

# Package ‘evobiR’

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**Type** Package

**Title** evobiR: evolutionary biology in R.

**Version** 1.0

**Date** 2013-10-08

**Author** Heath Blackmon

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**URL** <http://www.uta.edu/karyodb/evobiR/>

**Depends** seqinr, ape, stringr, geiger, taxize

**Description** evobiR is a collection of tools for use in evolutionary biology

**License** GPL (>= 2)

**NeedsCompilation** no

**Repository** CRAN

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evobiR-package

*evobiR: Evolutionary Biology in R*


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## Description

**evobiR** is a collection of tools for use in evolutionary biology. Some of the functions manipulate data in a way not implemented by other functions while others calculate sequence statistics or perform simulations, either of data across trees or genetic and genomic simulations.

## Details

Package: evobiR  
Type: Package  
Version: 1.0  
Date: 2013-10-08  
License: GPL (>=2)

More information on **evobiR** is available at <http://www.uta.edu/karyodb/evobiR/>

A full list of functions in the **evobiR** package is accessible by typing `library(help = evobiR)`

## Author(s)

Heath Blackmon

Maintainer: Heath Blackmon <coleoguy@gmail.com>

---

1.fasta                      *simulated SNP data*

---

**Description**

This file just contains some simulated SNP data

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

---

2.fasta                      *simulated SNP data*

---

**Description**

This file just contains some simulated SNP data

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

---

3.fasta                      *simulated SNP data*

---

**Description**

This file just contains some simulated SNP data

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

AnalyzeAssembly

*Analyze Genome Assembly*

---

**Description**

This function reports basic statistics for a genome assembly.

**Usage**

```
AnalyzeAssembly(genome, max_N = 25, plot = F)
```

**Arguments**

genome	a list of vectors with each element being a single string of the class "SeqFas-tadna".
max_N	Maximum number of consecutive N symbols. Scaffolds will be broken into contigs when this number is exceeded.
plot	When True an accumulation plot will be returned as well as the statistics

**Details**

If a standard FASTA file is read in with the function `read.fasta` from the package `seqinr` the argument `as.string` should set to `TRUE`. The genome should also be all lower case which is the default setting for `read.fasta`.

**Value**

A dataframe with the following rows:

- Number of Scaffolds
- Assembly Size Based on Scaffolds
- Number of Scaffolds over 1MB
- N50 Scaffold Size
- Number of Contigs
- Assembly Size Based on Contigs
- N50 Contig Size
- Minimum Contig Size
- Percent GC

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

**Examples**

```
## just a small simulated genome
data(genome)
## calculate summary statistics for the genome
AnalyzeAssembly(genome = genome, max_N = 25, plot = TRUE)
```

---

assembly.fasta	<i>a sample of a genome</i>
----------------	-----------------------------

---

**Description**

This is a small portion of Tribolium genome

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

---

CalcD	<i>Calculate Patterson's D-statistic</i>
-------	--

---

**Description**

These functions calculate Patterson's D-statistic to compare the frequencies of discordant SNP genealogies. These tests assume equal substitution rates and unlinked loci, D-statistics significantly different from 0 suggest that introgression has occurred.

**Usage**

```
CalcD(alignment = "alignment.fasta", boot = F, replicate = 1000)
```

```
CalcPartD(alignment = "alignment.fasta", boot = F, replicate = 1000, alpha = 0.05)
```

```
CalcPopD(alignment = "alignment.fasta")
```

**Arguments**

alignment	This is an alignment in fasta format
boot	This indicates whether or not bootstrapping should be performed to estimate variance
replicate	Number of replicates to be used in estimating variance
alpha	The desired alpha level, used in calculating significance of tests in CalcPartD after Bonferoni correction

**Details**

The functions CalcD and CalcPopD are implementations of the algorithm described in Durand et al. 2011. The function CalcPartD is an implementation of the extension that was reported by Eaton and Ree 2013 which extended the original ABBA-BABA test to include additional taxa providing the ability to identify the source population of introgression events when more than one extant lineage may have hybridized with the ingroup.

**Value**

Returns the number of each type of site, Z scores and p-values

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

Durand, Eric Y., et al. Testing for ancient admixture between closely related populations. *Molecular biology and evolution* 28.8 (2011): 2239-2252.

Eaton, D. A. R., and R. H. Ree. 2013. Inferring phylogeny and introgression using RADseq data: An example from flowering plants (Pedicularis: Orobanchaceae). *Syst. Biol.* 62:689-706

**Examples**

```
CalcD(alignment = system.file("1.fasta", package = "evobiR"), boot = TRUE, replicate=10)
```

```
CalcPartD(alignment = system.file("2.fasta", package = "evobiR"), boot = TRUE, 10)
```

```
CalcPopD(alignment = system.file("3.fasta", package = "evobiR"))
```

---

CoalSim

*Coalescence Simulations*

---

**Description**

This function simulates the coalescence of lineages through time. At the beginning of the simulation a number of unique lineages specified by the census size as we go back in time these coalesce until we have only a single lineage remaining.

**Usage**

```
CoalSim(census, lw, ln.col)
```

**Arguments**

- census            the size of population to use in the simulation.
- lw                line width factor passed to the plotting function.
- ln.col            specifies the color to be used in plotting the coalescence history

**Value**

Returns a plot showing the result of the coalescence simulation.

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

**Examples**

```
CoalSim(census = 15, lw = 2, ln.col = 'blue')
```

---

*data.taxonomy*            *taxonomic table*

---

**Description**

This file contains taxonomic information

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

---

`DriftSim`*Simulate Allele Frequencies*

---

**Description**

These functions simulate allele frequencies through time

**Usage**

```
SelSim(census, initial.freq, selection.coef, iter, generations)
```

```
DriftSim(census, initial.freq, iter, generations)
```

**Arguments**

<code>census</code>	The population size to be used for the simulation.
<code>initial.freq</code>	The proportion of individuals that should have the selected mutation at the beginning of the simulation.
<code>selection.coef</code>	The selection coefficient. 1 is a neutral mutation while values greater or less than 1 will be beneficial or deleterious respectively.
<code>iter</code>	The number of times that the simulation should be repeated
<code>generations</code>	How many generations each simulation should last.

**Value**

These functions were created to help in teaching population genetics. Each of these functions plot the frequency of a new mutation at each generation. Each iteration of the simulation is plotted in a new color and the functions also reports the number of iterations that resulted in fixation, loss or intermediate frequencies for the mutation. The function `DriftSim` is just a special case of `SelSim` where the selection coefficient is equal to 1.

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

**Examples**

```
set.seed(5)
SelSim(census = 200, initial.freq = .5, selection.coef = 1.01, iter = 10, generations = 200)

set.seed(5)
DriftSim(census = 200, initial.freq = .5, iter = 25, generations = 200)
```



---

**FuzzyMatch***Find Close Matches in a tree and dataset*

---

**Description**

When assembling data from different sources typos can sometimes cause a loss of perfect matches between trees and datasets. This function helps you find these close matches that can be hand curated to keep as many species as possible in your analysis.

**Usage**

```
FuzzyMatch(tree, data, max.dist)
```

**Arguments**

tree	a phylogenetic tree of the class "phylo".
data	character vector with the names from your dataset.
max.dist	This is the maximum number of characters that can differ between your tree and data and still be recognized as a close match.

**Value**

A dataframe with the following rows:

Name in data
Name in tree
Number of differences

**Author(s)**

Heath Blackmon and Matt Pennell

**References**

<http://www.uta.edu/karyodb/evobiR/>

**Examples**

```
data(hym.tree)
names <- c("Pepsis_elegans", "Plagiolepis_alluaudi", "Pheidele_lucreti",
          "Meliturgula_scriptifronsi", "Andrena_afimbriat")
FuzzyMatch(tree = hym.tree, data = names, max.dist=3)
```

---

 genome

*A small portion of the genome of Tribolium confusum*


---

**Description**

This is just little piece of the Tribolium confusum genome

**Format**

a list of vectors with each element being a single string of the class "SeqFastadna".

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

---

 GenomeSym

*Genome Simulation*


---

**Description**

THIS FUNCTION IS STILL IN THE DEVELOPMENT PHASE AND RESULTS FROM IT SHOULD NOT BE TRUSTED WITHOUT CONTACTING ME FIRST!!!

This function is designed to simulate the evolution of a diploid organism with chromosomal sex determination. It allows for the evolution of sexual antagonism, recombination suppression, aneuploidy, haplosufficiency.

**Usage**

```
GenomeSym(model = "fragileY", generations = 100, census = 50, gametes = 4,
  loci.a = 100, loci.s = 50, prob.point.mut = .0010,
  DFE = (c(rnorm(1000, mean = 0.9, sd = 0.06), rgamma(1000, rate = 2,
  shape = 0.4))), sex.bias.mut = .85, haplosuff.mut = 1, crossovers = 2,
  recom.mu = 10, aneuploi.mut = c(0, .0002), distance.mut = c(0, .000002),
  achiasmat.mut = c(0, .00002), sex.ant = c(.5, .3, .2),
  fragile.fact = 20, reporting = "None")
```

**Arguments**

model	the type of simulation to be run
generations	integer indicating the number of generations that the simulation should be run
census	integer indicating total census size
gametes	integer indicating number of gametes drawn from each individual
loci.a	integer indicating loci on autosomes
loci.s	integer indicating loci on sex chromosomes
prob.point.mut	probability that a new allele will arise
DFE	a vector of fitness effect for mutations
sex.bias.mut	proportion of mutations to arise in males
haplosuff.mut	a value of less than 1 from which mutations in haplosufficiency will be drawn from a uniform distribution between 0 and the specified value.
crossovers	the average number of crossovers per chromosome per meiosis in autosomes
recom.mu	maximum number of loci N that can quit recombining in a single mutational step
aneuploi.mut	probability of a Y aneuploidy mutation drawn from a uniform dist between X and Y
distance.mut	prob of evolving dist pairing uniform dist from X to Y
achiasm.mut	prob of evolving achiasmatic meiosis in males uniform from X to Y
sex.ant	numerical vector must sum to 1 indicating the probability of unbiased, male biased, female biased mutations
fragile.fact	this is the factor by which we increase aneuploidy based on par reduction
reporting	a character string indicating the desired type of insimulation reporting: 'PAR', 'XY.fitness', 'all.loci'.

**Details**

This function simulates a genome as a series of numerical vectors each chromosome in a genome has vectors to represent those genes fitness when in a female or when in a male additionally if they are sex chromosomes they have vectors describing the recombination characteristics and haplosufficiency data. All of these genome characteristics are allowed to evolve during the simulation through a variety of mutational mechanisms. The genomes perform in silico gametogenesis with recombination. The probability of gametes being used to create the next generation is in proportion to their parents relative fitness.

default values for the function are as follows:

```

model = "fragileY"
generations = 100
census = 50
gametes = 4
loci.a = 100
loci.s = 50

```

```

prob.point.mut = .3
DFE = (c(rnorm(1000,mean=.9,sd=.06),rgamma(1000, rate=2, shape=.4)))
sex.bias.mut = .85
haplosuff.mut = 1
crossovers = 2
recom.mu = 10
anneuploi.mut = c(0, .0002)
distance.mut = c(0, .000002)
achiasmat.mut = c(0, .00002)
sex.ant = c(.5, .3, .2)
fragile.fact = 20
reporting = "None"

```

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

**Examples**

```

## just a small simulated genome
Genome <- GenomeSym(census = 20, generations = 10, reporting = 'all.loci')

```

---

GetTaxonomy

*Create a taxonomy table*

---

**Description**

This function is an extension of the `tax_name` function from the package `taxize`. The input is a tree and the function will return a table with a description of the taxonomy of all species on the tree

**Usage**

```
GetTaxonomy(tree, database = "ncbi")
```

**Arguments**

<code>tree</code>	an object of class "phylo"
<code>database</code>	character string "ncbi" or "itis" indicating the database that should be used

**Value**

this function saves a csv file named: tree.taxonomy.csv

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

**Examples**

```
# data(hym.tree)
# GetTaxonomy(tree = hym.tree, database = 'ncbi')
```

---

HetLevels

*Calculates heterozygosity based on ambiguous IUPAC nucleotide codes*

---

**Description**

Some assemblers like AllPaths will use IUPAC codes to indicate polymorphic sites in an assembly. This function uses these to estimate the level of heterozygosity along a sequence.

**Usage**

```
HetLevels(fasta)
```

**Arguments**

fasta            A fasta formatted file that may contain 1 or more sequences.

**Details**

Ambiguity codes of "r", "y", "s", "w", "k", "m", "b", "d", "h", "v" are counted as polymorphic sites with an "n" are not counted since they are often indicative of scaffolding not polymorphism.

**Value**

A matrix is returned listing the length of each sequence the number of polymorphic sites and the number of "n" sites.

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

**Examples**

```
HetLevels(fasta = system.file("assembly.fasta", package = "evobiR"))
```

---

**HighLevelTree***Prune a phylogeny to a higher taxonomic levels.*

---

**Description**

This function takes a species level tree and prunes it so only a single representative of each clade, e.g. genus or family, is present.

**Usage**

```
HighLevelTree(taxa.table, tree, cur.tips, new.tips)
```

**Arguments**

<code>taxa.table</code>	a character matrix with columns containing the current tips of the tree and additional columns containing their assigned genera, family, tribe, etc.
<code>tree</code>	the current species level tree
<code>cur.tips</code>	a number indicating which column contains the current tip names
<code>new.tips</code>	a number indicating which column contains the names for the taxonomic level that you would like to prune the tree to.

**Value**

returns an object of the class "phylo"

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

**See Also**

[GetTaxonomy](#)

**Examples**

```
data(species.tree)
data(taxa.table)
```

```
family.tree <- HighLevelTree(taxa.table, species.tree, 1, 2)
genus.tree <- HighLevelTree(taxa.table, species.tree, 1, 3)
```

---

horn.beetle.csv	<i>Gnatocerus measurements</i>
-----------------	--------------------------------

---

**Description**

A csv file containing measurements of horn and body size for the beetle *Gnatocerus cornutus*.

---

hym.tree	<i>Phylogenetic tree</i>
----------	--------------------------

---

**Description**

This is a phylogenetic tree with 5 species of hymenoptera.

---

mcmc2	<i>mcmc log file</i>
-------	----------------------

---

**Description**

A 5 column by 100 row double matrix from a mcmc log file. The first column is the tree used during the iteration the remaining columns are the rate parameters of the Q matrix listed by column order.

---

mcmc3	<i>mcmc log file</i>
-------	----------------------

---

**Description**

A 10 column by 100 row double matrix from a mcmc log file. The first column is the tree used during the iteration the remaining columns are the rate parameters of the Q matrix listed by column order.

---

mite.data	<i>cytogenetic mite data</i>
-----------	------------------------------

---

**Description**

This file contains cytogenetic mite data available at the tree of sex database

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

---

mite.tree	<i>phylogenetic tree of mites</i>
-----------	-----------------------------------

---

**Description**

This file contains a phylogenetic tree of mites generated from phylota data by Nate Hardy

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

---

PPSDiscrete	<i>Create Simulated Datasets via PPS</i>
-------------	--

---

**Description**

This function performs posterior predictive simulations of discrete traits. The function is written to work with the output of bayesian programs that produce a collection of rate matrix parameter estimates based on either one or a collection of trees.

**Usage**

```
PPSDiscrete(trees, MCMC, states, N = 2)
```

**Arguments**

trees	an object of class "multiPhylo" or "phylo" containing the trees used in generating the rate estimates
MCMC	MCMC states bring this will normally be a log file that is brought into R with read.csv the columns for a three state character should be: tree, qAA, qBA, qCA, qAB, qBB, qCB, qAC, qBC, qCC. If your analysis involves only a single tree then the tree column should be excluded.
states	a vector of root probabilities
N	the number of PPS datasets desired

**Value**

A matrix is returned with the rownames being the species names from the tree and each column containing a result of a single PPS.



**Author(s)**

Heath Blackmon

**References**<http://www.uta.edu/karyodb/evobiR/>**Examples**

```

data(trees)
data(mcmc2)
data(mcmc3)
# 1 tree 100 q-mats 3 states
PPSDiscrete(trees[[1]], MCMC=mcmc3[,2:10], states=c(.5,.2,.3), N=2)
# 10 trees 100 q-mats 3 states
PPSDiscrete(trees, MCMC=mcmc3, states=c(.5,.2,.3), N=10)
# 10 trees 100 q-mats 2 states
PPSDiscrete(trees, MCMC=mcmc2, states=c(.5,.5), N=10)

```

---

ReplTable
*Maximize Retained Tips*


---

**Description**

These functions attempt to retain as many datapoints as possible during a comparative analysis.

**Usage**

```
ReplTable(tree, data.taxonomy, tree.taxonomy, levels=c("Genus", "Family"), verbose=TRUE)
```

```
MaxTips(tree, data, repl.table)
```

**Arguments**

tree	a phylogenetic tree of the class "phylo".
data.taxonomy	a dataframe containing taxonomic information for all species in the dataset see function <a href="#">GetTaxonomy</a>
tree.taxonomy	a dataframe containing taxonomic information for all species in the dataset see function <a href="#">GetTaxonomy</a>
data	a dataframe containing the data you would like to use in your comparative analysis
repl.table	the output of ReplTable and one of the inputs of MaxTips
levels	currently not implemented
verbose	turns on reports to the screen.

## Details

The goal of these two functions is to maximize the number of tips that are retained in a comparative phylogenetic analysis. This is done by searching for higher taxonomic level matches. This is accomplished by keeping one example of any available genera not matched at the species level and one available member of each family not matched at the genus or species level.

The taxonomy tables should have columns name of "Family", "Genus", and "Binomial". The dataframes (tree.taxonomy, data.taxonomy, and data) should all have row names corresponding to the binomial.

## Value

ReplTable produces a dataframe that is used as one of the inputs for MaxTips. It lists the tips on the tree that will be replaced and what taxonomic level matches have been found at. MaxTips returns a pruned dataset and tree.

## Author(s)

Heath Blackmon and Matt Pennell

## References

<http://www.uta.edu/karyodb/evobiR/>

## See Also

[GetTaxonomy](#)

## Examples

```
## Not run:
data('mite.data')
data('mite.tree')
data('data.taxonomy')
data('tree.taxonomy')

replacement.table <- ReplTable(tree = mite.tree, data.taxonomy = data.taxonomy,
                              tree.taxonomy = tree.taxonomy, levels=c("Genus", "Family"),
                              verbose=TRUE)
new.data <- MaxTips(mite.tree, mite.data, replacement.table)
## End(Not run)
```

---

ResSel	<i>Selection on Residuals</i>
--------	-------------------------------

---

**Description**

This function takes measurements of multiple traits and performs a linear regression and identifies those records with the largest and smallest residual. Originally it was written to perform a regression of horn size on body size allowing for high and low selection lines.

**Usage**

```
ResSel(data, traits, percent = 10, identifier = 1, model = "linear")
```

**Arguments**

data	this is a dataframe with subject identifiers and phenotypic trait values
traits	a numeric vector indicating the column containing the predictor and response variables in that order
percent	the percentage of highest and lowest residuals that should be identified
identifier	the column which contains the record numbers to identify individuals
model	currently this is not used

**Value**

This function returns a list

high line	the ID numbers for the individuals selected for the high line
low line	the ID numbers of the individuals selected for the low line

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

**Examples**

```
data <- read.csv(file = system.file("horn.beetle.csv", package = "evobiR"))  
ResSel(data = data, traits = c(2,3), percent = 15, identifier = 1, model = "linear")
```

---

SampleTrees                      *Select a random sample of trees*

---

### Description

This function takes as its input a large collection of trees from a program like MrBayes or Beast and allows the user to select the number of randomly drawn trees they wish to retrieve

### Usage

```
SampleTrees(trees, burnin, final.number, format, prefix)
```

### Arguments

trees	a nexus format file containing trees that the user wants to sample from
burnin	the proportion of trees to remove as burnin
final.number	the number of trees desired
format	options are "new" or "nex" indicating to save the trees in newick format or nexus format
prefix	a text string to assign to the new treefile name

### Value

an object of the class "multiPhylo" is returned

### Author(s)

Heath Blackmon

### References

<http://www.uta.edu/karyodb/evobiR/>

### Examples

```
SampleTrees(trees = system.file("trees.nex", package = "evobiR"),
            burnin = .1, final.number = 20, format = 'new', prefix = 'sample')
```

---

species.tree                      *Species Phylogenetic tree*

---

### Description

This is a simulated phylogenetic trees with 200 tips.

---

taxa.table	<i>A table with taxonomic information</i>
------------	---

---

**Description**

This character matrix contains species in one column followed by the genera and the family membership of each species

---

tree.taxonomy	<i>taxonomic table</i>
---------------	------------------------

---

**Description**

This file contains taxonomic information

**Author(s)**

Heath Blackmon

**References**

<http://www.uta.edu/karyodb/evobiR/>

---

trees	<i>10 Phylogenetic trees</i>
-------	------------------------------

---

**Description**

This is a collection of 10 simulated phylogenetic trees with 200 tips each.

---

trees.nex	<i>100 Phylogenetic trees</i>
-----------	-------------------------------

---

**Description**

This is a collection of 100 simulated phylogenetic trees with 10 tips each.

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