

The OMICs course

	Mon 8/5	Tue 9/5	Wed 10/5	Thu 11/5	Fri 12/5
09:00		<p>3) Yvonne Böttcher:</p> <p>"Multifactorial diseases/GWAS/disease association"</p> <p>"GWAS obesity": monogenic vs. polygenic/common obesity; gene-environment interactions; fat distribution, T2D; array based 5m DNA methylation; epigenome-wide analyses; pyrosequencing.</p> <p>These topics will be addressed together with an overview about the literature (lectures, "Green auditorium, Bygg 25, Ullevål")</p>	<p>5) Manuela Zucknick:</p> <p>"Statistical principles behind three basic data analysis tasks in omics"</p> <p>1) Screening for "interesting" features (e.g. differential gene expression) -> Multiple testing, gene group tests and pathway analyses</p> <p>2) Data visualisation and dimension reduction; "unsupervised learning" -> Cluster analysis and heatmaps, principal component analysis</p> <p>3) Prediction and classification of patients based on omics data; "supervised learning" -> Introduction to some popular prediction methods, correct assessment -> of prediction performance (lectures, "Green auditorium, Bygg 25, Ullevål")</p>	<p>7) Vessela Kristensen:</p> <p>"Genomics – from germ line to somatic genetic and epigenetics"</p> <p>SNPs and CNV -Normal genetic variation and disease GWAS/sequencing Genotype-phenotype associations Phenotypic changes in disease: usefulness for classification, prognosis and treatment response prediction At mRNA level mRNA arrays/RNAseq DNA copy number alterations DNA methylation/ DNA methylation arrays/vs sequencing miRNA/lncRNA regulation arrays/vs RNAseq DNA somatic mutations, dissemination the immunogenicity of somatic mutations. (lecture, "Auditorium 1, Kreftssenteret, Ullevål")</p>	<p>13) Johannes Hov:</p> <p>"Introduction to metagenomics and the gut microbiome in health and disease"</p> <p>Case/problem: Understanding the role of the gut microbiome in a chronic inflammatory disease Study designs (cross-sectional, longitudinal, interventions): discussion Metagenomics: - Choice of biological material - Phylogeny and databases - 16S rRNA-based sequencing - Complete metagenomic sequencing - Imputation methods - Analytical approach (overview) (lectures, "Auditorium 1, Kreftssenteret")</p>
10:00	<p>1) Eirik Frengen:</p> <p>* Introduction & Overview * Information about "student assignments"</p> <p>* "Genome anatomy" * Genetic variation * Monogenic disorders</p> <p>(lectures, "Green auditorium, Bygg 25, Ullevål")</p>			<p>8) Sigve Nakken:</p> <p>"DNA sequencing concept: tumor versus control"</p> <p>Bioinformatics: from raw data to variants (pipelines) Background noise: oxidation, contamination etc. (lecture, "Auditorium 1, Kreftssenteret")</p>	
11:00		<p>4) Tuula Nyman:</p> <p>"Proteomics lecture 1": experiment planning, different workflows including some basics on how mass spectrometry is used in proteomics, main applications (lectures, "Green auditorium")</p>		<p>9) Katja Elgstøen Metabolomics</p> <p>(lecture, "Auditorium 1, Kreftssenteret")</p>	<p>Initiation of the student assignments: Each student gets 7 min to present a project plan & 3 min for feedback from the teachers. (Auditorium 1, Kreftssenteret)</p>
12:00	Lunch	Lunch	Lunch	Lunch	Lunch
12:30		<p>Tuula Nyman:</p> <p>"Proteomics lecture 2": (lectures, "Green auditorium")</p>		<p>10) Alex Rowe:</p> <p>Integration of omics data (statistics & bioinformatics) (lecture, "Auditorium 1, Kreftssenteret")</p>	
13:30	<p>2) Magnus Vigeland:</p> <p>"Identification of disease-causing variants with exome sequencing data"</p> <p>(Hands-on analysis of WES data in the PC room: "PC-stue, 1.etg", Bygg 25")</p>	<p>Tuula Nyman:</p> <p>Protein identification, quantitative proteomics experiments, some post-translational modification (Hands-on analysis of proteomics results in PC room: "PC-stue, 1.etg", Bygg 25")</p>	<p>6) Robert Lyle:</p> <p>"High Throughput Sequencing (HTS)" Transcriptomics/RNAseq/lncRNA/miRNA Epigenetics (lectures, "Green auditorium")</p>	<p>11) Stig Ove Bøe:</p> <p>"The Human Cell Atlas project"</p> <p>"The Human cell atlas project"- a project aimed at identifying and mapping all types of cells in the human body. Technological advances in defining cell types and/or states at the single cell level. Use of high content imaging for analysis of gene function at the single cell level. (lecture, "Auditorium 1, Kreftssenteret")</p>	<p>Continue the student assignments: Each student gets 7 min to present a project plan & 3 min for feedback from the teachers. The students will write a small project as a home exam, which should include a discussion on strengths and weaknesses of the selected approach. This will be submitted two weeks after the course.</p>
14:30-16				<p>12) Tobias Kaufmann, Dennis van der Meer and Ida E. Sønnderby:</p> <p>"Connectomics – Imaging genetics"</p> <p>Introduction to principles, strengths and challenges in brain imaging Brain networks and their disruptions in severe mental illness Neurodevelopmental and neurodegenerative perspectives Linking connectomics with genetics: the role of genes in connectivity Methods keywords: multivariate statistics, data driven analysis, machine learning for classification and prediction, independent component analysis, multimodal fusion, network analysis/graph theory, polygenic risk (lectures, "Auditorium 1, Kreftssenteret")</p>	