# Positive selection in the chromosome 3 region 3p21.31

The 3p21.31 region on the human chromosome 3 spans about five megabases where positive selection seems to act recurrently. Previously published papers suggest that genes in the region have been under selection on multiple occasions in both African humans, in the ancestors of humans and chimpanzees, and more generally across primates. Why strong selection so often affects this region and which genes this selection affects is not not really known. With your newly aquired skills, you can apply the the most advanced population genomic methods and produce an updated inference of selection in Africans. For this project you have phased genotypes for chr3:46000000-54000000 individuals from the following populations:

```
YRI Yoruba Yoruba in Ibadan, Nigeria
LWK Luhya Luhya in Webuye, Kenya
GWD Gambian Gambian in Western Division, The Gambia
MSL Mende Mende in Sierra Leone
ESN Esan Esan in Nigeria
```

Make yourself familiar with the study populations. Where in Africa are they? How are they related?

Added all locations to favorites on google maps:

- 1. Gambian = GWD
  - a. Western Division is on the west coast of the Gambia (most west)
- 2. Mende = MSL
  - a. Sierra Leone us south of The Gambia (Second from West Coast
- 3. Yoruba = YRI
  - a. Ibadan is in South West Nigeria (3rd from the west coast)
- 4. Esan = ESN
  - a. Esan is in South Central Nigeria (East of Ibadan)
- 5. Luhya = LWK
  - a. Webuye is in Western Kenya (furthest east)

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## Investigate the following

Note: The data I am given is phased VCF files for all individuals and for each African population separately, so I do not need to do the week 3 exercise of using Beagle to to call the genotype phase.

Copy Data to folder on cluster
 #NOTE= using Cluster not Ucloud because I do not trust Ucloud.

#### Scp data

scp ~/populationgenomics/project\_data/chr3region/chr3\_460\_540\_phased.vcf.gz .

- ~/populationgenomics/project\_data/chr3region/chr3\_ESN\_460\_540\_phased.vcf.gz
- ~/populationgenomics/project\_data/chr3region/chr3\_GWD\_460\_540\_phased.vcf.gz
- ~/populationgenomics/project\_data/chr3region/chr3\_LWK\_460\_540\_phased.vcf.gz
- ~/populationgenomics/project data/chr3region/chr3 MSL 460 540 phased.vcf.gz
- ~/populationgenomics/project data/chr3region/chr3 YRI 460 540 phased.vcf.gz
- ~/populationgenomics/project\_data/chr3region/all\_inds.txt
- ~/populationgenomics/project data/chr3region/ESN inds.txt
- ~/populationgenomics/project\_data/chr3region/GWD\_inds.txt
- ~/populationgenomics/project data/chr3region/LWK inds.txt
- ~/populationgenomics/project data/chr3region/MSL inds.txt
- ~/populationgenomics/project\_data/chr3region/YRI\_inds.txt
- ~/populationgenomics/project data/chr3region/20140520.chr3.strict mask.fasta.gz
- ~/populationgenomics/project data/chr3region/human ancestor 3.fa

A. Perform an Fst scan between sets of populations in a sliding window of 100 SNP positions, comparing at least five pairs of populations. Identify the Fst outlier regions in each case.

#### Try using vcftools

ESN	Esan	Esan in Nigeria
GWD	Gambian	Gambian in Western Division, The Gambia
LWK	Luhya	Luhya in Webuye, Kenya
MSL	Mende	Mende in Sierra Leone
YRI	Yoruba	Yoruba in Ibadan, Nigeria

#### Remove non-relevant individuals from text file:

vcftools --gzvcf vcf/chr3\_460\_540\_phased.vcf.gz --keep txt/all\_inds.txt --recode --out filtered inds

#### NEW vcf file to use is called

vcf/filtered\_inds.recode.vcf.gz

## 1. ESN as compare

- a. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/ESN\_inds.txt--weir-fst-pop txt/GWD\_inds.txt --out output/ESN\_GWD
- b. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/ESN\_inds.txt --weir-fst-pop txt/LWK\_inds.txt --out output/ESN\_LWK
- c. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/ESN\_inds.txt --weir-fst-pop txt/MSL\_inds.txt --out output/ESN\_MSL
- d. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/ESN\_inds.txt --weir-fst-pop txt/YRI\_inds.txt --out output/ESN\_YRI
  - i. grep -v "\-nan" ESN\_YRI.weir.fst | awk '!/NaN/' > ESN\_YRI\_filtered1. Removes -nan values

#### 2. GWD as compare

- a. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/GWD\_inds.txt --weir-fst-pop txt/LWK inds.txt --out output/GWD LWK
- b. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/GWD\_inds.txt --weir-fst-pop txt/MSL\_inds.txt --out output/GWD\_MSL
- c. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/GWD\_inds.txt --weir-fst-pop txt/YRI\_inds.txt --out output/GWD\_YRI

#### 3. LWK as compare

- a. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/LWK\_inds.txt --weir-fst-pop txt/MSL\_inds.txt --out output/LWK\_MSL
- b. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/LWK\_inds.txt --weir-fst-pop txt/YRI\_inds.txt --out output/LWK\_YRI
- 4. MSL as compare
  - a. vcftools --gzvcf vcf/filtered\_inds.recode.vcf.gz --weir-fst-pop txt/MSL\_inds.txt --weir-fst-pop txt/YRI\_inds.txt --out output/MSL\_YRI

https://cloud.sdu.dk/app/files?path=%2F567654%2FPopgen%2FExercises%2F02\_F\_st atistics

**"F-statistics** (also known as **fixation indices**) describe the statistically expected level of heterozygosity in a population; more specifically the expected degree of (usually) a reduction in heterozygosity when compared to Hardy–Weinberg expectation."

./vcftools —vcf

B. Use Relate on all the individuals and visualize trees (using Relate or tskit) to get an impression of the relationship between the populations. How does this relate to your Fst results?

```
scp ~/populationgenomics/project_data/chr3region/20140520.chr3.strict_mask.fasta.gz .
```

- scp ~/populationgenomics/data/relate\_data/genetic\_map\_chr2\_combined\_b37.txt
- scp ~/populationgenomics/data/relate data/human ancestor 2.fa
- scp ~/populationgenomics/data/relate data/60 inds.txt
- scp ~/populationgenomics/data/relate\_data/chr2\_130\_145\_phased.vcf.gz

Remove non-relevant individuals from vcf file

vcftools --gzvcf vcf/chr3\_460\_540\_phased.vcf.gz --keep txt/all\_inds.txt --recode --out filtered\_inds

Relate does not accept the standard VCF file format, but instead uses a haps/sample format. You can read up on in the Relate documentation. The authors have been so kind as to supply a script to transform it. First, the vcf is converted to another file format (haplotype file format). If you want to know how it is structured, you can read about it here.

~/populationgenomics/software/relate/bin/RelateFileFormats --mode ConvertFromVcf --haps chr3.haps --sample chr3.sample -i filtered\_inds.recode

Then, repetitive (unreliably sequenced) regions must be masked to exclude them from our analysis. We also need to assign each variant as either ancestral or derived using the chimpanzee genome. We do both with this command:

NOTE: needed to gunzip all files first.

~/populationgenomics/software/relate/scripts/PrepareInputFiles/PrepareInputFiles.sh --haps chr3.haps --sample chr3.sample --ancestor human\_ancestor\_3.fa --mask 20140520.chr3.strict\_mask.fasta -o prep.chr3

Now, the input is fully prepared, and Relate can be run.

#### Initial

```
~/populationgenomics/software/relate/bin/Relate --mode All -m 1.25e-8 -N 30000 --haps prep.chr3.haps.gz --sample prep.chr3.sample.gz --map genetic map chr3 combined b37.txt -o chr3 relate
```

#### **ESN**

```
~/populationgenomics/software/relate/bin/Relate --mode All -m 1.25e-8 -N 30000 --haps prep.chr3_ESN.haps.gz --sample prep.chr3_ESN.sample.gz --map genetic map chr3 combined b37.txt -o chr3 relate
```

Relate outputs estimated mutation rate and coalescence times along the region

in which population size is estimated, and the population size is re-estimates branch lengths:

~/populationgenomics/software/relate/scripts/EstimatePopulationSize/EstimatePopulationSize.sh -i chr3\_relate -m 1.25e-8 --num\_iter 1 --poplabels all\_inds.txt -o popsize\_oneiter --threshold 0

~/populationgenomics/software/relate/scripts/EstimatePopulationSize/EstimatePopulationSize.sh -i chr3\_relate -m 1.25e-8 --num\_iter 2 --poplabels all\_inds.txt -o popsize twoiter --threshold 0

~/populationgenomics/software/relate/scripts/EstimatePopulationSize/EstimatePopulationSize.sh -i chr3\_relate -m 1.25e-8 --num\_iter 3 --poplabels all\_inds.txt -o popsize\_threeiter --threshold 0

When you are ready, you should have a command prompt from srun. Use that terminal to activate the pg-relate environment.

```
conda activate pg-relate
```

The Relate command below detects positive selection. At this point, we are detecting selection based on 5 distinct populations.

```
~/populationgenomics/software/relate/scripts/DetectSelection/DetectSelection.sh -i popsize_oneiter -m 1.25e-8 --poplabels all_inds.txt -o selection_relate_oneiter 
~/populationgenomics/software/relate/scripts/DetectSelection/DetectSelection.sh -i popsize -m 1.25e-8 --poplabels 60_inds.txt -o selection_relate
```

#### PLAN: USE MY ONE ITER TO ID REGION THAT I WILL BUILD TREES ON

```
relate/one_iter/selection_relate_oneiter.sele
-n 10

sort -k35r relate/one_iter/selection_relate_oneiter.sele | cut -f 1,2,35 -d ' ' | head
-n 10

Output:

pos rs_id when_mutation_has_freq2
46707244 rs17079122 -9.93441e-05
52381322 rs563971947 -7.16044e-05
52386137 rs76451298 -7.16044e-05
52210700 rs352152 -6.73561
46374436 rs35465430 -6.69821
46051363 rs58139750 -6.40215
46051532 rs55743140 -6.40215
51836739 rs113885732 -6.08449
```

Try plotting some of the sites with the following command. Use -o to determine name, and bp\_of\_interest for the position (remember to chagne both each time you run the command).

52196241 rs6786592 -5.9645

```
~/populationgenomics/software/relate/scripts/TreeView/TreeView.sh --haps relate/chr3.haps --sample relate/chr3.sample --anc relate/chr3_relate.anc --mut relate/one_iter/popsize_oneiter.mut --poplabels relate/all_inds.txt --years_per_gen 28 -o tree_52196241 --bp_of_interest 52196241

~/populationgenomics/software/relate/scripts/TreeView/TreeView.sh --haps relate/chr3.haps --sample relate/chr3.sample --anc relate/chr3_relate.anc --mut relate/one_iter/popsize_oneiter.mut --poplabels relate/all_inds.txt --years_per_gen 28 -o tree_468300001 --bp_of_interest 46830000
```

#### C. Use Relate on each population separately to infer positive selection.

~/populationgenomics/software/relate/bin/RelateFileFormats --mode ConvertFromVcf --haps chr3.haps --sample chr3.sample -i filtered\_inds.recode

#### ESN:

- ~/populationgenomics/software/relate/bin/Relate --mode All -m 1.25e-8 -N 30000 --haps prep.chr3\_ESN.haps.gz --sample prep.chr3\_ESN.sample.gz --map genetic\_map\_chr3\_combined\_b37.txt -o chr3\_relate\_ESN
- 2. ~/populationgenomics/software/relate/scripts/EstimatePopulationSize/EstimatePopulationSize.sh -i chr3\_relate\_ESN -m 1.25e-8 --poplabels all\_inds.txt -o popsize --threshold 0
- 3. ~/populationgenomics/software/relate/scripts/DetectSelection/DetectSelection.sh -i popsize ESN -m 1.25e-8 --poplabels ESN inds.txt -o selection relate ESN
- sort -k35r selection\_relate\_ESN.sele | cut -f 1,2,35 -d ' ' | head -n 10 > top\_SNPs\_Relate\_ESN

#### LWK:

- 1. ~/populationgenomics/software/relate/bin/Relate --mode All -m 1.25e-8 -N 30000 --haps prep.chr3\_LWK.haps.gz --sample prep.chr3\_LWK.sample.gz --map genetic map chr3 combined b37.txt -o chr3 relate LWK
- ~/populationgenomics/software/relate/scripts/EstimatePopulationSize/EstimatePopulationSize.sh -i chr3\_relate\_LWK -m 1.25e-8 --poplabels LWK\_inds.txt -o popsize\_LWK --threshold 0
- 3. ~/populationgenomics/software/relate/scripts/DetectSelection/DetectSelection.sh -i popsize LWK -m 1.25e-8 --poplabels LWK inds.txt -o selection\_relate\_LWK
- sort -k35r selection\_relate\_LWK.sele | cut -f 1,2,35 -d ' ' | head -n 10 > top SNPs Relate LWK

#### GWD:

- ~/populationgenomics/software/relate/bin/Relate --mode All -m 1.25e-8 -N 30000 --haps prep.chr3\_GWD.haps.gz --sample prep.chr3\_GWD.sample.gz --map genetic\_map\_chr3\_combined\_b37.txt -o chr3\_relate\_GWD
- ~/populationgenomics/software/relate/scripts/EstimatePopulationSize/EstimatePo
- 3. ~/populationgenomics/software/relate/scripts/DetectSelection/DetectSelection.sh -i popsize GWD -m 1.25e-8 --poplabels GWD inds.txt -o selection relate GWD
- sort -k35r selection\_relate\_GWD.sele | cut -f 1,2,35 -d ' ' | head -n 10 > top\_SNPs\_Relate\_GWD

#### MSL

- 1. ~/populationgenomics/software/relate/bin/Relate --mode All -m 1.25e-8 -N 30000 --haps prep.chr3\_MSL.haps.gz --sample prep.chr3\_MSL.sample.gz --map genetic map chr3 combined b37.txt -o chr3 relate MSL
- 2. ~/populationgenomics/software/relate/scripts/EstimatePopulationSize/EstimatePopulationSi

- 3. ~/populationgenomics/software/relate/scripts/DetectSelection/DetectSelection.sh -i popsize\_MSL -m 1.25e-8 --poplabels MSL\_inds.txt -o selection\_relate\_MSL
- sort -k35r selection\_relate\_MSL.sele | cut -f 1,2,35 -d ' ' | head -n 10 > top SNPs Relate MSL

#### YRI

- ~/populationgenomics/software/relate/bin/Relate --mode All -m 1.25e-8 -N 30000 --haps prep.chr3\_YRI.haps.gz --sample prep.chr3\_YRI.sample.gz --map genetic map chr3 combined b37.txt -o chr3 relate YRI
- ~/populationgenomics/software/relate/scripts/EstimatePopulationSize/EstimatePo
- 3. ~/populationgenomics/software/relate/scripts/DetectSelection/DetectSelection.sh -i popsize\_YRI -m 1.25e-8 --poplabels YRI\_inds.txt -o selection\_relate\_YRI
- sort -k35r selection\_relate\_YRI.sele | cut -f 1,2,35 -d ' ' | head -n 10 > top\_SNPs\_Relate\_YRI

https://github.com/standard-aaron/clues/wiki/Sampling-coalescence-times-with-Relate

D. Run one or more additional methods for selection inference. If possible this should be another tree sequence based method such as CLUES. Compare the results to those obtained using Relate.

#### ESN SNP -> 46659530

~/populationgenomics/software/relate/scripts/SampleBranchLengths/SampleBranchLeng ths.sh -i chr3\_relate\_ESN -o chr3\_ESN\_relate\_resample\_46659530\_100 -m 1.25e-8 --coal popsize\_ESN.coal --format b --num\_samples 100 --first-bp 46659530 --last-bp 46659530

LWK SNP -> 46659530

~/populationgenomics/software/relate/scripts/SampleBranchLengths/SampleBranchLeng ths.sh -i chr3\_relate\_LWK -o chr3\_LWK\_relate\_resample\_46659530\_100 -m 1.25e-8 --coal popsize\_LWK.coal --format b --num\_samples 100 --first-bp 46659530 --last-bp 46659530

#### GWD SNP -> 46659530

~/populationgenomics/software/relate/scripts/SampleBranchLengths/SampleBranchLeng ths.sh -i chr3\_relate\_GWD -o chr3\_GWD\_relate\_resample\_46659530\_100 -m 1.25e-8 --coal popsize\_GWD.coal --format b --num\_samples 100 --first-bp 46659530 --last-bp 46659530

#### YRI SNP -> 46659530

~/populationgenomics/software/relate/scripts/SampleBranchLengths/SampleBranchLeng ths.sh -i chr3\_relate\_YRI -o chr3\_YRI\_relate\_resample\_46659530\_100 -m 1.25e-8 --coal popsize\_YRI.coal --format b --num\_samples 100 --first-bp 46659530 --last-bp 46659530

#### MSL SNP -> 46659530

~/populationgenomics/software/relate/scripts/SampleBranchLengths/SampleBranchLeng ths.sh -i chr3\_relate\_MSL -o chr3\_MSL\_relate\_resample\_46659530\_100 -m 1.25e-8 --coal popsize\_MSL.coal --format b --num\_samples 100 --first-bp 46659530 --last-bp 46659530

## 52196241

## ESN SNP -> 52196241

 $\label{lem:continuous_software/relate/scripts/SampleBranchLengths/SampleBranchLengths.sh-i-chr3\_relate\_ESN-o-chr3\_ESN\_relate\_resample\_52196241\_100-m\\ 1.25e-8--coal\ popsize\_ESN.coal\ --format\ b\ --num\_samples\ 100\ --first-bp\ 52196241\\ --last-bp\ 52196241\\ \end{aligned}$ 

#### python inference.py --times

- ${\it $\sim$} / population genomics/students/tess div/Final Project Pop Genome/relate/ESN/chr3_ESN_relate\_resample\_52196241\_100-coal$
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/ESN/popsize\_ESN.coal --burnin 60 --out
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/CLUES/selection\_clues\_ESN\_

# 52196241\_100\_burnin\_coal

Log 10 p value = -2.72972

## LWK SNP -> 52196241

 $\label{lem:continuous} $$ \sim \operatorname{populationgenomics/software/relate/scripts/SampleBranchLengths/SampleBranchLengths.sh -i chr3_relate_LWK -o chr3_LWK_relate_resample_52196241_100 -m \\ 1.25e-8 --coal popsize_LWK.coal --format b --num_samples 100 --first-bp 52196241 \\ --last-bp 52196241 \\ \end{aligned}$ 

#### python inference.py --times

- $\label{lem:continuous} $$ \sim \text{populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/LWK/chr3\_LWK\_relate\_r esample\_52196241\_100-coal} $$$
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/LWK/popsize\_LWK.coal --burnin 60 --out
- ${\it \sim} / population genomics/students/tess div/Final Project Pop Genome/CLUES/selection\_clues\_LWK\_52196241\_100\_burnin\_coal$

## GWD SNP -> 52196241

~/populationgenomics/software/relate/scripts/SampleBranchLengths/SampleBranchLeng ths.sh -i chr3\_relate\_GWD -o chr3\_GWD\_relate\_resample\_52196241\_100 -m 1.25e-8 --coal popsize\_GWD.coal --format b --num\_samples 100 --first-bp 52196241 --last-bp 52196241

## YRI SNP -> 52196241

~/populationgenomics/software/relate/scripts/SampleBranchLengths/SampleBranchLeng ths.sh -i chr3\_relate\_YRI -o chr3\_YRI\_relate\_resample\_52196241\_100 -m 1.25e-8 --coal popsize\_YRI.coal --format b --num\_samples 100 --first-bp 52196241 --last-bp 52196241

## python inference.py --times

- ${\it ~~/} population genomics/students/tess div/Final Project Pop Genome/relate/YRI/chr3\_YRI\_relate\_resample\_52196241\_100--coal$
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/YRI/popsize\_YRI.coal --burnin 60 --out

~/populationgenomics/students/tessdiv/FinalProjectPopGenome/CLUES/selection\_clues\_YRI\_ 52196241 100 burnin coal

-1.5974

## MSL SNP -> 52196241

~/populationgenomics/software/relate/scripts/SampleBranchLengths/SampleBranchLeng ths.sh -i chr3\_relate\_MSL -o chr3\_MSL\_relate\_resample\_52196241\_100 -m 1.25e-8 --coal popsize\_MSL.coal --format b --num\_samples 100 --first-bp 52196241--last-bp 52196241

**MSL** 

python inference.py --times

- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/MSL/chr3\_MSL\_relate\_r esample 52196241 100 --coal
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/MSL/popsize\_MSL.coal --burnin 60 --out
- ${\it $^{\prime}$ population genomics/students/tess div/Final Project Pop Genome/CLUES/selection\_clues\_MSL\_52196241\_100\_burnin\_coal}$

Log 10 p- value = -2.65713

#### Run CLUES from CLUES folder

ESN

python inference.py --times

- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/ESN/chr3\_ESN\_relate\_r esample\_46659530\_100 --coal
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/ESN/popsize\_ESN.coal --burnin 60 --out
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/CLUES/selection\_clues\_ESN\_ 46659530 100 burnin coal

YRI

python inference.py --times

~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/YRI/chr3\_YRI\_relate\_res ample\_46659530\_100 --coal

- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/YRI/popsize\_YRI.coal --burnin 60 --out
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/CLUES/selection\_clues\_YRI\_4 6659530\_100\_burnin\_coal

LWK

#### python inference.py --times

- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/LWK/chr3\_LWK\_relate\_r esample 46659530 100 --coal
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/LWK/popsize\_LWK.coal --burnin 60 --out
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/CLUES/selection\_clues\_LWK\_ 46659530\_100\_burnin\_coal

#### **GWD**

python inference.py --times

- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/GWD/chr3\_GWD\_relate \_resample\_46659530\_100 --coal
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/GWD/popsize\_GWD.coa I --burnin 60 --out
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/CLUES/selection\_clues\_GWD \_46659530\_100\_burnin\_coal

#### **MSL**

python inference.py --times

- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/MSL/chr3\_MSL\_relate\_r esample 46659530 100 --coal
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/relate/MSL/popsize\_MSL.coal --burnin 60 --out
- ~/populationgenomics/students/tessdiv/FinalProjectPopGenome/CLUES/selection\_clues\_MSL\_46659530\_100\_burnin\_coal

```
python plot_traj.py selection_clues_ESN_46659530_100_burnin_coal plots/ESN_traj.png --ext png

python plot_traj.py selection_clues_GWD_46659530_100_burnin_coal plots/GWD_traj.png --ext png

python plot_traj.py selection_clues_LWK_46659530_100_burnin_coal plots/LWK_traj.png --ext png

python plot_traj.py selection_clues_MSL_46659530_100_burnin_coal plots/MSL_traj.png --ext png

python plot_traj.py selection_clues_MSL_46659530_100_burnin_coal plots/MSL_traj.png --ext png

python plot_traj.py selection_clues_YRI 46659530_100_burnin_coal plots/YRI traj.png --ext png
```

https://github.com/standard-aaron/clues/wiki

E. Identify genes potentially under selection and any known function of these genes. Consider what may drive recurrent selection in this region.

# **Papers**

Patterns of Ancestry, Signatures of Natural Selection, and Genetic Association with Stature in Western African Pygmies

Perhaps: An approximate full-likelihood method for inferring selection and allele frequency trajectories from DNA sequence data