

Package ‘MKomics’

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Title Omics Data Analysis

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Imports stats, utils, graphics, grDevices, RColorBrewer, robustbase,
limma, grid, circlize, ComplexHeatmap

Suggests knitr, rmarkdown

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Description Similarity plots based on correlation and median absolute deviation (MAD); adjusting colors for heatmaps; aggregate technical replicates; calculate pairwise fold-changes and log fold-changes; compute one- and two-way ANOVA; simplified interface to package 'limma' (Ritchie et al. (2015), <[doi:10.1093/nar/gkv007](https://doi.org/10.1093/nar/gkv007)>) for moderated t-test and one-way ANOVA; Hamming and Levenshtein (edit) distance of strings as well as optimal alignment scores for global (Needleman-Wunsch) and local (Smith-Waterman) alignments with constant gap penalties (Merkl and Waack (2009), ISBN:978-3-527-32594-8).

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MKomics-package	<i>Omic Data Analysis.</i>
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Description

Similarity plots based on correlation and median absolute deviation (MAD); adjusting colors for heatmaps; aggregate technical replicates; calculate pairwise fold-changes and log fold-changes; compute one- and two-way ANOVA; simplified interface to package 'limma' (Ritchie et al. (2015), <doi:10.1093/nar/gkv007>) for moderated t-test and one-way ANOVA; Hamming and Levenshtein (edit) distance of strings as well as optimal alignment scores for global (Needleman-Wunsch) and local (Smith-Waterman) alignments with constant gap penalties (Merkl and Waack (2009), ISBN:978-3-527-32594-8).

Details

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 Suggests: knitr, rmarkdown
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 URL: <https://www.stamats.de/>

```
library(MKomics)
```

Author(s)

Matthias Kohl <https://www.stamats.de>

Maintainer: Matthias Kohl <matthias.kohl@stamats.de>

corDist *Correlation Distance Matrix Computation*

Description

The function computes and returns the correlation and absolute correlation distance matrix computed by using the specified distance measure to compute the distances between the rows of a data matrix.

Usage

```
corDist(x, method = "pearson", diag = FALSE, upper = FALSE, abs = FALSE,  
        use = "pairwise.complete.obs", ...)
```

Arguments

x	a numeric matrix or data frame
method	the correlation distance measure to be used. This must be one of "pearson", "spearman", "kandall", "cosine", "mcd" or "ogk", respectively. Any unambiguous substring can be given.
diag	logical value indicating whether the diagonal of the distance matrix should be printed by 'print.dist'.
upper	logical value indicating whether the upper triangle of the distance matrix should be printed by 'print.dist'.
abs	logical, compute absolute correlation distances
use	character, corresponds to argument use of function cor
...	further arguments to functions covMcd or covOGK , respectively.

Details

The function computes the Pearson, Spearman, Kendall or Cosine sample correlation and absolute correlation; confer Section 12.2.2 of Gentleman et al (2005). For more details about the arguments we refer to functions [dist](#) and [cor](#). Moreover, the function computes the minimum covariance determinant or the orthogonalized Gnanadesikan-Kettenring estimator. For more details we refer to functions [covMcd](#) and [covOGK](#), respectively.

Value

corDist returns an object of class "dist"; cf. [dist](#).

Note

A first version of this function appeared in package SLMisc.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

- Gentleman R. Ding B., Dudoit S. and Ibrahim J. (2005). Distance Measures in DNA Microarray Data Analysis. In: Gentleman R., Carey V.J., Huber W., Irizarry R.A. and Dudoit S. (editors) Bioinformatics and Computational Biology Solutions Using R and Bioconductor. Springer.
- P. J. Rousseeuw and A. M. Leroy (1987). Robust Regression and Outlier Detection. Wiley.
- P. J. Rousseeuw and K. van Driessen (1999) A fast algorithm for the minimum covariance determinant estimator. Technometrics 41, 212-223.
- Pison, G., Van Aelst, S., and Willems, G. (2002), Small Sample Corrections for LTS and MCD, Metrika, 55, 111-123.
- Maronna, R.A. and Zamar, R.H. (2002). Robust estimates of location and dispersion of high-dimensional datasets; Technometrics 44(4), 307-317.
- Gnanadesikan, R. and John R. Kettenring (1972). Robust estimates, residuals, and outlier detection with multiresponse data. Biometrics 28, 81-124.

Examples

```
## only a dummy example
M <- matrix(rnorm(1000), ncol = 20)
D <- corDist(M)
```

corPlot

Plot of similarity matrix based on correlation

Description

Plot of similarity matrix. This function is a slight modification of function `plot.cor` of the archived package "sma".

Usage

```
corPlot(x, new = FALSE, col, minCor,
        labels = FALSE, lab.both.axes = FALSE, labcols = "black",
        title = "", cex.title = 1.2,
        protocol = FALSE, cex.axis = 0.8,
        cex.axis.bar = 1, signifBar = 2, ...)

corPlot2(x, new = FALSE, col, minCor = 0.5, labels = FALSE,
         row.width = 6, column.height = 6, lab.both.axes = TRUE,
         fontsize.axis = 12, title = "", fontsize.title = 16,
         signifBar = 2, use_raster = FALSE)
```

Arguments

x	data or correlation matrix, respectively
new	If new=FALSE, x must already be a correlation matrix. If new=TRUE, the correlation matrix for the columns of x is computed and displayed in the image.
col	colors palette for image. If missing, the RdYlGn palette of RColorBrewer is used.
minCor	numeric value in [-1,1], used to adjust col
labels	vector of character strings to be placed at the tickpoints, labels for the columns of x.
lab.both.axes	logical, display labels on both axes
labcols	colors to be used for the labels of the columns of x. labcols can have either length 1, in which case all the labels are displayed using the same color, or the same length as labels, in which case a color is specified for the label of each column of x.
title	character string, overall title for the plot.
cex.title	numerical value giving the amount by which plotting text and symbols should be magnified relative to the default; cf. par , cex.main.
fontsize.title	numerical value giving the fontsize of the title.
protocol	logical, display color bar without numbers.
cex.axis	The magnification to be used for axis annotation relative to the current setting of 'cex'; cf. par .
fontsize.axis	numerical value giving the fontsize of the axis labels.
cex.axis.bar	The magnification to be used for axis annotation of the color bar relative to the current setting of 'cex'; cf. par .
signifBar	integer indicating the precision to be used for the bar.
row.width	numerical value giving width of the row in centimeters; i.e., can be used to change space available for the labels.
column.height	numerical value giving the height of the column in centimeters; i.e., can be used to change space available for the labels.
use_raster	see Heatmap .
...	graphical parameters may also be supplied as arguments to the function (see par). For comparison purposes, it is good to set <code>zlim=c(-1,1)</code> .

Details

This functions generates the so called similarity matrix (based on correlation) for a microarray experiment.

If $\min(x)$, respectively $\min(\text{cor}(x))$ is smaller than minCor, the colors in col are adjusted such that the minimum correlation value which is color coded is equal to minCor.

The newer function corPlot2 is based on function Heatmap of Bioconductor package **ComplexHeatmap**.

Value

invisible()

Note

A first version of function corPlot appeared in package SLMisc.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

Sandrine Dudoit, Yee Hwa (Jean) Yang, Benjamin Milo Bolstad and with contributions from Natalie Thorne, Ingrid Loennstedt and Jessica Mar. sma: Statistical Microarray Analysis. <http://www.stat.berkeley.edu/users/terry/zarray/Software/smacode.html>

Examples

```
## only a dummy example
M <- matrix(rnorm(1000), ncol = 20)
colnames(M) <- paste("Sample", 1:20)
M.cor <- cor(M)

corPlot(M.cor, minCor = min(M.cor))
corPlot(M.cor, minCor = min(M.cor), lab.both.axes = TRUE)
corPlot(M.cor, minCor = min(M.cor), protocol = TRUE)
corPlot(M.cor, minCor = min(M.cor), signifBar = 1)

corPlot2(M.cor, minCor = min(M.cor))
corPlot2(M.cor, minCor = min(M.cor), lab.both.axes = FALSE)
corPlot2(M.cor, minCor = min(M.cor), signifBar = 1)
```

heatmapCol

Generate colors for heatmaps

Description

This function modifies a given color vector as used for heatmaps.

Usage

```
heatmapCol(data, col, lim, na.rm = TRUE)
```

Arguments

data	matrix or data.frame; data which shall be displayed in a heatmap; ranging from negative to positive numbers.
col	vector of colors used for heatmap.
lim	constant colors are used for data below <code>-lim</code> resp. above <code>lim</code> .
na.rm	logical; remove NA values.

Details

Colors below and above a specified value are kept constant. In addition, the colors are symmetrized.

Value

vector of colors

Note

A first version of this function appeared in package `SLmisc`.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

Examples

```
data.plot <- matrix(rnorm(100*50, sd = 1), ncol = 50)
colnames(data.plot) <- paste("patient", 1:50)
rownames(data.plot) <- paste("gene", 1:100)
data.plot[1:70, 1:30] <- data.plot[1:70, 1:30] + 3
data.plot[71:100, 31:50] <- data.plot[71:100, 31:50] - 1.4
data.plot[1:70, 31:50] <- rnorm(1400, sd = 1.2)
data.plot[71:100, 1:30] <- rnorm(900, sd = 1.2)
nrcol <- 128

require(RColorBrewer)
myCol <- rev(colorRampPalette(brewer.pal(10, "RdBu"))(nrcol))
heatmap(data.plot, col = myCol, main = "standard colors")
myCol2 <- heatmapCol(data = data.plot, col = myCol,
                    lim = min(abs(range(data.plot)))-1)
heatmap(data.plot, col = myCol2, main = "heatmapCol colors")
```

`madMatrix`*Compute MAD between columns of a matrix or data.frame*

Description

Compute MAD between columns of a matrix or data.frame. Can be used to create a similarity matrix for a microarray experiment.

Usage

```
madMatrix(x)
```

Arguments

`x` matrix or data.frame

Details

This functions computes the so called similarity matrix (based on MAD) for a microarray experiment; cf. Buness et. al. (2004).

Value

matrix of MAD values between columns of `x`

Note

A first version of this function appeared in package SLmisc.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

Andreas Buness, Wolfgang Huber, Klaus Steiner, Holger Sueltmann, and Annemarie Poustka. arrayMagic: two-colour cDNA microarray quality control and preprocessing. *Bioinformatics Advance Access* published on September 28, 2004. doi:10.1093/bioinformatics/bti052

See Also

plotMAD

Examples

```
## only a dummy example  
madMatrix(matrix(rnorm(1000), ncol = 10))
```

madPlot

Plot of similarity matrix based on MAD

Description

Plot of similarity matrix based on MAD between microarrays.

Usage

```
madPlot(x, new = FALSE, col, maxMAD = 3, labels = FALSE,
        labcols = "black", title = "", protocol = FALSE, ...)
```

```
madPlot2(x, new = FALSE, col, maxMAD = 3, labels = FALSE,
         row.width = 6, column.height = 6,
         lab.both.axes = TRUE, fontsize.axis = 12,
         title = "", fontsize.title = 16, signifBar = 2)
```

Arguments

x	data or correlation matrix, respectively
new	If new=FALSE, x must already be a matrix with MAD values. If new=TRUE, the MAD matrix for the columns of x is computed and displayed in the image.
col	colors palette for image. If missing, the RdYlGn palette of RColorBrewer is used.
maxMAD	maximum MAD value displayed
labels	vector of character strings to be placed at the tickpoints, labels for the columns of x.
labcols	colors to be used for the labels of the columns of x. labcols can have either length 1, in which case all the labels are displayed using the same color, or the same length as labels, in which case a color is specified for the label of each column of x.
title	character string, overall title for the plot.
fontsize.title	numerical value giving the fontsize of the title.
protocol	logical, display color bar without numbers
lab.both.axes	logical, display labels on both axes
fontsize.axis	numerical value giving the fontsize of the axis labels.
signifBar	integer indicating the precision to be used for the bar.
row.width	numerical value giving width of the row in centimeters; i.e., can be used to change space available for the labels.
column.height	numerical value giving the height of the column in centimeters; i.e., can be used to change space available for the labels.
...	graphical parameters may also be supplied as arguments to the function (see par). For comparison purposes, it is good to set zlim=c(-1,1).

Details

This function generates the so called similarity matrix (based on MAD) for a microarray experiment; cf. Bunes et. al. (2004). The function is similar to [corPlot](#).

Note

A first version of this function appeared in package SLmisc.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

Sandrine Dudoit, Yee Hwa (Jean) Yang, Benjamin Milo Bolstad and with contributions from Natalie Thorne, Ingrid Loennstedt and Jessica Mar. sma: Statistical Microarray Analysis. <http://www.stat.berkeley.edu/users/terry/zarray/Software/smacode.html>

Andreas Bunes, Wolfgang Huber, Klaus Steiner, Holger Suetmann, and Annemarie Poustka. arrayMagic: two-colour cDNA microarray quality control and preprocessing. Bioinformatics Advance Access published on September 28, 2004. doi:10.1093/bioinformatics/bti052

See Also

[corPlot](#)

Examples

```
## only a dummy example
set.seed(13)
x <- matrix(rnorm(1000), ncol = 10)
x[1:20,5] <- x[1:20,5] + 10
madPlot(x, new = TRUE, maxMAD = 2.5)
madPlot2(x, new = TRUE, maxMAD = 2.5)
## in contrast
corPlot2(x, new = TRUE, minCor = -0.5)
```

mod.oneway.test

Moderated 1-Way ANOVA

Description

Performs moderated 1-Way ANOVAs based on Bioconductor package limma.

Usage

```
mod.oneway.test(x, group, repeated = FALSE, subject, adjust.method = "BH",
               sort.by = "none")
```

Arguments

x a (non-empty) numeric matrix of data values.
group an optional factor representing the groups.
repeated logical indicating whether there are repeated-measures.
subject factor with subject IDs; required if repeated = TRUE.
adjust.method see [p.adjust](#)
sort.by see [toptable](#)
 , where "logFC" corresponds to difference in means.

Details

The function uses Bioconductor package *limma* to compute moderated 1-way ANOVAs. For more details we refer to [ebayes](#).

Value

A data.frame with the results.

References

Ritchie, M.E., Phipson, B., Wu, D., Hu, Y., Law, C.W., Shi, W., and Smyth, G.K. (2015). *limma* powers differential expression analyses for RNA-sequencing and microarray studies. *Nucleic Acids Research* 43(7), e47.

See Also

[oneway.test](#), [mod.t.test](#)

Examples

```

set.seed(123)
X <- rbind(matrix(rnorm(5*20), nrow = 5, ncol = 20),
            matrix(rnorm(5*20, mean = 1), nrow = 5, ncol = 20))
gr <- factor(c(rep("A1", 5), rep("B2", 5), rep("C3", 5), rep("D4", 5)))
mod.oneway.test(X, gr)

## Welch 1-Way ANOVA (not moderated)
ow.test <- function(x, g){
  res <- oneway.test(x ~ g)
  c(res$statistic, res$p.value)
}
ow.res <- t(apply(X, 1, ow.test, g = gr))
colnames(ow.res) <- c("F", "p.value")
ow.res

## repeated measures
X <- rbind(matrix(rnorm(6*18), nrow = 6, ncol = 18),
            matrix(rnorm(6*18, mean = 1), nrow = 6, ncol = 18))
gr <- factor(c(rep("T1", 6), rep("T2", 6), rep("T3", 6)))

```

```
subjectID <- factor(c(rep(1:6, 3)))
mod.oneway.test(X, gr, repeated = TRUE, subject = subjectID)
```

mod.t.test

Moderated t-Test

Description

Performs moderated t-tests based on Bioconductor package limma.

Usage

```
mod.t.test(x, group = NULL, paired = FALSE, subject, adjust.method = "BH",
           sort.by = "none", na.rm = TRUE)
```

Arguments

`x` a (non-empty) numeric matrix of data values.

`group` an optional factor representing the groups.

`paired` a logical indicating whether you want a paired test.

`subject` factor with subject IDs; required if `paired = TRUE`.

`adjust.method` see [p.adjust](#)

`sort.by` see [toptable](#)

, where "logFC" corresponds to difference in means.

`na.rm` logical. Should missing values (including NaN) be omitted from the calculations of group means?

Details

The function uses Bioconductor package limma to compute moderated t-tests. For more details we refer to [ebayes](#).

Value

A data.frame with the results.

References

Ritchie, M.E., Phipson, B., Wu, D., Hu, Y., Law, C.W., Shi, W., and Smyth, G.K. (2015). limma powers differential expression analyses for RNA-sequencing and microarray studies. *Nucleic Acids Research* 43(7), e47.

See Also

[t.test](#)

Examples

```

## One-sample test
X <- matrix(rnorm(10*20, mean = 1), nrow = 10, ncol = 20)

mod.t.test(X)
## corresponds to
library(limma)
design <- matrix(1, nrow = ncol(X), ncol = 1)
colnames(design) <- "A"
fit1 <- lmFit(X, design)
fit2 <- eBayes(fit1)
topTable(fit2, coef = 1, number = Inf, confint = TRUE, sort.by = "none")[,-4]

## Two-sample test
set.seed(123)
X <- rbind(matrix(rnorm(5*20), nrow = 5, ncol = 20),
            matrix(rnorm(5*20, mean = 1), nrow = 5, ncol = 20))
g2 <- factor(c(rep("group 1", 10), rep("group 2", 10)))

mod.t.test(X, group = g2)
## corresponds to
design <- model.matrix(~ 0 + g2)
colnames(design) <- c("group1", "group2")
fit1 <- lmFit(X, design)
cont.matrix <- makeContrasts(group1vsgroup2="group1-group2", levels=design)
fit2 <- contrasts.fit(fit1, cont.matrix)
fit3 <- eBayes(fit2)
topTable(fit3, coef = 1, number = Inf, confint = TRUE, sort.by = "none")[,-4]

## Paired two-sample test
subjID <- factor(rep(1:10, 2))
mod.t.test(X, group = g2, paired = TRUE, subject = subjID)

```

oneWayAnova

A function for Analysis of Variance

Description

This function is a slight modification of function Anova of package **genefilter**.

Usage

```
oneWayAnova(cov, na.rm = TRUE, var.equal = FALSE)
```

Arguments

cov The covariate. It must have length equal to the number of columns of the array that the result of oneWayAnova will be applied to.

<code>na.rm</code>	a logical value indicating whether NA values should be stripped before the computation proceeds.
<code>var.equal</code>	a logical variable indicating whether to treat the variances in the samples as equal. If TRUE, then a simple F test for the equality of means in a one-way analysis of variance is performed. If FALSE, an approximate method of Welch (1951) is used, which generalizes the commonly known 2-sample Welch test to the case of arbitrarily many samples.

Details

The function returned by `oneWayAnova` uses `oneway.test` to perform a one-way ANOVA, where `x` is the set of gene expressions. The F statistic for an overall effect is computed and the corresponding p-value is returned.

The function `Anova` of package **genefilter** instead compares the computed p-value to a prespecified p-value and returns TRUE, if the computed p-value is smaller than the prespecified one.

Value

`oneWayAnova` returns a function with bindings for `cov` that will perform a one-way ANOVA.

The covariate can be continuous, in which case the test is for a linear effect for the covariate.

Note

A first version of this function appeared in package `SLmisc`.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

R. Gentleman, V. Carey, W. Huber and F. Hahne (2006). `genefilter`: methods for filtering genes from microarray experiments. R package version 1.13.7.

See Also

[oneway.test](#), [twoWayAnova](#)

Examples

```
set.seed(123)
af <- oneWayAnova(c(rep(1,5),rep(2,5)))
af(rnorm(10))
```

pairwise.fc	<i>Compute pairwise fold changes</i>
-------------	--------------------------------------

Description

This function computes pairwise fold changes. It also works for logarithmic data.

Usage

```
pairwise.fc(x, g, ave = mean, log = TRUE, base = 2, mod.fc = TRUE, ...)
```

Arguments

x	numeric vector.
g	grouping vector or factor
ave	function to compute the group averages.
log	logical. Is the data logarithmic?
base	If log = TRUE, the base which was used to compute the logarithms.
mod.fc	logical. Return modified fold changes? (see details)
...	optional arguments to ave.

Details

The function computes pairwise fold changes between groups, where the group values are aggregated using the function which is given by the argument ave.

The fold changes are returned in a slightly modified form if mod.fc = TRUE. Fold changes FC which are smaller than 1 are reported as to $-1/FC$.

The implementation is in certain aspects analogously to [pairwise.t.test](#).

Value

Vector with pairwise fold changes.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

See Also

[pairwise.t.test](#)

Examples

```

set.seed(13)
x <- rnorm(100) ## assumed as log2-data
g <- factor(sample(1:4, 100, replace = TRUE))
levels(g) <- c("a", "b", "c", "d")
pairwise.fc(x, g)

## some small checks
res <- by(x, list(g), mean)
2^(res[[1]] - res[[2]]) # a vs. b
-1/2^(res[[1]] - res[[3]]) # a vs. c
2^(res[[1]] - res[[4]]) # a vs. d
-1/2^(res[[2]] - res[[3]]) # b vs. c
-1/2^(res[[2]] - res[[4]]) # b vs. d
2^(res[[3]] - res[[4]]) # c vs. d

```

pairwise.logfc

Compute pairwise log-fold changes

Description

The function computes pairwise log-fold changes.

Usage

```
pairwise.logfc(x, g, ave = mean, log = TRUE, base = 2, ...)
```

Arguments

x	numeric vector.
g	grouping vector or factor
ave	function to compute the group averages.
log	logical. Is the data logarithmic?
base	If log = TRUE, the base which was used to compute the logarithms.
...	optional arguments to ave.

Details

The function computes pairwise log-fold changes between groups, where the group values are aggregated using the function which is given by the argument ave.

The implementation is in certain aspects analogously to [pairwise.t.test](#).

Value

Vector with pairwise log-fold changes.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

See Also

[pairwise.t.test](#)

Examples

```
set.seed(13)
x <- rnorm(100) ## assumed as log2-data
g <- factor(sample(1:4, 100, replace = TRUE))
levels(g) <- c("a", "b", "c", "d")
pairwise.logfc(x, g)

## some small checks
res <- by(x, list(g), mean)
res[[1]] - res[[2]] # a vs. b
res[[1]] - res[[3]] # a vs. c
res[[1]] - res[[4]] # a vs. d
res[[2]] - res[[3]] # b vs. c
res[[2]] - res[[4]] # b vs. d
res[[3]] - res[[4]] # c vs. d
```

pairwise.mod.t.test *Pairwise Moderated t-Tests*

Description

Performs pairwise moderated t-tests (unpaired) based on Bioconductor package limma.

Usage

```
pairwise.mod.t.test(x, group, adjust.method = "BH", sort.by = "none")
```

Arguments

`x` a (non-empty) numeric matrix of data values.
`group` an optional factor representing the groups.
`adjust.method` see [p.adjust](#)
`sort.by` see [tortable](#)
, where "logFC" corresponds to difference in means.

Details

The function uses Bioconductor package limma to compute pairwise moderated t-tests. For more details we refer to [ebayes](#).

Value

A data.frame with the results.

References

Ritchie, M.E., Phipson, B., Wu, D., Hu, Y., Law, C.W., Shi, W., and Smyth, G.K. (2015). limma powers differential expression analyses for RNA-sequencing and microarray studies. *Nucleic Acids Research* 43(7), e47.

See Also

[oneway.test](#), [mod.t.test](#)

Examples

```
set.seed(123)
X <- rbind(matrix(rnorm(5*20), nrow = 5, ncol = 20),
             matrix(rnorm(5*20, mean = 1), nrow = 5, ncol = 20))
gr <- factor(c(rep("A1", 5), rep("B2", 5), rep("C3", 5), rep("D4", 5)))
mod.oneway.test(X, gr)
pairwise.mod.t.test(X, gr)
```

 repMeans

Compute mean of replicated spots

Description

Compute mean of replicated spots where additionally spot flags may incorporated.

Usage

```
repMeans(x, flags, use.flags = NULL, ndups, spacing, method, ...)
```

Arguments

x	matrix or data.frame of expression values
flags	matrix or data.frame of spot flags; must have same dimension as x
use.flags	should flags be included and in which way; cf. section details
ndups	integer, number of replicates on chip. The number of rows of x must be divisible by ndups
spacing	the spacing between the rows of 'x' corresponding to replicated spots, spacing = 1 for consecutive spots; cf. function unwrapdups in package "limma"
method	function to aggregate the replicated spots. If missing, the mean is used.
...	optional arguments to method.

Details

The incorporation of spot flags is controlled via argument `use.flags`.

NULL: flags are not used; minimum flag value of replicated spots is returned

"max": only spots with flag value equal to the maximum flag value of replicated spots are used

"median": only spots with flag values larger or equal to median of replicated spots are used

"mean": only spots with flag values larger or equal to mean of replicated spots are used

Value

LIST with components

<code>exprs</code>	mean of expression values
<code>flags</code>	flags for mean expression values

Note

A first version of this function appeared in package `SLmisc`.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

See Also

[unwrapdups](#)

Examples

```
## only a dummy example
M <- matrix(rnorm(1000), ncol = 10)
FL <- matrix(rpois(1000, lambda = 10), ncol = 10) # only for this example
res <- repMeans(x = M, flags = FL, use.flags = "max", ndups = 5, spacing = 20)
```

simPlot

Plot of a similarity matrix.

Description

Plot of similarity matrix.

Usage

```
simPlot(x, col, minVal, labels = FALSE, lab.both.axes = FALSE,
        labcols = "black", title = "", cex.title = 1.2,
        protocol = FALSE, cex.axis = 0.8,
        cex.axis.bar = 1, signifBar = 2, ...)

simPlot2(x, col, minVal, labels = FALSE,
         row.width = 6, column.height = 6, lab.both.axes = TRUE,
         fontsize.axis = 12, title = "", fontsize.title = 16,
         signifBar = 2)
```

Arguments

x	quadratic data matrix.
col	colors palette for image. If missing, the RdYlGn palette of RColorBrewer is used.
minVal	numeric, minimum value which is display by a color; used to adjust col
labels	vector of character strings to be placed at the tickpoints, labels for the columns of x.
lab.both.axes	logical, display labels on both axes
labcols	colors to be used for the labels of the columns of x. labcols can have either length 1, in which case all the labels are displayed using the same color, or the same length as labels, in which case a color is specified for the label of each column of x.
title	character string, overall title for the plot.
cex.title	A numerical value giving the amount by which plotting text and symbols should be magnified relative to the default; cf. par , <code>cex.main</code> .
fontsize.title	numerical value giving the fontsize of the title.
protocol	logical, display color bar without numbers
cex.axis	The magnification to be used for axis annotation relative to the current setting of 'cex'; cf. par .
fontsize.axis	numerical value giving the fontsize of the axis labels.
cex.axis.bar	The magnification to be used for axis annotation of the color bar relative to the current setting of 'cex'; cf. par .
signifBar	integer indicating the precision to be used for the bar.
row.width	numerical value giving width of the row in centimeters; i.e., can be used to change space available for the labels.
column.height	numerical value giving the height of the column in centimeters; i.e., can be used to change space available for the labels.
...	graphical parameters may also be supplied as arguments to the function (see par). For comparison purposes, it is good to set <code>zlim=c(-1,1)</code> .

Details

This functions generates a so called similarity matrix.

If `min(x)` is smaller than `minVal`, the colors in `col` are adjusted such that the minimum value which is color coded is equal to `minVal`.

Value

`invisible()`

Note

The function is a slight modification of function `corPlot`.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

Sandrine Dudoit, Yee Hwa (Jean) Yang, Benjamin Milo Bolstad and with contributions from Natalie Thorne, Ingrid Loennstedt and Jessica Mar. *sma: Statistical Microarray Analysis*.
<http://www.stat.berkeley.edu/users/terry/zarray/Software/smacode.html>

Examples

```
## only a dummy example
M <- matrix(rnorm(1000), ncol = 20)
colnames(M) <- paste("Sample", 1:20)
M.cor <- cor(M)

simPlot(M.cor, minVal = min(M.cor))
simPlot(M.cor, minVal = min(M.cor), lab.both.axes = TRUE)
simPlot(M.cor, minVal = min(M.cor), protocol = TRUE)
simPlot(M.cor, minVal = min(M.cor), signifBar = 1)

simPlot2(M.cor, minVal = min(M.cor))
simPlot2(M.cor, minVal = min(M.cor), lab.both.axes = FALSE)
simPlot2(M.cor, minVal = min(M.cor), signifBar = 1)
```

stringDist

Function to compute distances between strings

Description

The function can be used to compute distances between strings.

Usage

```
stringDist(x, y, method = "levenshtein", mismatch = 1, gap = 1)
```

Arguments

x	character vector, first string
y	character vector, second string
method	character, name of the distance method. This must be "levenshtein" or "hamming". Default is the classical Levenshtein distance.
mismatch	numeric, distance value for a mismatch between symbols
gap	numeric, distance value for inserting a gap

Details

The function computes the Hamming and the Levenshtein (edit) distance of two given strings (sequences).

In case of the Hamming distance the two strings must have the same length.

In case of the Levenshtein (edit) distance a scoring and a trace-back matrix are computed and are saved as attributes "ScoringMatrix" and "TraceBackMatrix". The characters in the trace-back matrix reflect insertion of a gap in string y (d: deletion), match (m), mismatch (mm), and insertion of a gap in string x (i).

Value

stringDist returns an object of S3 class "stringDist" inherited from class "dist"; cf. [dist](#).

Note

The function is mainly for teaching purposes.

For distances between strings and string alignments see also Bioconductor package **Biostrings**.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

R. Merkl and S. Waack (2009). Bioinformatik Interaktiv. Wiley.

See Also

[dist](#), [stringSim](#)

Examples

```
x <- "GACGGATTATG"
y <- "GATCGGAATAG"
## Levenshtein distance
d <- stringDist(x, y)
d
attr(d, "ScoringMatrix")
attr(d, "TraceBackMatrix")
```

```
## Hamming distance
stringDist(x, y)
```

stringSim *Function to compute similarity scores between strings*

Description

The function can be used to compute similarity scores between strings.

Usage

```
stringSim(x, y, global = TRUE, match = 1, mismatch = -1, gap = -1, minSim = 0)
```

Arguments

x	character vector, first string
y	character vector, second string
global	logical; global or local alignment
match	numeric, score for a match between symbols
mismatch	numeric, score for a mismatch between symbols
gap	numeric, penalty for inserting a gap
minSim	numeric, used as required minimum score in case of local alignments

Details

The function computes optimal alignment scores for global (Needleman-Wunsch) and local (Smith-Waterman) alignments with constant gap penalties.

Scoring and trace-back matrix are computed and saved in form of attributes "ScoringMatrix" and "TraceBackMatrix". The characters in the trace-back matrix reflect insertion of a gap in string y (d: deletion), match (m), mismatch (mm), and insertion of a gap in string x (i). In addition stop indicates that the minimum similarity score has been reached.

Value

stringSim returns an object of S3 class "stringSim" inherited from class "dist"; cf. [dist](#).

Note

The function is mainly for teaching purposes.

For distances between strings and string alignments see also Bioconductor package **Biostrings**.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

R. Merkl and S. Waack (2009). Bioinformatik Interaktiv. Wiley.

See Also

[dist](#), [stringDist](#)

Examples

```
x <- "GACGGATTATG"
y <- "GATCGGAATAG"

## optimal global alignment score
d <- stringSim(x, y)
d
attr(d, "ScoringMatrix")
attr(d, "TraceBackMatrix")

## optimal local alignment score
d <- stringSim(x, y, global = FALSE)
d
attr(d, "ScoringMatrix")
attr(d, "TraceBackMatrix")
```

traceBack

Function to trace back

Description

Function computes an optimal global or local alignment based on a trace back matrix as provided by function [stringDist](#) or [stringSim](#).

Usage

```
traceBack(D, global = TRUE)
```

Arguments

D	object of class "stringDist"
global	logical, global or local alignment

Details

Computes one possible optimal global or local alignment based on the trace back matrix saved in an object of class "stringDist" or "stringSim".

Value

matrix: pairwise global/local alignment

Note

The function is mainly for teaching purposes.

For distances between strings and string alignments see Bioconductor package **Biostrings**.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

R. Merkl and S. Waack (2009). Bioinformatik Interaktiv. Wiley.

See Also

[stringDist](#)

Examples

```
x <- "GACGGATTATG"
y <- "GATCGGAATAG"

## Levenshtein distance
d <- stringDist(x, y)
## optimal global alignment
traceBack(d)

## Optimal global alignment score
d <- stringSim(x, y)
## optimal global alignment
traceBack(d)

## Optimal local alignment score
d <- stringSim(x, y, global = FALSE)
## optimal local alignment
traceBack(d, global = FALSE)
```

twoWayAnova

A function for Analysis of Variance

Description

This function is a slight modification of function Anova of package **genefilter**.

Usage

```
twoWayAnova(cov1, cov2, interaction, na.rm = TRUE)
```

Arguments

cov1	The first covariate. It must have length equal to the number of columns of the array that the result of twoWayAnova will be applied to.
cov2	The second covariate. It must have length equal to the number of columns of the array that the result of twoWayAnova will be applied to.
interaction	logical, should interaction be considered
na.rm	a logical value indicating whether 'NA' values should be stripped before the computation proceeds.

Details

The function returned by twoWayAnova uses `lm` to fit a linear model of the form `lm(x ~ cov1*cov2)`, where `x` is the set of gene expressions. The F statistics for the main effects and the interaction are computed and the corresponding p-values are returned.

Value

twoWayAnova returns a function with bindings for cov1 and cov2 that will perform a two-way ANOVA.

Note

A first version of this function appeared in package SLmisc.

Author(s)

Matthias Kohl <Matthias.Kohl@stamats.de>

References

R. Gentleman, V. Carey, W. Huber and F. Hahne (2006). `genefilter`: methods for filtering genes from microarray experiments. R package version 1.13.7.

See Also

[lm](#), [oneWayAnova](#)

Examples

```
set.seed(123)
af1 <- twoWayAnova(c(rep(1,6),rep(2,6)), rep(c(rep(1,3), rep(2,3)), 2))
af2 <- twoWayAnova(c(rep(1,6),rep(2,6)), rep(c(rep(1,3), rep(2,3)), 2),
                  interaction = FALSE)
x <- matrix(rnorm(12*10), nrow = 10)
apply(x, 1, af1)
apply(x, 1, af2)
```

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