Overview of HPViewer

Contents

| Description \dots | 1 |
|---------------------|---|
| Installation | 1 |
| Parameters | 1 |
| Results | 2 |
| Basic Usage (demo) | 2 |
| Workflow | |

Description

HPViewer is a tool for **genotyping and quantification of HPV from metagenomic or human genomic shotgun sequencing data.** We designed it to improve performance by masking nonspecific sequences from reference genomes and directly identifying HPV short DNA reads. It contains two HPV databases with different masking strategies, repeat-mask and homology-mask and one homology distance matrix to choose between those two databases.

If you use the HPViewer software, please cite our manuscript:

Yuhan Hao, Liying Yang, Antonio Galvao Neto, Milan R. Amin, Dervla Kelly, Stuart M. Brown, Ryan C. Branski, Zhiheng Pei. "HPViewer: Sensitive and specific genotyping of human papillomavirus in metagenomic DNA" (Submitted)

Installation

```
$ git clone https://github.com/yuhanH/HPViewer.git
```

Pre-requisites

Python (2.7+)

Python packages (sys, getopt, subprocess)

Bowtie2: http://bowtie-bio.sourceforge.net/bowtie2/manual.shtml

SAMtools: http://www.htslib.org/

Bedtools: http://bedtools.readthedocs.io/en/latest/

Parameters

Required

- a) input files (**-U** or **-1 -2**): fastq files (or fastq.gz), unpaired (**-**U unpaired.fastq) or R1,R2 paired (**-**1 R1.fastq **-**2 R2.fastq)
- b) output file name (**-o**)

Optional

- a) database mask type (-m): hybrid-mask(default), repeat-mask, homology-mask. repeat-mask is a more sensitive mode; and homology-mask is suggested when some types of HPV are present in large abundance which may lead to false positive of other types of HPV.
- b) number of threaded used in bowtie2 alignment (-p)
- c) minimal coverage threshold to determine HPV present (-cov), default is 150 bp (1.5 x average length of your reads).

Results

a) output_HPV_summary.txt has three coloumns with types of HPV present, number of reads per kilobase (RPK) for the matching HPV, and number of reads of the matching HPV.

$$average\ coverage = \frac{RPK*average\ reads\ length}{1000}$$

b) alignment results after bowtie2: output.sam, output.bam

Basic Usage (demo)

```
python HPViewer2.py -U test_unpaired.fastq -o TEST
more TEST/TEST_HPV_profile.txt
## HPV_type RPK(reads_per_kilobase) count_of_reads
## HPV5 114.3199
                    859
## HPV7 109.0264
                    860
## HPV21
            120.3347
                         906
## HPV49
            109.3479
                         820
## HPV66
            123.1417
                         936
## HPV69
            108.1513
                         812
## HPV73
            117.2339
                         868
## HPV88
            126.1261
                         924
## HPV101
            115.1348
                         833
## HPV117
            116.1049
                         899
```

Workflow

