# **Practice: Using Python on the Unix Command Line**

BIOS 274: Introductory Python Programming for Genomics

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### WINDOWS USERS ONLY

### 1. Open Mobaxterm

2. Install python3 for MobaXterm

```
apt-get install python3
This will take quite a while to finish.
```

3. Install nano for MobaXterm:

/bin/apt-cyg install nano

You can open a new tab within MobaXterm and run this command while python3 is installing.

- 4. Settings → Configuration → Terminal (it's a tab at the top) → Check the box near the bottom next to "Use Windows PATH"
- 5. Close MobaXterm
- 6. Re-open MobaXterm

### WINDOWS USERS ONLY

### 7. Test whether python has been correctly installed

python3 --version
Python 3.6.8 should appear

### 8. Test whether nano has been correctly installed

#### nano

A different screen should pop up To exit nano, do 'ctrl' + X

# **Basic Unix Commands**

Print working directory (your current directory) pwd Change directory to specified PATH cd PATH cd .. Move up one directory level List all the files/folders in the directory ls Copy FILE to DIRECTORY CD FILE DIRECTORY Move FILE to DIRECTORY mv FILE DIRECTORY Delete/Remove FILE rm FTLE Delete/Remove DIRECTORY rm -r DIRECTORY Find all instances of 'STRING' in FILE grep 'STRING' FILE Count the number of words in FILE WC FILE wc -1 FILE Count the number of lines in FILE Open a simple plaintext editor nano

### **REMEMBER:**

Type man COMMAND (i.e. man grep) to see usage information!

# In Terminal (Mac) or MobaXterm (Windows):

1.	Change directory	to wherever v	you downloaded v	vour files for Dav	7 from Canvas
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2. Print out the names of all the files in the directory

3. From the terminal, view the contents of utils.py

4. Enter an interactive version of python on the command line

5. Use the functions in utils.py to find the complement and reverse complement of a DNA sequence

6. Exit the interactive version of python

Open sys.argv\_example.py in Atom.

Follow the directions at the top of the script to run it.

Open sumLinesOfFile.py in Atom.

Follow the directions at the top of the script.

Open shortStatureGenes.py in Atom.

Follow the directions at the top of the script.

## Can we find the diseases associated with a gene with basic Unix commands?

1. Find the diseases in shortStatureDisorders.tsv associated with FLNB. Save them to a file called FLNB.txt

2. How many diseases are associated with FLNB?

Hint: Use a flag with either grep or wc
Do man grep or man wc to see usage information.

3. Can you do step 1 for both FLNB and COL1A1 at the same time?

4. Display only the names of the disorders in FLNB.txt (without the column of genes).