Package: simboot (via r-universe)

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Type Package

Title Simultaneous Inference for Diversity Indices

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Description Provides estimation of simultaneous bootstrap and asymptotic confidence intervals for diversity indices, namely the Shannon and the Simpson index. Several pre--specified multiple comparison types are available to choose. Further user--defined contrast matrices are applicable. In addition, simboot estimates adjusted as well as unadjusted p--values for two of the three proposed bootstrap methods. Further simboot allows for comparing biological diversities of two or more groups while simultaneously testing a user-defined selection of Hill numbers of orders q, which are considered as appropriate and useful indices for measuring diversity.

License GPL (>= 2)

URL https://github.com/shearer/simboot, http://shearer.github.io/simboot/

BugReports https://github.com/shearer/simboot/issues

Depends boot, mvtnorm

LazyLoad yes

Repository https://shearer.r-universe.dev

RemoteUrl https://github.com/shearer/simboot

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Contents

`simboot-package` *Simultaneous inference for diversity indices.*

Description

Package **simboot** provides estimation of simultaneous bootstrap and asymptotic confidence intervals for diversity indices, namely the Shannon and the Simpson index. Several pre-specified multiple-comparison types are available. Further user-defined contrast matrices are applicable. In addition, **simboot** estimates adjusted as well as unadjusted p–values for two of the three proposed bootstrap methods. Further simboot allows for comparing biological diversities of two or more groups with simultaneously testing a user-defined selection of Hill numbers of orders q, which are considered appropriate and useful indices for measuring diversity.

Details

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License: GPL (>= 2)
LazyLoad: yes

Author(s)

Ralph Scherer\ Philip Pallmann\

References

Scherer, R. and Schaarschmidt, F. (2013) Simultaneous confidence intervals for comparing biodiversity indices estimated from overdispersed count data. *Biometrical Journal* 55, 246–263.

Evaluation of the methods in `sbdiv`

Pallmann, P. et al. (2012) Assessing group differences in biodiversity by simultaneously testing a user-defined selection of diversity indices. *Molecular ecology resources* 12, 1068–??78.

Evaluation of the methods in `mcpHill`

Westfall, P. H. and Young, S. S. (1993) Resampling-Based Multiple Testing: Examples and Methods for p–Value Adjustment. New York: Wiley.

Corresponding method `sbdiv` with method `WYht`

Besag, J., Green, P. J., Higdon, D., Mengersen, K. (1995) Bayesian computation and stochastic systems (with discussion) . *Statistical Science*, 10, 3–66.

Corresponding method `sbdiv` with method `rpht`

Beran, R. (1988) Balanced simultaneous confidence sets. *Journal of the American Statistical Association*, 83, 679–686.

Corresponding method `sbdiv` with method `tsht`

Fritsch, K. S., Hsu, J. C. (1999) Multiple comparison of entropies with application to dinosaur biodiversity. *Biometrics*, 55, 4, 1300–1305.

Rogers, J. A., Hsu, J. C. (2001) Multiple comparisons of biodiversity. *Biometrical Journal*, 43, 5, 617–625.

Corresponding method `sbdiv` with method `asht`

Jost, L. (2008) G(ST) and its relatives do not measure differentiation. *Molecular Ecology*, 17, 4015-4026.

Corresponding method `mcpHill`

asht *Internal function for simultaneous asymptotic intervals*

Description

Internal function for simultaneous asymptotic intervals

Note

Only internal function. Use function sbdiv instead

References

Fritsch, K. S., Hsu, J. C. (1999) Multiple comparison of entropies with application to dinosaur biodiversity. *Biometrics*, 55, 4, 1300–1305.

Rogers, J. A., Hsu, J. C. (2001) Multiple comparisons of biodiversity. *Biometrical Journal*, 43, 5, 617–625.

Bacteria *Relative Abundances of Soil Bacteria*

Description

Relative abundances of soil bacteria from 27 samples collected in nine forest and 18 grassland sites in Germany. The data set includes abundances of 18 bacterial phyla (including three candidate phyla) and five proteobacterial classes.

Usage

```
data(Bacteria)
```

Format

A data frame with 27 observations on the following 24 variables.

`Land use type` a factor with levels `forest` `grassland`

`Acidobacteria` a numeric vector

`Actinobacteria` a numeric vector

`Bacteroidetes` a numeric vector

`Chloroflexi` a numeric vector

`Cyanobacteria` a numeric vector

`Deinococcus-Thermus` a numeric vector

`Fibrobacteres` a numeric vector

`Firmicutes` a numeric vector

`Fusobacteria` a numeric vector

`Gemmatimonadetes` a numeric vector

`Nitrospira` a numeric vector

`OP11` a numeric vector

`Planctomycetes` a numeric vector

`Spirochaetes` a numeric vector

`Tenericutes` a numeric vector

`TM7` a numeric vector

`Verrucomicrobia` a numeric vector

`WS3` a numeric vector

`Alphaproteobacteria` a numeric vector

`Betaproteobacteria` a numeric vector

`Deltaproteobacteria` a numeric vector

`Gammaproteobacteria` a numeric vector

`Epsilonproteobacteria` a numeric vector

Details

Relative abundances of 18 bacterial phyla (including three candidate phyla) and five proteobacterial classes (alpha, beta, gamma, delta and epsilon) from two ecological metagenomics studies (Will et al. 2010, Nacke et al. 2011). There are 27 observations altogether, nine of which stem from forest and 18 from grassland plots in Germany.

One goal of these investigations was to unravel differences in bacterial diversity and community composition between the land use types forest and grassland.

The bacteria's relative abundances were determined by analyzing the V2-V3 region of the 16S rRNA gene via pyrosequencing-based DNA techniques.

Source

Will, C., Thuermer, A., Wollherr, A., et al. (2010) Horizon- specific bacterial community composition of German grassland soils, as revealed by pyrosequencing-based analysis of 16S rRNA genes. *Applied and Environmental Microbiology*, 76, 6751–6759.

Nacke, H., Thuermer, A., Wollherr, A., et al. (2011) Pyrosequencing- based assessment of bacterial community structure along different management types in German forest and grassland soils. *PLoS One*, 6, e17000.

Examples

```
data(Bacteria)
str(Bacteria)

### Assess whether there is a difference in biodiversity and
### community composition species richness (Shannon index,
```

```
### Simpson index) between grassland and forest.
### Bootstrap times set to 50 due to example time settings

library(simboot)
mcpHill(dataf=Bacteria[,2:24], fact=Bacteria[,1], boots=50, qval=c(0,1,2))
```

`Boutrp` *Internal function*

Description

Internal function for method `rpht` in function `sbdiv`

Note

Only for internal use.

`CCdrp` *Internal function*

Description

Internal function for method rpht in `sbdiv`

`contrMat` *Contrast Matrices*

Description

Computes contrast matrices for several multiple comparison procedures.

Usage

```
contrMat(n, type = c("Dunnett", "Tukey", "Sequen", "AVE",
                     "Changepoint", "Williams", "Marcus",
                     "McDermott", "UmbrellaWilliams", "GrandMean"),
         base = 1)
```

Arguments

`n`	a (possibly named) vector of sample sizes for each group.
`type`	type of contrast.
`base`	an integer specifying which group is considered the baseline group for Dunnett contrasts.

Details

Computes the requested matrix of contrasts for comparisons of mean levels.

Value

The matrix of contrasts with appropriate row names is returned.

Note

Function `contrMat` is adapted from package **multcomp**

References

Frank Bretz, Alan Genz and Ludwig A. Hothorn (2001), On the numerical availability of multiple comparison procedures. *Biometrical Journal*, **43**(5), 645–656.

Examples

```
n <- c(10,20,30,40)
names(n) <- paste("group", 1:4, sep="")
contrMat(n) # Dunnett is default
contrMat(n, base = 2) # use second level as baseline
contrMat(n, type = "Tukey")
contrMat(n, type = "Sequen")
contrMat(n, type = "AVE")
contrMat(n, type = "Changepoint")
contrMat(n, type = "Williams")
contrMat(n, type = "Marcus")
contrMat(n, type = "McDermott")
### Umbrella-protected Williams contrasts, i.e. a sequence of
### Williams-type contrasts with groups of higher order
### stepwise omitted
contrMat(n, type = "UmbrellaWilliams")
### comparison of each group with grand mean of all groups
contrMat(n, type = "GrandMean")
```

`corrmatgen` *Internal function.*

Description

Correlation matrix for confidence intervals assuming multivariate standard normal distribution. Calculates the correlation matrix for method asci in function `sbdiv`

Usage

```
corrmatgen(CM, varp)
```

Arguments

CM	a matrix of contrast coefficients, dimension MxI, where M=number of contrasts, and I=number of groups in a oneway layout
varp	a numeric vector of groupwise variance estimates (length = I)

Value

A matrix of dimension MxM.

---

estShannon *Estimator for Shannon's index*

Description

Estimation function for Shannon's index. Internal use in estShannonf.

Usage

```
estShannon(x)
```

Arguments

x	Vector of discrete-scaled numerical values.

Details

Estimator of Shannon-Wiener index with bias correction. Number of Species S in the bias correction does not take zeros into account.

Value

Shannon-Wiener index with bias correction

estShannonf *Estimator for Shannon's index odered by a factorial variable f.*

Description

Estimation function for Shannon's index. Internal use in sbdiv for methods rpht, tsht, asht. Sums up species counts in each columns for every treatment group and estimates Shannon's index with bias correction on the resulting vectors of summed up species counts.

$\widehat{HBC}_i = \hat{H}_i + (S_i - 1)/(2N_{i\bullet}) - (1 - \sum(1/\hat{p}_{i\bullet s}))/(12N_{i\bullet}^2) - \sum((1/\hat{p}_{i\bullet s}) - (1/(\hat{p}_{i\bullet s}^2)))/(12N_{i\bullet}^3);$

$i = 1, ..., k; s = 1, ..., S; p_{i\bullet s} = \frac{\sum_{j=1}^{n} x_{sj}}{N_{i\bullet}};$

$$\hat{H}_i = (-1)\sum_{s=1}^{S}(\hat{p}_{i\bullet s} log(\hat{p}_{i\bullet s}))$$

$N_{i\bullet} = \sum_{j=1}^{n} N_{ij}$ Number of observed individuals in treatment i.

Usage

```
estShannonf(X, f)
```

Arguments

X	n times p matrix containing species in p columns and replicates in n rows.
f	Factor variable containing treatment groups. Must be of length: replicates times treatment groups.

Value

estimate	Estimated Shannon-Wiener index for treatment groups
varest	Estimated variance of Shannon-Wiener index for treatment groups

estShannonWY *Estimator for Shannon's index row wise.*

Description

Estimation function for Shannon's index. Internal use in WYht. Calculates Shannon-Wiener index with bias correction

$$\widehat{HBC}_{ij} = \hat{H}_{ij} + (S_{ij} - 1)/(2N_{ij}) - (1 - \sum_{s=1}^{S}(1/\hat{p}_{ijs}))/(12N_{ij}^2) - \sum_{s=1}^{S}((1/\hat{p}_{ijs}) - (1/(\hat{p}_{ijs}^2)))/(12N_{ij}^3);$$

$$\hat{H}_{ij} = (-1)\sum_{s=1}^{S}(\hat{p}_{ijs} log(\hat{p}_{ijs}))$$

$i = 1, ..., k; j = 1, ..., n; s = 1, ..., S;$

$S_j =$ Number of observed species in replicate j;

$N_j =$ Number of observed individuals in replicate j

for every row in a $n \times p$ matrix.

Usage

```
estShannonWY(x)
```

Arguments

x	Vector of p numerical species counts.

Value

Shannon-Wiener index with bias correction

`estSimpson` *Estimator for Simpson's index*

Description

Estimation function for Simpson's index $1 - p^2 * n/(n-1)$. Internal use in estSimpsonf.

Usage

```
estSimpson(x)
```

Arguments

x	Vector of discrete-scaled numerical values.

Value

Estimator of Simpson's index

estSimpsonf *Estimator for Simpson's index odered by a factorial variable f.*

Description

Estimation function for Simpson's index. Internal use in sbdiv for methods rpht, tsht, asht. Sums up species counts in each columns for every treatment group and estimates Simpson's index on the resulting vectors of summed up species counts.

Usage

```
estSimpsonf(X, f)
```

Arguments

X	n times p matrix containing species in p columns and replicates in $n$rows.
f	Factor variable comtaining treatment groups. Must be of length: replicates times treatment groups.

Value

estimate	Estimated Simpson index for treatment groups
varest	Estimated variance of Simpson's index for treatment groups

estThetaRow *Internal function*

Description

Internal function for method WYht in function sbdiv. Calculates the specified diversity index for every replicated sample in each treatment group.

Usage

```
estThetaRow(X, f, theta)
```

Arguments

X	Matrix with dimension $n \times p$.
f	Factorial variable containing treatment groups.
theta	Shannon or Simpson index

mcpHill *Multiplicity-adjusted p-values for comparing biodiversity via simultaneous inference of a user-defined selection of diversity indices*

Description

The function mcpHill allows for comparing biological diversities of two or more groups. It simultaneously tests a user-defined selection of Hill numbers of orders q, which are considered appropriate and useful indices for measuring diversity (Jost 2008). As an output mcpHill gives p-values adjusted for multiplicity according to the method of Westfall & Young (1993).

Usage

```
mcpHill(dataf, fact, align = FALSE, block, boots = 5000, udmat
 = FALSE, usermat, mattype = "Dunnett", dunbase = 1, qval = seq(-1, 3),
 opt = "two.sided")
```

Arguments

dataf	Data frame containing numerical values (e.g. species counts or relative abundances). Rows represent repeated observations of the (two or more) groups, columns represent taxonomic units (usually species, or phyla, classes etc.).
fact	Vector assigning (two or more) factor levels to the observations, i.e. the groups to be compared. The length of fact must equal the number of rows in dataf.
align	Logical indicating whether a block alignment should be carried out. If TRUE, the blocks must be specified as a vector in block. Default is FALSE.
block	Vector assigning which block an observation belongs to. Only required if align=TRUE. The length of block must equal the number of rows in dataf.
boots	Number of bootstrap replications. Values lower than 999 are rejected. Default is 5000.
udmat	Logical indicating whether used-defined contrasts are applied for multiple testing. If TRUE, a contrast matrix has to be specified via usermat. Default is FALSE, meaning that the contrast matrix is specified by a catchword (e.g. "Tukey", "Dunnett" etc.).
usermat	Matrix specifying user-defined multiple testing contrasts. Only required if udmat=TRUE. The row sums in the matrix must equal zero.
mattype	Type of contrast matrix for multiple comparisons of groups. Hence only required for comparisons of more than two groups. Can be specified by the catchwords used in function contrMat (e.g. "Dunnett", "Tukey", "GrandMean", "AVE", "Williams", "Changepoint" etc.). Default is "Dunnett".
dunbase	Integer determining the factor group (in alphanumerical order) to be considered the baseline or control and therefore only needed for Dunnett-type multiple contrasts. Default is 1.
qval	Vector containing the requested selection of q-values in order to specify the Hill numbers of orders q to be investigated. Default is seq(-1,3).

opt "greater" performs an upper-tailed test, "less" a lower-tailed test and "two.sided" a two-tailed test. Default is "two.sided".

Value

The output of mcpHill is a matrix containing the chosen selection of Hill numbers (their orders q) in the first column. The multiplicity-adjusted p-values for each hypothesis tested are in the second column. The names of the rows denote which groups are being compared.

Author(s)

Philip Pallmann

References

Pallmann, P. et al. (2012) Assessing group differences in biodiversity by simultaneously testing a user-defined selection of diversity indices. Molecular ecology resources 12, 1068–78.

Jost, L. (2008) G(ST) and its relatives do not measure differentiation. Molecular Ecology, 17, 4015–4026.

Westfall, P.H. and Young S.S. (1993) Resampling-based multiple testing: examples and methods for p-value adjustment. New York: Wiley.

Examples

```
### Multiple testing with user-defined contrasts after block alignment

data(predatGM)

mymat <- rbind( "GM - S1" = c(1,-1,0,0), "GM - S2" = c(1,0,-1,0), "GM -
  S3" = c(1,0,0,-1), "S1 - S2" = c(0,1,-1,0), "S1 - S3" = c(0,1,0,-1) )

# example runs with only 100 bootstrap steps. For estimation use 2000 or more.
mcpHill(dataf=predatGM[,3:35], fact=predatGM[,2], align=TRUE,
block=predatGM[,1], boots=100, udmat=TRUE, usermat=mymat, qval=seq(-1,
3, by=0.5))

# with Dunnett-type contrast matrix
mcpHill(dataf=predatGM[,3:35], fact=predatGM[,2], align=TRUE,
block=predatGM[,1], boots=100, udmat=FALSE, mattype = "Dunnett", qval=seq(-1,
3, by=0.5))
```

---

predatGM *Abundance data of predatory insects*

Description

In a field trial with 8 complete blocks, one genetically modified crop variety and three varieties without genetical modification (S1, S2, S3) have been cultivated. Note that S1 is genetically closely related to the GM variety, and mainly differs from GM by not containing the transformation, while S2 and S3 are conventional varieties, which are genetically not closely related to GM and S1. In each of the 24 plots, a certain taxonomic group of predatory insects has been trapped. Trapped individuals have been classified to the species level. A total of 33 different species has been observed. For each plot, the summed counts of each species over one cultivation period is given in the variables Sp1, Sp2,...Sp33. Among others, one question in research was: Does the genetic modified variety effect biodiversity of the (ecologically important, non-target) species?

Usage

```
data(predatGM)
```

Format

A data frame with 32 observations on the following 35 variables.

`Block` a numeric vector, values 1,...,8 indicate the blocks of the trial

`Variety` a factor distinguishing the four varieties in the field trial, with levels `GM` (the genetically modified variety), `S1` (the near-isogenic, conventional variety), `S2` and `S3` (further conventional varieties)

`Sp1` a numeric vector, observed counts of species 1

`Sp2` a numeric vector, ...

`Sp3` a numeric vector

`Sp4` a numeric vector

`Sp5` a numeric vector

`Sp6` a numeric vector

`Sp7` a numeric vector

`Sp8` a numeric vector

`Sp9` a numeric vector

`Sp10` a numeric vector

`Sp11` a numeric vector

`Sp12` a numeric vector

`Sp13` a numeric vector

`Sp14` a numeric vector

`Sp15` a numeric vector

`Sp16` a numeric vector

`Sp17` a numeric vector

`Sp18` a numeric vector

`Sp19` a numeric vector

`Sp20` a numeric vector

`Sp21` a numeric vector

`Sp22` a numeric vector

`Sp23` a numeric vector

`Sp24` a numeric vector

`Sp25` a numeric vector

`Sp26` a numeric vector

`Sp27` a numeric vector

`Sp28` a numeric vector

`Sp29` a numeric vector

`Sp30` a numeric vector

`Sp31` a numeric vector

`Sp32` a numeric vector

`Sp33` a numeric vector

Source

Data set provided by Kai U. Priesnitz, Bavarian State Research Center for Agriculture, Institute for Plant Protection, Freising, Germany.

Examples

```
data(predatGM)

str(predatGM)

# Display data as a mosaicplot

# load("D:/Mueller/Biodiv/data/predatGM.rda")

# Matrix of counts with appropriate names
COUNTS<-as.matrix(predatGM[,3:35])
SPECNAM<-names(predatGM)[3:35]
colnames(COUNTS)<-SPECNAM
rownames(COUNTS)<-predatGM[,"Variety"]

# Assign colors and order by decreasing total abundance
COL<-grey(c(0,2,4,6,8,1,3,5,7)/8)
DMO<-COUNTS[,order(colSums(COUNTS), decreasing=TRUE)]
colnames(DMO)[15:33]<-"."

# Mosaicplot
par(mar=c(4,2,1,1))
mosaicplot(DMO, col=COL, las=2, off=15, main="", cex=1.1)
mtext("A", side=3, line=-1.5, adj=0, cex=2)
```

rpht *Internal function for simultaneous bayesian bootstrap intervals*

Description

Internal function for simultaneous bayesian bootstrap intervals

Note

Only internal function. Use function `sbdiv` instead

References

Besag, J., Green, P. J., Higdon, D., Mengersen, K. (1995) Bayesian computation and stochastic systems (with discussion) . *Statistical Science*, 10, 3–66.

saproDipGM *Abundance data of Diptera with saprophagous larvae*

Description

In a field trial with 6 complete blocks, three treatments have been applied: a genetically modified crop variety was cultivated without insecticide treatment (GM), its near-isogenic counterpart (i.e. not genetically modified but otherwise genetically closely related to the GM crop) has been cultivated without insecticide treatment (Iso), and the near-isogenic variety has been cultivated with insecticide treatment (Ins). In each of the 18 plots, two emergence traps have been placed and Diptera with saprophagous larvae were classified to the species level and counted. A total number of 25 different species has been observed and included in the present data set. For each plot, the summed counts of each species over one cultivation period (in 2002) and the two traps is given in the columns Acor, ..., Tnud. Among others, one question in this trial was: Does the genetic modified variety effect biodiversity of the (ecologically important, non-target) species in comparison to the isogenic variety (as a negative control) and in comparison to the insecticide treated plants (as a positive control)?

Usage

```
data(saproDipGM)
```

Format

A data frame with 18 observations on the following 27 variables.

`Block` a numeric vector, values 1,...,6 indicate the blocks of the trial

`Variety` a factor, distinguishing the 3 treatment levels: `GM` (genetically modified, no insecticide), `Ins` (not genetically modified, insecticide treatment) , and `Iso` (not genetically modified, no insecticide)

`Acor` a numeric vector of counts of the first species

`Arub` a numeric vector...

`Aaph` a numeric vector

`Bbre` a numeric vector

`Btri` a numeric vector

`Burt` a numeric vector

`Bvag` a numeric vector

`Bill` a numeric vector

`Ccru` a numeric vector

`Cmir` a numeric vector

`Cvag` a numeric vector

`Dnit` a numeric vector

`Dand` a numeric vector

`Lcin` a numeric vector

`Lcas` a numeric vector

`Malt` a numeric vector

`Moli` a numeric vector

`Mluc` a numeric vector

`Mtox` a numeric vector

`Ppha` a numeric vector

`Sato` a numeric vector

`Spal` a numeric vector

`Sate` a numeric vector

`Sleu` a numeric vector

`Tnud` a numeric vector

Source

Data set provided by Dr. Sabine Prescher,Institute for Biosafety of Genetically Modified Plants, Julius-Kuehn-Institut, Braunschweig, Germany

Examples

```
data(saproDipGM)

str(saproDipGM)

# load("D:/Mueller/Biodiv/data/saproDipGM.rda")

# Display data as a mosaicplot

# Matrix of counts with appropriate names
```

```
COUNTS<-as.matrix(saproDipGM[,3:27])
SPECNAM<-names(saproDipGM)[3:27]
colnames(COUNTS)<-SPECNAM
rownames(COUNTS)<-saproDipGM[,"Variety"]

# Assign colors and order by decreasing total abundance
COL<-grey(c(0,2,4,6,8,1,3,5,7)/8)
DMO<-COUNTS[,order(colSums(COUNTS), decreasing=TRUE)]

# Mosaicplot
par(mar=c(4,2,1,1))
mosaicplot(DMO, col=COL, las=2, off=15, main="", cex=1.1)
mtext("A", side=3, line=-1.5, adj=0, cex=2)
```

---

sbdiv *Perform simultaneous confidence intervals or adjusted p–values for the Shannon and the Simpson index.*

Description

Function sbdiv estimates simultaneous confidence intervals for the Shannon or the Simpson index. This function provides calculation of several pre–defined contrasts for confidence intervals.Further self-defined contrast are applicable. Simultaneous resampling confidence intervals are estimated according to the Algorithm of Besag et al. (1995) using method rpht, Westfall et al. (1993) using method WYht or similar to Beran (1988) using method tsht. Further estimation of simultaneous asymptotic intervals adjusting for heterogeneous variances is provided by method asht according to Fritsch and Hsu (1999) and Rogers and Hsu (2001). However, estimation of asymptotic intervals may make no sense in data sets with replicated samples due to overdispersion.

Usage

```
sbdiv(X, f, theta = c("Shannon", "Simpson"),
type = c("Dunnett", "Tukey", "Sequen", "AVE",
                    "Changepoint", "Williams", "Marcus",
                    "McDermott", "UmbrellaWilliams", "GrandMean"),
cmat = NULL, method = c("WYht", "tsht", "rpht", "asht"), conf.level =
0.95, alternative = c("two.sided", "less", "greater"), R = 2000, base =
1, ...)
```

Arguments

X	Data frame containing numerical values for counts in columns. Every column represents on species.
f	Vector of factorial variables for treatment groups. Vector length must be equal to the length of treatment groups multiplicated with sample replications.
theta	Biodiversity index. Options are Shannon and Simpson index.

type	Type of comparison. Options are Dunnett, Tukey, Sequen, AVE, Changepoint, Williams, Marcus, McDermott, UmbrellaWilliams, GrandMean intervals. We tested only Dunnett and Tukey contrasts in simulations.
cmat	Optional self-defined contrast matrix. In case of using this argument, the type argument is not considered.
method	Possible methods are simultaneous bootstrap confidence intervals: WYht, tsht, rpht and asymptotic simultaneous confidence intervals: asht. Adjusted and unadjusted p–values are estimated with method WYht and method tsht.
conf.level	Pre-defined overall confidence level. Default is 0.95, while two-sided inference is estimated with $(1 - conf.level)/2$ for each side and one-sided inference is estimated with $1 - conf.level$ for the side of interest.
alternative	Specified type of interval. Could be "one-sided" or "two.sided".
R	Number of bootstrap steps. Default is 2000, which is a good compromise between accuracy and computing time
base	Control group. base = 1 uses the first group in alphabetical order.
...	Further optional arguments for the internal used function boot from package **boot**. Most importantly, the number of Bootstrap samples can be chosen via the parameter R (default is R=2000); see ?boot for further options.

Details

sbdiv is the main function for estimating the different multiplicity adjusted confidence intervals. Different methods are called from internal functions.

Value

conf.int	estimate: Estimated difference between groups. Estimators differ between the methods due to calculation. lower: Lower bounds of estimated intervals. upper: Upper bounds of estimated intervals.
p.value	adj. p: multiplicity adjusted p-values. raw p: unadjusted p-values
conf.level	Pre-specified confidence level
alternative	Pre-specified alternative

Author(s)

Ralph Scherer

References

Scherer, R. and Schaarschmidt, F. (2013) Simultaneous confidence intervals for comparing biodiversity indices estimated from overdispersed count data. *Biometrical Journal* 55, 246–263.

Evaluation of the methods in sbdiv

Westfall, P. H. and Young, S. S. (1993) Resampling-Based Multiple Testing: Examples and Methods for p–Value Adjustment. New York: Wiley.

Corresponding method sbdiv with method WYht

Besag, J., Green, P. J., Higdon, D., Mengersen, K. (1995) Bayesian computation and stochastic systems (with discussion) . *Statistical Science*, 10, 3–66.

Corresponding method sbdiv with method rpht

Beran, R. (1988) Balanced simultaneous confidence sets. *Journal of the American Statistical Association*, 83, 679–686.

Corresponding method sbdiv with method tsht

Fritsch, K. S., Hsu, J. C. (1999) Multiple comparison of entropies with application to dinosaur biodiversity. *Biometrics*, 55, 4, 1300–1305.

Rogers, J. A., Hsu, J. C. (2001) Multiple comparisons of biodiversity. *Biometrical Journal*, 43, 5, 617–625.

Corresponding method sbdiv with method asht

Examples

```
## For plots of the datasets see the help files for the data sets.

## First dataset
data(predatGM)

## structure of data
str(predatGM)

## remove block variable
datspec_1 <- predatGM[, -1]
str(datspec_1)

## Order of factorial variable
datspec_1$Variety

## argument base = 1 uses GM as control group. Not directly executable
## due to intensive computing time
# sbdiv(X = datspec_1[, 2:length(datspec_1)], f = datspec_1[, 1], theta =
# "Shannon", type = "Dunnett", method = "WYht", conf.level = 0.95,
# alternative = "two.sided", R = 2000, base = 1)

## Directly executable but senseless value for boot steps R
sbdiv(X = datspec_1[, 2:length(datspec_1)], f = datspec_1[, 1], theta =
"Shannon", type = "Dunnett", method = "WYht", conf.level = 0.95,
alternative = "two.sided", R = 100, base = 1)


## Second dataset
data(saproDipGM)

## structure
str(saproDipGM)

## remove block variable
datspec_2 <- saproDipGM[, -1]
str(datspec_2)
```

```
## Order of factor variable
datspec_2$Variety

## argument base = 2 uses Ins as control group. Not directly executable
## due to intensive computing time
# sbdiv(X = datspec_2[, 2:length(datspec_2)], f = datspec_2[, 1], theta =
# "Shannon", type = "Dunnett", method = "rpht", conf.level = 0.95,
# alternative = "two.sided", R = 2000, base = 2)

## Directly executable but senseless value for boot steps R
sbdiv(X = datspec_2[, 2:length(datspec_2)], f = datspec_2[, 1], theta =
"Shannon", type = "Dunnett", method = "rpht", conf.level = 0.95,
alternative = "two.sided", R = 100, base = 2)
```

`SCIrp` *Internal function*

Description

Interval estimation in method rpci in function sbci

Note

Internal function. Use `sbdiv` instead.

`Simpson` *Internal function for Simpson estimator*

Description

Calculates Simpson's index on probability vector p

Usage

```
Simpson(p)
```

Arguments

`p`	Probability vector x_s/n

Value

Simpson's index

Note

Only for internal use

tsht *Internal function for simultaenous bootstrap intervals*

Description

Internal function for simultaenous bootstrap intervals based on summed up counts for every species.

Note

Only internal function. Use function sbdiv instead

References

Beran, R. (1988) Balanced simultaneous confidence sets. *Journal of the American Statistical Association*, 83, 679–686.

waldci *Internal function for Wald intervals*

Description

Internal function for wald intervals in method asht in function sbdiv

Note

Internal function. Use function sbdiv instead.

WYht *Internal function for simultaneous bootstrap confidence intervals*

Description

Internal function for simultaneous bootstrap confidence intervals based on resampled residuals

Note

Only internal function. Use function sbdiv instead

References

Westfall, P. H. and Young, S. S. (1993) Resampling-Based Multiple Testing: Examples and Methods for p–Value Adjustment. New York: Wiley.

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