## **Command line Practical**

### **Basic Unix Command Line Skill**

**Note:** Lines that start with a \$\\$ should be entered on the command line without the \$\\$

#### 1. Using terminal in Virtual Box

- Open the Terminal application in the Virtual Machine window
- Set up Shared Clipboard in virtual box. Navigate to the Devices tab on the menu bar and set it to bidirectional.

#### 2. Change to /home/<virtual\_box\_name>/Desktop/ and make a folder

- \$ cd /home/<virtual\_box\_name>/Desktop/
- \$ mkdir workshop This will be where your workshop documents will be stored.
- \$ 1s to your folder and everyone else's
- \$ cd workshop change into your directory

#### 3. Download and Copy the sample data

• Today's sample data can be downloaded here:

```
https://github.com/gonzalezvl/Microbial_Metagenomics_Workshop_UCLA_2015/
```

- use cp to copy the directory Command\_line\_practical and its contents to your new folder /home/<virtual box name>/workshop/
  - We're not giving the exact command line here. Hint: Find the cp option to copy an entire directory. The format of the cp command is cp [options] source destination For the destination use . to copy to the current folder. Did you use an absolute path or a relative path for source and destination?
- \$ 1s -1 to confirm that the folder copied successfully. You should see the Day1 folder.
- Error about ...omitting directory... ? Check the cheat sheet for the cp option to copy a whole directory.

#### 4. Viewing file contents

- Use cd to change the directory to Command\_line\_practical and then change again into data which has the sequence files we'll be using. Use \$ ls -1 to confirm that the two sequences.fa and sequences.fq are there.
- Try all four of these commands to examine the file sequences.fa
  - a) \$ cat sequences.fa
  - b) \$ head sequences.fa
  - c) \$ tail sequences.fa
  - d) \$ less sequences.fa (Remember, q is quit in less)
  - e) Use less to *find the sequences* that contains the description EAS20\_8\_6\_1\_5\_388 (Type h in less for help screen, look for "Search forward..." in the "SEARCHING" section.).
  - f) Use head to display the first sequence only (first two lines). Type \$ man head (or Google) to find the option to limit the number of lines head displays (man pages open in less, use the arrow keys to navigate and q to quit).
  - g) Use tail and wildcard globbing to display the last 10 lines of the two sequences files (sequences.fa and sequences.fq) with one command.

#### 5. Edit a file with nano

- Open sequences.fa with nano
- Change some bases in one of the sequences. We'll need this file in the afternoon so keep it valid fasta.
- Save and close the file Remember in nano shortcuts are at the bottom of the screen and ^ is the control kev.

#### 6. Create a new text file and delete it

- \$ nano newfile If the filename doesn't exist, an empty document will be opened.
- Enter some text into the file, save it and exit.
- Use one of the text viewing commands we used in Step 4 to view your new file's contents.
- After viewing your file delete it with the rm command

# Digging deeper (optional for more advanced users):

1. Want to learn the vi editor? Enter the command \$ vimtutor on Hydra or your Mac for a tutorial. To exit

type :q

- 2. Want to learn the emacs editor? Do the emacs tutorial by starting \$ emacs and then <control+h> and then t . To exit type <control+x> and then <control+c> .
- 3. Using \$ 1s -1 Compare the creation date of the original file and the one you copied. Check the cp man page or online resources to learn how to copy the file and preserve the original create date.
- 4. When you use less to view the fasta file, the lines automatically wrap ("fold" in less lingo) which can be annoying for fasta files with long sequences. What flag for less will stop this so the lines don't wrap ("chop" in less lingo)? Hint: this is an option when starting less rather than a command when less is already running.

**UCLA Microbial Metagenomics Workshop 2015** 

\*Modified from Hydra Workshop 2015\*