



Innovating Epigenetics Solutions

Epigenomics Profiling Services

Chromatin analysis

- Histone modifications via ChIP-qPCR/ChIP-seq
- Transcription factor binding via ChIP-qPCR/ChIP-seq
- Chromatin accessibility (ATAC-seq)

RNA sequencing and analysis

- mRNA and small RNA sequencing
- Low and difficult input RNA-seq
- Whole transcriptome analysis and gene expression profiling

DNA Methylation analysis

- **Genome-wide:** Whole genome bisulfite-seq, EPIC array, RRBS, MeDIP/MeDIP-seq
- **Targeted** DNA methylation assays
- **DNA hydroxymethylation** analysis

Bioinformatics data mining

- Machine learning for enhanced analysis of complex data
- Robust deep learning to uncover underlying biological significance of your data
- Establish new biomarkers for disease diagnostics

	Techniques	Standard analysis	Advanced analysis
Chromatin analysis	<ul style="list-style-type: none"> •ChIP-seq (histones/TFs) •ChIP-qPCR (histones/TFs) 	<ul style="list-style-type: none"> • Read filtering and trimming • Read mapping to the genome • Peak calling 	<ul style="list-style-type: none"> • Identification of differentially bound sites between groups of samples • Genomic annotation • Gene ontology enrichment • Pathway enrichment • Specific genomic region visualization
	<ul style="list-style-type: none"> • ATAC-seq 	<ul style="list-style-type: none"> • Read filtering and trimming • Read mapping to the genome • Identification of open chromatin regions 	<ul style="list-style-type: none"> • Differentially open chromatin region analysis • Genomic annotation • Nucleosome positioning • Motif recognition • Gene ontology enrichment • Pathway enrichment
DNA methylation analysis	<ul style="list-style-type: none"> • RRBS • WGBS 	<ul style="list-style-type: none"> • Read filtering and trimming • Read mapping to the genome • Methylation calling 	<ul style="list-style-type: none"> • CpG and region level differential methylation • Genomic annotation • Gene ontology enrichment • Pathway enrichment
	<ul style="list-style-type: none"> • MeDIP/hMeDIP-seq 	<ul style="list-style-type: none"> • Read filtering and trimming • Read mapping to the genome • Relative methylation quantification 	<ul style="list-style-type: none"> • Differentially-methylated region identification • Genomic annotation • Gene ontology enrichment • Pathway enrichment
	<ul style="list-style-type: none"> • EPIC arrays 	<ul style="list-style-type: none"> • Raw β- values • Data normalization • Probe filtering • Background correction 	<ul style="list-style-type: none"> • Differentially-methylated region identification • Genomic annotation • Gene ontology enrichment • Pathway enrichment
	<ul style="list-style-type: none"> • Targeted analysis 	<ul style="list-style-type: none"> • Read filtering and trimming • Read mapping to the genome 	<ul style="list-style-type: none"> • Differential methylation
RNA analysis	<ul style="list-style-type: none"> • RNA-seq 	<ul style="list-style-type: none"> • Read filtering and trimming • Read mapping to the genome or transcriptome • Gene and transcript quantification 	<ul style="list-style-type: none"> • Differentially-expressed gene identification • Annotation • Gene ontology enrichment • Pathway enrichment
Data mining	<ul style="list-style-type: none"> • Probabilistic modeling • Deep learning approaches • Integrative analyses of different -omics datasets 	<ul style="list-style-type: none"> • Feasibility phase - validation of the data set • Training and testing • Data mining analysis • Interpretation of results 	<ul style="list-style-type: none"> • Reporting • Model validation • Scientific support



www.diagenode.com Please contact us for more information

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