

Sambar workflow

Load SambaR into R

```
source('C:/path/to/dir/SAMBAR.txt')
```

SambaR can be downloaded from Github:
<https://github.com/mennodejong1986/SambaR>

Install and load dependencies

```
getpackages(mylib=NULL)
```

Import data (binary PED/MAP) into R

```
setwd('C:/path/to/workdir/')
```

```
importdata(inputprefix='prefix',samplefile=NULL,  
sumstatsfile=NULL,depthfile=FALSE)
```

Or instead convert from an existing genlight object:

```
genlight2sambar(genlight_object=NULL)
```

Filter data

```
filterdata(indmiss=0.25,snpmiss=0.1,min_mac=2  
,doefilter=TRUE,snpdepthfilter=TRUE,  
min_spacing=500,nchrows=NULL,silent=TRUE)
```

Included: quality control, F, HWE, relatedness

Analyze data

population structure

```
findstructure(Kmax=6,add_legend=TRUE,  
legend_pos='bottomright',legend_cex=3,  
symbol_size=3)
```

Included: PCoA, PCA, DAPC, CA, MDS,
 admixture analyses (LEA), Tess3r

population differentiation

```
calcdistance(nchrows=NULL)
```

Included: Nei's D, (sliding window) Wright's Fst,
 Nei's Fst, Weir & Cockerham Fst, ABBA-BABA

genetic diversity

```
calcdiversity(nrsites=NULL,legend_cex=2.5)
```

Included: SFS, heterozygosity, nucleotide
 diversity, Watterson's theta, Tajima's D, private
 alleles, proportion segregating sites

selection analyses

```
selectionanalyses(do_pcadapt=TRUE,do_outflank  
=TRUE,do_fsthet=TRUE,export='pdf')
```

Included: GWDS, FstHet, OutFLANK, PCadapt

Sambar output

mygenlight

	snp1	snp2	snp3	...
ind1				
Ind2				
Ind3				
...				

inds

	name	pop	nr	filter	...
	Ind1			FALSE	
	Ind2			TRUE	
	Ind3			TRUE	
	...				

snps

	snp1	snp2	snp3	...
name				
chr				
pos				
miss				
filter	TRUE	TRUE	FALSE	
maf				
...				

output folders

- Demography
- Divergence
- Diversity
- Inputfiles
- QC
- Selection
- Structure

