

ANGSD-wrapper: utilities for analyzing next generation sequencing data

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Abstract

High throughput sequencing has changed many aspects of population genetics, molecular ecology, and related fields, affecting both experimental design and data analysis. The software package ANGSD allows users to perform a number of population genetic analyses on high-throughput sequencing data. The package is specifically designed to produce more accurate results for samples with low sequencing depth and makes use of full genome data while handling a wide array of sampling and experimental designs. Here we present ANGSD-wrapper, a user-friendly interface for running ANGSD and visualizing results. ANGSD-wrapper includes a number of 'wrapper' scripts that facilitate configuration and execution of multi-step analyses. ANGSD-wrapper also provides interactive graphing of ANGSD results to enhance data exploration. We demonstrate the usefulness of ANGSD-wrapper by analyzing resequencing data from populations of wild and domesticated *Zea*. ANGSD-wrapper is freely available from <https://github.com/mojaveazure/angsd-wrapper>.

Introduction

High throughput sequencing has revolutionized evolutionary genetics, allowing researchers to quickly assay large numbers of individuals or survey fine-scale patterns of variation across the genome. Application of these approaches has led to changes in both experimental design and data analysis [1]. Many popular software packages used by researchers for analysis of comparative resequencing data [see 2] were not designed to handle these novel data types or efficiently analyze the large volumes of data now being generated. In particular, short read sequencing has brought new challenges, including highly variable coverage, missing data, and high per-nucleotide error rates.

A number of tools have recently been published to estimate population genetic descriptive statistics using high throughput sequencing data [3, 4, 5, 6], but the majority of these either make limiting assumptions about the data (e.g., all sites have been sequenced, all genomes are haploid, sequencing is to sufficient depth, all individuals are outcrossing) or are specialized tools offering a narrow set of analysis options. Korneliussen et al. [7] recently published the software package ANGSD, which enables users to flexibly perform a large number of common population genetic analyses, including estimation of diversity statistics, admixture analysis including Patterson's D statistic [8], site frequency spectrum estimation [9], and calculation of neutrality test statistics [10]. One of the most important features of ANGSD is that most analyses are performed directly on genotype likelihoods, freeing users from the requirement of calling variants or genotypes and permitting analysis of low-coverage data or sequences with large amounts of missing data.

Here we present ANGSD-wrapper, a user-friendly interface to ANGSD. ANGSD-wrapper takes the form of a set of configuration files and 'wrapper' scripts (Figure S1) that streamline the execution of

Method	Calculations	Interactive Graphing
SFS	Site frequency spectrum	Yes
2DSFS	Joint site frequency spectrum, F_{ST}	Yes
ABBA/BABA	Patterson’s D statistic	Yes
Ancestral	Extract ancestral sequence from BAM file	No
Genotypes	Genotype likelihood estimations	No
PCA	Principal component analysis	Yes
Thetas	Diversity statistics (θ_w , θ_π , Fu and Li’s θ , Fay’s θ) and neutrality tests (Tajima’s D, Fu and Li’s D, Fu and Li’s F, Fay and Wu’s H, Zeng’s E)	Yes
Inbreeding	Calculate per-individual inbreeding coefficients with ngsF	No
Admixture	Perform admixture analysis	Yes

Table 1: Table of methods implemented in ANGSD-wrapper

multi-step pipelines required for data analysis in ANGSD. ANGSD-wrapper also eases the configuration of ANGSD-related programs such as ngsPopGen [11], ngsF[12], and ngsAdmix [13]. Because the large volume of data associated with high throughput sequence analysis is often difficult to visualize, ANGSD-wrapper also provides a suite of interactive visualization tools to plot results and explore patterns at multiple scales. We demonstrate some of the analyses possible using ANGSD-wrapper when applied to low-coverage whole-genome data from domesticated maize and two related wild teosinte subspecies. ANGSD-wrapper is freely available from <https://github.com/mojaveazure/angsd-wrapper>.

Methods

ANGSD-wrapper is a set of configuration files and scripts written primarily in the Bash UNIX shell. The scripts can be run either on a standalone computer with a UNIX terminal, or on computing clusters where they can be submitted to a queuing system such as SGE [14], Slurm [15] or TORQUE [16]. An installation of the statistical software R [17] is required to make use of the visualization tools incorporated in ANGSD-wrapper. The visualization portion of ANGSD-wrapper also requires installation of the R packages shiny [18], genomeIntervals [19], and ape [20].

ANGSD-wrapper is divided into scripts associated with analytical approaches implemented in ANGSD and associated software. ANGSD-wrapper provides a common configuration file, “common.conf,” which holds variables that are likely to remain constant across analyses, including identifiers for chromosomal regions and the paths to project directories. In ANGSD-wrapper, each method is self-contained in a shell script which uses information from the common configuration file and a method-specific configuration file. Each analysis is run using a simple command:

```
$ angsd-wrapper <method> <configuration_file>
```

Analyses supported by ANGSD-wrapper are shown in Table 1. A detailed flowchart of each of these workflows is shown in Figure S1, and additional details, documentation, a tutorial, and a wiki can be found on the GitHub page: <https://github.com/mojaveazure/angsd-wrapper/wiki>.

The visualization software included with ANGSD-wrapper is contained within it’s own directory called “shinyGraphing.” This application must be started in R and can be accessed locally from a web browser. This software provides a graphical user interface (GUI) to quickly and interactively plot results obtained from ANGSD-wrapper. Each tab in the GUI contains plots for different ANGSD methods.

In order to use the plotting software, the user navigates to the desired tab and uploads the appropriate results file. The Shiny server automatically parses ANGSD output files and creates the resulting plot(s) (Figure 1), which can be saved using the browser’s built in image saving capabilities.

ANGSD-wrapper graph

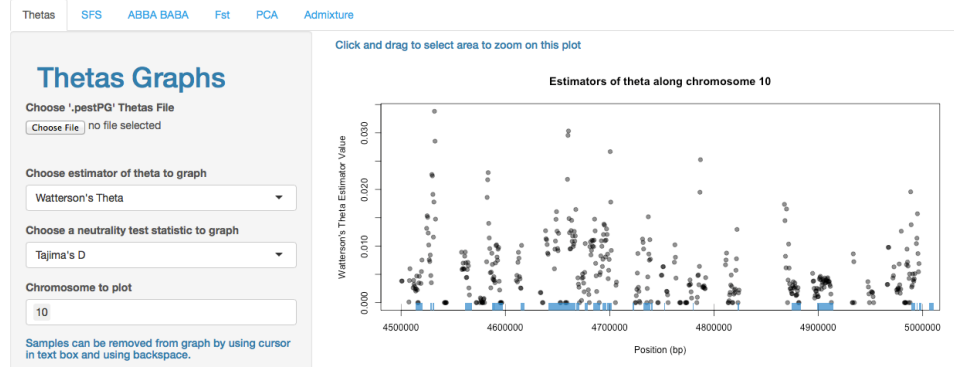


Figure 1: Visualization of Watterson's θ estimated by ANGSD across a 1.5Mb region of chromosome 10 in *Zea mays* ssp. *mays* using ANGSD-wrapper. Blue boxes indicate genic regions provided by a GFF annotation.

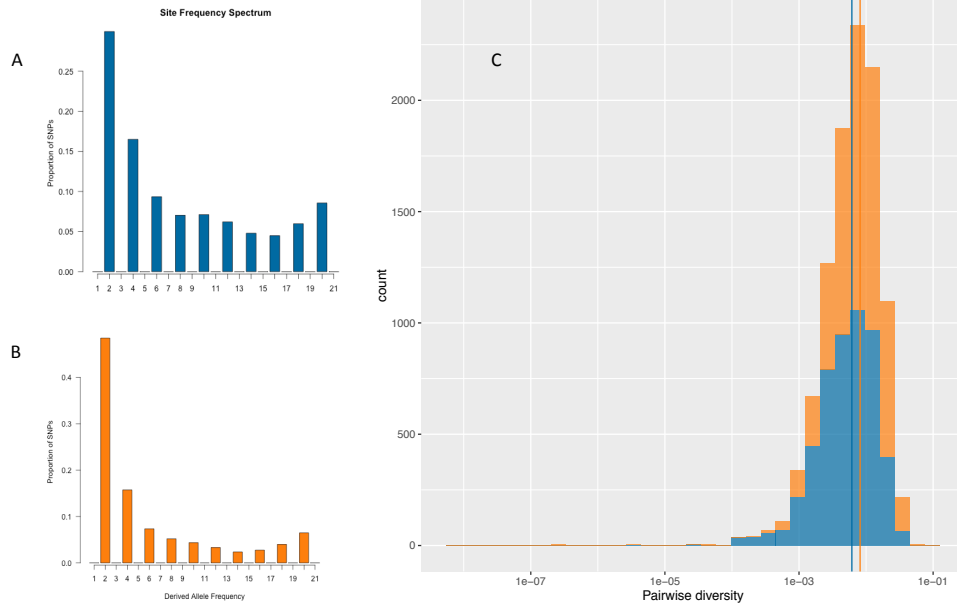


Figure 2: Summary statistics for *Zea mays*. Site frequency spectra for A. maize and B. teosinte. C. distribution of pairwise differences for maize (blue) and teosinte (orange). Pairwise diversity results are visualized separately from the interactive graphics and colors were added to A and B using a custom script.

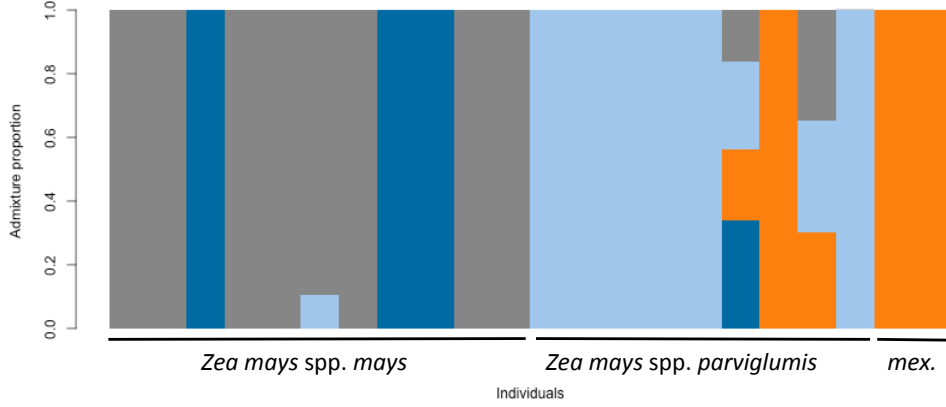


Figure 3: Admixture analysis for *Zea mays* spp. *mays*, *Zea mays* spp. *parviglumis*, and *Zea mays* spp. *mexicana* (*mex*) with K=4 source populations. Individuals are shown in the same order as Table S1.

As a demonstration of analyses in ANGSD-wrapper, we explore patterns of diversity in a single genomic region of domesticated maize and wild teosinte. We used a subset of the resequenced samples from the Maize HapMap2 project (Table S1) and calculated summary statistics using a 10Mb region on chromosome 10 [21]. The data are available at https://figshare.com/articles/Example_Data_tar_bz2/2063442. In the following we refer to methods listed in Table 1 when describing analyses.

We first use the “SFS” method to estimate the site frequency spectrum (SFS) of both maize and its wild progenitor *Zea mays* spp. *parviglumis*, assuming an inbreeding coefficient of $F = 1$ for these highly inbred samples. The SFS and diversity statistics were calculated using ancestral states inferred in the “Ancestral Sequence” method from a single resequenced genome of *Tripsacum dactyloides*. We show that the maize SFS is skewed towards more intermediate-frequency variants (Figure 2A-B, Tajima’s D of 0.2 and 0.0085 in maize and teosinte, respectively), likely a result of the bottleneck associated with maize domestication [22] [23]. Using the “Thetas” method we find further evidence of the domestication bottleneck, with mean levels of nucleotide diversity in this region in maize $\approx 25\%$ lower than in teosinte (0.0061 and 0.0082, respectively; Figure 2C). Using the “2DSFS method,” which includes an F_{ST} calculation, we find a mean F_{ST} in this region of 0.116, nearly identical to the genome-wide value of 0.11 reported in [24]. There are no genes in this region that have been identified as potential domestication candidates, consistent with the lack of extended regions of high F_{ST} in our analysis (Figure S3).

Finally, we include two samples of the related wild teosinte *Zea mays* spp. *mexicana* to assess evidence for admixture. Using the “Admixture” method, which implements an estimate of admixture proportion from genotype likelihoods [13], we identify structure within domesticated maize separating three high-latitude temperate landraces from the other tropical accessions (Figure 3). We find no evidence of admixture between these lowland maize samples and spp. *mexicana*, consistent with an independent analysis from SNP genotyping [25]. *Zea mays* spp. *mexicana* clusters into its own group, along with a single accession of spp. *parviglumis* collected from a region in which many teosinte populations appear to be the result of admixture between the two subspecies [26]. A single spp. *parviglumis* accession from the Northern extent of the range does not appear to be well-classified with these data, likely due to our relatively limited geographic and genomic sampling.

Conclusions

ANGSD-wrapper provides an easy-to-use interface that simplifies many population genetic analyses implemented in ANGSD [7] and permits the exploration of genome-scale results through interactive visualization. ANGSD-wrapper is under active development to incorporate updates to the ANGSD software package.

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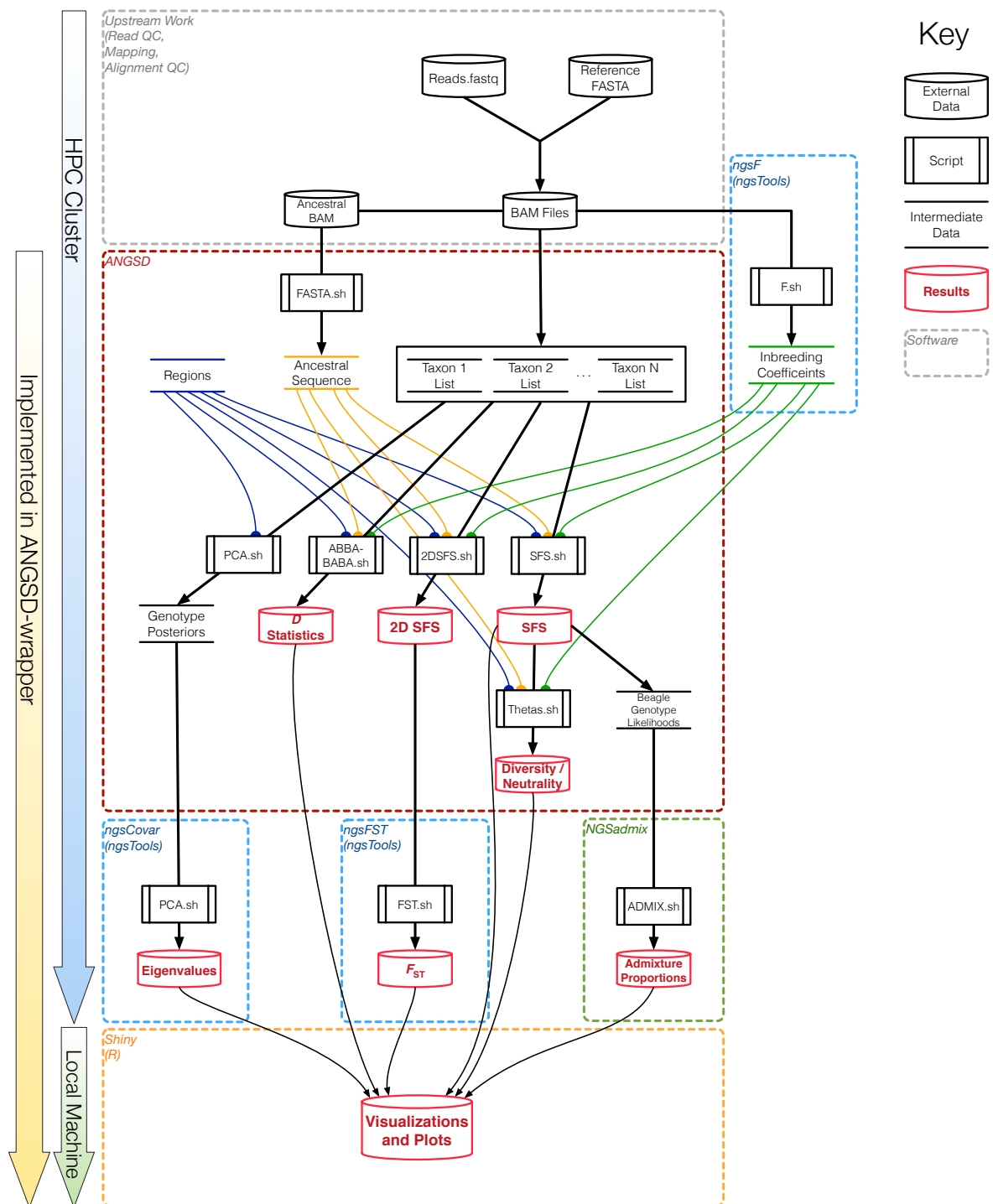


Figure S1: Workflow diagram for all methods available in ANGSD-wrapper.

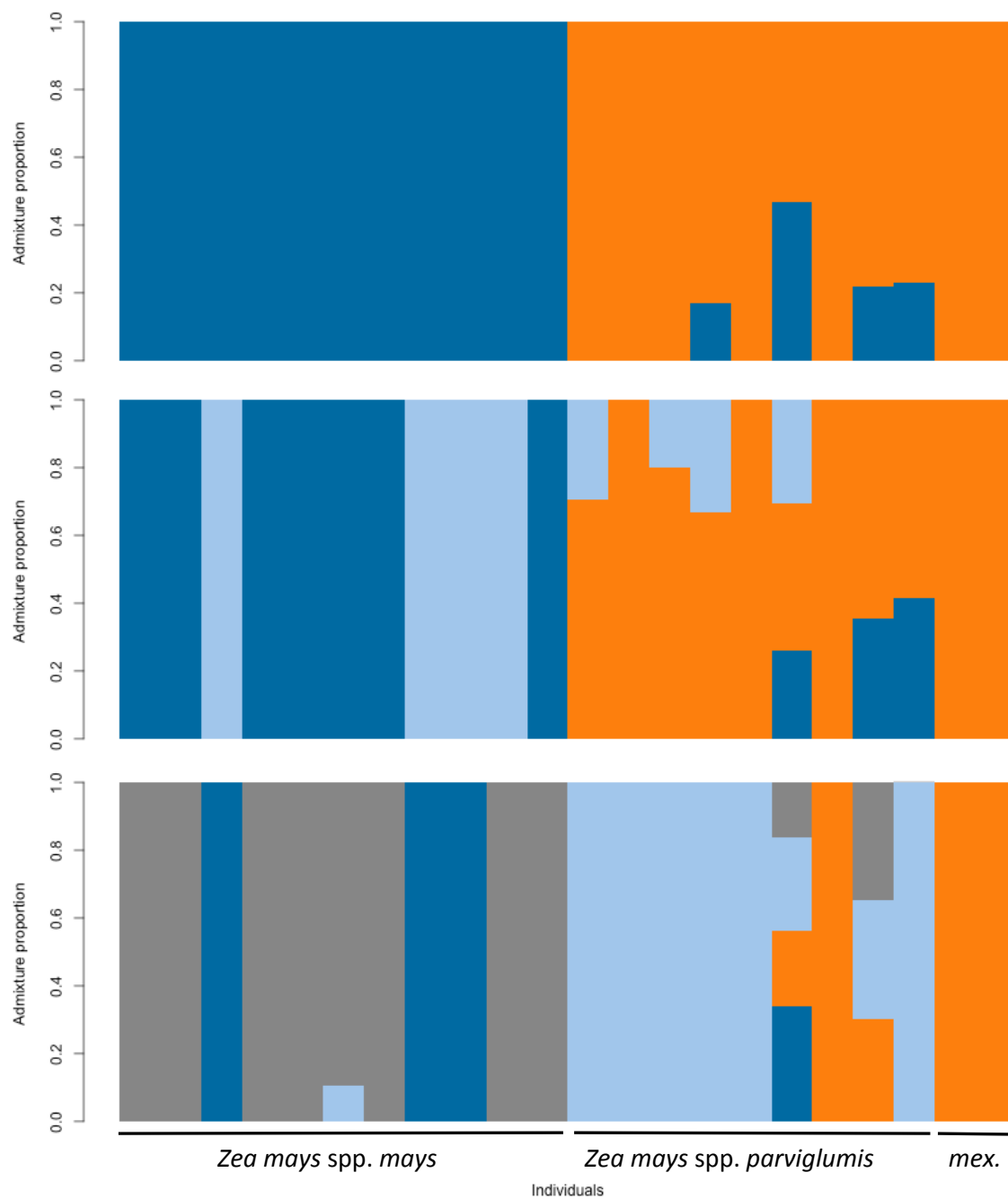


Figure S2: Admixture analysis for K=2 (top), K=3 (middle), and K=4 (bottom).

Sample	Mean Depth	Species
BKN009	7.00194	<i>Zea mays</i> spp. <i>mays</i>
BKN011	6.9777	<i>Zea mays</i> spp. <i>mays</i>
BKN014	6.78829	<i>Zea mays</i> spp. <i>mays</i>
BKN015	3.739	<i>Zea mays</i> spp. <i>mays</i>
BKN019	3.71944	<i>Zea mays</i> spp. <i>mays</i>
BKN022	3.71268	<i>Zea mays</i> spp. <i>mays</i>
BKN025	3.53799	<i>Zea mays</i> spp. <i>mays</i>
BKN026	3.91798	<i>Zea mays</i> spp. <i>mays</i>
BKN027	7.05576	<i>Zea mays</i> spp. <i>mays</i>
BKN033	3.85336	<i>Zea mays</i> spp. <i>mays</i>
BKN035	3.74839	<i>Zea mays</i> spp. <i>mays</i>
TIL01	3.60133	<i>Zea mays</i> spp. <i>parviglumis</i>
TIL03	4.07518	<i>Zea mays</i> spp. <i>parviglumis</i>
TIL04	6.09163	<i>Zea mays</i> spp. <i>parviglumis</i>
TIL07	5.11419	<i>Zea mays</i> spp. <i>parviglumis</i>
TIL09	5.29004	<i>Zea mays</i> spp. <i>parviglumis</i>
TIL11	3.15477	<i>Zea mays</i> spp. <i>parviglumis</i>
TIL15	6.87873	<i>Zea mays</i> spp. <i>parviglumis</i>
TIL16	2.67186	<i>Zea mays</i> spp. <i>parviglumis</i>
TIL17	2.61892	<i>Zea mays</i> spp. <i>parviglumis</i>
TIL08	6.09453	<i>Zea mays</i> ssp. <i>mexicana</i>
TIL25	13.1566	<i>Zea mays</i> ssp. <i>mexicana</i>
TDD39103	8.62	<i>Tripsacum dactyloides</i>

Table S1: Table of samples used in analysis with mean depth over the region 15000000-25000000 on chromosome 10.

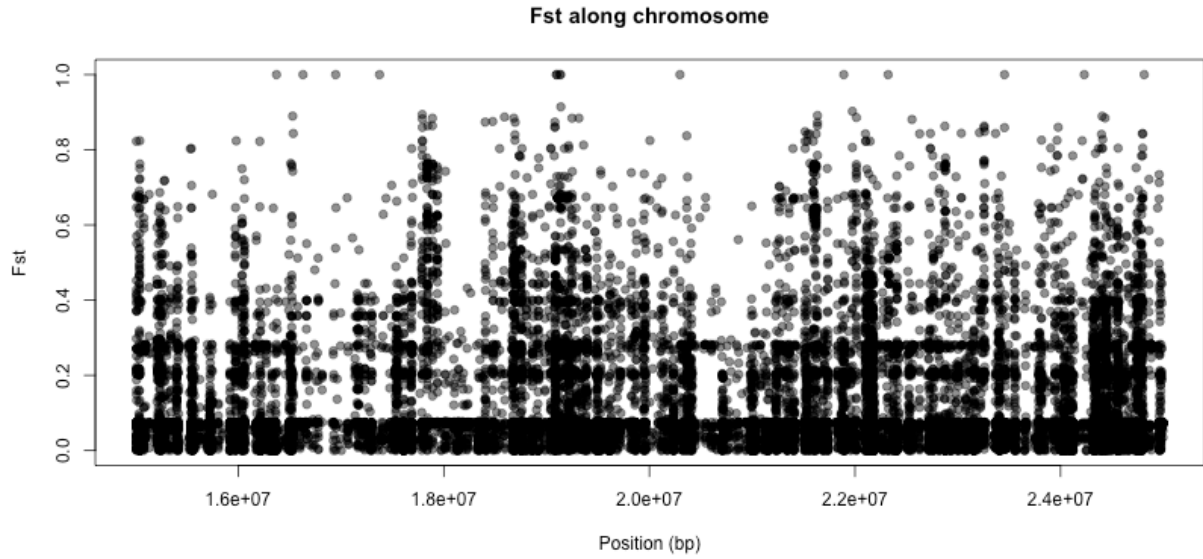


Figure S3: F_{ST} values plotted against base pair position on chromosome 10 of maize. F_{ST} is calculated between the maize and teosinte samples.