

```
#####  
####
```

**(c) 2015-2017 Merly Escalona**  
**[merlyescalona@uvigo.es](mailto:merlyescalona@uvigo.es)**

---

**Phylogenomics Lab. University of  
Vigo.**

---

**Description:**

---

**=====**

---

**Pipelines for data analysis**

---

**Running [@triploid.uvigo.es](mailto:@triploid@uvigo.es)**

---

```
#####  
####  
#!/bin/bash -l  
#####  
####
```

**Folder paths**

---

```
#####  
####  
source $HOME/src/vc-benchmark-cesga/src/vcs.variables.sh
```

```
#####
####
simphyReplicateID=1
pipelinesName="ssp"
replicatesNumDigits=5
replicateID="$(printf "%0${replicatesNumDigits}g" $simphyReplicateID)"
ngsphyReplicatePath="$HOME/data/NGSphy_${pipelinesName}.${replicateID}"
simphyReplicatePath="$HOME/data/${pipelinesName}.${replicateID}"
referencesReplicatePath="$HOME/data/references/references.${pipelinesName}.${replicateID}"
#####
####
```

## 1. REFERENCE INDEXING WITH BWA

---

```
#####
####
for fastaFile in $(find $referencesReplicatePath -name *.fasta); do
qsub $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.1.sh $fastaFile
done
#####
####
```

## 2. GENERATION OF BWA COMMAND LINESs

---

```
#####
####
```

**qsub -t \$simphyReplicateID  
\$HOME/src/vc-benchmark-  
cesga/jobs/analysis/ssp.analysis.2.sh  
PE150OWN HiSeq2500**

---

```
qsub -t $simphyReplicateID $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.2.sh
```

```

PE150DFLT HiSeq2500
qsub -t $simphyReplicateID $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.2.sh
SE150DFLT HiSeq2500
qsub -t $simphyReplicateID $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.2.sh
PE250DFLT MiSeqV3
qsub -t $simphyReplicateID $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.2.sh
SE250DFLT MiSeqV3
#####
####

```

## 3. MAPPINGS

---

```

#####
####
profiles=("PE250DFLT" "SE250DFLT") # ("SE150DFLT") # ("PE150DFLT") # ("PE150OWN") #
for profileFOLDER in ${profiles[]};do numJobs=$(find
"$HOME/data/mappings/${pipelinesName}.${replicateID}/scripts/" -name
"${pipelinesName}.${replicateID}.${profileFOLDER}.bwa.commands." -type f | wc -l );
echo $numJobs
qsub -t 1-$numJobs $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.3.sh
"$HOME/data/mappings/${pipelinesName}.${replicateID}/scripts/${pipelinesName}.${replicateID}
.${profileFOLDER}.bwa.commands"
done
#####
####

```

## 4. Generating BAMMING SORTING commands

---

```

#####
####
qsub -t $simphyReplicateID $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.4.sh
PE150DOWN HiSeq2500
qsub -t $simphyReplicateID $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.4.sh
PE150DFLT HiSeq2500
qsub -t $simphyReplicateID $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.4.sh
SE150DFLT HiSeq2500
qsub -t $simphyReplicateID $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.4.sh

```

PE250DFLT MiSeqV3

qsub -t \$simphyReplicateID \$HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.4.sh

SE250DFLT MiSeqV3

```
#####  
####
```

## 5. BAMMING SORTING

---

```
#####  
####  
profiles=("PE250DFLT" "SE250DFLT") # ("SE150DFLT") # ("PE150DFLT") # ("PE150OWN") #  
for profileFOLDER in ${profiles[]};do numJobs=$(find  
"$HOME/data/mappings/${pipelinesName}.${replicateID}/scripts/" -name  
"${pipelinesName}.${replicateID}.${profileFOLDER}.samtools.commands." -type f | wc -l );  
echo $numJobs  
qsub -t 1-$numJobs $HOME/src/vc-benchmark-cesga/jobs/analysis/ssp.analysis.5.sh  
"$HOME/data/mappings/${pipelinesName}.${replicateID}/scripts/${pipelinesName}.${replicateID}  
.${profileFOLDER}.samtools.commands"  
done  
  
#####  
####
```

## 6. INFORMATION ON THE MAPPING

---

```
#####  
####
```

To ask the view command to report solely “proper pairs” we use the -f option and ask for alignments where the second bit is true (proper pair is true).

samtools view -f 0x2 sample.sorted.bam

How many properly paired alignments are there?

samtools view -f 0x2 sample.sorted.bam | wc -l

Now, let’s ask for alignments that are NOT properly paired. To do this, we use the -F option (note the capitalization to denote “opposite”).

```
samtools view -F 0x2 sample.sorted.bam
```

How many improperly paired alignments are there?

```
samtools view -F 0x2 sample.sorted.bam | wc -l
```