



## **HPC Core: Bioinformatics Analysis Rate 2025 Fall**

Category	Analysis Type	Pricing Basis	Internal Rate (USD)	NextLab Rate (USD)	External Profit Rate (USD)	Description	Pipeline Outline
<b>Bioinformatics Consulting</b>	Custom project	Per hour	\$55	\$83	\$165	Bioinformatics guidance, support, and solution	Experimental design, literature review, determination of best practices, methods recommendations, results interpretation
<b>AI/ML Consulting</b>	Custom project	Per hour	\$55	\$83	\$165	Artificial Intelligence (Machine learning and deep learning) guidance, support, and solution	Experimental design, literature review, determination of best practices, methods recommendations, results interpretation
<b>Data Analysis Consulting</b>	Custom project	Per hour	\$55	\$83	\$165	Data analysis guidance, support, and solution	Experimental design, literature review, determination of best practices, methods recommendations, results interpretation
<b>Genomics</b>	Whole Genome Sequencing (WGS)	Per sample	\$250	\$375	\$750	Comprehensive detection of genetic variants across the entire genome, including single-nucleotide changes and larger structural alterations.	FASTQ QC → adapter trimming → BWA-MEM alignment & duplicate marking → post-alignment QC → GATK variant calling → basic CNV/SV screening → VEP/ANNOVAR annotation → QC report & consult
	Whole Exome Sequencing (WES)	Per sample	\$230	\$345	\$690	Identification and annotation of coding-region variants	FASTQ QC → trimming → alignment (BWA-MEM) → post-alignment QC → GATK variant calling (exome)

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						in your sample's exome.	targets) → annotation → summary report & consult
	Targeted Gene Sequencing	Per sample	\$250	\$375	\$750	Focused analysis of specified gene panels to detect known or novel mutations in selected genes.	FASTQ QC → trimming → alignment to panel regions → variant calling on target genes → annotation → QC/variant summary
	Copy Number Variation (CNV) Analysis	Per sample	\$200	\$300	\$600	Detection of genomic regions with gains or losses (copy number changes) from your sequencing data.	Alignment (or use provided BAM) → CNVkit/ExomeDepth CNV calling → manual review → gene annotation → QC metrics & report
	Structural Variant (SV) Analysis	Per sample	\$210	\$315	\$630	Identification of larger genomic rearrangements such as deletions, duplications, inversions, and translocations.	Alignment (or use provided BAM) → Manta/LUMPY SV calling → filter high-confidence SVs → annotate breakpoints → QC report
	Genome-Wide Association Study (GWAS)	Per hour	\$55	\$83	\$166	Statistical association analysis linking genetic variants to traits or diseases in large cohorts.	Genotype QC (PLINK) → PCA for stratification → optional imputation (Minimac/Beagle) → association testing → multiple testing corrections → Manhattan/QQ plots
	Whole Genome Assembly	Per sample	\$250	\$375	\$750	Bioinformatics guidance, support, and solution with literature Review, Methods Survey, & Tool Recommendation	Generation of a high-quality draft genome sequence, including annotation of genes and genomic features.
<b>Epigenomics &amp; Accessibility</b>	Methylation & Epigenomics Analysis	Per sample	\$250	\$375	\$750	Analysis of DNA methylation or histone	WGBS alignment (Bismark) or array processing → DMR analysis (BSseq/DSS) → peak

						modification patterns to study gene regulation and epigenetic changes.	calling for ChIP-Seq → DMR/peak annotation → QC reports & interpretation
	Chromatin Accessibility (ATAC-Seq)	Per sample	\$250	\$375	\$750	Mapping open chromatin regions to identify regulatory elements and transcription factor binding sites.	FASTQ QC → trimming → alignment (Bowtie2/STAR) → Tn5 cut-site & peak calling (MACS2) → differential accessibility (DiffBind) → QC metrics → report
<b>Transcriptomics</b>	Bulk RNA-Seq Analysis	Per sample	\$250	\$375	\$750	Quantitative profiling of gene expression across whole samples to identify differentially expressed genes.	FASTQ QC → trimming → alignment (STAR/Hisat2) → gene quantification (featureCounts/RSEM) → differential expression (DESeq2/edgeR) → QC plots → summary report
	Single-Cell RNA-Seq (scRNA-Seq) Analysis	Per sample	\$260	\$390	\$780	High-resolution, cell-by-cell gene expression profiling to identify distinct cell populations and their marker genes.	FASTQ QC → Cell Ranger/STARsolo quantification → filtering & QC → normalization (SCTransform) → PCA+UMAP → clustering & annotation → differential expression → QC report
	De Novo Transcriptome Assembly & Annotation	Per sample	\$200	\$300	\$600	Construction of a reference transcriptome for organisms without a reference genome, including functional annotation of transcripts.	Read QC → de novo assembly (Trinity/Velvet) → transcript clustering (CD-HIT) → functional annotation (Trinotate) → assembly quality (TransRate) → summary report
	Spatial Transcriptomics Analysis	Per sample	\$280	\$420	\$840	Mapping gene expression patterns to	bustools) → image registration (Giotto/STUtility) → spot QC

						their spatial locations within tissue sections to reveal tissue architecture and microenvironments.	→ normalization & clustering (Seurat/Scanpy) → spatial gene detection → deconvolution → visualization → pathway enrichment → report & consult
	MicroRNA (miRNA-Seq) Analysis	Per sample	\$220	\$330	\$660	Profiling of small non-coding RNAs (microRNAs) to study their expression and regulatory roles.	Adapter trimming → alignment (miRDeep2/STAR) → miRNA quantification → differential analysis (DESeq2) → QC metrics → summary report
	Long Non-Coding RNA (lncRNA-Seq) Analysis	Per sample	\$230	\$345	\$690	Measurement of long non-coding RNA expression levels to study their potential functions in biology.	FASTQ QC → alignment to transcriptome (STAR) → quantification (featureCounts/RSEM) → differential analysis (DESeq2) → QC report
	Alternative Splicing Analysis	Per hour	\$55	\$110	\$220	Identification and quantification of different splice isoforms to understand transcript diversity.	STAR 2-pass alignment → isoform assembly (StringTie/Cufflinks) → junction quantification (rMATS/LeafCutter) → differential splicing → sashimi plots → report
<b>Metabolomics &amp; Lipidomics</b>	Untargeted Metabolomics Analysis	Per sample	\$250	\$375	\$750	Comprehensive profiling of small-molecule metabolites to discover differences between sample groups.	Feature detection (XCMS/MZmine) → peak alignment & annotation (HMDB) → statistical analysis (MetaboAnalyst) → pathway enrichment → QC → summary report
	Targeted Metabolomics Analysis	Per sample	\$250	\$375	\$750	Quantitative measurement of predefined metabolites using targeted assays.	Targeted feature extraction → calibration curve integration → quantification → QC (internal standards) → statistical summary → pathway annotation
	Lipidomics Analysis	Per sample	\$250	\$375	\$750	Identification and	Lipid feature detection → annotation (LIPID MAPS) →

						quantification of lipid species to study lipid metabolism and related pathways.	differential abundance → QC (calibration) → pathway & network analysis → summary report
<b>Metagenomics &amp; Microbiome</b>	Shotgun Metagenomics Analysis	Per sample	\$280	\$420	\$840	Genome-scale profiling of microbial community composition and function in complex samples.	FASTQ QC → host read removal (KneadData) → taxonomic profiling (Kraken2/MetaPhlAn3) → optional assembly → functional profiling (HUMAnN3) → diversity metrics → summary report
	16S rRNA Sequencing Analysis	Per sample	\$280	\$420	\$840	Targeted sequencing of bacterial 16S rRNA genes to analyze microbial community structure and diversity.	Demultiplex & QC (QIIME2/DADA2) → ASV/OTU calling → taxonomic assignment (SILVA/Greengenes) → alpha/beta diversity → differential abundance → summary report
	Metatranscriptomics Analysis	Per sample	\$250	\$375	\$750	Profiling of active microbial gene expression to understand functional activities in environmental or host-associated samples.	rRNA depletion QC → FASTQ QC → alignment (Salmon) → functional annotation (eggNOG-mapper) → differential expression → pathway enrichment (KEGG) → summary report
<b>Proteomics &amp; PTM</b>	Mass Spectrometry Proteomics Analysis	Per hour	\$55	\$83	\$166	Global analysis of protein abundance and differential expression using mass spectrometry data.	Raw file to mzML → database search (MaxQuant/Proteome Discoverer) → FDR filtering & quantification → differential protein analysis → pathway enrichment → summary report
	Post-Translational Modification (PTM) Analysis	Per hour	\$55	\$83	\$166	Identification and quantification of protein modifications such as	Variable modification search → site localization scoring (PTMP prophet) → quantification → functional annotation → summary report

						phosphorylation to study signaling pathways.	
	Protein-Protein Interaction (PPI) Analysis	Per hour	\$55	\$83	\$166	Mapping and analysis of protein interaction networks to identify key partners and functional modules.	Integrate PPI databases (STRING/BioGRID) → network construction & analysis (Cytoscape) → module detection → pathway identification → report
<b>Structural Bioinformatics</b>	Homology Modeling	Per protein	\$55	\$83	\$166	Building 3D models of proteins based on known structures to predict structure-function relationships.	Template search & alignment (BLAST/HHblits) → model building (MODELLER/SWISS-MODEL) → refinement & energy minimization → validation (Ramachandran/QMEAN) → report
	Molecular Docking	Per protein	\$55	\$83	\$166	Predicting how small molecules bind to target proteins for virtual screening or lead optimization.	Receptor prep → ligand library prep → docking runs (AutoDock Vina/Glide) → rescoring (MM-GBSA) → visualization of top hits → summary report
	Molecular Dynamics (MD) Simulation	Per protein	\$55	\$66	\$73	Simulating atomic motions of biomolecules to study dynamics and stability over time.	System setup (solvation/ion placement) → parameterization → equilibration & production (GROMACS/NAMD) → trajectory analysis → report
	QM/MM Simulations	Per protein	\$55	\$83	\$166	High-accuracy quantum/classical simulations to investigate enzyme mechanisms or reaction energetics.	Partition QM/MM → QM setup (basis set) → QM/MM run (Gaussian/ORCA with GROMACS) → energy profile analysis → report
<b>Functional Genomics</b>	CRISPR Screening Analysis	Per sample	\$110	\$165	\$330	Identifying genes essential for a	FASTQ QC → mapping to sgRNA library (MAGeCK) → hit calling & statistics → off-

						phenotype via pooled CRISPR knockout or activation screens.	target checks → pathway enrichment → summary report
	RNAi Screening Analysis	Per sample	\$110	\$165	\$330	Screening gene knockdowns via RNA interference to discover gene functions or drug targets.	FASTQ QC → mapping to shRNA library → dropout score calculation → hit calling → summary report
	Gene Expression Perturbation Analysis	Per sample	\$110	\$165	\$330	Assessing gene expression changes after treatments or perturbations to understand biological responses.	Differential expression (DESeq2) → pathway enrichment (clusterProfiler) → optional regulatory network inference → summary report
	Gene Regulatory Network (GRN) Analysis	Per hour	\$250	\$375	\$750	Constructing and analyzing gene networks to uncover regulatory relationships between genes.	Data normalization (voom/SCTransform) → network inference (ARACNe/WGCNA/GENIE3) → module detection → validation → report
<b>Clinical Trial &amp; Statistics</b>	Survival Analysis	Per hour	\$55	\$83	\$166	Analyzing time-to-event data to estimate survival probabilities and risk factors.	Data cleaning/censoring → Kaplan–Meier curves → Cox proportional hazards modeling → assumption checks → report
	Bayesian Statistical Analysis	Per hour	\$55	\$83	\$166	Performing statistical inference with Bayesian methods to quantify uncertainty in model parameters.	Model specification (priors/likelihood) → MCMC sampling (Stan/JAGS) → convergence diagnostics → posterior predictive checks → report
	RCT Analysis (Randomized Controlled Trials)	Per hour	\$55	\$83	\$166	Statistical analysis of clinical trial data to compare treatment	Data cleaning/randomization checks → ITT vs PP modeling → mixed-effects models for longitudinal outcomes → multiplicity adjustments → report

						effects and account for study design complexities.	
	Adaptive Clinical Trial Analysis	Per hour	\$55	\$83	\$166	Planning and analyzing adaptive trial designs that allow modifications based on interim results.	Interim analysis design & simulations → type I error control & sample size recalculation → interim execution → operating characteristics simulation → report

#### **Charge Per Sample with Biological Replicates & Time Points**

- Biological Replicates: Each independent sample (e.g., different individuals, cell lines) counts as one sample for analysis pricing.
- Time Points: If analyzing the same sample across multiple time points, each time point is considered as a separate dataset.

#### **Example Calculation:**

- Experiment: 3 replicates × 3 time points = 9 data files
- RNA-Seq Analysis (Bulk RNA-Seq: \$250/sample) →  $9 \times \$250 = \$2250$  total
- Single-Cell RNA-Seq Analysis (\$300/sample) →  $9 \times \$260 = \$2340$  total

**Notes:**

- All rates assume the client provides raw reads from sequencing platforms, raw FASTQ files, or aligned BAMs (and TIFFs for spatial). Wet-lab (extraction, library prep, sequencing, imaging) completed separately.
- Compute costs are absorbed in these rates. External nonprofit rates are 20% added to internal rates, and an extra 10% added for external profit rates.
- The pipeline description of the column is a potential list of tools and open-source packages. It does not contain any pricing based on any proprietary software, and pipeline modification is determined as per request.