Lab 3 Example

1 Fst outliers

Let's take a visual approach to detecting Fst outliers (non-rigorous)

- Download the repens_150.stru file.
- This contains a subset of 187 individuals and 150 snp loci
- Read repens_150.stru into R using the adegenet package and read.structure() function, save it to object called repens
 - Use str() to look at repens object
- Call the pegas library and use the code Fst(as.loci(repens)) to look at locus-by-locus Fst values, and save the output as a data.frame
- Make a histogram of the Fst values. You may want to increase the number of breaks.
- Do there appear to be any outliers? What do these mean?

2 Run Structure

- Call the strataG library and convert and save repens to a gtypes object using genind2gtypes
- Use the runStructure() function to run Structure through strataG
 - num.k.rep = 3
 - k = 2:19 (will take some time)
 - burnin = 2000, numreps = 2000
- Make plots using evanno()
- Choose a few values of K and make bar plots using structurePlot()
- How does your choice of K compare to those of Prunier et al.? To Project #2?

3 Calculate Tajima's D on sequence data

Now let's calculate Tajima's D on a small sample of *Protea repens* sequence data Each sequence contains up to 274,405 bp

- Download the repens.fasta data file and read it into R using read.fasta, saving it as the object repens.fasta, then convert and save it as a matrix
- Esimate the average per nucleotide diversity $\hat{\theta}_{\pi}$ using: pws.diff <- dist.dna(repens.fasta, model = "N", pairwise.deletion = TRUE, as.matrix = TRUE) pi <- mean(pws.diff[lower.tri(pws.diff)])
- Estimate the number of segregating sites k using: S <- ncol(variableSites(repens.fasta)\$sites)
- Now estimate $\hat{\theta}_k$ n <- nrow(repens.fasta) n.vec <- 1:(n-1) a1 <- sum(1/n.vec)
- Finally, look at the difference between $\hat{\theta}_{\pi}$ and $\hat{\theta}_{k}$ pi (S/a1)
- What do you get? What a crazy number!
- There are other pieces going on here that need to be corrected for. Use the function tajimasD() in strataG to calculate Tajima's D on repens.fasta. Look at the tajimasD() source code for details!
- What value do you get? Is it significant? What does this value imply about the evolutionary history of these samples?