kiel, Germany  
*Mapping the functional epigenome of Ulcerative Colitis in monozygotic twins*

5:00 - 5:30 pm **Coffee Break**

5:30 - 5:45 pm **Anke Hinney**, University of Duisburg-Essen, Germany  
*Childhood ADHD and obesity: Evidence for a common genetic link*

5:45 - 6:00 pm **Ralf Herwig**, Max Planck Institute for molecular Genetics, Berlin, Germany  
*An integrated catalogue of genome, methylome and gene expression variations in colon cancer*

#### Evening Lecture:

6:00 - 7:00 pm **Randolph Nesse**, The University of Michigan, Ann Arbor, USA  
*Evolutionary Medicine and Molecular Medicine: Synergistic Siblings*

7:00 - 10:00 pm **Get-Together (Wine, Cheese, Live Music)**

### Thursday, December 13, 2012

#### Symposium IV Genomics of Common Disease II

9:00 - 9:30 am **Keynote: Hugo A. Katus**, Heidelberg University Hospital, Germany  
*Innovations in translational research – Successes of NGFN*

9:30 - 9:45 am **Christina Loley**, University of Lübeck, Germany  
*Association of X-chromosomal variants with coronary heart disease: Results from a meta-analysis*

9:45 - 10:00 am **Martin Peifer**, University of Cologne, Germany  
*Toward a novel genomics-based taxonomy of lung cancer: The Clinical Lung Cancer Genome Project*

10:00 - 10:15 am **Christel Herold-Mende**, University of Heidelberg, Germany  
*Aberrant self-renewal and quiescence contribute to Glioblastoma aggressiveness*

10:15 - 10:45 am **Coffee Break**

#### Symposium V Functional Genomics

10:45 - 11:00 am **Jan Korbel**, The European Molecular Biology Laboratory, Heidelberg, Germany  
*Genome sequencing of childhood medulloblastoma brain tumors links chromothripsis with TP53 mutations*

11:00 - 11:15 am **Gerrit Erdmann**, German Cancer Research Center, Heidelberg, Germany  
*Secretion of Wnts is required for Wnt/ $\beta$ -catenin pathway activity in colorectal cancer (CRC) despite APC or  $\beta$ -catenin mutations*

- 11:15 - 11:30 am **Sven Lindner**, University Hospital Essen, Germany  
*LIN28B induces neuroblastoma and enhances MYCN levels via let-7 suppression*
- 11:30 - 11:45 am **Sören Westphal**, University Clinic Ulm, Germany  
*Human BCL2-associated athanogene 3 (BAG3) mutations lead to dilated cardiomyopathy in zebrafish*
- 11:45 - 12:00 pm **Zouhair Aherrahrou**, University of Lübeck, Germany  
*Cyp17a1 deficient mice display increased body weight, visceral/subcutaneous fat deposition and altered lipid metabolism*
- 12:00 - 2:00 pm **Lunch Break and Poster Session II**  
12:00 – 1:00 pm odd numbers  
01:00 – 2:00 pm even numbers
- Company Satellite Session**  
*Life Technologies - Roche Diagnostics - SEQUENOM - Nanostring Technologies*
- 2:00 - 2:15 pm **Ceremony: „Annemarie Poustka Poster Award 2012“**  
sponsored by Roche Diagnostics GmbH  
**Christine Kuch**, Roche Diagnostics Deutschland GmbH  
**Hugo A. Katus**, Heidelberg University Clinics, Project Committee of NGFN-Plus / NGFN-Transfer in the Program of Medical Genome Research

## Conference Organization

**NGFN Management Office**  
c/o German Cancer Research Center - DKFZ  
Im Neuenheimer Feld 280, V025, D-69120 Heidelberg

Dr. Silke Argo  
E-mail: s.argo@dkfz.de; Phone: +49-6221-424649  
Dr. Cornelia Depner; Dr. Tanja Jutzi;  
Dr. Johanna Lampert; Anita Miehle

## Venue

**German Cancer Research Center - DKFZ  
Communication Center**  
Im Neuenheimer Feld 280  
69120 Heidelberg  
www.dkfz.de

With special thanks to:



## Symposium VI Personalized Medicine

- 2:15 - 2:45 pm **Keynote: Rudi Balling**, Luxembourg Centre for Systems Biomedicine, Luxembourg  
*Systems approaches to Parkinson's disease*
- 2:45 - 3:00 pm **Nicole Teichmann**, University Hospital rechts der Isar, Munich, Germany  
*A promising therapy strategy for PDAC: MEK1/2 inhibition with the novel chemotherapeutic drug BAY 86-9766 (RDEA119)*
- 3:00 - 3:15 pm **Margret Hoehle**, Max Planck Institute for Molecular Genetics, Berlin, Germany  
*Haplotype-resolving multiple human genomes: Key to personalized medicine and genome biology*
- 3:15 - 3:30 pm **Jeanette Erdmann**, University of Lübeck, Germany  
*The risk of myocardial infarction is increased by digenic mutation in GUCY1A3 and CCT7 - identified by exome sequencing in an extended family*
- 3:30 - 3:45 pm **Brajesh Kaistha**, University of Marburg, Germany  
*High-throughput cell-based assays identify Placenta-specific 8 (Plac8; Onzin) as a key regulator of proliferation and survival in pancreatic cancer cells*
- 3:45 - 4:00 pm **Concluding Remarks: Markus Nöthen**, Friedrich-Wilhelms University, Bonn, Spokesperson for the Project Committee of NGFN-Plus / NGFN-Transfer in the Program of Medical Genome Research
- 4:00 - 4:30 pm **Closing Coffee**

## 5th Annual Meeting of NGFN-Plus and NGFN-Transfer in the Program of Medical Genome Research

**December 11-13, 2012  
DKFZ, Heidelberg**

[www.ngfn-meeting.de](http://www.ngfn-meeting.de)

## Scientific Program Committee

Prof. Dr. Jürgen Brosius  
University of Münster

Prof. Dr. Martin Hrabě de Angelis  
Helmholtz Zentrum München

Prof. Dr. Hugo Katus  
Heidelberg University Hospital

Prof. Dr. Peter Lichter  
German Cancer Research Center (DKFZ), Heidelberg

Prof. Dr. Markus Nöthen  
University of Bonn

Prof. Dr. Matthias Riemenschneider  
Saarland University Medical Center

Prof. Dr. Stefan Schreiber  
University Medical Center Schleswig-Holstein

Prof. Dr. Roman Thomas  
University of Cologne

Prof. Dr. Wolfgang Wurst  
Helmholtz Zentrum Munich

SPONSORED BY THE