Finding Breakpoints of Viral Genomes using recan

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1 Introduction

In many different group of viruses recombination of genetic material is an important evolutionary process that generates much of genetic diversity.The patterns of recombination evident within the genomes of viruses can reveal lots of detail about their evolution and biology.Recombination is responsible for the evolution of viruses in response to selective forces present in a host environment [1]. A better understanding of recombination events taking place in the genomes of viruses are useful for gaining in-sites about the biology of the viruses.The patterns of sequence exchange between viruses in different species can reveal otherwise undetectable ecological links between some species and barriers between others.[2]. The breakpoints of genomes can reveal details about mechanistic and biochemical processes regarding the recombination process.[2].This project makes use of the python package 'recan'[3](recombination analyzer) for constructing genetic distance plots and exploring them.Also, we make use of bio-python[4] package for plotting fasta sequences.

2 Methods:

2.1 recan Python Package:

recan is a python package developed for the recombination analysis and it provides a mean for constructing genetic distance plots and exploring them interactively.recan is based on python packages like Biopython, Pandas, Matplotlib, and Plotly libraries.recan requires the input to be in the form of alligned fasta sequence.There are options for changing the window size,adjusting the sequence of interest(for example; sequence where the breakpoints occurs), changing the method of distance calculations etc.

2.2 Input Data:

The fasta sequence of viral genomes can be obtained from any of the protein data banks. RCSB protein data bank [5](https://www.rcsb.org/) is most popular among bioinformatic scientists. We can directly obtain the fasta sequence of the genome from RCSB.Since the recan only takes aligned fasta format of the genome, the fasta sequence downloaded from the RCSB protein data bank needs to be alligned.This can be achieved by using NCBI Multiple SEquence Allignment Viewer(https://www.ncbi.nlm.nih.gov/projects/msaviewer/).In this project, i chose lumky skin disease virus's genome [6] for finding out the breakpoints. I have also used the genome sequence of BCRA1 genes, which are the genes that produce tumor suppressor proteins, for demonstrating the visualization of the fasta sequence.
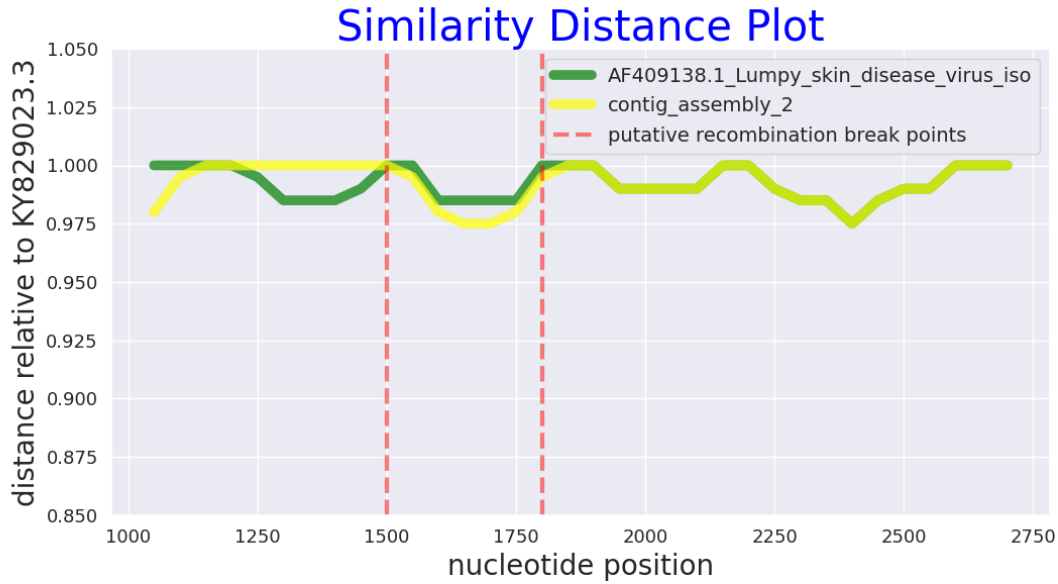
Figure 1: Distance Plot

3 Results and Discussion:

3.1 Breakpoints:

In genetics, the rearrangement of chromosomes is called mutation.It is an abnormality in the chromosome involving a change in the native chromosome [7].Theses changes may involve several events such as, duplication, inversion, deletion and translocation.These events are usually caused by a breakage in the helices of DNA and rejoining of the broken ends to form a new gene,different from the gene order of the chromosomes before they were broken [8].Some regions of chromosomes are more susceptible for rearrangements than others and thus are sources of genetic diseases and cancer.The Figure1 shows the breakpoints of lumky skin disease viral genome. We can see that, the recombination breakpoints are at around nucleotide number 1500 and at 1800. This is the clear evidence that recombination is taking place and the genome is undergoing mutation.

3.2 FASTA Sequence Plotting

The FASTA sequences of genomes can be visualized in several ways. We can make use of the Biopython package of renac for plotting the sequence of BCRA1 gene having multiple sequences.The Figure2 shows the line plot and Figure3 shows the histogram of BCRA1 gene.

One of the most commonly used analytical data to compare different sequence is GC Percentage. In genetics and molecular biology, GC content(or guanine-cytosine content) is the s the percentage of nitrogenous bases in a DNA or RNA molecule that are either guanine (G) or cytosine (C)[9].The GC percentage indicates the proportion of bases out of an implied four total bases, also including adenine and thymine in DNA and adenine and uracil in RNA.Figure4shows the GC percentage of BCRA1 gene.
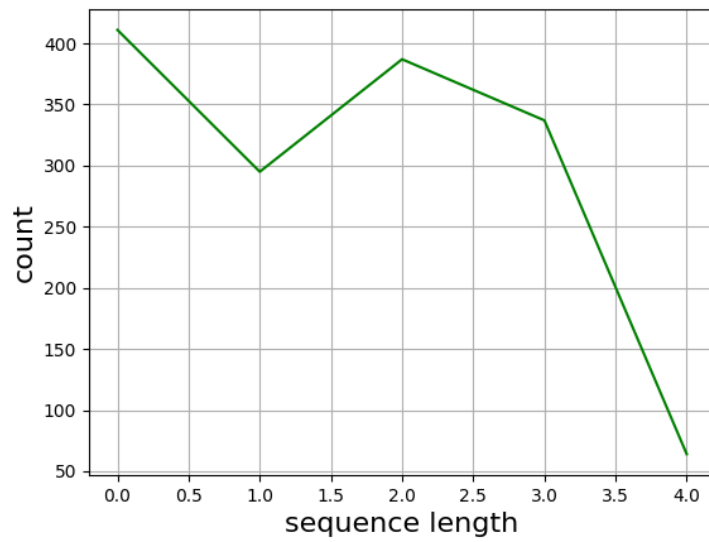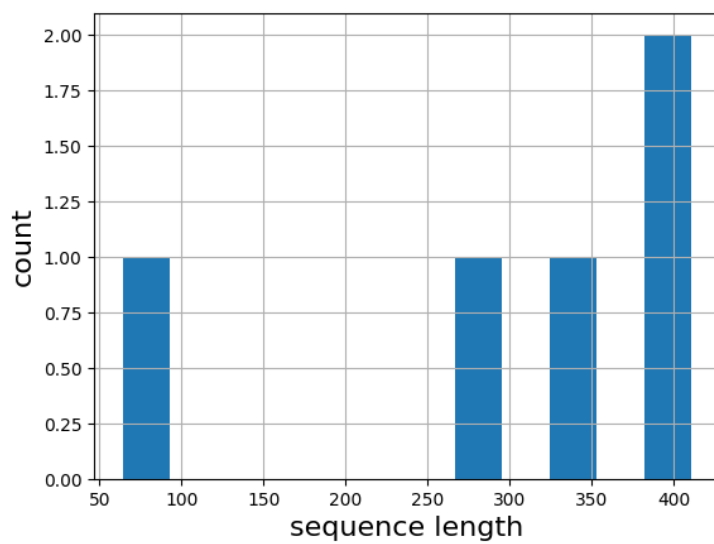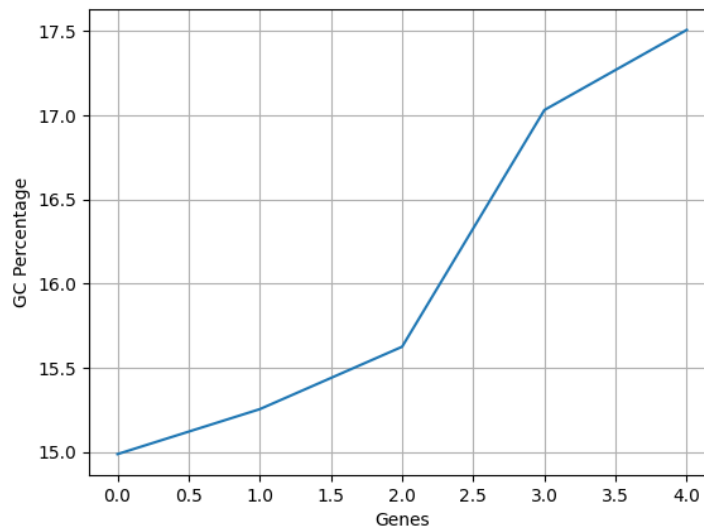
Figure 2: Line plot

Figure 3: Histogram

Figure 4: GC Content

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