Linux Command Line

Rachel Wood

2023-03-22

For this portfolio I will use the example of bioinformatics to showcase the utility of the Linux command line. The data we will be using is the genome and protein datasets from the SARS-CoV-2 virus, which can be found at https://www.ncbi.nlm.nih.gov/genome/?term=SARS-CoV-2.

Working with Files and File Systems

We start in a folder containing the files downloaded from the link above:

```
ls
GCF_009858895.2_ASM985889v3_genomic.fna.gz
GCF_009858895.2_ASM985889v3_protein.faa.gz
```

We first create a folder to work within using the mkdir command and use the cd command to move into this new file. Finally we can use the ls command to view the contents of this folder:

```
mkdir bioinf_ex
```

As expected, the directory is empty, but we can move the genome and protein files to our new folders. We use the cd .. to go one 'step back' in the directory and return to our original directory. We use the ls command again and then the mv command to move our two files into the new folder we have created:

```
mv GCF_009858895.2_ASM985889v3_protein.faa.gz -t bioinf_ex/
mv GCF_009858895.2_ASM985889v3_genomic.fna.gz -t bioinf_ex/
cd bioinf_ex/
```

We now check these files have been successfully moved:

```
ls
```

```
GCF_009858895.2_ASM985889v3_genomic.fna.gz
GCF_009858895.2_ASM985889v3_protein.faa.gz
```

We notice the .gz file extension, meaning we need to unzip the files. We can do this with the gzip command:

```
gzip -d GCF_009858895.2_ASM985889v3_genomic.fna.gz
gzip -d GCF_009858895.2_ASM985889v3_protein.faa.gz
```

าร

```
GCF_009858895.2_ASM985889v3_genomic.fna
GCF_009858895.2_ASM985889v3_protein.fa
```

We finally rename the files to something more convenient with the mv command:

```
mv GCF_009858895.2_ASM985889v3_genomic.fna genomic.fna
mv GCF_009858895.2_ASM985889v3_protein.faa protein.faa
```

Genomic Data

This section focuses on the genomic.fna file. We first use the head command to view the first few lines of the file:

We might want to know how long a genome is, this is often measured by the number of base pairs, here each pair is represented by a character - a base pair is either A-T or G-C. We use the grep command using the -o flag instead combined with the wc command to get the count of A, C, G or T characters in the file (excluding the header line):

```
tail -n+2 genomic.fna | grep -o [ATCG] | wc -1
```

We can use similar commands to obtain the individual counts of the letters:

```
tail -n+2 genomic.fna | grep -o A | wc -l
8954
tail -n+2 genomic.fna | grep -o T | wc -l
9594
tail -n+2 genomic.fna | grep -o C | wc -l
5492
tail -n+2 genomic.fna | grep -o G | wc -l
5863
```

From this we can see the genome is A-T enriched.

The start codon Methionine is represented by ATG, we can see how many times this appears.

For this we first need to remove the header line and all new lines characters (denoted \n) and save the changes in a new file:

```
tail -n+2 genomic.fna > genomic_new.fna
echo -n $(tr -d "\n" < genomic_new.fna) > genomic_new.fna
```

Note the use of echo, this creates a subshell, which allows us to read from and write to the same file.