



### **Certificate of Achievement**

# Ankit singh

has completed the following course:

# BIOINFORMATICS FOR BIOLOGISTS: ANALYSING AND INTERPRETING GENOMICS DATASETS WELLCOME CONNECTING SCIENCE

The course covered NGS sequencing, file formats used, Linux commands for sequence quality control, mapping, and variant calling. Same tasks were performed using Nextflow, applied to viralrecon pipeline with data visualization of variants obtained from genomic data as part of the downstream analysis.

3 weeks, 6 hours per week

Dr Treasa Creavin

Head of Scientific Meetings and Digital Learning Wellcome Connecting Science

**Dr Fatma Guerfali**Researcher
Institut Pasteur de Tunis







### Ankit singh

has completed the following course:

## BIOINFORMATICS FOR BIOLOGISTS: ANALYSING AND INTERPRETING GENOMICS DATASETS

WELLCOME CONNECTING SCIENCE



The course covered Next Generation Sequencing and its significance in genomics, the installation of widely-used bioinformatics tools, the use of different file formats, and Linux commands for sequence quality control, mapping, and variant calling. The analysis was also performed using the workflow management system Nextflow, and applied to viralrecon, an existing bioinformatics pipeline. Downstream analysis included RStudio data visualization of variants obtained from genomic data sample.

### STUDY REQUIREMENT

3 weeks, 6 hours per week

#### LEARNING OUTCOMES

- Use software managers to install and run reproducible bioinformatics tools
- Handle and analyse sequence datasets through hands-on exercises
- Analyse quality control metrics for sequencing data
- Modify existing workflows to suit specific task requirements and optimise analysis processes
- Interrogate and interpret the results obtained from running bioinformatics pipelines
- Perform downstream analyses of pipeline outputs using R, enabling data visualisation and further exploration

#### **SYLLABUS**

- Brief introduction to sequencing technologies and an overview of current sequencing outputs
- Sequence quality control to ensure data accuracy and reliability
- Bioinformatics workflows utilising tools such as Nextflow using nf-core pipelines

- Mapping sequencing data to a reference genome for alignment and variant calling
- Working with pipeline outputs in R to perform downstream analysis and visualisation.

