

Bioinformatics Unit: Analysis Services

The unit provides data processing and analysis for genomic, epigenomic and transcriptomic data from microarrays and high-throughput sequencing

Microarrays

Genome

- SNV panels
- QC, Variant calling
 - Custom genotype clustering

Epigenome

- DNA methylation (450K, EPIC)
- QC, Normalization
 - Methylation calling (beta/M values)
 - Clustering
 - Differential methylation (DM/DMR)
 - Gene + regulatory elements association

Transcriptome

- Gene expression arrays
- QC, Normalization
 - Quantification
 - Clustering
 - Differential expression
 - Gene set enrichment analysis
- microRNA arrays
- QC, Normalization
 - Quantification
 - Clustering
 - Differential expression
 - Target prediction
 - Correlation miRNA-gene expression

High-throughput sequencing

Genome

- Whole Genome Sequencing (WGS)
- Exome sequencing
 - QC, Alignment
 - Variant calling

Epigenome

- ChIP-seq, ATAC-seq, DNase-seq
- QC, Alignment
 - Peak calling
 - Differential binding
 - Functional analysis
- Whole Genome Bisulfite Sequencing (WGBS),
Reduced Representation Analysis Bisulfite
sequencing (RRBS)
- QC, Mapping
 - Methylation calling
 - Variant calling
 - Differential analysis (DMR/ methylome
segmentation)
 - Functional annotation

Transcriptome

- RNAseq (poly +/-, total)
- QC, mapping
 - Quantification
 - Alternative splicing
 - Clustering
 - Differential expression
 - Gene set enrichment analysis

Other services:

❑ Consultancy

- Experimental Design
- Statistical advice
- Recommendation analysis workflows and tools

❑ Training

- Internships
- Seminars
- Workshops