

CODING FOR BIOLOGIST PROJECT TOPICS

- 1 **Genomic Data Visualization Tool** Develop an integrated bioinformatics tool for visualizing and exploring genomic data, including gene expression profiles with its mutations
- 2 **3D Protein Structure and Drug Visualizer and Analysis Tool** Develop an interactive platform for 3D protein structure visualization, allowing detailed analysis of protein-ligand interactions
- 3 Microbiome Diversity and Functional Analysis Tool Build a comprehensive tool for analyzing microbiome datasets, providing rich visualizations of microbial diversity, composition, and functional capabilities
- 4 Cancer Mutation Signature Analysis App Develop a user-friendly software tool to analyze mutational signatures in cancer genomics, uncovering patterns that can guide therapeutic strategies
- 5 **Gene Expression Classification Web App** Design a Streamlit app to upload gene expression data and classify disease vs. normal using ML models (SVM, Random Forest).
- 6 **Protein Stability Predictor Based on Mutations** Use Biopython and ML to predict the effect of mutations on protein stability using sequence and structure info.
- 7 Rare Disease Variant Prioritizer Using NLP + ML Use text mining and ML to prioritize candidate variants in rare disease cases based on literature co-occurrence.
- 8 **miRNA Target Prediction using ML** Use sequence features to train a model that predicts miRNA-mRNA interactions.
- 9 **Single-cell RNA-seq Clustering and Cell Type Classification** Perform clustering of scRNA-seq data to uncover cellular heterogeneity.
- 10 **Epigenetic Pattern Prediction Using ML** Predict DNA methylation or histone modification sites using sequence and structural features.
- 11 RNA-Seq Analysis Pipeline (Snakemake/Nextflow) Build an automated RNA-seq pipeline including trimming, alignment, and expression quantification.
- 12 Variant Calling Pipeline for NGS Data Create a complete script-based pipeline from FASTQ to VCF using tools like STAR, GATK.
- **Automatic Literature Curation from PubMed** Build a PubMed scraper using Python and NCBI Entrez API to collect articles by keywords.
- 14 **In-Silico Mutation Analysis on Disease Genes** Study the structural and functional impact of known SNPs in disease-related genes.
- 15 **Host–Pathogen Interaction Analysis** Map interactions between host and viral proteins in datasets (e.g., SARS-CoV-2, HIV).
- 16 **RNA Editing Site Exploration in Neurodegeneration** Identify RNA editing sites in ALS or Alzheimer's patient RNA-seg datasets.
- 17 Al-Based Drug Side Effect Prediction from Molecular Data Use ML to predict potential adverse drug reactions based on chemical descriptors and targets.

- 18 **Al Model for Identifying Functional IncRNAs** Use Al to distinguish functional vs. non-functional long non-coding RNAs using multi-feature inputs.
- 19 **Prediction of Cancer Progression Using Multi-Omics Data** Combine transcriptomics, proteomics, and metabolomics data to predict cancer progression using ensemble machine learning models
- 20 Analysis of Tumor Microenvironment Using Single-Cell Data Utilize unsupervised learning to identify cell populations and interactions in the tumor microenvironment
- 21 **Identifying De Novo SCN1A Mutations in Dravet Syndrome** Combining AlphaMissense Deleteriousness Scores with dbSNP Absence and Parental Confirmation
- 22 **Pharmacogenomic Variant Identification in CYP2C19** Utilizing AlphaMissense to Predict Functional Impact of Rare Variants Beyond dbSNP-Curated Alleles
- 23 **Population-Specific Pathogenic Variant Burden in PAH for Phenylketonuria** Integrating AlphaMissense, dbSNP Population Data, and ClinVar
- 24 Developing Allosteric Inhibitors of Plasmodium falciparum DHODH via Fragment Linking for Malaria
- 25 Leveraging Cryo-EM Structures of the MC4R-Gs Complex for Agonist Design in Obesity
- 26 Structure-Based Optimization of RSV Fusion (F) Protein Inhibitors for Respiratory Syncytial Virus Infection
- 27 CFTR Gene Variant Analysis for Cystic Fibrosis using ClinVar API and Python
- 28 Virtual Screening + ML Re-Ranking for MC4R Agonists
- 29 Deep Learning for Predicting Type 2 Diabetes from Gut Microbiota
- 30 Multi-Class Disease Prediction Using Microbiome Data
- 31 Time Series Analysis of Cell Growth or Infection Progression
- 32 Codon Optimization Tool for Synthetic Genes Create a codon usage bias calculator for optimizing gene sequences for different expression hosts (e.g., E. coli vs. human).