

Analysis 1. Computing fixation index (Fst)

Fst is a measure of genetic differentiation between two populations. For this reason it can be a useful indicator of selection (as well as other things!).

As this is a scan, it is data, not hypothesis driven, and we compute Fst over a whole chromosome.

We use data from the ag1000 genomes project phase 2 data release. We use a VCF file, this is ok as we do not need phased haplotypes to compute Fst.

This analysis is carried out in windows, what might be an appropriate window size?

This analysis will be carried out on the lstm cluster.

NOTES: - The VCF file we are using has been downsampled to 10% to allow the analysis to be done in a short time. - The commands below can't be copied and pasted verbatim, you will need to change parts of them.

```
ssh yourusername@opteron.lstmed.ac.uk
```

Once you have logged into the cluster, we are going to create a directory in your home for this analysis. Good organisation of your analyses is *very* important.

```
cd # make sure you are in your home directory.  
mkdir fst_analysis  
cd fst_analysis/  
mkdir output
```

Now we create some *symlinks*, these are shortcuts to the data we will be using. They are not strictly required, but make finding files much easier.

```
ln -s /GAARD/selection/samples  
ln -s /GAARD/selection/scripts  
ln -s /home/elucas/galaxy_stuff/phase2.AR1
```

Now we have created 3 symlinks to:

- *samples*. This links to txt files describing which samples in the project are in which population.
- *scripts*. This links to scripts we will use to compute Fst.
- *phase2.AR1*. This links to the ag1000G data.

NOTE: These are just links to the data. Deleting your link will not delete the data itself.

This is a table describing the files in the `samples/` directory.

File	Population
phase2.ar1.AOcol.txt	Angola <i>coluzzii</i>
phase2.ar1.CIcol.txt	Cote d'Ivoire <i>coluzzii</i>
phase2.ar1.GAgam.txt	Gabon <i>gambiae</i>
phase2.ar1.GM.txt	The Gambia
phase2.ar1.GQgam.txt	Equatorial Guinea <i>gambiae</i>
phase2.ar1.UGgam.txt	Uganda <i>gambiae</i>
phase2.ar1.BFcol.txt	Burkina Faso <i>coluzzii</i>
phase2.ar1.CMgam.txt	Cameroon <i>gambiae</i>
phase2.ar1.GHcol.txt	Ghana <i>coluzzii</i>
phase2.ar1.GNcol.txt	Guinea <i>coluzzii</i>
phase2.ar1.GW.txt	Guinea Bissau
phase2.ar1.BFgam.txt	Burkina Faso <i>gambiae</i>
phase2.ar1.FRgam.txt	French Mayotte - <i>gambiae</i>
phase2.ar1.GHgam.txt	Ghana <i>coluzzii</i>
phase2.ar1.GNgam.txt	Guinea <i>gambiae</i>
phase2.ar1.KE.txt	Kenya (kilifi)

Task: Use the `wc` command to count how many samples are in each population. The `wc` command counts the number of lines, words, and characters in a file.

Try something like:

```
wc samples/phase2.ar1.CMgam.txt
```

What does `wc -l` do?

Next we need to run the script. The `Fst` script is written in python, but we will call it using bash. Copy the template for the script, as follows.

```
cp scripts/run_fst.sh my_run_fst.sh
```

Open the script for editing using `nano`, and follow the instructions in the file.

```
nano my_run_fst.sh
```

When you have edited you file, you can run the script using:

```
bash my_run_fst.sh
```

Expect this to take several minutes.

If it finishes successfully. Look at the file you have created in `output/!`

If you get this far, using what you learned on Wednesday to download this file from the cluster for you to plot these data.

~ nick harding