# Package 'DeadCanMove'

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Title Assess how spatial roadkill patterns change with temporal

Type Package

sampling scheme	
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Description  From a baseline data frame of dead individuals recorded daily at different road stretches, simulate varying sub-sampling schemes, calculate roadkill hotspots based on each scheme, and compare hotspot patterns between sampling schemes and the baseline data.  License GPL-3	
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Dead Can Move-package Assess how spatial roadkill patterns change with temporal sampling scheme

## **Description**

From a baseline data frame of dead individuals recorded daily at different road stretches, simulate varying sub-sampling schemes, calculate roadkill hotspots based on each scheme, and compare hotspot patterns between sampling schemes and the baseline data.

#### **Details**

Package: DeadCanMove

Type: Package Version: 0.2

Date: 2015-05-8 License: GPL-3

## Author(s)

Barbosa A.M., Marques J.T., Santos S.M., Lourenco A., Medinas D., Beja P., Mira A.

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#### References

Santos S.M., Marques J.T., Lourenco A., Medinas D., Barbosa A.M., Beja P., Mira A. (under revision) Sampling effects on the identification of roadkill hotspots and implications for survey design and mitigation

#### **Examples**

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binary.comp.methods Binary comparison methods

## Description

This function provides the methods implemented for calculating binary similarity between hotspots obtained from different sampling schemes and those obtained from the baseline (reference) sampling scheme. It is used by functions binary.comparison, sequential.corr and hotspots.comparison.

## Usage

binary.comp.methods()

#### Value

A character vector naming the methods implemented.

#### Author(s)

A. Marcia Barbosa

#### See Also

binary.comparison, sequential.corr

## Examples

binary.comp.methods()

binary.comparison

Binary comparison

## **Description**

Compares two binary vectors using the coefficient specified in method.

## Usage

binary.comparison(x, y, method)

#### **Arguments**

x a binary (0-1) vector

y a binary (0-1) vector to compare with x

method the comparison measure to use. Current options are "Phi", "Mathews", "Yule",

"Jaccard", "Baroni", "kappa", "CCR", "TSS", "gain", "loss", and "balance".

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#### Value

A numeric value indicating the cofficient of association specified in method.

#### Author(s)

A. Marcia Barbosa

#### See Also

binary.comp.methods

## **Examples**

```
\begin{array}{l} bin1 <- \ sample(c(0,\,1),\,100,\,replace = TRUE) \\ bin2 <- \ sample(c(0,\,1),\,100,\,replace = TRUE) \\ \\ binary.comparison(bin1,\,bin2,\,method = "Phi") \\ binary.comparison(bin1,\,bin2,\,method = "kappa") \\ \end{array}
```

 ${\it getBoxplots}$ 

Get boxplots

## Description

Get boxplots

#### Usage

```
getBoxplots(corrs.list, ...)
```

## Arguments

corrs.list a list of corrs.tables given by function schemeCorrs additional arguments for the boxplot function (e.g. ylim = c(0, 1), las = 2, main = as.character(bquote(corrs.list)))

#### Value

Box plots

## Author(s)

A. Marcia Barbosa

#### See Also

schemeCorrs

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hotspot.numbers

Hotspot numbers

#### **Description**

This function calculates the total numbers of events and the hotspot thresholds for each group and sampling interval.

## Usage

```
hotspot.numbers(hotspots.list, sampl.intervals, groups, include.all.together = TRUE, min.total.events = 0, min.hotspot.threshold = 2)
```

## **Arguments**

hotspots.list results of the sequential.hotspots function

sampl.intervals integer vector of the sampling intervals to analyse (at the moment, these intervals

must be consecutive and start with one)

groups taxa or groups to analyse separately (e.g. as.character(unique(dataset\$group)))

include.all.together

logical, whether to run the analysis also for all groups combined

min.total.events minimum total number of events to calculate hotspots for a group

min.hotspot.threshold

minimum number of events for a region to be considered a hotspot

#### Value

A list of the following matrices:

N.events

HS.threshold

N.hotspots

events.in.HS

## Author(s)

A. Marcia Barbosa, J. Tiago Marques, Sara M. Santos

## See Also

hotspots, sequential.hotspots

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#### **Examples**

```
\label{eq:condition} $$\operatorname{submats} < -\operatorname{sequential.submatrix}(\operatorname{dataset} = \operatorname{roadkills}, \\ \operatorname{sampl.columns} = 4:\operatorname{ncol}(\operatorname{roadkills}), \\ \operatorname{window.size} = 1, \\ \operatorname{gap.size} = 0:2, \\ \operatorname{group.column} = "\operatorname{taxon}", \\ \operatorname{include.all.together} = \operatorname{TRUE}, \\ \operatorname{remove.zeros} = \operatorname{TRUE}, \\ \operatorname{keep.nonsampl.columns} = \operatorname{TRUE}, \\ \operatorname{n.subsampl.columns} = 120) \\ \\ \operatorname{hsl} < -\operatorname{sequential.hotspots}(\operatorname{dataset} = \operatorname{roadkills}, \\ \operatorname{submats} = \operatorname{submats}, \\ \operatorname{region.column} = "\operatorname{segment}", \\ \operatorname{first.subsampl.col} = 4, \\ \operatorname{confidence} = 0.95) \\ \\ \operatorname{hsn} < -\operatorname{hotspot.numbers}(\operatorname{hotspots.list} = \operatorname{hsl}, \\ \operatorname{sampl.intervals} = 1:3, \\ \\ \operatorname{groups} = \operatorname{as.character}(\operatorname{unique}(\operatorname{roadkills\$taxon})), \\ \operatorname{include.all.together} = \operatorname{TRUE}, \\ \\ \operatorname{min.hotspot.threshold} = 2) \\ \\ \operatorname{hsn}
```

hotspots

Calculate roadkill hotspots

#### **Description**

This function identifies the hotspot regions in a dataset, or in a submatrix compared to the total dataset, using an adaptation of the method of Malo et al. (2004).

## Usage

```
hotspots(dataset, submat = NULL, region.column, subsampl.columns, hotspots = TRUE, confidence = 0.95, min.total.events = 0, min.hotspot.threshold = 2)
```

#### **Arguments**

dataset name of the matrix or dataframe containing the complete data

submat name of the matrix or dataframe containing the data of the group and sampling

window/gap for which to calculate hotspots

region.column name or index number of the column containing the regions (road sectors, sites)

to classify as hotspots or non-hotspots

subsampl.columns

index numbers of the consecutive columns of submat (or, if there is no submat,

of the dataset) containing the (daily) sampling data, e.g. 4:180

hotspots logical, whether to calculate the hotspots confidence confidence threshold to consider hotspots

min.total.events minimum total number of events to calculate hotspots. Not totally implemented

yet!

min.hotspot.threshold

minimum number of events for a region to be considered a hotspot. If the Malo method says that regions with less than this value are hotspots, the value returned is NA. The default threshold is 2.

hotspots.comparison 7

#### Value

A list with elements threshold (an integer value indicating the number of deaths obtained as a threshold for considering a site a roadkill hotspot) and hotspots (a data frame showing the total number of deaths per region and whether or not it was considered a hospot.)

#### Author(s)

A. Marcia Barbosa, J. Tiago Marques, Sara M. Santos

#### References

Malo, J.E., Suarez, F., Diez, A. (2004) Can we mitigate animal-vehicle accidents using predictive models? J. Appl. Ecol. 41, 701-710 (doi: 10.1111/j.0021-8901.2004.00929.x)

#### See Also

```
sequential.hotspots
```

## Examples

```
\label{eq:data} $\operatorname{data}(\operatorname{roadkills})$$ hs <- hotspots(\operatorname{dataset} = \operatorname{roadkills}, \operatorname{submat} = \operatorname{NULL}, \operatorname{region.column} = "\operatorname{segment}", \\ \operatorname{subsampl.columns} = 4:\operatorname{ncol}(\operatorname{roadkills}), \operatorname{confidence} = 0.95)$$ hs
```

hotspots.comparison

Hotspots comparison

#### **Description**

This is a wrapper for most of the functions in this package (one function to rule them all). You'll probably only need to use this one, which in turn calls each of the other functions and does all the calculations in one step.

## Usage

```
hotspots.comparison(dataset, sampl.columns, sampl.intervals, region.column, group.column, include.all.together = TRUE, confidence = 0.95, min.total.events = 80, min.hotspot.threshold = 2, comp.method = "Phi", plot = TRUE, sep.plots = FALSE, omit.baseline.interval = TRUE, ...)
```

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#### **Arguments**

dataset name of the matrix or dataframe to analyze sampl.columns index numbers of the columns containing the (daily) sampling data, e.g. 4:180 intervals at which to extract sampling data, e.g. 1:30; currently must be consecsampl.intervals utive and start with 1 name or index number of the column containing the regions (road segments, region.column sites) to classify as hotspots or non-hotspots group.column name or index number of the column containing the taxa or groups to analyse separately, e.g. 3 or "Family"; if NULL, all records will be used together include.all.together logical, whether to get subsampling matrices also for the complete data (including all groups combined) confidence confidence threshold to consider hotspots (see Malo et al. 2004); defaults to 0.95 min.total.events minimum total number of events (e.g. deaths) to calculate hotspots for a group min.hotspot.threshold minimum number of events for a region to be considered a hotspot the method with which to compare the hotspots obtained with increasing sampl.intervals comp.method with those of the baseline scenario; type binary.comp.methods() for available options logical, whether to plot the correlations between subsamples and baseline for plot each group (may cause function to fail if sep. plots = FALSE and figure margins are too large for the number of resulting plots) sep.plots logical, whether to present the plots in separate windows rather than all in the same window omit.baseline.interval logical, whether to omit the first column (correlation of baseline hotspots with themselves) from calculations and results

additional arguments to pass to the plot function

## Value

#### A list with 9 elements:

hotspots.list

N.events

HS.threshold

N.hotspots

events.in.HS

event.corrs

event.loss

event.gain

event.balance

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#### Author(s)

A. Marcia Barbosa, J. Tiago Marques, Sara M. Santos

#### References

Malo, J.E., Suarez, F., Diez, A. (2004) Can we mitigate animal-vehicle accidents using predictive models? J. Appl. Ecol. 41, 701-710 (doi: 10.1111/j.0021-8901.2004.00929.x)

#### See Also

hotspots

## **Examples**

```
data(roadkills)
```

```
\label{eq:comparison} $$hc <-$hotspots.comparison(dataset = roadkills, sampl.columns = 4:ncol(roadkills), sampl.intervals = 1:5, region.column = "segment", group.column = "taxon", include.all.together = TRUE, confidence = 0.95, min.total.events = 80, min.hotspot.threshold = 2, comp.method = "Phi", plot = TRUE, sep.plots = FALSE, omit.baseline.interval = TRUE, ylim = c(0, 1))
```

jumping.window

Jumping window

## **Description**

This function extracts a moving (a.k.a. running, rolling, sliding) window but with no overlap between windows and with the option for gaps between windows.

#### Usage

```
jumping.window(sampl.columns, window.size, gap.size, start.column = 1, J = FALSE)
```

## Arguments

sampl.columns	index numbers of the consecutive columns with the sampling data (e.g. 3:180) from which to extract the jumping windows. Can also be any vector from which to extract a jumpting window.
${\it window.size}$	size of each sampling window/season (consecutive sampling days each time)
gap.size	size of the gap between sampling windows. Can be zero or a positive integer.
start.column	column of sampl.columns where to actually start the sampling windows. The default is 1, but e.g. with a gap size of 1 between windows, the start column can be either 1 or 2.
J	logical, whether to provide the results in the form of J for function ${\it carcass}$ ::etterson. Defaults to FALSE.

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#### **Details**

This function is now integrated within submatrix to provide for additional sampling scheme options

#### Value

This function returns a vector containing the elements of sampl.columns that are included in the extracted windows; elements falling within the gaps are left out.

## Author(s)

A. Marcia Barbosa, J. Tiago Marques

## **Examples**

```
\label{eq:coadkills} $$ names(roadkills)$ $$ names(roadkills)$ jumping.window(4:ncol(roadkills), window.size = 1, gap.size = 0)$ $$ jumping.window(4:ncol(roadkills), window.size = 1, gap.size = 1)$ $$ w3g5 <- jumping.window(4:ncol(roadkills), window.size = 3, gap.size = 5)$ $$ w3g5$ $$ sm <- submatrix(dataset = roadkills, sampl.columns = w3g5, sampl.interval = 1, group.column = "taxon", group.names = NULL, remove.zeros = TRUE, keep.nonsampl.columns = TRUE)$ $$
```

plotEventCorrs

Plot correlations between events in each subsampling dataset and the baseline dataset

## Description

This function plots the correlation with baseline against sampling interval for each group

## Usage

```
plotEventCorrs(event.corrs, sep.plots = FALSE, ...)
```

#### **Arguments**

event.corrs a matrix of correlations resulting from the sequential.corr function sep.plots logical, whether to place each plot in a separate window additional arguments to pass to the plot function

roadkills 11

#### Author(s)

A. Marcia Barbosa

#### See Also

plot, binary.comparison, sequential.corr

#### **Examples**

```
\label{eq:data} $$\operatorname{data}(\operatorname{roadkills})$$ submats <- \operatorname{sequential.submatrix}(\operatorname{dataset} = \operatorname{roadkills}, \\ \operatorname{sampl.columns} = 4 : \operatorname{ncol}(\operatorname{roadkills}), \\ \operatorname{window.sizes} = 1, \\ \operatorname{gap.sizes} = 0 : 2, \\ \operatorname{group.column} = "\operatorname{taxon}", \\ \operatorname{include.all.together} = \operatorname{TRUE}, \\ \operatorname{remove.zeros} = \operatorname{TRUE}, \\ \operatorname{keep.nonsampl.columns} = \operatorname{TRUE}, \\ \operatorname{n.subsampl.columns} = 80)$$ $$ hsl <- \operatorname{sequential.hotspots}(\operatorname{dataset} = \operatorname{roadkills}, \\ \operatorname{submats} = \operatorname{submats}, \\ \operatorname{region.column} = "\operatorname{segment}", \\ \operatorname{first.subsampl.col} = 4, \\ \operatorname{confidence} = 0.95)$$ $$ hsn <- \\ \operatorname{hotspot.numbers}(\operatorname{hotspots.list} = \operatorname{hsl}, \\ \operatorname{sampl.intervals} = 1 : 3, \\ \operatorname{groups} = \operatorname{as.character}(\operatorname{unique}(\operatorname{roadkills\$taxon})), \\ \operatorname{include.all.together} = \operatorname{TRUE}, \\ \operatorname{min.total.events} = 0, \\ \operatorname{min.hotspot.threshold} = 2)$$ $$ \operatorname{seqcorr} <- \\ \operatorname{sequential.corr}(\operatorname{hotspots.list} = \operatorname{hsl}, \\ \operatorname{hotspots.thresholds} = \operatorname{hsn\$HS.threshold}, \\ \operatorname{comp.method} = "\operatorname{Phi"}, \\ \operatorname{baseline.gap} = 0, \\ \operatorname{messages} = "\operatorname{TRUE}")$$ $$ plotEventCorrs}(\operatorname{event.corrs} = \operatorname{seqcorr}, \\ \operatorname{sep.plots} = \operatorname{FALSE}, \\ \operatorname{ylim} = \operatorname{c}(0, 1), \\ \operatorname{pch} = 20)$$ $$
```

roadkills

Imaginary roadkill data

## **Description**

An imaginary dataset of roadkill data for 5 "taxonomic" groups.

#### Usage

```
data(roadkills)
```

#### **Format**

A data frame with 900 observations on the following variables:

individ an integer vector attributing an identifier to each recorded individual segment a numeric vector identifying the road segment at which each individual was recorded group a character vector indicating the "taxonomic" group to which each individual belongs day1 a numeric vector indicating whether the individual was found (1) or not (0) on that sampling day (the same for all remaining days in the data frame)

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#### **Details**

Each row corresponds to an individual recorded at a particular road stretch (segment), with a 1 if it was present and a 0 if it was not present at that segment on each of the sampling days. Individuals were not manually removed from the road, so each individual has value 1 in all days in which its body was detected on the road.

#### Source

Freely modified from data collected by: Santos S.M., Carvalho F., Mira A. (2011) How long do the dead survive on the road? Carcass persistence probability and implications for road-kill monitoring surveys. PLoS ONE 6(9), e25383 (doi:10.1371/journal.pone.0025383)

## **Examples**

```
data(roadkills)
roadkills[1:20, 1:10]
```

schemeCorrs

Scheme correlations

#### **Description**

Get correlation between each sampling scheme and the corresponding baseline

#### Usage

schemeCorrs(dataset, submats, submats.baseline, region.column, group.column, first.subsampl.col)

## Arguments

dataset

name of the matrix or dataframe containing the complete data submats a list of the submatrices for which to calculate the correlation (result of the sequential.submatrix function) submats.baseline a list of the submatrices corresponding to the baseline sampling scheme for each group region.column name or index number of the column containing the regions (road segments, sites) to classify as hotspots or non-hotspots group.column name or index number of the column containing the taxonomic groups

first.subsampl.col

index number of the first column containing subsampling data

#### Value

This function returns a list of corrs.tables.

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## Author(s)

A. Marcia Barbosa

#### See Also

getBoxplots

sequential.corr

Sequential correlation

## **Description**

This function calculates the correlation between the hotspots obtained from each submatrix and those of the baseline (sub)matrix of the corresponding group.

#### Usage

```
sequential.corr(hotspots.list, hotspots.thresholds, comp.method = "Phi", baseline.interval = 1, baseline.gap = 0, messages = "TRUE")
```

#### **Arguments**

hotspots.list a list of hotspot tables resulting from the sequential.hotspots function hotspots.thresholds

a matrix of hotspots thresholds (element 2 of the results of the hotspot.numbers

function)

comp.method characer value indicating the correlation coefficient to use; type binary.comp.methods()

for available options

baseline.interval the sampling interval with which to correlate all the other sampling intervals for

each group; defaults to 1 (take every sample)

baseline.gap the sampling gap with which to correlate all other sampling schemes for each

group; defaults to 0 (no gap between samples)

messages logical, whether to display messages

#### Value

A matrix of correlations (or whatever index was defined in method) between the hotspots obtained for each group and sampling scheme, and the hotspots obtained from the baseline data for the group under analysis.

#### Author(s)

A. Marcia Barbosa

#### See Also

binary.comparison

14 sequential.estimateN

#### **Examples**

```
\label{eq:data} $\operatorname{data}(\operatorname{roadkills})$ submats <- sequential.submatrix(dataset = \operatorname{roadkills}, sampl.columns = 4:\operatorname{ncol}(\operatorname{roadkills}), \operatorname{window.size} = 1, \operatorname{gap.size} = \operatorname{c}(0,\ 1,\ 2), \operatorname{group.column} = "\operatorname{taxon}", \operatorname{include.all.together} = \operatorname{TRUE}, \operatorname{remove.zeros} = \operatorname{TRUE}, \operatorname{keep.nonsampl.columns} = \operatorname{TRUE}, \operatorname{n.subsampl.columns} = 120) $$ hsl <- sequential.hotspots(dataset = \operatorname{roadkills}, \operatorname{submats} = \operatorname{submats}, \operatorname{region.column} = "\operatorname{segment}", \operatorname{first.subsampl.col} = 4, \operatorname{confidence} = 0.95) $$ hsn <- \operatorname{hotspot.numbers}(\operatorname{hotspots.list} = \operatorname{hsl}, \operatorname{sampl.intervals} = 1:3, \operatorname{groups} = \operatorname{as.character}(\operatorname{unique}(\operatorname{roadkills\$taxon})), \operatorname{include.all.together} = \operatorname{TRUE}, \operatorname{min.total.events} = 0, \operatorname{min.hotspot.threshold} = 2) $$$ seqcorr <- \operatorname{sequential.corr}(\operatorname{hotspots.list} = \operatorname{hsl}, \operatorname{hotspots.thresholds} = \operatorname{hsn\$HS.threshold}, \operatorname{comp.method} = "\operatorname{Phi}", \operatorname{baseline.gap} = 0, \operatorname{messages} = "\operatorname{TRUE}") $$$$ seqcorr $$
```

sequential.estimateN

Sequential estimate N

## Description

This function estimates the actual numbers of animal casualties given the observed numbers and a set of estimators, sequentially for all given submats. Requires package **carcass**.

#### Usage

```
sequential.estimateN(submats, submats.N, first.subsampl.col, region.column, persist, effic, estimators = c("korner", "huso", "erickson", "etterson"), margin = 0.01, ...)
```

#### **Arguments**

 $\begin{array}{ll} submats & result of the \ sequential. submatrix \ function. \\ submats. N & result of the \ sequential. Nevents \ function. \end{array}$ 

first.subsampl.col

index number of the first column containing the (sub)sampling data in submats

region.column name or index number of the column containing the regions (road segments,

sites) to classify as hotspots or non-hotspots

persist named vector of persistence per group; group names must match those in the

data

effic named vector of detection efficiency per group; group names must match those

in the data

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estimators character vector of the estimator(s) to use. The default is all estimators available.

margin numeric value to subtract from and add to the estimator(s) to get p.lower and

p.upper, respectively, when using function estimateN in package **carcass** 

... currently not in use

#### Value

This function returns a list.

#### Author(s)

A. Marcia Barbosa, J. Tiago Marques, Sara Santos

#### See Also

function estimateN in package carcass

sequential.hotspots

Calculate roadkill hotspots for a series of (sub)sampling datasets

#### **Description**

This function applies hotspots sequencially to a given set of submatrices to identify the hotspot regions in each dataset, using an adaptation of the method of Malo et al. (2004).

#### Usage

 $sequential.hotspots(dataset,\,submats,\,region.column,\,first.subsampl.col,\,confidence=0.95)$ 

#### **Arguments**

dataset name of the matrix or dataframe containing the complete data

submats a list of the submatrices for which to calculate the hotspots (result of the sequential.submatrix

function)

region column name or index number of the column containing the regions (road segments,

sites) to classify as hotspots or non-hotspots

first.subsampl.col

index number of the first column containing subsampling data

confidence confidence threshold to consider hotspots. The default is 0.95

#### Value

A list of 2 elements:

hotspots.thresholds

A named integer vector

hotspots.maps A list of data frames, each showing the total number of events (deaths) per region

and whether or not it was considered a hospot.

16 sequential.Nevents

#### Author(s)

A. Marcia Barbosa

#### References

Malo, J.E., Suarez, F., Diez, A. (2004) Can we mitigate animal-vehicle accidents using predictive models? J. Appl. Ecol. 41, 701-710 (doi: 10.1111/j.0021-8901.2004.00929.x)

#### See Also

hotspots

#### **Examples**

```
data(roadkills)

submats <- sequential.submatrix(dataset = roadkills,
sampl.columns = 4:ncol(roadkills), window.sizes = 1, gap.sizes = 1:3,
group.column = "taxon", include.all.together = TRUE, remove.zeros = TRUE,
keep.nonsampl.columns = TRUE, n.subsampl.columns = 85)

shs <- sequential.hotspots(dataset = roadkills, submats = submats,
region.column = "segment", first.subsampl.col = 4, confidence = 0.95)

shs
str(shs)
```

sequential. Nevents

Sequential numbers of events

#### **Description**

Applies function link{hotspots} (with hotspots=FALSE) sequentially to a given set of submatrices

### Usage

```
sequential.Nevents(dataset, submats, region.column, first.subsampl.col, estimate = FALSE)
```

## **Arguments**

dataset name of the matrix or dataframe containing the complete data

submats a list of the submatrices for which to calculate the hotspots (result of the sequen-

tial.submatrix function)

region.column name or index number of the column containing the regions (road sectors, sites)

to classify as hotspots or non-hotspots

first.subsampl.col

index number of the first column containing subsampling data

estimate logical, whether to add estimates from package carcass

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#### Author(s)

A. Marcia Barbosa

#### See Also

link{hotspots}, link{sequential.hotspots}

sequential.submatrix

Extract subsampling matrices for a series of subsampling schemes

## Description

This function applies submatrix sequencially to all specified sampling schemes and taxonomic groups.

## Usage

```
sequential.submatrix (dataset, sampl.columns, sampl.intervals = NULL,\\ window.sizes = NULL, gap.sizes = NULL, start.columns = 1, all.combinations = TRUE,\\ group.column = NULL, include.all.together = TRUE, remove.zeros = TRUE,\\ keep.nonsampl.columns = TRUE, n.subsampl.columns = NULL)
```

## **Arguments**

dataset	name of the matrix or dataframe to analyze		
$_{ m sampl.columns}$	numbers of the consecutive columns with the (daily) sampling data, e.g. 4:180		
sampl.intervals	a vector of the intervals at which to extract sampling data, e.g. 5 to take one every five samples		
window.sizes	the size (in sampling time units, e.g. days) of the sampling periods		
gap.sizes	the size of the gaps between sampling periods		
start.columns	vector of start columns, see submatrix		
all.combinations	logical, whether to use all window x gap size combinations (the default, TRUE) or just the number corresponding to the length of window.sizes and gap sizes (in which case window.sizes and gap.sizes must have the same length)		
group.column	name or index number of the column containing the taxa or groups to analyse separately, e.g. 3 or "Family"; if NULL, all records will be used together		
include.all.together			
	logical, whether to get subsampling matrices also for the complete data (including all groups together)		
remove.zeros	logical, whether to remove rows where all extracted samples have zero observations		
keep.nonsampl.columns			
	logical, whether to include also the non-sampling columns of dataset in the resulting submatrices)		
${\it n.subsampl.columns}$			

number of subsampl.columns

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#### Value

A list of submatrices

#### Author(s)

A. Marcia Barbosa

#### See Also

```
submatrix, subset
```

### **Examples**

submatrix

Extract a submatrix for a given taxomic group and/or sampling scheme

## **Description**

Given a baseline dataset, this function extracts a sub-dataset for a given taxomic group and/or sampling scheme, defined either by a sampling interval (periodicity) or by a window size (consecutive sampling days each time) and a gap size (gaps between sampling windows).

#### Usage

```
submatrix(dataset, sampl.columns, sampl.interval = NULL, window.size = NULL, \\ gap.size = NULL, start.column = 1, group.column = NULL, group.names = NULL, \\ remove.zeros = TRUE, keep.nonsampl.columns = TRUE)
```

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#### **Arguments**

dataset	name of the matrix or dataframe to analyze	
sampl.columns	index numbers of the (consecutive) columns containing the baseline (daily) sampling data, e.g. $3:180$	
sampl. interval	interval at which to extract sampling data, e.g. 5 (to take one every five samples)	
window.size	instead of sampl.interval (for one sampling every so many time units), size (in sampling time units, e.g. days) of each sampling window (e.g. 3 for 3 consecutive days sampling each time); must be complemented with gap.size, for the gap between sampling windows.	
gap.size	size (in sampling time units, e.g. days) of the gaps between sampling windows $-$ e.g. 1 for 1 gap (non-sampled) day between sampling windows.	
start.column	column of sampl.columns where to actually start the sampling (e.g. with a gap size of 1 between sampling windows, the start column can be either 1 or 2)	
group.column	name or index number of the column containing the taxa or groups to analyse, e.g. 3 or "Family" $$	
group.names	$name(s) \ of \ the \ group(a) \ to \ extract, e.g. \ c("Mustelidae", "Procyonidae"); if \ NULL \ (the \ default), \ all \ groups \ in \ group.names \ are \ extracted$	
remove.zeros	logical indicating whether to remove rows where all extracted days have zero observations	
keep.nonsampl.columns		
	logical indicating whether to keep the non-sampling columns in the extracted result	

## Value

This function returns a subset of dataset containing the taxonomic groups and sampling columns resulting from the given sampling scheme.

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## See Also

sequential.submatrix, subset

## **Examples**

```
data(roadkills)

submat1 <- submatrix(dataset = roadkills, sampl.columns = 4:ncol(roadkills),
sampl.interval = 3, start.column = 1, group.column = "taxon", group.names = NULL)

head(submat1)

submat2 <- submatrix(dataset = roadkills, sampl.columns = 4:ncol(roadkills),
window.size = 5, gap.size = 2, start.column = 1, group.column = "taxon",
```

20 submatrix

 $\begin{aligned} & group.names = NULL) \\ & head(submat2) \end{aligned}$ 

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