

# Workbook

## Basic Bioinformatics for Biologist

### 1. Linux basic commands

Command	Function
pwd	Print the current working directory (where you are)
ls	List files and directories in the current directory
cd	Change directory (move to another folder)
mkdir	Create a new directory
rmdir	Remove an empty directory
cp	Copy files or directory
mv	Move or rename files or directories
rm	Remove (delete) files
cat	Display the entire content of a file
less	View file content page by page (scrollable)
head	Show the first lines of a file
tail	Show the last lines of a file
grep	Search for patterns/keyword inside files
wc -l	Count the number of lines in a file
cut	Extract specific columns or fields from text
sort	Sort lines of a file
uniq	Remove duplicate lines (work after sort)
echo	Print a message or variable to the terminal
man	Show the manual/help page of command

\*) put `-h` or `--help` option to show the short instruction for each command

## 2. Obtain training data

### 2.1 Update package list

```
$ sudo apt-get update
```

### 2.2 Upgrade general packages

```
$ sudo apt-get upgrade
```

### 2.3 Install git using apt package manager

```
$ sudo apt-get install git
```

### 2.4 Clone training data from github

```
$ git clone https://github.com/reditama/bioinformatics-workshop.git
```

### 2.5 Check if the cloning is complete

```
$ ls -la
```

## 3. Basic Linux operation

### 3.1 Change directory to bioinformatics-workshop

```
$ cd bioinformatics-workshop
```

### 3.2 List files and folder in the directory

```
$ ls
```

### 3.3 Show the file size of files

```
$ ls -la
```

### 3.4 View the content of ecoli\_500kb.fasta

```
$ less ecoli_500kb.fasta
```

### 3.5 View the content of ecoli\_500kb.fastq.gz

```
$ less ecoli_500kb_hifi.fastq.gz
```

### 3.6 Use the -S option to unwrap the content

```
$ less -S ecoli_500kb.fastq.gz
```

### 3.7 View the content of gff file using head

```
$ head ecoli_K12_MG1655_example.gff3
```

### 3.8 Find gene annotation using grep

```
$ grep gene ecoli_K12_MG1655_example.gff3
```

### 3.8 Count the number of gene (Use pipe to perform commands simultaneously)

```
$ grep gene ecoli_K12_MG1655_example.gff3 | wc -l
```

## 4. Software installation

### 4.1 Seqkit installation

Hint: ask chatgpt

### 4.2 fastp installation

Hint: ask chatgpt

### 4.3 flye installation

```
$ cd
$ git clone https://github.com/fenderglass/Flye
$ cd Flye
$ make
```

## 5. Assembly preparation

### 5.1 Change directory to home (cd default argument is home/username)

```
$ cd
```

### 5.2 Create new folder called assembly and move inside (use && to perform a series of commands)

```
$ mkdir assembly && cd assembly
```

### 5.3 Copy ecoli\_500kb.fasta to assembly folder

```
$ cp ../bioinformatics-workshop/ecoli_500kb_hifi.fastq.gz .
```

### 5.4 Inspect the statistics of fastq

```
$ seqkit stats ecoli_500kb_hifi.fastq.gz
```

### 5.5 Put the statistics in a file

```
$ seqkit stats ecoli_500kb_hifi.fastq.gz > ecoli_500kb_hifi.stats.txt
```

### 5.6 Perform qc using fastp, put your cleaned fastq in a new file (ecoli\_500kb\_hifi\_clean.fastq.gz)

Hint: use fastp --help to learn how to.

### 5.7 Inspect the statistics of cleaned fastq

```
$ seqkit stats ecoli_500kb_hifi_clean.fastq.gz >
ecoli_500kb_hifi_clean.stats.txt
```

### 5.8 Assembly the clean reads using flye

```
$ /home/reditama/Flye/bin/flye --pacbio-raw
ecoli_500kb_reads_hifi_clean.fastq.gz --genome-size 5m --out-dir
flye_output
```

### 5.9 Inspect the output directory

```
$ cd flye_output  
$ ls -la
```

### 5.10 Inspect the statistics of assembled contigs

```
$ seqkit stats assembly.fasta > assembly.stats.txt
```

\*\*\*