

DELLA

```
ssh vonholdt@della.princeton.edu
```

Managing scripts:
to_run_mapping.sh
to_run_methylCall.sh
to_run_qseq2fastq.sh

```
cd /tigress/VONHOLDT  
cd /tigress/vonholdt
```

```
to_run_qseq2fastq.sh
```

```
### Job Name  
#PBS -N qseq2fastq  
#PBS -l nodes=1;ppn=1;walltime=0:10:00  
#PBS -M vonholdt@princeton.edu  
#PBS -m abe
```

```
cd /tigress/VONHOLDT/scripts  
./batchqseq2fastq.sh
```

```
ssh netID@della.princeton.edu
```

Managing scripts:
to_run_parser.sh
to_run_R_library1.sh

```
cd /tigress/VONHOLDT  
cd /tigress/netID
```

```
ssh netID@della.princeton.edu
```

Managing scripts:
to_run_dmrAnalysis.sh
to_run_plotDMR.sh

```
cd /tigress/VONHOLDT  
cd /tigress/netID
```

TIGRESS

```
/tigress/vonholdt  
/tigress/netID1  
/tigress/netID2
```

/tigress/VONHOLDT # this is the shared folder!
/tigress/VONHOLDT/data # do not move/edit this data
You must know the path of the data files so you can direct your script.sh (in /tigress/VONHOLDT/scripts) to the right input file and write your output to the correct place.

Example: /tigress/VONHOLDT/data contains
maleWolf.qseq
femaleWolf.qseq

/tigress/VONHOLDT/scripts # these are the working scripts
Store all of your scripts here and call them in a batch.sh file, analogous to a pipeline. Then point your to_run.sh to this batch.sh which actually contains all of your real commands.

Example: I want to run a perl script that converts the two qseq files to the fastq format. So I write a batch.sh file that is:

```
batchqseq2fastq.sh
```

```
#!/bin/bash  
  
./qseq2fastq.pl <maleWolf.qseq  
./qseq2fastq.pl <femaleWolf.qseq  
  
exit 0;
```