

Variant interpretation and sharing of genetic data

November 7-8 2017

Ibsenhuset, Skien

Tuesday November 7th

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| 0830-0930 | Registration |
| 0930-0935 | Welcome
Asbjørg Stray-Pedersen, Chair, Norwegian Society for Medical Genetics (NFMG) |
| 0935-1050 | Variant interpretation and sharing of sequence variants
Heidi Rehm, Medical Director, Broad Institute Clinical Research Sequencing Platform
Principal investigator, Clinical Genome Resource (ClinGen) |
| 1050-1110 | Break |
| 1110-1210 | Challenges in variant interpretation – how to minimize inter and intra- laboratory inconsistencies
Teresa Neuhan, MD, Clinical Geneticist, MGZ - Medical Genetics Center, Munich |
| 1210-1315 | Lunch |
| 1315-1415 | Workshops <ul style="list-style-type: none">- Variant interpretation, TBA- Pharmacogenetics, TBA |
| 1415-1445 | Break |
| 1445-1530 | Presentation of submitted abstracts (“frie foredrag”).
12 minutes + 3 minutes for questions per presentation |
| 1530-1545 | Genetikkportalen |
| 1545-1600 | Closing remarks |
| 1600 | General Assembly, Norwegian Society for Human Genetics (NSHG)
Årsmøte for NSHG (In Norwegian) |
| 1700 | General Assembly, Norwegian Society for Medical Genetics (NFMG)
Årsmøte for NFMG (In Norwegian) |
| 1930 | Dinner |

Wednesday November 8th

- 0830-0840 Registration
- 0840-0845 Welcome
- Wenche Listøl, Chair, Norwegian Society for Human Genetics, Haukeland University Hospital, Bergen, Norway
- 0845-0945 Copy number alterations
- Nicole de Leeuw, Head team Intellectual Disability, Multiple Congenital Abnormalities & Endocrine disorders, Radboud UMC, Nijmegen
- 0945-1030 Presentation of submitted abstracts ("frie foredrag").
- 12 minutes + 3 minutes for questions per presentation
- 1030-1100 Break
- 1100-1200 Workshops
- HTS/CNVs, TBA
 - Prenatal diagnostics, TBA
- 1200-1300 Lunch
- 1300-1345 Can IVF influence human evolution?
- Hans Ivar Hanevik, Overlege Fertilitetsklinikken Sør, Chair, Norwegian Society of Human Reproduction and Embryology
- 1345-1405 Break
- 1405-1505 Workshops
- Rare syndromes and inborn errors of metabolism, TBA
 - Cancer genetics, inherited cancer, TBA
- 1505-1520 CLG
- 1520-1530 Closing remarks