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EXPERIENCE

PROJECT- TRANSCRIPTOME DATA ANALYSIS TO IDENTIFY THE RELATION BETWEEN ADIPOGENESIS AND ANGIOGENESIS

Present

SELF PROJECT IN VARIANT CALLING

VARIANT CALLING-germline variants in a human WGS paired end reads using GATK Good Practice Workflow

Developed a complete variant calling workflow for identifying genomic variants from NGS data. The pipeline involved raw sequence quality assessment, adapter and low-quality base trimming, alignment of reads to the reference genome (hg38), sorting and indexing of BAM files, and variant detection using GATK. Post-processing included variant filtration and annotation to identify SNPs and INDELs. The project strengthened understanding of data preprocessing, alignment strategies, and variant interpretation in genomics.

DEGREE PROJECT WORK

EDUCATION

M.Sc. Computational Biology 7

08/2024 - Present

Department of Computational Biology and Bioinformatics, University of Kerala

Specialization: Next-Generation Sequencing (NGS) Data Analytics.Relevant Coursework: Bioinformatics, Genomics, Big Data in Biology, and Machine Learning for Biological Systems.

B.Sc. Biochemistry 7 06/2021 - 06/2024

Government of College Kariavattom, University of Kerala

SKILLS

Leadership PCR

Communication DNA Isolation

RNA seq analysis

Variant interpretation

Clinical Biochemistry

Genome Alignment

Gel Electrophoresis

SPAdes and QUAST NGS tools

Shell Scripting Using Bash Variant Calling using GATK

Perl Programming Language HTML,CSS,JS

Python R Programing language

LANGUAGES

English Hindi

1/2

Tamil Malayalam