

# More Command Line

# We've gone through many essential commands

- `ls`
- `wc`
- `pwd`
- `cd`
- `mkdir`
- `man`
- `rm`
- `touch`
- `mv`
- `echo`
- `less`
- `cat`
- `>> vs >`
- `grep`
- `sort`
- `| (pipe)`
- `nano`

# | (pipes)

You cannot be a productive command line user until you really understand the power of pipes

```
grep TP53 genes.txt | grep "missense" | wc -l
```

This kind of construction allows you to get answers quickly!

# I'm stuck!

- **Ctrl-C** to interrupt/kill a running process
- **q** quits some interactive commands (e.g. less)
- editing a file with vim?
  - press **Escape**
  - type **:q!**
  - press **Return**

# It's not working!

- Did you check case?
  - capital vs lowercase matters!
- Are you in the right directory?
  - use ``ls`` all the time!
- typos
  - tab-complete is your friend!

# First, let's download some data

## make a directory, then move into it

```
mkdir ~/workspace/commandline
```

```
cd ~/workspace/commandline
```

## download the files from the links posted in slack

```
wget <URL>
```

## use ls to list the files

```
ls -l
```

# Examine the files

genes1.txt = mutations identified in patient set 1

genes2.txt = mutations identified in patient set 1

# Working with `sort` | `uniq`

```
## Sorts genes in genes1.txt  
cat genes1.txt | sort  
  
## ... is equivalent to ...  
sort genes1.txt
```

Genes are sorted  
alphabetically, great!

```
CEBPA  
CEBPA  
DNMT3A  
FLT3  
IDH1  
IDH2  
IDH2  
NPM1  
NRAS  
RUNX1  
TET2  
TP53
```



# Working with `sort` | `uniq`

```
## Sorts genes in genes1.txt
```

```
cat genes1.txt | sort
```

```
## ... is equivalent to ...
```

```
sort genes1.txt
```

```
## Get the unique genes... right?
```

```
cat genes1.txt | uniq
```

DNMT3A

FLT3

NPM1

TET2

**IDH2**

RUNX1

TP53

**IDH2**

IDH1

**CEBPA**

NRAS

**CEBPA**

Why are IDH2 and  
CEBPA repeated?

# Working with `sort` | `uniq`

```
## Sorts genes in genes1.txt
```

```
cat genes1.txt | sort
```

```
## ... is equivalent to ...
```

```
sort genes1.txt
```

```
## Sort then unique
```

```
cat genes1.txt | sort | uniq
```

```
CEBPA  
DNMT3A  
FLT3  
IDH1  
IDH2  
NPM1  
NRAS  
RUNX1  
TET2  
TP53
```

`This is the way!`

(`uniq` only identifies matching values when they are immediately next to one another)

## More **sort** | **uniq** combinations

## Report only duplicate values (uniq -d flag)

```
cat genes1.txt | sort | uniq -d
```

## Report values that are in the file a single time (uniq -u)

```
cat genes1.txt | sort | uniq -u
```

## Count the number of occurrences for each value (uniq -c)

```
cat genes1.txt | sort | uniq -c
```

Unsorted data:

DNMT3A

FLT3

NPM1

TET2

**IDH2**

RUNX1

TP53

**IDH2**

IDH1

**CEBPA**

NRAS

**CEBPA**

# Performing set operations with `sort | uniq`

Question 1: how can we find if a gene is present in both genes1.txt and genes2.txt?

Question 2: how can we find genes only found in genes1.txt and not in genes2.txt?

# Performing set operations with `sort` | `uniq`

Question 1: how to find if a gene is present in both genes1.txt and genes2.txt?

```
cat genes1.txt | sort | uniq -u > genes1.uniq.txt
cat genes2.txt | sort | uniq -u > genes2.uniq.txt
cat genes1.uniq.txt genes2.uniq.txt | sort | uniq -d
```

Question 2: how do we find genes only found in genes1.txt and not in genes2.txt?

# Performing set operations with `sort` | `uniq`

Question 1: how to find if a gene is present in both genes1.txt and genes2.txt?

```
cat genes1.txt | sort | uniq -u > genes1.uniq.txt
cat genes2.txt | sort | uniq -u > genes2.uniq.txt
cat genes1.uniq.txt genes2.uniq.txt | sort | uniq -d
```

Question 2: how do we find genes only found in genes1.txt and not in genes2.txt?

```
cat genes1.uniq.txt genes1.uniq.txt genes2.uniq.txt | sort | uniq -u
```

# Examine the mutation file

```
less tcga.tsv
```

```
#wrap long lines
```

```
less -S tcga.tsv
```

```
#wrap long lines and set tab spacing to 20 characters
```

```
less -S -x20 tcga.tsv
```

# Examine the mutation file

```
less tcga.tsv
```

```
#wrap long lines
```

```
less -S tcga.tsv
```

```
#wrap long lines and set tab spacing to 20 characters
```

```
less -S -x20 tcga.tsv
```

```
type "q" to exit
```



# Sort the mutation file

```
sort tcga.tsv | less
```

```
#sort by chromosome (second column)
```

```
sort -k 2 tcga.tsv | less
```

```
#sort by chromosome, and then position numerically
```

```
sort -k 2,2 -k 3,3n tcga.tsv | less
```

# Extract info from the mutation file

```
#cut the 8th column (gene names)  
cut -f 8 tcga.tsv | less
```

```
#cut multiple columns  
cut -f 2-4,8,10 tcga.tsv | head
```

```
#find the most frequently mutated genes in this cohort  
cut -f 8 tcga.tsv | sort | uniq -c | sort -nrk 1 | head -n 20
```

# Some useful UNIX commands

- **head**      print the first 10 lines of a file
- **tail**      print the last 10 lines of a file  
getting fancy: **tail -n +2** (start at the second line of a file)
- **wc**      count the number of characters/words/lines in a file  
**wc -l** for only lines
- **less**      because you don't want 3 million lines scrolling through your terminal  
**q** to exit, **-S** to wrap lines (lots more useful options here)
- **grep**      to search through a file (**-v** to search for lines *without* pattern)

# Working with compressed data

- **tar**      work with a “bundle” of data

create:      **tar -cvf output.tar infile1 infile2**

extract:      **tar -xvf output.tar**

- **gzip**      compress a single file

create:    **gzip mydata.txt**      (creates mydata.txt.gz)

extract:    **gunzip mydata.txt.gz**      (creates mydata.txt)

Often these operations are combined

```
tar -czvf myfile.tar.gz <list of files>
```

```
tar -xzvf myfile.tar.gz
```

# sed and awk

sed is most commonly used for find and replace operations:

```
cat file.txt | sed 's/foo/bar/g' >file_fixed.txt
```

Awk can be used to reorder particular columns (here, third, first, then second):

```
awk '{print $3,$1,$2}' file.txt >file2.txt
```

Or to print only certain lines of a file - here, every third line, starting at line 0

```
awk 'NR % 3 == 0' file > file2.txt
```

(both are very powerful, if somewhat opaque tools, this is just scratching the surface!)

# Working with FASTQs

- <https://gist.github.com/chrisamiller/230cf13c1ee0ca10a5535279957f48a5>