## Calculation of evolutionary rates and rate shifts with the RRphylo method

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**Depends** ape, phytools, geiger, stats4, foreach, doParallel, pvclust, mvMORPH, lmtest, parallel, phangorn, rlist, scales, R.utils, binr, RColorBrewer, nlme, penalized, smatr, data.tree, car, outliers, emmeans, picante, vegan, plotrix, cluster, tseries, ddpcr, geomorph

# 1. RRphylo

**Title** *Evolutionary rates computation along phylogenies* 

## **Description**

The function RRphylo performs the phylogenetic ridge regression. It takes a tree and a vector of tip data (phenotypes) as entries, calculates the penalization factor  $\lambda$ , produces the matrices of tip to root, and node to root distances (matrix **L** and **L**'), the vector of ancestral state estimates, the vector of predicted phenotypes, and the rates vector  $\hat{\beta}$  for all the branches of the

tree. For multivariate data, rates are given as both one entry per variable, and as a multivariate vector obtained by computing the Euclidean Norm of the rate entries per observation.

### **Usage**

RRphylo(tree, y, cov=NULL, rootV=NULL, aces=NULL, clus=0.5)

## **Arguments**

tree a phylogenetic tree. The tree needs not to be ultrametric or fully dichotomous y either a single vector variable or a multivariate dataset of class 'matrix' cov the covariate to be indicated if its effect on the rates must be accounted for. In this case residuals of the covariate versus the rates are used as rates. 'cov' must be as long as the number of nodes plus the number of tips of the tree, which can be obtained by running RRphylo on the covariate as well, and taking the vector of ancestral states and tip values to form the covariate, as in the example below.

**rootV** phenotypic value (values if multivariate) at the tree root. If rootV=NULL the function 'learns' about the root value from the 10% tips being closest in time to the tree root, weighted by their temporal distance from the root itself (close tips phenotypes weigh more than more distant tips).

**aces** a named vector (or matrix if y is multivariate) of ancestral character values at nodes. Names correspond to the nodes in the tree.

x1 the additional predictor to be indicated to perform the multiple version of RRphylo. 'x1' vector must be as long as the number of nodes plus the number of tips of the tree, which can be obtained by running RRphylo on the predictor as well, and taking the vector of ancestral states and tip values to form the x1.

aces.x1 a named vector of ancestral character values at nodes for 'x1'. It must be indicated if both 'aces' and 'x1' are specified. Names correspond to the nodes in the tree.clus the proportion of clusters to be used in parallel computing (only if y is multivariate).

#### **Details**

Details about RRphylo are in paragraph 1.1.

#### Value

**tree** the tree used by RRphylo. The fully dichotomous version of the tree argument. For trees with polytomies, the tree is resolved by using multi2di function in the package ape. If the latter is a dichotomous tree, the two trees will be identical **tip.path** a n x m matrix, where n=number of tips and m=number of branches (i.e. n\*2-1). Each row represents the branch lengths along a root-to-tip path.

**node.path** a n x n matrix, where n=number of internal branches. Each row represents the branch lengths along a root-to-node path.

**rates** single rate values computed for each branch of the tree. If y is a single vector variable, rates are equal to multiple.rates. If y is a multivariate dataset, rates are computed as the square root of the sum of squares of each row of multiple.rates. **aces** the phenotypes reconstructed at nodes.

**predicted.phenotypes** the vector of estimated tip values. It is a matrix in the case of multivariate data.

**multiple.rates** a n x m matrix, where n= number of branches (i.e. n\*2-1) and m = number of variables. For each branch, the column entries represent the evolutionary rate.

**lambda** the penalization factor fitted within RRphylo by the inner function optL. With multivariate data, several optL runs are performed. Hence, the function provides a single lambda for each individual variable.

**ace.values** if aces are specified, the function returns a data frame containing the corresponding node number on the RRphylo tree for each node, along with estimated values.

**x1.rate** if x1 is specified, the function returns the partial regression coefficient for x1.

## **Examples**

```
## data("DataOrnithodirans")
## DataOrnithodirans$treedino->treedino
## DataOrnithodirans$massdino->massdino

## Case 1. RRphylo without accounting for the effect of a covariate
# RRphylo(tree=treedino,y=massdino)

## Case 2. RRphylo without accounting for the effect of a covariate
## performing RRphylo on the covariate (body mass itself) in order
to retrieve ancestral state values
# RRphylo(tree=treedino,y=massdino)->RRcova
# c(RRcova$aces,massdino)->cov.values
# c(rownames(RRcova$aces),names(massdino))->names(cov.values)
```

# RRphylo(tree=treedino,y=massdino,cov=cov.values)

```
## Case 3. "RRphylo" specifying the ancestral states
# data("DataCetaceans")
# DataCetaceans$treecet->treecet
# DataCetaceans$masscet->masscet
# DataCetaceans$brainmasscet->brainmasscet
# DataCetaceans$aceMyst->aceMyst
# RRphylo(tree=treecet, y=masscet, aces=aceMyst)
## Case 4. Multiple "RRphylo"
# drop.tip(treecet, treecet$tip.label[-match(names(brainmasscet),
treecet$tip.label)])->treecet.multi
# masscet[match(treecet.multi$tip.label,names(masscet))]
->masscet.multi
# RRphylo(treecet.multi, masscet.multi) ->RRmass.multi
# RRmass.multi$aces[,1]->acemass.multi
# c(acemass.multi, masscet.multi) ->x1.mass
# RRphylo(treecet.multi,y=brainmasscet,x1=x1.mass)
```

## 1.1. Phylogenetic Ridge Regression penalization within RRphylo

Under phylogenetic Ridge Race regression, the vector of rates at individual branches 'betas' is calculated as:

```
betas <- (solve(t(L) %*% L + lambda * diag(ncol(L))) %*% t(L)) \  \  \, \%*\% \  \  \, as.matrix(y)
```

and the vector of predicted phenotypes as:

```
y <- L %*% betas
```

Where **L** is the matrix of root to tip distances provided by RRphylo, and lambda is the penalization factor fitted within RRphylo. Note that without penalization, the vector of rates is calculated as to compute the phenotypic vector perfectly:

```
L%*%t(L)%*%solve(L%*%t(L))%*%y
```

Kratch and McHardy (2014) used L2 (quadratic) penalization to find the value of the parameter  $\lambda$ . Within RRphylo,  $\lambda$  is fitted by an inner function optL, which is written as to minimize the rate variation along individual branches of the tree, thereby acting conservatively in terms of the chance to find rate shifts. We illustrate here the effect of assuming different  $\lambda$  values in RRphylo. We produced a random, non-ultrametric tree by using rtree function in the R package ape. Secondly, we simulated a phenotype evolving on such tree under the Brownian motion model (BM) of evolution. Then, we produced a uniform distribution of  $\lambda$  values, ranging from 0 to 10, and computed the phenotypes predicted by

RRphylo regression under the values of the uniform distribution of lambdas. Eventually, we calculated the resulting phylogenetic signal K (Blomberg et al. 2003) for each of the simulated phenotype. The plot of Ks versus the corresponding  $\lambda$  values is reported in figure 1.

```
## produce the tree and phenotype
library(ape)
library(lmtest)
rtree(100)->tree
setBM(tree,1,type="brown")->y
## perform RRphylo
RRphylo(tree,y)->RR
RR$tip.path->L
RR$node.path->L1
## compute the phylogenetic signal at different values of lambda, taken
from a uniform distribution of lambda values
seq(0,10, length.out=10000)->lambdaD
f=function(lambda) {
require(phytools)
betas <- (solve(t(L) %*% L + lambda * diag(ncol(L))) %*% t(L)) %*%
as.matrix(y)
aceRR <- L1 %*% betas[1:Nnode(tree), ]</pre>
y.hat <- L %*% betas
return(phylosig(tree,y.hat,test=TRUE))
}
```

```
library(parallel)
library(pbapply)
cl<-makeCluster(detectCores()-1)</pre>
clusterExport(cl, c("f", "L", "L1", "y", "tree"))
pbsapply(lambdaD, f, cl=cl) ->Ks
unlist (Ks[1,]) \rightarrow KS
unlist (Ks[2,]) \rightarrow Ps
## plot the phylogenetic signals versus lambdas. Eventually, the real
(fitted) lambda is superimposed on the plot
library(phytools)
col<-rep("black",length(KS))</pre>
col[which(KS>phylosig(tree,y)*0.75 & KS<phylosig(tree,y)*1.25)]<-"red"</pre>
col[which(Ps>.05)]<-"green"</pre>
plot(lambdaD, KS, col=col, xlab="fixed lambda value", ylab="fitted
phylogenetic signal K", cex=1.5, cex.lab=1.8)
abline(v=RR$lam, col="purple", lwd=2.5)
```

From the simulations it is empirically possible to note that the Brownian motion model of evolution (which is expected to produce values of phylogentic signal K close to 1) corresponds to small absolute values of  $\lambda$ . Yet, the phylogenetic signal becomes non-significant (i.e. phylogenetic effects disappear) for even smaller absolute values of  $\lambda$  (the green dots in Figure 1). It is important to notice that when the value of  $\lambda$  is fitted to the original tree and phenotype (rather than taken from a uniform distribution of  $\lambda$  values as in the simulations), RRphylo produces phenotypes evolving according to BM, as expected (the

fitted  $\lambda$  value is represented by the purple vertical line in figure 1), which implies the function works properly.

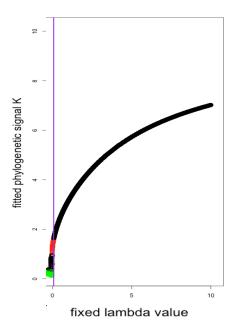
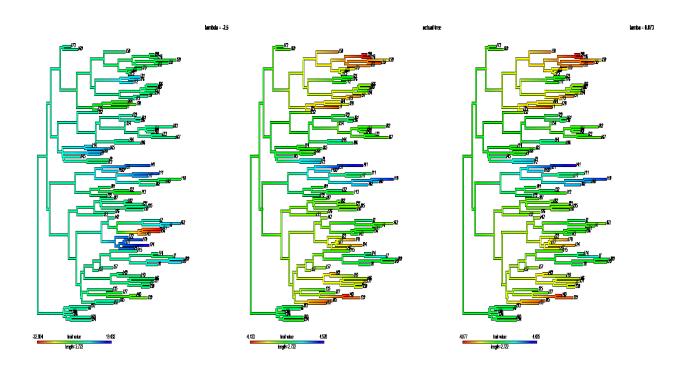


Figure 1 The phylogenetic signal K at different values of the penalization factor  $\lambda$ . The initial phenotype was simulated according to the Brownian motion model of evolution. As expected, the fitted  $\lambda$  value (indicated by the purple vertical line) produces a phylogenetic signal close to 1. The small box on the right represents the same plot as in the larger figure, yet cut down to the region of small  $\lambda$  values. Please notice that since version 1.1 the estimation of  $\lambda$  is restricted in between 0 to 10

However, values of K consistently close to 1 appear at very low values of  $\lambda$  values. To better appreciate the effect of assuming different  $\lambda$  values, we plotted the original tree with the branches colored according to the phenotypes simulated under BM by using the function

contMap in the R package phytools (Figure 2, center), produced with the fitted  $\lambda$  (Figure 2, right), and with a  $\lambda$  values producing a phylogenetic signal close to 1 (see the red dots in the negative realm of  $\lambda$  in Figure 1). Under these small values of  $\lambda$ , the phenotypes deviate significantly from the BM expectation (Figure 2, left). This means that RRphylo produces, beyond a legitimate value of K, a distribution of phenotypes at the tips very close to the original distribution.



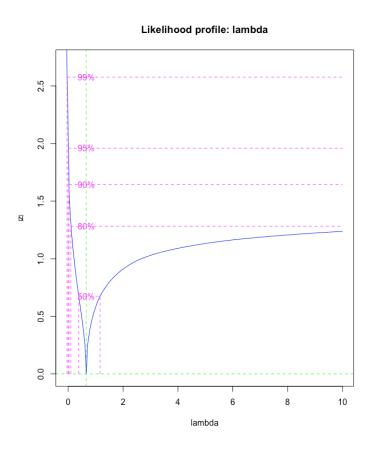
**Figure 2** The distribution of phenotypes produced with  $\lambda$  values giving a K value close to the original, Brownian motion simulation. The original phenotypes are reported in the tree at the center of the figure. The phenotypes with the  $\lambda$  value fitted in RRphylo are to the right. The tree to the left represent the phenotypes produced with a small value of  $\lambda$  that still gives a K value close to 1.

We next calculated the likelihood surface for the uniform distribution of  $\lambda$  values, by using the package bbmle (Bolker 2017). Of course, the likelihood peaks (in terms of deviance) at the

fitted value of  $\lambda$ , where deviance is the difference in negative log-likelihood from the likelihood at the MLE  $\lambda$ . The y-axis in Figure 2 represents the signed square root of the deviance difference (z), while the x-axis is the uniform distribution of starting  $\lambda$  values as used in the simulations. Please notice that the right end of the confidence interval is rather flat below the alpha level (95%). This could be adjusted by changing the standard error manually. Please refer to the bbmle2 package vignette for full explanation.

```
t<-tree
library(bbmle)
 optL <- function(lambda) {</pre>
  y < - scale(y)
 betas <- (solve(t(L) %*% L + lambda * diag(ncol(L))) %*%
        t(L)) %*% as.matrix(y)
  aceRR <- L1 %*% betas[1:Nnode(t), ]</pre>
  y.hat <- L %*% betas
  Rvar <- array()</pre>
  for (i in 1:Ntip(t)) {
   ace.tip <- betas[match(names(which(L[i, ] != 0)),</pre>
                rownames(betas)), ]
   mat = as.matrix(dist(ace.tip))
   Rvar[i] <- sum(mat[row(mat) == col(mat) + 1])
  abs(1 - (var(Rvar) + (mean(as.matrix(y))/mean(y.hat))))
 }
```

```
logLik(mle2(optL, start = list(lambda = 10), method = "L-BFGS-B"))->LL1
mle2(optL, start = list(lambda = 10), method = "L-BFGS-B")->f1
profile(f1,maxsteps=5000,prof.upper=10,trace=TRUE)->pf1
plot(pf1)
```



**Figure 3** The likelihood profile of the *optL* function, used within *RRphylo* for the purpose of penalization.

As with version 2.2.0, RRphylo can perform multiple regression. If a predictor is specified by the argument x1 the the partial regression coefficient calculated for x1 is reported as \$x1.rate. The effect of \$x1.rate is easy to assess graphically.

```
# given a tree (tree) and a phenotype (y) produce a predictor x1 as:

### create a predictor x1 related to y
x1 <- (y+(runif(length(y), -0.5*sd(y), 0.5*sd(y))))*-1
plot(y,x1) ### this plot shows the original relationship between y and the predictor x1

### compute ancestral state estimates for x1
RRphylo(tree, x1)$aces[,1]->ace1
### perform multiple regression RRphylo
RRphylo(tree,y,x1=c(ace1,x1))->RR

plot(y,RR$x1.rate*x1) ### this plot shows the effect of the partial regression coefficient $x1.rate$ on the relationship between y and the predictor x1
```

## 2. Locating shifts with the search.shift function

**Title** *Locating shifts in phenotypic evolutionary rates* 

## **Description**

The function search.shift tests whether individual clades or isolated tips dispersed through the phylogeny evolve at different 'RRphylo' rates as compared to the rest of the tree. Instances of rate shifts may be automatically found.

## Usage

```
search.shift(RR, status.type = c("clade", "sparse"), node = NULL,
state = NULL, cov = NULL, nrep = 1000, f = NULL, foldername)
```

### **Arguments**

**RR** an object fitted by RRphylo.

**status.type** whether the "clade" or "sparse" condition must be tested.

**node** under the "clade" condition, the node (clade) to be tested for the rate shift. When multiple nodes are tested, they need to be written as in the example below. If 'node' is left unspecified, the function performs under the 'auto-recognize' feature, meaning it will automatically test individual clades for deviation of their rates from the background rate of the rest of the tree (see details).

**state** the state of the tips specified under the "sparse" condition.

**cov** the covariate to be indicated if its effect on rate values must be accounted for.

Contrary to RRphylo, 'cov' needs to be as long as the number of tips of the tree.

**nrep** the number of simulations to be performed for the rate shift test, by default nrep is set at 1000.

**f** the size of the smallest clade to be tested. By default, nodes subtending to one tenth of the tree tips are tested.

**foldername** the path of the folder where plots are to be found.

### **Details**

The function search shift takes the object produced by RRphylo. Two different conditions of rate change can be investigated. Under the "clade" condition the vector of node or nodes subjected to the shift must be provided. Alternatively, under the "sparse" case the (named) vector of states (indicating which tips are or are not evolving under the rate shift according to the tested hypothesis) must be indicated. In the "clade" case, the function may perform an 'auto-recognize' feature. Under such specification, the function automatically tests individual clades (from clades as large as one half of the tree down to a specified clade size) for deviation of their rates from the background rate of the rest of the tree, which is identical to the "clade" case. An inclusive clade with significantly high rates is likely to include descending clades with similarly significantly high rates. Hence, with 'auto-recognize' the search shift function is written as to scan clades individually and to select only the node subtending to the highest difference in mean absolute rates as compared to the rest of the tree. A plot of the tree highlighting nodes subtending to significantly different rates is automatically produced. Caution must be put into interpreting the 'auto-recognize' results. For instance, a clade with small rates and another with large rates could be individuated even under BM. This does not mean these clades are actual instances for rate shifts. Clades must be tested on their own without the 'auto-recognize' feature, which serves as guidance to the investigator, when no strong a priori hypothesis to be tested is advanced. The 'auto-recognize' feature is not meant to provide a test for a specific hypothesis. It serves as an optional guidance to understand whether and which clades show significantly large or small rates as compared to the rest of the tree. Individual clades are tested at once, meaning that significant instances of rate variation elsewhere on the tree are ignored. Conversely, running the "clade" condition without including the 'auto-recognize' feature, multiple clades presumed to evolve under the same shift are tested together, meaning that their rates are collectively contrasted to the rest of the tree, albeit they can additionally be compared to each other upon request. Under both the "clade" and "sparse" conditions, multiple clades could be specified at once, and optionally tested individually (for deviation of rates) against the rates of the rest of the tree and against each other. The histogram of random differences of mean rates distribution along with the position of the real difference of means is automatically generated by search.shift. Regardless of which condition is specified, the function output produces the real difference of means, and their significance value.

### Value

Under all circumstances, search.shift provides a vector of **\$rates**. If 'cov' values are provided, rates are regressed against the covariate and the residuals of such regression form the vector **\$rates**. Otherwise, **\$rates** are the same rates as with the RR argument.

Under "clade" case without specifying nodes (i.e. 'auto-recognize') a list including: \$ **\$all.clades** for each detected node, the data-frame includes the average rate difference (computed as the mean rate over all branches subtended by the node minus the average rate for the rest of the tree) and the probability that it do represent a real shift. Probabilities are contrasted to simulations shuffling the rates across the tree branches for a number of replicates specified by the argument nrep. Note that the p-values refer to the number of times the real average rates are larger (or smaller) than the rates averaged over the rest of the tree, divided by the number of simulations. Hence, large rates are significantly larger than the rest of the tree (at alpha = 0.05), when the probability is > 0.975; and small rates are significantly small for p < 0.025.

**\$single.clades** the same as with 'all.clades' but restricted to the largest/smallest rate values along a single clade (i.e. nested clades with smaller rate shifts are excluded). Large rates are significantly larger than the rest of the tree (at alpha = 0.05), when the probability is > 0.975; and small rates are significantly small for p < 0.025. Under "clade" condition by specifying the node argument:

**\$all.clades.together** if more than one node is tested, this specifies the average rate difference and the significance of the rate shift, by considering all the specified nodes as evolving under a single rate. As with the 'auto-recognize' feature, large rates are significantly larger than the rest of the tree (at alpha = 0.05), when the probability is > 0.975; and small rates are significantly small for p < 0.025.

**\$single.clades** this gives the significance for individual clades, tested separately. As previously, large rates are significantly larger than the rest of the tree (at alpha = 0.05), when the probability is > 0.975; and small rates are significantly small for p < 0.025.

Under the "sparse" condition:

**\$state.results** for each state, the data-frame includes the average rate difference (computed as the mean rate over all leaves evolving under a given state, minus the average rate for each other state or the rest of the tree) and the probability that the shift is real. Large rates are significantly larger (at alpha = 0.05), when the probability is > 0.975; and small rates are significantly small for p < 0.025. States are compared pairwise.

### **Examples**

data("DataOrnithodirans")

DataOrnithodirans\$treedino->treedino

DataOrnithodirans\$massdino->massdino

```
RRphylo(tree=treedino,y=massdino)->dinoRates
# Case 1. Without accounting for the effect of a covariate
# Case 1.1 "clade" condition
# with auto-recognize
search.shift(dinoRates, status.type= "clade", foldername=tempdir())
# testing two hypothetical clades
search.shift(dinoRates, status.type= "clade", node=c(697,748),
foldername=tempdir())
# Case 1.2 "sparse" condition
# testing the sparse condition.
search.shift(dinoRates, status.type= "sparse", state=statedino,
foldername=tempdir())
# Case 2. Accounting for the effect of a covariate
# Case 2.1 "clade" condition
search.shift(dinoRates, status.type= "clade", cov=massdino,
foldername=tempdir())
# Case 2.2 "sparse" condition
search.shift(dinoRates, status.type="sparse", state=statedino,
cov=massdino, foldername=tempdir())
```

DataOrnithodirans\$statedino->statedino

3. Producing phenotypes with desired trends in rate over time or phenotypic mean over time with setBM

**Title** *Producing simulated phenotypes with trends* 

## **Description**

The function setBM is wrapper around 'phytools' *fastBM* function, which generates BM simulated phenotypes with or without a trend.

### Usage

```
setBM(tree, nY = 1, s2 = 1, a = 0, type = c("", "brown",
    "trend", "drift"), trend.type = c("linear", "stepwise"), tr = 5, t.shift
= 0.5, es=2, ds=1)
```

## **Arguments**

**tree** a phylogenetic tree.

**nY** the number of phenotypes to simulate.

**s2** value of the Browian rate to use in the simulations.

**a** the phenotype at the tree root.

**type** the type of phenotype to simulate. With the option "brown" the phenotype will have no trend in the phenotypic mean or in the rate of evolution. A variation in the phenotypic mean over time (a phenotypic trend) is obtained by selecting the option "drift". A trend in the rate of

evolution produces an increased variance in the residuals over time. This is obtained by specifying the option "trend".

**trend.type** two kinds of heteroscedastic residuals are generated under the "trend" type. The option "linear" produces an exponential increase (or decrease) in heteroscedasticity, whereas the "stepwise" option produces an increase (or decrease) after a specified point in time.

**tr** the intensity of the trend with the "stepwise" option is controlled by the tr argument. The scalar tr is the multiplier of the branches extending after the shift point as indicated by "t.shift". **t.shift** the relative time distance from the tree root where the stepwise change in the rate of evolution is indicated to apply.

**es** when trend.type="linear", es is a scalar representing the exponent at which the evolutionary time (i.e. distance from the root) scales to change to phenotypic variance. With es=1 the phenotypic rate will be trendless, with es<1 the variance of the phenotypes will decrease exponentially towards the present and the other way around with es>1.

**ds** a scalar indicating the change in phenotypic mean in the unit time, in type="drift" case. With ds = 0 the phenotype will be trendless, with ds < 0 the phenotypic mean will decrease towards the present and the other way around with ds > 0.

## **Details**

setBM differs from fastBM in that the produced phenotypes are checked for the existence of a temporal trend in the phenotype. The user may specify whether she wants trendless data (option "brown"), phenotypes trending in time (option "drift"), or phenotypes whose variance increases/decreases exponentially over time, consistently with the existence of a trend in the rate of evolution (option "trend"). In the latter case, the user may indicate the intensity of the

trend (by applying different values of *es*), and whether it should occur after a given proportion of the tree height (hence a given point back in time, specified by the argument *t.shift*).

Trees in setBM are treated as non ultrametric. If an ultrametric tree is fed to the function, setBM alters slightly the leaf lengths multiplying randomly half of the leaves by  $1 * 10^{-3}$  in order to make it non-ultrametric.

### **Value**

Either an object of class array containing a single phenotype or an object of class matrix of n phenotypes as columns, where n is indicated as nY = n

## **Examples**

```
## data("DataOrnithodirans")
## DataOrnithodirans$treedino->treedino

## setBM(tree=treedino, nY= 1, type="brown")
## setBM(tree=treedino, nY= 1, type="drift", ds=2)
## setBM(tree=treedino, nY= 1, type="trend", trend.type="linear", es=2)
```

# 4. Find a node subtending to a clade of desired size

The function *sizedsubtree* scans a phylogentic tree to randomly find nodes subtending to a subtree of desired minimum size, up to one half of the tree size (number of tips).

## **Usage**

```
sizedsubtree(tree, Size=NULL, time.limit=10)
```

# **Arguments**

**tree** a phylogenetic tree

**Size** the desired size of the tree subtending to the extracted node. By default, the minimu tree size is set at one tenth of the tree size (i.e. number of tips).

**time.limit** specifies a limit to the searching time, a warning message is thrown if the limit is reached.

### **Details**

The argument time.limit sets the searching time. The algorithm stops if that limit is reached, avoiding recursive search when no solution is in fact possible.

### **Value**

A node subtending to a subtree of desired minimum size.

# **Examples**

```
## data("DataOrnithodirans")
## DataOrnithodirans$treedino->treedino
## sizedsubtree(tree=treedino)
## library(geiger)
## tips(treedino,sizedsubtree(treedino,size=50)) # to view the tip
extracted
## extract.clade(treedino,sizedsubtree(treedino,size=50))# to extract the
subtree
```

## 5. evo.dir

**Title** *Phylogenetic vector analysis of phenotypic change* 

# **Description**

This function quantifies direction, size and rate of evolutionary change of multivariate traits along node-to-tip paths and between species.

## **Usage**

```
evo.dir(RR, angle.dimension=c("rates", "phenotypes"), y.type=c("original", "
RR"), y=NULL, pair.type=c("node", "tips"), pair=NULL, node=NULL, random=c("yes
", "no"), nrep=100)
```

## **Arguments**

**RR** an object produced by *RRphylo*.

angle.dimension specifies whether "rates" or "phenotypes" vectors are used.

y.type must be indicated when angle.dimension = "phenotypes". If "original", it uses the phenotypes provided by the user, if "RR" it uses RR\$predicted.phenotypes as variable.
y specifies the phenotypes to be provided if y.type = "original".

**pair.type** either "node" or "tips". Angles are computed between each possible couple of species descending from a specified node ("node"), or between a given couple of species ("tips").

pair species pair to be specified if pair.type = "tips". It needs to be written as in the example
below.

**node** node number to be specified if pair.type = "node". Notice the node number must refer to the dichotomic version of the original tree, as produced by *RRphylo*.

**random** whether to perform randomization test ("yes"/"no").

**nrep** number of replications must be indicated if random = "yes". It is set at 100 by default.

#### **Details**

The way *evo.dir* computes vectors depends on whether phenotypes or rates are used as variables. *RRphylo* rates along a path are aligned along a chain of ancestor/descendant relationships. As such, each rate vector origin coincides to the tip of its ancestor, and the resultant of the path is given by vector addition. In contrast, phenotypic vectors are computed with reference to a common origin (i.e. the consensus shape in a geometric morphometrics). In this latter case, vector subtraction (rather than addition) will define the resultant of the species evolutionary direction.

It is important to realize that resultants could be at any angle even if the species (the terminal vectors) have similar phenotypes, because path resultants, rather than individual phenotypes, are being contrasted. However, the function also provides the angle between individual phenotypes as 'angle.between.species'.

To perform randomization test, the evolutionary directions of the two species are collapsed together. Then, for each variable, the median is found, and random paths of the same size as the original paths are produced sampling at random from the 47.5th to the 52.5th percentile around the medians. This way, a random distribution of angles is obtained under the hypothesis that the two directions are actually parallel. The 'angle.direction' represents the angle formed by the species phenotype and a vector of 1s (as long as the number of variables representing the phenotype). This way, each species phenotype is contrasted to the same vector. The 'angle.direction' values could be inspected to test whether individual species phenotypes evolve towards similar directions.

#### Value

Under all specs, *evo.dir* returns a list object. The length of the list is one if pair.type = "tips". If pair.type = "node", the list is as long as the number of all possible species pairs descending from the node. Each element of the list contains:

angle.path.A angle of the resultant vector of species A to MRCA
vector.size.species.A size of the resultant vector of species A to MRCA
angle.path.B angle of the resultant vector of species B to MRCA
vector.size.species.B size of the resultant vector of species B to MRCA
angle.between.species.to.mrca angle between the species paths resultant vectors to
the MRCA

**angle.between.species** angle between species vectors (as they are, without computing the path)

MRCA the node identifying the most recent common ancestor of A and B angle.direction.A angle of the vector of species A (as it is, without computing the path) to a fixed reference vector (the same for all species)

vec.size.direction.A size of the vector of species A

**angle.direction.B** angle of the vector of species B (as it is, without computing the path) to a fixed reference vector (the same for all species)

**vec.size.direction.B** size of the vector of species B

If random = "yes", results also include p-values for the angles.

# **Examples**

```
# data("DataApes")
# DataApes$PCstage->PCstage
# DataApes$Tstage->Tstage

# RRphylo(tree=Tstage,y=PCstage)->RR

## Case 1. Without performing randomization test

# Case 1.1 Computing angles between rate vectors
## for each possible couple of species descending from node 72
```

# evo.dir(RR,angle.dimension="rates",pair.type="node",node=72,

```
random="no")
## for a given couple of species
# evo.dir(RR,angle.dimension="rates",pair.type="tips",pair=c("Sap 1",
"Tro 2"), random="no")
## Case 1.2 computing angles between phenotypic vectors provided by the
user
## for each possible couple of species descending from node 72
# evo.dir(RR,angle.dimension="phenotypes",y.type="original",y=PCstage,
pair.type="node", node=72, random="no")
## for a given couple of species
# evo.dir(RR,angle.dimension="phenotypes",y.type="original",y=PCstage,
pair.type="tips",pair=c("Sap 1","Tro 2"),random="no")
## Case 1.3 computing angles between phenotypic vectors produced by RRphylo
## for each possible couple of species descending from node 72
# evo.dir(RR,angle.dimension="phenotypes",y.type="RR",pair.type="node",
node=72, random="no")
### for a given couple of species
# evo.dir(RR,angle.dimension="phenotypes",y.type="RR",pair.type="tips",
pair=c("Sap 1","Tro 2"),random="no")
## Case 2. Performing randomization test
## Case 2.1 Computing angles between rate vectors
```

## for each possible couple of species descending from node 72

```
# evo.dir(RR,angle.dimension="rates",pair.type="node",node=72,
random="yes", nrep=100)
## for a given couple of species
# evo.dir(RR,angle.dimension="rates",pair.type="tips",pair=c("Sap 1",
"Tro 2"), random="yes", nrep=100)
## Case 2.2 computing angles between phenotypic vectors provided by the
user
## for each possible couple of species descending from node 72
# evo.dir(RR,angle.dimension="phenotypes",y.type="original",y=PCstage,
pair.type="node", node=72, random="yes", nrep=100)
## for a given couple of species
# evo.dir(RR, angle.dimension="phenotypes", y.type="original", y=PCstage,
pair.type="tips",pair=c("Sap_1","Tro_2"),random="yes",nrep=100)
## Case 2.3 computing angles between phenotypic vectors produced by RRphylo
## for each possible couple of species descending from node 72
# evo.dir(RR,angle.dimension="phenotypes",y.type="RR",pair.type="node",
node=72, random="yes", nrep=100)
## for a given couple of species
# evo.dir(RR,angle.dimension="phenotypes",y.type="RR",pair.type="tips",
pair=c("Sap 1","Tro 2"), random="yes", nrep=100)
```

### 6. retrieve.angles

**Title** *Extracting a user-specified subset of the evo.dir results* 

## **Description**

This function takes the result list produced by *evo.dir* as the input and extracts a specific subset of it. The user may specify whether to extract the set of angles between species resultant vectors and the MRCA, the size of resultant vectors, or the set of angles between species.

## **Usage**

```
retrieve.angles(angles.res, wishlist=c("anglesMRCA", "angleDir", "angles.be
tween.species"), random=c("yes", "no"), focus=c("node", "species", "both", "no
ne"), node=NULL, species=NULL, write2csv=c("yes", "no"))
```

# **Arguments**

**angles.res** the object resulting from *evo.dir* function.

wishlist specifies whether to extract angles and sizes ("anglesMRCA") of resultant vectors between individual species and the MRCA, angles and sizes ("angleDir") of vectors between individual species and a fixed reference vector (the same for all species), or angles between species resultant vectors ("angles.between.species").

random it needs to be "yes" if angles.res object contains randomization results.

**focus** it can be "node", "species", "both", or "none", whether the user wants the results for a focal node, or for a given species, for both, or just wants to visualize everything.

**node** must be indicated if focus= "node" or "both". As for *evo.dir*, the node number must refer to the dichotomic version of the original tree, as produced by *RRphylo*.

**species** must be indicated if focus= "species" or "both".

write2csv if "yes" results are saved to a .csv file in your working directory.

#### **Details**

retrieve.angles allows to focalize the extraction to a particular node, species, or both. Otherwise it returns the whole dataset.

### Value

Irrespective of the cases, retrieve.angles outputs an object of class 'data.frame'.

If wishlist = "anglesMRCA", the data frame includes:

**MRCA** the most recent common ancestor the angle is computed to.

**species** species ID.

**angle** the angle between the resultant vector of species and the MRCA.

**vector.size** the size of the resultant vector computed from species to MRCA.

If wishlist = "angleDir", the data frame includes:

**MRCA** the most recent common ancestor the vector is computed to.

**species** species ID.

**angle.direction** the angle between the vector of species and a fixed reference.

**vector.size** the size of the vector of species.

If wishlist = "angles.between species", the data frame includes:

**MRCA** the most recent common ancestor the vector is computed from.

species pair IDs of the species pair the "angle between species" is computed for.
angleBTWspecies2MRCA angle between species resultant vectors to MRCA
anglesBTWspecies angle between species resultant vectors

## **Examples**

```
# data("DataApes")
# DataApes$PCstage->PCstage
# DataApes$Tstage->Tstage
# RRphylo(tree=Tstage, y=PCstage) ->RR
## Case 1. "evo.dir" without performing randomizatin
# evo.dir(RR, angle.dimension="rates", pair.type="node", node=57,
random="no") ->evo.p
## Case 1.1 retrieve angles and sizes of resultant vectors between
individual species and the MRCA:
## for a focal node
   retrieve.angles(evo.p, wishlist="anglesMRCA", random="no", focus="node",
node=68, write2csv="no")
## for a fcal species
# retrieve.angles(evo.p, wishlist="anglesMRCA", random="no", focus=
"species", species="Sap", write2csv="no")
## for both focal node and species
   retrieve.angles(evo.p, wishlist="anglesMRCA", random="no", focus="both",
node=68, species="Sap", write2csv="no")
```

```
## without any specific requirement
   retrieve.angles(evo.p, wishlist="anglesMRCA", random="no", focus="none",
write2csv="no")
## Case 1.2 retrieve angles and sizes of vectors between individual species
and a fixed reference vector:
## for a focal node
     retrieve.angles(evo.p, wishlist="angleDir", random="no", focus="node",
node=68, write2csv="no")
## for a focal species
# retrieve.angles(evo.p, wishlist="angleDir", random="no", focus=
"species", species="Sap", write2csv="no")
## for both focal node and species
     retrieve.angles(evo.p, wishlist="angleDir", random="no", focus="both",
node=68, species="Sap", write2csv="no")
## without any specific requirement
     retrieve.angles(evo.p, wishlist="angleDir", random="no", focus="none",
write2csv="no")
## Case 1.3 retrieve angles between species resultant vectors:
 ## for a focal node
    retrieve.angles(evo.p, wishlist="angles.between.species", random="no",
focus="node", node=68, write2csv="no")
## for a focal species
    retrieve.angles(evo.p, wishlist="angles.between.species", random="no",
focus="species", species="Sap", write2csv="no")
## for both focal node and species
```

```
retrieve.angles(evo.p, wishlist="angles.between.species", random="no",
focus="both", node=68, species="Sap", write2csv="no")
## without any specific requirement
    retrieve.angles(evo.p, wishlist="angles.between.species", random="no",
focus="none", write2csv="no")
## Case 2. "evo.dir" with performing randomization
# evo.dir (RR, angle.dimension="rates", pair.type="node", node=57,
random="yes")->evo.p
## Case 2.1 retrieve angles and sizes of resultant vectors between
individual species and the MRCA:
## for a focal node
# retrieve.angles(evo.p,wishlist="anglesMRCA",random="yes",focus="node"
, node=68, write2csv="no")
## for a focal species
# retrieve.angles(evo.p, wishlist="anglesMRCA", random="yes", focus=
"species", species="Sap", write2csv="no")
## for both focal node and species
# retrieve.angles(evo.p, wishlist="anglesMRCA", random="yes", focus=
"both", node=68, species="Sap", write2csv="no")
## without any specific requirement
# retrieve.angles(evo.p, wishlist="anglesMRCA", random="yes", focus=
"none", write2csv="no")
```

```
## Case 2.2 retrieve angles and sizes of resultant vectors between
individual species and the MRCA:
## for a focal node
# retrieve.angles(evo.p, wishlist="angleDir", random="yes", focus="node"
, node=68, write2csv="no")
## for a focal species
# retrieve.angles(evo.p,wishlist="angleDir",random="yes",focus=
"species", species="Sap", write2csv="no")
## for both focal node and species
# retrieve.angles(evo.p, wishlist="angleDir", random="yes", focus=
"both", node=68, species="Sap", write2csv="no")
## without any specific requirement
# retrieve.angles(evo.p, wishlist="angleDir", random="yes", focus=
"none", write2csv="no")
## Case 2.3 retrieve angles between species resultant vectors:
## for a focal node
# retrieve.angles(evo.p, wishlist="angles.between.species", random="yes",
focus="node", node=68, write2csv="no")
## for a focal species
   retrieve.angles(evo.p, wishlist="angles.between.species", random="yes",
focus="species", species="Sap", write2csv="no")
## for both focal node and species
   retrieve.angles(evo.p, wishlist="angles.between.species", random="yes",
focus="both", node=68, species="Sap", write2csv="no")
## without any specific requirement
```

```
# retrieve.angles(evo.p, wishlist="angles.between.species", random="yes",
focus="none", write2csv="no")
```

## 8. angle.matrix

**Title** *Ontogenetic shape vectors analysis* 

## **Description**

This function computes ontogenetic vectors for each species and compares them between species pairs.

## **Usage**

```
angle.matrix(RR, node, Y=NULL, select.axes=c("no", "yes"), type=c("phenotypes
", "rates"), cova=NULL)
```

## **Arguments**

RR an object produced by RRphylo.

**node** the number of the node subtending species to compute ontogenetic vectors on. Notice the node number must refer to the dichotomic version of the original tree, as it is produced by *RRphylo*.

Y multivariate trait values at tips.

**select.axes** if "yes", Y variables are individually regressed against developmental stages and only significant phenotypes are retained to compute ontogenetic vectors. If "no", all variables are retained.

**type** specifies weather to perform the analysis on phenotypic ("phenotypes") or rate ("rates") vectors.

**cova** the covariate to be indicated if its effect on rate values must be accounted for. Contrary to *RRphylo*, 'cova' needs to be as long as the number of tips of the tree. As the covariate only affects rates computation, there is no covariate to provide when type = "phenotypes".

#### **Details**

The *angle.matrix* function takes as objects a phylogenetic tree (deriving directly from an RR object), including the different ontogenetic stages of each species as polytomies. Names at tips must be written as species ID and stage number separated by the underscore (for example "Sap\_1", that is *Homo sapiens* at stage 1).

The RRphylo object *angle.matrix* is fed with is just used to extract the dichotomized version of the phylogeny. This is useful because node number changes randomly at dichotomizing non-binary trees. However, when performing *angle.matrix* with the covariate there is no need for the RR object to be produced with the argument covariate set to TRUE. Furthermore, as the covariate only affects the rates computation, it makes no sense to use it when computing vectors for phenotypic variables.

Once angles and vectors are computed, *angle.matrix* performs two tests by means of standard major axis (SMA) regression. For each species pair, the "biogenetic test" verifies whether the angle between species grows during development, meaning that the two species becomes less similar to each other during growth, in keeping with the biogenetic law. The

"paedomorphosis test" tells whether there is heterochronic shape change in the data. Under paedomorphosis, the adult stages of one (paedomorphic) species will resemble the juvenile stages of the other (peramorphic) species. The test regresses the angles formed by the shapes at different ontogenetic stages of a species to the shape at the youngest stage of the other in the pair, against age. Then, it tests whether the two regression lines (one per species) have different slopes, and whether they have different signs. If the regression lines point to different directions and differ from each other, it means that one of the two species in the pair resembles, with age, the juveniles of the other, indicating paedomorphosis. Ontogenetic vectors of individual species are further computed, in reference to the MRCA of the pair, and to the first stage of each species (i.e. intraspecifically). Importantly, the size of the ontogenetic vectors of rates tell whether the two species differ in terms of developmental rate, which is crucial to understand which process is behind paedomorphosis, where it applies.

While performing the analysis, the function prints messages on-screen. If select.axes = "yes", it informs the user about which phenotypic variables are used. Secondly, it specifies whether ontogenetic vectors to MRCA, and intraspecific ontogenetic vectors significantly differ in angle or size between species pairs. Then, for each species pair, it indicates if the biogenetic law and paedomorphosis apply.

#### Value

A *list* containing 4 objects.

**\$regression.matrix:** a 'list' object including "angles between species" and "angles between species to MRCA" matrices for all possible combinations of species pairs from the two sides descending from the MRCA. For each matrix, corresponding biogenetic and paedomorphosis tests are reported.

```
$`angles between species`

$`Species1/Species2`

$matrix

$paedomorphosis test

$biogenetic test

$`Species1/Species3`

$matrix

$paedomorphosis test
```

\$biogenetic test

\$`angles between species 2 MRCA`

\$`Species1/Species2`

\$matrix

\$paedomorphosis test

\$biogenetic test

\$`Species1/Species3`

\$matrix

\$paedomorphosis test

\$biogenetic test

**\$angles.2.MRCA.and.vector.size:** a 'data.frame' including angles between the resultant vector of species and the MRCA and the size of the resultant vector computed from species to MRCA, per stage per species.

**\$ontogenetic.vectors2MRCA:** a 'data.frame' including angle, size, and corresponding x and y components, of ontogenetic vectors between each species and the MRCA. For both angle and size, the p-value for the difference between species pairs is reported.

**\$ontogenetic.vectors.to.1st.stage:** a 'list' object containing:

**\$matrices:** for all possible combinations of species pairs from the two sides descending form the MRCA, the upper triangle of the matrix contains angles between different ontogenetic stages for the first species (i.e. Species1). The same applies to the lower triangle for the second species (i.e. Species2 or Species3).

\$ `Species1/Species2`

\$`Species1/Species3`

**\$vectors**: for all possible combinations of species pairs from the two sides descending form the MRCA, angle and size of ontogenetic vectors computed to the first stage of each species. For both angle and size, the p-value for the difference between the species pair is reported.

\$`Species1/Species2`

\$`Species1/Species3`

#### **Examples**

# data("DataApes")

# DataApes\$PCstage->PCstage

# DataApes\$Tstage->Tstage

# DataApes\$CentroidSize->CS

# RRphylo(tree=Tstage,y=PCstage)->RR

```
## Case 1. "angle.matrix" without accounting for te effect of a
covariate
## Case 1.1 selecting only shape variables that show significant
relationship with age
## on phenotypic vectors
# angle.matrix(RR, node=72, Y=PCstage, select.axes="yes", type="phenotypes")
## on rates vectors
# angle.matrix(RR, node=72, Y=PCstage, select.axes="yes", type="rates")
## Case 1.2 using all shape variables
## on phenotypic vectors
# angle.matrix(RR, node=72, Y=PCstage, select.axes="no", type="phenotypes")
## on rates vectors
# angle.matrix(RR, node=72, Y=PCstage, select.axes="no", type="rates")
## Case 2. "angle.matrix" accounting for the effect of a covariate (on
rates vectors only)
## Case 2.1 selecting only shape variables that show significant
relationship with age
# angle.matrix(RR, node=72, Y=PCstage, select.axes="yes", type="rates",
cova=CS)
```

## Case 2.2 using all shape variables

```
# angle.matrix(RR, node=72, Y=PCstage, select.axes="no", type="rates",
cova=CS)
```

#### 9. makeL

**Title** *Matrix of branch lengths along root-to-tip paths* 

# **Description**

This function produces a  $n \times m$  matrix, where n=number of tips and m=number of branches (i.e. n + number of nodes). Each row represents the branch lengths along a root-to-tip path.

## **Usage**

makeL(tree)

# **Arguments**

**tree** a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous.

#### Value

The function returns a n x m matrix of branch lengths along a root-to-tip path.

#### **Examples**

```
# data("DataApes")
```

# DataApes\$Tstage->Tstage

```
# makeL(tree=Tstage)
```

#### 10.makeL1

**Title** *Matrix of branch lengths along a root-to-node path* 

## **Description**

This function produces a  $n \times n$  matrix, where n=number of internal branches. Each row represents the branch lengths along a root-to-node path.

## **Usage**

```
makeL1(tree)
```

## **Arguments**

**tree** a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous.

#### Value

The function returns a n x n matrix of branch lengths along a root-to-node path.

## **Examples**

```
# data("DataApes")
# DataApes$Tstage->Tstage
# makeL1(tree=Tstage)
```

#### 11. getMommy

**Title** Upward tip or node to root path

## **Description**

Thi function is a wrapper around phytools *getDescendants* (Revell 2012). It returns the node path from a given node or species to the root of the phylogeny.

## **Usage**

getMommy(tree,N,curr=NULL)

### **Arguments**

**tree** a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous.

**N** the number of node or tip to perform the function on. Notice the function only works with number, not tip labels.

curr has not to be provided by the user.

#### **Details**

The object 'curr' is created inside the function in order to produce an array of nodes on the path.

#### Value

The function produces a vector of node numbers as integers, collated from a node or a tip towards the tree root.

# **Examples**

```
# data("DataApes")
# DataApes$Tstage->Tstage
# getMommy(tree=Tstage,12)
```

#### 12. makeFossil

**Title** *make fossil species on a phylogeny* 

## **Description**

This function takes an object of class 'phylo' and randomly changes the lengths of the leaves.

#### **Usage**

```
makeFossil(tree, p=0.5, ex=0.5)
```

## **Arguments**

tree a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous.
p the proportion of tips involved. By default it is half of the number of tips.
ex the multiplying parameter to change the leaf lengths. It is set at 0.5 by default.

#### Value

The function produces a phylogeny having the same backbone of the original one.

# **Examples**

```
# data("DataApes")
# DataApes$Tstage->Tstage
# makeFossil(tree=Tstage,p=0.5,ex=0.5)
```

## 13. swap.phylo

**Title** test the effect of phylogenetic uncertainty on rate shifts found at a particular node

#### **Description**

The function uses a number of alternative phylogenies with altered (as compared to the reference tree) topology and branch lengths to tests whether the tips descending from the specified node ('node') have statistically different rates from the rest of the tree. A phenotypic vector 'y' must be supplied. Eventually, the effect of a covariate could be included.

#### **Usage**

```
swap.phylo(tree, si=0.5, si2=0.5, node, y, rts, nrep=100, cov=NULL)
```

### **Arguments**

**tree** a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous.

**si** the proportion of tips whose topologic arrangement will be swapped.

**si2** the proportion of nodes whose age will be changed.

**node** the focal node to be tested.

**y** the phenotype under testing.

**rts** the rates found by RRphylo on the original tree.

**nrep** the number of simulated trees to be produced.

**cov** the covariate to be indicated if its effect on rate values must be accounted for. Contrary to *RRphylo*, 'cov' needs to be as long as the number of tips of the tree.

#### **Details**

swap.phylo changes the tree topology and branch lengths to a level specified by the user. Up to half of the tips, and half of the branch lengths can be changed randomly. The function provides a 'swapped' tree, yet, importantly, once a shift in the rate of evolution has been found by RRphylo, this function can be used to test whether the shift depends on the tree topology and branch lengths. It runs RRphylo on swapped trees (default is 100) and then calculates the absolute rate difference between all the branches of the shifted node and the rest of the tree. A t-test is eventually performed to assess significance.

#### Value

The function returns a 'list' object containing:

**\$p.swap** the probability that the rates at 'node' are different from rates at the rest of the tree.

**\$rates** the distribution of rates per branch as calculated by RRphylo on 'swapped' phylogenies.

### **Examples**

```
# data("DataApes")
# DataApes$PCstage->PCstage
# DataApes$Tstage->Tstage
# DataApes$CentroidSize->CS
## Case 1. swap.phylo without accounting for the effect of a
covariate
# RRphylo(tree=Tstage,y=PCstage)->RR
# RR$rates->rr
# swap.phylo(tree=Tstage, node=72, y=PCstage, rts=rr)
## Case 2. swap.phylo accounting for the effect of a covariate
# RRphylo(tree=Tstage,y=CS)->RRcova
# c(RRcova$aces,CS)->cov.values
# c(rownames(RRcova$aces), names(CS)) ->names(cov.values)
# RRphylo(tree=Tstage, y=PCstage, cov=cov.values) ->RR
# RR$rates->rr
# swap.phylo(tree=Tstage, node=72, y=PCstage, rts=rr, cov=CS)
```

## 14. swap.one

**Title** create alternative phylogenies from a given tree

#### **Usage**

swapONE (tree, si=0.5, si2=0.5)

## **Description**

The function produces an alternative phylogeny with altered topology and branch lengths, and computes the Kuhner-Felsenstein (Kuhner & Felsenstein 1994) distance between original and 'swapped' tree.

## **Arguments**

tree a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous.si the proportion of tips whose topologic arrangement will be swapped.si2 the proportion of nodes whose age will be changed.

#### **Details**

swap.one changes the tree topology and branch lengths. Up to half of the tips, and half of the branch lengths can be changed randomly. Each randomly selected node is allowed to move up to 2 nodes apart from its original position.

#### Value

The function returns a list containing the 'swapped' version of the original tree, and the Kuhner-Felsenstein distance between the trees.

### **Examples**

## data("DataOrnithodirans")

```
## DataOrnithodirans$treedino->treedino
```

```
# swapONE(tree = treedino)
```

### 15. plotRates

**Title** plot RRphylo rates at a specified node

### **Usage**

```
plotRates(RR, node, export.tiff = TRUE , foldername)
```

# **Description**

The function *plotRates* plots the *RRphylo* rates computed for a given clade as compared to the rates computed for the rest of the tree.

## **Arguments**

**RR** an object produced by RRphylo.

**node** the node subtending the clade of interest.

**export.tiff** if TRUE the function save a "rate\_bars.tiff" file inside the working directory. It is TRUE by default.

**foldername** the path of the folder where plots are to be found.

#### Value

The function produces two barplots. On the left side, the rates (in absolute values) computed for the focal clade (blue) are plotted against the rates of the rest of the tree (green). On the right side, the absolute rates of individual branches of the focal clade are collated in increasing rate value (blue bars), and contrasted to the average rate computed over the rest of the tree branches (the vertical red line). It also returns the rate values for both nodes and species descending from the focal node.

## **Examples**

```
# data("DataApes")
# DataApes$PCstage->PCstage
# DataApes$Tstage->Tstage
# RRphylo(tree=Tstage, y=PCstage)->RR
# plotRates(RR, node=72, export.tiff = FALSE, foldernam=tempdir())
```

### 16. search.trend

**Title** *searching for evolutionary trends in phenotypes and rates* 

#### **Description**

This function searches for evolutionary trends in phenotypic mean and evolutionary rates.

### **Usage**

```
search.trend(RR,y,node=NULL,nsim=100,clus=.5,foldername,cov=NULL,
ConfInt=FALSE)
```

### **Arguments**

**RR** an object produced by RRphylo.

**y** the named vector (or matrix if multivariate) of phenotypes.

x1 the additional predictor to be specified if the RR object has been created using an additional predictor (i.e. multiple version of RRphylo). 'x1' vector must be as long as the number of nodes plus the number of tips of the tree, which can be obtained by running RRphylo on the predictor as well, and taking the vector of ancestral states and tip values to form the 'x1'.

**node** number of individual clades to be specifically tested and contrasted to each other. It is NULL by default. Notice the node number must refer to the dichotomic version of the original tree, as produced by RRphylo.

**nsim** number of simulations to be performed. It is set at 100 by default.

**clus** the proportion of clusters to be used in parallel computing.

**foldername** the path of the folder where plots are to be found.

**cov** the covariate values to be specified if the RR object has been created using a covariate for rates calculation. As for RRphylo, 'cov' must be as long as the number of nodes plus the number of tips of the tree, which can be obtained by running RRphylo on the covariate as well, and taking the vector of ancestral states and tip values to form the covariate (see the example below).

**Confint** if TRUE, the function returns 95% confidence intervals around phenotypes and rates produced according to the Brownian motion model of evolution. It is FALSE by default.

#### **Details**

The function simultaneously returns the regression of phenotypes and phenotypic evolutionary rates against age (in terms of distance from the tree root) tested against Brownian motion simulations to assess significance. It stores the rates (absolute values) versus age regression and the phenotype versus age regression plots as .pdf files. In the plots, the 95% confidence intervals of phenotypes and rates simulated under the Brownian motion for each node are plotted as shaded areas. Regression lines are printed for all regressions. To assess significance, slopes are compared to a family of simulated slopes (BM<sub>slopes</sub>, where the number of simulations is equal to 'nsim'), generated under the Brownian motion, using the *fastBM* function in the package phytools.

Individual nodes are compared to the rest of the tree in different ways depending on whether phenotypes or rates versus age regressions are tested. With the former, the regression slopes for individual clades and the slope difference between clades is contrasted to slopes obtained through Brownian motion simulations. For the latter, regression models are tested and contrasted to each other referring to estimated marginal means, by using the *emmeans* function in the package emmeans.

### Value

The function returns a 'list' object including:

**\$rbt:** for each branch of the tree, the RRphylo rates and the distance from the tree root (age). If y is multivariate, it also includes the multiple rates for each y vector. If node is specified, each branch is classified as belonging or not to the indicated clades.

**\$pbt:** a data frame of phenotypic values and their distance from the tree root for each node (i.e. ancestral states) and tip of the tree.

**\$phenotypic.regression**: results of phenotype versus age regression. It reports a p-value for the regression slope between the variables (p.real), a p-value computed contrasting the real slope to Brownian motion simulations (p.random), and a parameter indicating the deviation of the phenotypic mean from the root value in terms of the number of standard deviations of the trait distribution (dev). dev is 0 under Brownian Motion. Only p.random should be inspected to assess significance.

**\$rate.regression**: results of the rates (absolute values) versus age regression. It reports a p-value for the regression between the variables (p.real), a p-value computed contrasting the real slope to Brownian motion simulations (p.random), and a parameter indicating the ratio between the range of phenotypic values and the range of such values halfway along the tree height, divided to the same figure under Brownian motion (spread). spread is 1 under Brownian Motion. Only p.random should be inspected to assess significance.

**\$Confints**: the 95% confidence intervals around phenotypes and rates produced according to the Brownian motion model of evolution.

**\$node.phenotypic.regression** results of phenotype versus age regression through node. It reports the slope for the regression between the variables at node (slope), a p-value computed contrasting the real slope to Brownian motion simulations (p.random), the difference between estimated marginal means predictions for the group and for the rest of the tree (emm.difference), and a pvalue for the emm.difference (p.emm).

**\$node.rate.regression** results of the rates (absolute values) versus age regression through node. It reports the difference between estimated marginal means predictions for the group and for the rest of the tree (emm.difference), a p-value for the emm.difference (p.emm), the difference between regression slopes for the group and for the rest of the tree (slope.difference), and a p-value for the slope.difference (p.slope).

If more than one node is specified, the object **\$group.comparison** reports the same results as \$node.phenpotypic.regression and \$node.rate.regression obtained by comparing individual clades to each other.

### **Examples**

```
## data("DataOrnithodirans")
## DataOrnithodirans$treedino->treedino
## DataOrnithodirans$massdino->massdino
### Extract the Pterosaurs tree and data
## extract.clade(treedino,748)->treeptero
## massdino[match(treeptero$tip.label, names(massdino))] ->massptero
## massptero[match(treeptero$tip.label,names(massptero))]->massptero
### Case 1. "RRphylo" whitout accounting for the effect of a covariate
# RRphylo(tree=treeptero,y=log(massptero))->RRptero
## Case 1.1. "search.trend" whitout indicating nodes to be specifically
tested for trends
# search.trend(RR=RRptero, y=log(massptero), nsim=100, clus=0.5,
foldername=tempdir(),cov=NULL,ConfInt=FALSE)
## Case 1.2. "search.trend" indicating nodes to be specifically tested
for trends
# search.trend(RR=RRptero, y=log(massptero), nsim=100, node=143,
clus=0.5, foldername=tempdir(),cov=NULL,ConfInt=FALSE)
```

```
### Case 2. "RRphylo" accounting for the effect of a covariate
## "RRphylo" on the covariate in order to retrieve ancestral state
values
# RRphylo(tree=treeptero,y=log(massptero))->RRptero
# c(RRptero$aces,log(massptero))->cov.values
# names(cov.values)<-c(rownames(RRptero$aces), names(massptero))</pre>
# RRphylo(tree=treeptero,y=log(massptero),cov=cov.values)->RRpteroCov
## Case 2.1. "search.trend" whitout indicating nodes to be specifically
tested for trends
# search.trend(RR=RRpteroCov, y=log(massptero), nsim=100, clus=0.5,
foldername=tempdir(),ConfInt=FALSE,cov=cov.values)
## Case 2.2. "search.trend" indicating nodes to be specifically tested
for trends
# search.trend(RR=RRpteroCov, y=log(massptero), nsim=100, node=143,
clus=0.5, foldername=tempdir(),ConfInt=FALSE,cov=cov.values)
## Case 3. "search.trend" on multiple "RRphylo"
# data("DataCetaceans")
# DataCetaceans$treecet->treecet
# DataCetaceans$masscet->masscet
# DataCetaceans$brainmasscet->brainmasscet
# DataCetaceans$aceMyst->aceMyst
# drop.tip(treecet, treecet$tip.label[-
match(names(brainmasscet), treecet$tip.label)]) ->treecet.multi
```

```
# masscet[match(treecet.multi$tip.label,names(masscet))]->masscet.multi
# RRphylo(treecet.multi,masscet.multi)->RRmass.multi
# RRmass.multi$aces[,1]->acemass.multi
# c(acemass.multi,masscet.multi)->x1.mass
# RRphylo(treecet.multi,y=brainmasscet,x1=x1.mass)->RRmulti
# search.trend(RR=RRmulti, y=brainmasscet,x1=x1.mass,
clus=0.5,foldername=tempdir())
```

### 17. getSis

Title Get sister clade

## **Description**

The function identifies and returns the sister clade of a given node/tip.

### **Usage**

```
getSis(tree,n,printZoom= TRUE)
```

## **Arguments**

**tree** a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous.

**n** number of focal node or name of focal tip.

**printZoom** if TRUE the function plots the tree section of interest.

#### Value

The sister node number or sister tip name. In case of polytomies, the function returns a vector.

## **Examples**

```
# data(DataOrnithodirans)
```

```
# DataOrnithodirans$treedino->treedino
```

```
# getSis(tree=treedino,n=678,printZoom=FALSE)
```

```
# getSis(tree=treedino,n="Shenzhoupterus chaoyangensis",printZoom=FALSE)
```

#### 18. distNodes

**Title** *Finding distance between nodes and tips* 

## **Description**

The function computes the distance between pairs of nodes, pairs of tips, or between nodes and tips. The distance is meant as both patristic distance and the number of nodes intervening between the pair.

#### **Usage**

```
distNodes(tree, node=NULL, clus=0.5)
```

## **Arguments**

**tree** a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous. **node** either a single node/tip or a pair of nodes/tips.

**clus** the proportion of clusters to be used in parallel computing.

### Value

If 'node' is specified, the function returns a data frame with distances between the focal node/tip and the other nodes/tips on the tree (or for the selected pair only). Otherwise, the function returns a matrix containing the number of nodes intervening between each pair of nodes and tips.

### **Examples**

```
# data("DataApes")
# DataApes$Tstage->Tstage

# distNodes(tree=Tstage)
# distNodes(tree=Tstage, node=64)
# distNodes(tree=Tstage, node="Tro_2")
# distNodes(tree=Tstage, n=c(64,48))
# distNodes(tree=Tstage, n=c(64, "Tro 2"))
```

#### 19. search.conv

**Title** Searching for morphological convergence among species and clades

## **Description**

The function scans a phylogenetic tree looking for morphological convergence between entire clades or species evolving under specific states.

#### Usage

```
search.conv(RR=NULL, tree=NULL, y, nodes=NULL, state=NULL, aceV=NULL,
min.dim=NULL, max.dim=NULL, min.dist=NULL, PGLS=TRUE, nsim=1000, rsim=1000,
clus=.5, foldername=NULL)
```

#### **Arguments**

**RR** an object produced by RRphylo. This is not indicated if convergence among states is tested. **tree** a phylogenetic tree. The tree needs not to be ultrametric or fully dichotomous. This is not indicated if convergence among clades is tested.

**y** a multivariate phenotype. The object y should be either a matrix or dataframe with species names as rownames.

**nodes** node pair to be tested. If unspecified, the function automatically searches for convergence among clades. Notice the node number must refer to the dichotomic version of the original tree, as produced by RRphylo.

state the state of the tips. If a single convergent state is indicated species within the state are tested. Otherwise, species belonging to different parts of the tree could be tested for convergence on each other by indicating different states. This latter case is especially meant to test for iterative evolution (i.e. the appearance of repeated morphotypes into different clades). The state for non-focal species (i.e. not belonging to any convergent group) must be indicated as "nostate". The state for non-focal species (i.e. not belonging to any convergent group) must be indicated as "nostate".

**aceV** phenotypic values at internal nodes. The object 'aceV' should be either a matrix

or dataframe with nodes (referred to the dichotomic version of the original tree, as produced by RRphylo) as rownames. If 'aceV' are not indicated, ancestral phenotypes are estimated via RRphylo.

**min.dim** the minimum size of the clades to be compared. When node is indicated, it indicates the minimum size of the smallest clades in nodes, otherwise it is set at one tenth of the tree size.

**max.dim** the maximum size of the clades to be compared. When node is indicated, it is 'min.dim'\*2 if the largest clade in 'nodes' is smaller than this value, otherwise it corresponds to the size of the largest clade. Whitout nodes it is set at one third of the tree size.

min.dist the minimum distance between the clades to be compared. When nodes is indicated, it is the distance between the pair. Under the automatic mode, the user can choose whether time distance or node distance (i.e. the number of nodes intervening between the pair) should be used. If time distance has to be considered, min.dist should be a character argument containing the word "time" and then the actual time distance to be used. The same is true for node distance, but the word "node" must preced the node distance to be used. For example, if the user want to test only clades more distant than 10 time units, the argument should be "time10". If clades separated by more than 8 nodes has to be tested, the argument min.dist should be "node8". If left unspecified, it automatically searches for convergence between clades separated by a number of nodes bigger than one tenth of the tree size.

**PGLS** if the argument is set to TRUE, the function tests whether states have phylogenetic structure, by running a runs test. If the states are phylogenetically structured, a procD.pgls (package geomorph) is performed using states as the predictor variables. Pgls residuals will be used to test for convergence. Please notice, this is not suited for ultrametric trees or for variables which are not ordinal (i.e. the categories possess a natural ordering).

**nsim** number of simulations to perform sampling within the theta random distribution. It is set at 1000 by default.

**rsim** number of simulations to be performed to produce the random distribution of theta values. It is set at 1000 by default.

**clus** the proportion of clusters to be used in parallel computing.

**foldername** the path of the folder where plots are to be found.

#### **Details**

Regardless the case (either 'state' or 'clade'), the function stores a plot into the folder specified by 'foldername'. If convergence among clades is tested, the clade pair plotted corresponds to those clades with the smallest **\$average distance** from group centroid. The figure shows the Euclidean distances computed between the MRCAs of the clades and the mean Euclidean distance computed between all the tips belonging to the converging clades, as compared to the distribution of these same figures across the rest of the tree. Furthermore, the function stores the PC1/PC2 plot obtained by PCA of the species phenotypes. Convergent clades are indicated by colored convex hulls. Large colored dots represent the mean phenotypes per clade (i.e. their group centroids). Eventually, a modified traitgram plot is produced, highlighting the branches of the clades found to converge. In both PCA and traitgram, asterisks represent the ancestral phenotypes of the individual clades. If convergence among states is tested, the function produces a PC plot with colored convex hulls enclosing species belonging to different states. Furthermore, it generates circular plots of the mean angle between states (blue lines) and the range of random angles (gray shaded area). The pvalue for the convergence test is printed within the circular plots.

#### Value

If convergence between clades is tested, the function returns a list including:

- **\$node pairs**: a dataframe containing for each pair of nodes:
  - ang.bydist.tip: the mean theta angle between clades divided by the time distance.
  - ang.conv: the mean theta angle between clades plus the angle between aces, divided by the time distance.
  - ang.ace: the angle between aces.
  - ang.tip: the mean theta angle between clades.
  - nod.dist: the distance intervening between clades in terms of number of nodes.
  - time.dist: the time distance intervening between the clades.
  - p.ang.bydist: the p-value computed for ang.bydist.tip.
  - p.ang.conv: the p-value computed for ang.conv.
  - clade.size: the size of clades.
- **\$node pairs comparison**: pairwise comparison between significantly convergent pairs (all pairs if no istance of significance was found) performed on the distance from group centroids (the mean phenotype per clade).
- **\$average distance from group centroids**: smaller average distances mean less variable phenotypes within the pair.

If convergence between (or within a single state) states is tested, the function returns a dataframe including for each pair of states (or single state):

- **\$ang.state**: the mean theta angle between species belonging to different states (or within a single state).
- **\$ang.state.time**: the mean of theta angle between species belonging to different states (or within a single state) divided by time distance.

- **\$p.ang.state**: the p-value computed for ang.state.
- **\$p.ang.state.time**: the p-value computed for ang.state.time.

## **Examples**

```
# data("DataFelids")
# DataFelids$PCscoresfel->PCscoresfel
# DataFelids$treefel->treefel
# DataFelids$statefel->statefel
# RRphylo(treefel, PCscoresfel) ->RRfel
#
## searching convergence within a single state
# search.conv(tree=treefel, y=PCscoresfel, state=statefel,
# foldername = tempdir())
## searching convergence between clades by setting min.dist as node
## distance
# search.conv(RR=RRfel, y=PCscoresfel, min.dim=5, min.dist="node9",
# foldername = tempdir())
## searching convergence between clades by setting min.dist as time
# distance
# search.conv(RR=RRfel, y=PCscoresfel, min.dim=5, min.dist="time38",
# foldername = tempdir())
```

## 20. PGLS\_fossil

**Title** Phylogenetic Generalized Least Square with fossil phylogenies

# **Description**

The function performs pgls for non-ultrametric trees using either Pagel's lambda transform,

Brownian Motion or RRphylo rates to change the correlation structure.

#### **Usage**

PGLS fossil(modform,data,tree,RR=NULL)

#### **Arguments**

**modform** the formula for the regression model.

**data** a list of named vectors including response and predictor variables as named in modform.

**tree** a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous.

RR the result of RRphylo performed on the response variable. If NULL (default), the function fits Pagel's lambda in the regression for univariate data or uses the tree variance covariance matrix in the multivariate case. If RR is specified, tree branches are rescaled to the absolute branch-wise rate values calculated by RRphylo to transform the variance-covariance matrix.

#### Value

Fitted pgls parameters and significance.

#### **Details**

With univariate data, the user may want to use either Pagel's lambda or RRphylo rates to transform the correlation structure. In the former case, the lambda transform is fitted to the data (Revell, 2010). In the latter case, branch lengths are multiplied by absolute rates as computed by RRphylo to accommodate rate variation across the tree. In the multivariate case, the variance-covariance structure is either left unaltered by keeping RR = NULL (Adams and Collyer, 2015) or changed according to the norm-2 vector of rates computed for each phenotype by RRphylo.

# **Examples**

```
# library(ape)
# library(phytools)
# rtree(100) -> tree
# fastBM(tree) -> y
# fastBM(tree) -> x
# RRphylo(tree, y) -> RR
#
# PGLS_fossil(y~x,RR=RR,type="Pagel")
# PGLS_fossil(y~x,RR=RR,type="RRphylo")
# library(ape)
# library(phytools)
# rtree(100) -> tree
# fastBM(tree, nsim=3) -> resp.multi
```

```
# fastBM(tree) ->pred1
# fastBM(tree) ->pred2
# PGLS_fossil (modform=y1~x1+x2, data=list(y1=resp, x2=pred1, x1=pred2),
# tree=tree)
# RRphylo::RRphylo(tree, resp) ->RR
# PGLS_fossil (modform=y1~x1+x2, data=list(y1=resp, x2=pred1, x1=pred2),
# tree=tree, RR=RR)
#
# PGLS_fossil (modform=y1~x1+x2,
# data=list(y1=resp.multi, x2=pred1, x1=pred2), tree=tree)
# RRphylo::RRphylo(tree, resp.multi) ->RR
# PGLS_fossil (modform=y1~x1+x2,
# data=list(y1=resp.multi, x2=pred1, x1=pred2), tree=tree, RR=RR)
```

#### 21. StableTraitsR

**Title** Run StableTraits from within R

#### **Description**

This function runs StableTraits and StableTraitsSum (Elliot and Mooers 2014) from within the R environment and returns its output into the workspace.

#### **Usage**

StableTraitsR(tree,y,path,output=NULL,aces=NULL,argST=NULL,argSTS=NULL)

### **Arguments**

**tree** a phylogenetic tree. The tree needs not to be either ultrametric or fully dichotomous.

**y** a named vector of phenotypic trait.

**path** the folder path where the StableTraits output will be stored. Notice that the input tree and data (modified automatically if the original tree is not fully dichotomous or if aces are specified) will be stored in this folder as well.

**output** name of the output to be returned, if unspecified it will be named "output".

**aces** a named vector of ancestral character values at nodes specified in advance. Names correspond to the nodes in the tree.

**argST** a list of further arguments passed to StableTraits. If the argument has no value (for example "brownian") it must be specified as TRUE.

**argSTS** list of further arguments passed to StableTraitsSum. If the argument has no value (for example "brownian") it must be specified as TRUE.

#### **Details**

The StableTraits software is available at http://www.michaelelliot.net/stabletraits/, along with instructions for compilation. Once it is installed, the user must set as R working directory the folder where the StableTraits software are installed. Further information about the arguments and outputs of StableTraits and StableTraitsSum can be found at <a href="http://www.michaelelliot.net/stabletraits/">http://www.michaelelliot.net/stabletraits/</a>. StableTraitsR automatically recognizes which Operating System is running on the computer (it has been tested successfully on MacOS and Windows machines).

#### Value

The function returns a 'list' containing the output of StableTraits and StableTraitsSum.

**\$progress** a table reporting the DIC and PRSF diagnostics.

**\$rates\_tree** a copy of the original tree with branch lengths set to the evolutionary rate imputed by the stable reconstruction. Specifically, each branch length is equal to the absolute difference in the stable reconstruction occurring on that branch divided by the square root of the input branch length.

**\$rates** the original branch lengths, evolutionary rates, node height and (optionally) scaled branch lengths.

**\$aces** the median estimates of ancestral states and stable parameters along with the 95% credible interval.

**\$brownian\_tree** if "brownian" is TRUE in 'argSTS', a copy of the original tree with branch lengths set such that the Brownian motion reconstruction of the character on this tree is approximately the same as the stable ancestral reconstruction.

**\$ace.prior.values** if aces is specified, the function returns a dataframe containing the corresponding node number on the RRphylo tree for each node, the original (preset) and the estimated values, and the 95% credible interval.

#### **Examples**

```
# library(ape)
# library(phytools)
# setwd("~/Software/StableTraits")
# dir.create("Analyses")
# rtree(100) ->tree
```

```
# fastBM(tree) ->y
# c(1,2,3) ->acev
# sample(Ntip(tree) + seq(1:Nnode(tree)),3) ->names(acev)
# StableTraitsR(tree,y,path="Analyses/",output="my_output",aces=acev,
# argST=list(iterations=500000,chains=4),argSTS=list(brownian=TRUE))->ST
```

#### 22. overfitRR

**Title** *Testing RRphylo methods overfit* 

## **Description**

Testing the robustness of search.trend (Castiglione et al. 2019) and search.shift (Castiglione et al. 2018) to sampling effects and phylogenetic uncertainty.

### **Usage**

```
overfitRR(RR,y,s=0.25,trend=FALSE,shift.node=NULL,shift.state=NULL,
trend.node=NULL,aces=NULL,cov=NULL,rootV=NULL,nsim=100,clus=.5)
```

## **Arguments**

**RR** an object produced by RRphylo.

**y** a named vector of phenotypes.

**s** the percentage of tips to be cut off. It is set at 25% by default.

**trend** if TRUE, the function performs search.trend on the subsampled tree, searching for trends in either phenoypes or rates through time. If the argument trend.node is specified (see below), the 'trend' argument is automatically set to TRUE.

**shift.node** the nodes to be tested in search.shift under the 'clade' condition. Notice that the conditions on search.shift are exclusive, they cannot be tested together.

**shift.state** the named vector of states to be tested in search.shift under the 'sparse' condition. Notice that the conditions on search.shift are exclusive, they cannot be tested together.

**trend.node** the nodes to be tested by search.trend for trends in phenotypes or rates through time.

**aces** if used to produce the 'RR' object, the vector of those ancestral character values at nodes known in advance must be specified. Names correspond to the nodes in the tree.  $\mathbf{x1}$  the additional predictor to be specified if the RR object has been created using an additional predictor (i.e. multiple version of RRphylo). 'x1' vector must be as long as the number of nodes plus the number of tips of the tree, which can be obtained by running RRphylo on the predictor as well, and taking the vector of ancestral states and tip values to form the 'x1'.

aces.x1 a named vector of ancestral character values at nodes for 'x1'. It must be indicated if the RR object has been created using both aces and 'x1'. Names correspond to the nodes in the tree.

**cov** if used to produce the 'RR' object, the covariate must be specified. As in RRphylo, the covariate vector must be as long as the number of nodes plus the number of tips of the tree, which can be obtained by running RRphylo on the covariate as well, and taking the vector of ancestral states and tip values to form the covariate.

**rootV** if used to produce the 'RR' object, the phenotypic value at the tree root must be specified.

**nsim** number of simulations to be performed. It is set at 100 by default.

**clus** the proportion of clusters to be used in parallel computing.

#### **Details**

Methods using a large number of parameters risk being overfit. This usually translates in poor fitting with data and trees other than the those originally used. With RRphylo methods this risk is usually very low. However, the user can assess how robust the results got by applying search.shift or search.trend are by running overfitRR. With the latter, the original tree and data are subsampled by specifying a s parameter, that is the proportion of tips to be removed from the tree. Internally, overfitRR further shuffles the tree by using the function swapONE. Thereby, both the potential for overfit and phylogenetic uncertainty are accounted for straight away.

#### Value

The function returns a 'list' containing:

**\$rootCI** the 95% confidence interval around the root value.

**\$ace.regressions** the results of linear regression between ancestral state estimates before and after the subsampling.

**\$shift.results** if 'shift.state' is specified, a list including for each state the p-values at each simulation (\$states) and the percentage of simulations producing significant p-value (\$p.states). If the argument 'shift.node' is specified, the list contains for each node the p-values at each simulation (\$single clades) and the percentage of simulations producing

significant p-value (\$p.single.clades), and the same figures by considering all the specified nodes as evolving under a single rate (\$all.caldes.togheter and \$p.all.clades.together). **\$trend.results** if 'trend' = TRUE, a list including slopes and p-values of phenotypes versus age and absolute rates versus age regressions for the entire tree at each simulation

(\$tree.phenotype and \$tree.rates) and the percentage of simulations showing significant p-values for both variables (\$tree.p). If 'trend.node' is specified, the list also includes the same results at nodes (\$node.phenotype, \$node.rates and \$node.p) and the results for comparison between nodes (\$comparison.phenotype, \$comparison.rates and \$comparison.p).

## **Examples**

```
# data("DataOrnithodirans")
# DataOrnithodirans$treedino->treedino
# DataOrnithodirans$massdino->massdino
# DataOrnithodirans$statedino->statedino

## Extract Pterosaurs tree and data
# library(ape)
# extract.clade(treedino,748)->treeptero
# massdino[match(treeptero$tip.label,names(massdino))]->massptero
# massptero[match(treeptero$tip.label,names(massptero))]->massptero
# RRphylo(tree=treedino,y=massdino)->dinoRates
# RRphylo(tree=treeptero,y=log(massptero))->RRptero

## Case 1 search.shift under "clade" condition
# search.shift(RR=dinoRates, status.type= "clade",
```

```
# foldername=tempdir())->SSnode
# overfitRR(RR=dinoRates, y=massdino,
# shift.node=rownames(ss$single.clades))->overfit.SSnode
## Case 2 search.shift under "sparse" condition
# search.shift(dinoRates, status.type= "sparse", state=statedino,
# foldername=tempdir())->SSstate
# overfitRR(RR=dinorates, y=massdino,
# shift.state=statedino)->overfit.SSstate
## Case 3 search.trend on the entire tree
# search.trend(RR=RRptero, y=log(massptero),foldername=tempdir())->STtree
# overfitRR(RR=RRptero,y=log(massdino),trend=TRUE)->overfit.STtree
## Case 4 search.trend at specified nodes
# search.trend(RR=RRptero,y=log(massptero),node=143,
# foldername=tempdir())->STnode
# overfitRR(RR=RRptero,y=log(massdino),trend.node=143)->overfit.STnode
## Case 5 overfitRR on multiple RRphylo
# data("DataCetaceans")
# DataCetaceans$treecet->treecet
# DataCetaceans$masscet->masscet
# DataCetaceans$brainmasscet->brainmasscet
# DataCetaceans$aceMyst->aceMyst
# drop.tip(treecet,treecet$tip.label[-match(names(brainmasscet),
treecet$tip.label)])->treecet.multi
```

```
# masscet[match(treecet.multi$tip.label,names(masscet))]->masscet.multi
# RRphylo(treecet.multi,masscet.multi)->RRmass.multi
# RRmass.multi$aces[,1]->acemass.multi
# c(acemass.multi,masscet.multi)->x1.mass
# RRphylo(treecet.multi,y=brainmasscet,x1=x1.mass)->RRmulti
# search.trend(RR=RRmulti,y=brainmasscet,x1=x1.mass,clus=0.5,
foldername=tempdir())->STcet
# overfitRR(RR=RRmulti,y=brainmasscet,trend=TRUE,x1=x1.mass)-
>overfit.STcet
```

#### 23. scaleTree

**Title** *Phylogenetic tree calibration* 

### **Description**

The function is a wrapper around the functions "scalePhylo", "assign.ages", and "assign.brlen" written by Gene Hunt (http://paleobiology.si.edu/staff/individuals/hunt.cfm). It rescales tree branch lengths according to given calibration dates.

## **Usage**

```
scaleTree(tree, tip.ages, node.ages=NULL, min.branch=0.1)
```

## **Arguments**

**tree** a phylogenetic tree. The tree needs not to be ultrametric and fully dichotomous. **tip.ages** a named vector of tip ages in terms of distance from the youngest tip.

node.ages a named vector of node ages in terms of distance from the youngest tip. If no calibration date for nodes is available except for the tree root, the vector must contain as only entry the tree height with the root number as name.
min.branch the minimum branch length that will be imposed when no age is specified.

#### Value

Rescaled phylogentic tree.

# **Examples**

```
# library(ape)
# library(phytools)
# set.seed(76)
# rtree(100)->tree
#
# max(nodeHeights(tree))->H
#
# Example 1
# H-diag(vcv(tree))->tipAges
# tipAges[tips(tree,160)]<-min(tipAges)
# H->nodeAges
# names(nodeAges)<-Ntip(tree)+1
#
# scaleTree(tree,tipAges,nodeAges)->treeS1
#
```

```
# edge.col<-rep("black",nrow(tree$edge))</pre>
# edge.col[which(treeS1$edge[,2]%in%getDescendants(treeS1,160))]<-"red"</pre>
# layout(2:1)
# plot(tree,edge.color = edge.col,show.tip.label=F)
# plot(treeS1,edge.color = edge.col,show.tip.label=F)
## Example 2
\# nodeAges<-c(H, 4.5, 6, 6)
# names(nodeAges) <-c(101,143,171,121)</pre>
# tipAges<-H-diag(vcv(tree))</pre>
# scaleTree(tree, tipAges, nodeAges) ->treeS2
# edge.col<-rep("black",nrow(tree$edge))</pre>
# edge.col[which(treeS1$edge[,2]%in%c(getDescendants(treeS1,143),
#
                                       getDescendants(treeS1,171),
                                       getDescendants(treeS1,121)))]<-"red"</pre>
# layout(2:1)
# plot(tree,edge.color = edge.col,show.tip.label=F)
# plot(treeS2,edge.color = edge.col,show.tip.label=F)
```

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