Package ‘timecourse’

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Title Statistical Analysis for Developmental Microarray Time Course Data

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Description Functions for data analysis and graphical displays for developmental microarray time course data.

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R topics documented:

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abs2ratio

Convert log-values to log-ratios

Description

For a single gene, computes the log ratios between time courses from two paired biological conditions.

Usage

abs2ratio(x, mn, k, c.grp, reference)

Arguments

x
a numeric vector giving the log-values of a gene with two paired biological conditions, sorted in ascending order by biological condition, replicate, and time groups.

mn
a numeric matrix giving the sample sizes for the two biological conditions.

k
a positive integer giving the number of time points.

c.grp
an numeric or character vector with length equals to that of x, giving the biological condition group for each element of x.

reference
a numeric value or character assigning the reference biological condition.

Details

This function is for internal use only and is not to be called by the user.

Value

a numeric vector containing log-ratios between two paired biological conditions.

Author(s)

Yu Chuan Tai <yuchuan@stat.berkeley.edu>

See Also

mb.paired.
fruitfly  

\textit{Drosophila microarray time course data in Tomancak et al. (2002)}

\textbf{Description}

This is a subset of the Drosophila microarray time course data in Tomancak et al. (2002).

\textbf{Usage}

\texttt{data(fruitfly)}

\textbf{Format}

A matrix of $\log_2$ gene expression values for 2000 probesets.

\textbf{Source}

Tomancak et al. (2002) \textit{Systematic determination of patterns of gene expression during Drosophila embryogenesis}. Genome Biology 2002, 3:research0088.1-0088.14 \url{http://genomebiology.com/2002/3/12/research/0088.1} The complete dataset can be downloaded from the following website \url{http://www.fruitfly.org/cgi-bin/ex/insitu.pl}

\textbf{MArrayTC-class}  

\textit{Microarray Time Course Object-class}

\textbf{Description}

A list-based class for storing the analysis results from the multivariate empirical Bayes models of differential expression for longitudinal replicated developmental microarray time course data. Objects are normally created by \texttt{mb.long} and \texttt{mb.MANOVA}.

\textbf{Slots/Components}

\texttt{MArrayTC} objects do not contain any slots (apart from \texttt{.Data}) but they should contain the following list components:

- \texttt{M}: input matrix of log-ratios or log-values of expression for a series of microarrays.

Objects may also contain the following optional components:

- \texttt{prop}: numeric value giving the proportion of differentially expressed genes.
- \texttt{nu}: numeric value containing the estimated amount of moderation.
- \texttt{Lambda}: the estimated Lambda.
- \texttt{Lambda1}: the estimated Lambda1.
- \texttt{eta}: the estimated prior scale parameter.
- \texttt{alpha}: the estimated common mean of the expected time course vector under the null.
- \texttt{alpha.d}: the estimated condition-specific means of the expected time course vectors under the alternative.
matrix.cov

beta: the estimated scale parameter for the common covariance matrix of the common expected
time course vector under the null.

beta.d: the estimated condition-specific scale parameters for the common covariance matrix of
the expected time course vectors under the alternative.

percent: numeric matrix containing the percent of moderation corresponding to each sample size
for the longitudinal one- and two- sample problems.

size: numeric vector or matrix containing the sample sizes for all genes corresponding to different
biological conditions, when the latter are sorted in ascending order.

con.group: numeric or character vector giving the biological condition group of each array. The
i_th element of con.group corresponds to the biological condition of the i_th column of M.

rep.group: numeric or character vector giving the replicate group of each array. The i_th ele-
ment of rep.group corresponds to the replicate of the i_th column of M.

time.group: numeric vector giving the time group of each array. The i_th element of time.group
corresponds to the time of the i_th column of M.

HotellingT2: numeric vector giving the $\tilde{T}^2$ statistics of differential expression.

MB: numeric vector giving the MB-statistics of differential expression.

pos.HotellingT2: numeric vector whose i_th element corresponds to the index of the gene with
ranking i in HotellingT2.

pos.MB: numeric vector whose i_th element corresponds to the index of the gene with ranking i in
MB.

geneNames: character vector giving gene names.

descriptions: character vector giving gene descriptions.

Methods

MArrayTC extends the LargeDataObject class in package limma, and inherits a show method from
there.

The function plotProfile takes MArrayTC as the input argument.

Author(s)

Yu Chuan Tai <yuchuan@stat.berkeley.edu>

matrix.cov Covariance

Description

For a single gene, computes the transformed or untransformed sample covariance matrix if one
biological condition, or pooled sample covariance matrix if two or more biological conditions.

Usage

matrix.cov(x, k, trans = TRUE, c.grp = NULL, use = "complete.obs")
Arguments

- **x**: a numeric vector giving the log-ratios or log-values for a gene, sorted in ascending order by biological condition, replicate, and time groups.
- **k**: a positive integer giving the number of time points.
- **trans**: logical. Should the Helmert transformation be performed?
- **c.grp**: a numeric vector corresponding to the biological condition group for each element of \( x \).
- **use**: character. The same as the use in stats function `cov`. The default uses complete observations.

Details

This function is for internal use only and is not to be called by the user.

Value

A numeric matrix.

Author(s)

Yu Chuan Tai <yuchuan@stat.berkeley.edu>

References


See Also

cov, ot.helmert.

Description

Computes the \( \tilde{T}^2 \) statistics and/or the MB-statistics of differential expression for longitudinal replicated developmental microarray time course data by multivariate empirical Bayes shrinkage of gene-specific sample variance-covariance matrices towards a common matrix.

Usage

```r
mb.long(object, method = c("1D", "paired", "2D"), type = c("none", "robust"),
times, reps, prior.df = NULL, prior.COV = NULL,
prior.eta = NULL, condition.grp = NULL, rep.grp = NULL, time.grp = NULL,
one.sample = FALSE, ref = NULL, p = 0.02, out.t = FALSE,
tuning = 1.345, HotellingT2.only=TRUE)
```
Arguments

object Required. An object of class matrix, MAList, marrayNorm, or ExpressionSet containing log-ratios or log-values of expression for a series of microarrays.

method a character string, "1D" for the one-sample case where genes of interest are those which change over time, "paired" for the one-sample case where genes of interest are those whose expected temporal profiles do not stay 0, for example, cDNA microarrays, or the paired two-sample case where genes of interest are those with different expected temporal profiles across 2 biological conditions, "2D" for the independent two-sample case where genes of interest are those with different expected temporal profiles across 2 biological conditions. The default is "1D".

type a character string, indicating whether possible outliers should be down-weighted.

times Required. A positive integer giving the number of time points.

reps Required. A numeric vector or matrix corresponding to the sample sizes for all genes across different biological conditions, when biological conditions are sorted in ascending order. If a matrix, rows represent genes while columns represent biological conditions.

prior.df an optional positive value giving the degrees of moderation.

prior.COV an optional numeric matrix giving the common covariance matrix to which the gene-specific sample covariances are smoothed toward.

prior.eta an optional numeric value giving the scale parameter for the covariance matrix for the expected time course profile.

condition.grp a numeric or character vector with length equals to the number of arrays, assigning the biological condition group of each array. Required if method=2D.

rep.grp an optional numeric or character vector with length equals to the number of arrays, assigning the replicate group of each array.

time.grp an optional numeric vector with length equals to the number of arrays, assigning the time point group of each array.

one.sample Is it a one-sample problem? Only specify this argument when method=paired. The default is FALSE which means it is a paired two-sample problem.

ref an optional numeric value or character specifying the name of reference biological condition. The default uses the first element of condition.grp. Only specify this argument when method=paired and one.sample is FALSE.

p a numeric value between 0 and 1, assumed proportion of genes which are differentially expressed.

out.t logical. Should the moderated multivariate t-statistics be outputed? The default is FALSE.

tuning the tuning constant for the Huber weight function with a default 1.345.

HotellingT2.only logical. Should only the HotellingT2 statistics be outputed? This should be set as TRUE (default) when the sample size(s) are the same across genes, in order to reduce computational time.

Details

This function implements the multivariate empirical Bayes statistics described in Tai and Speed (2004), to rank genes in the order of interest from longitudinal replicated developmental microarray
time course experiments. It calls one of the following functions, depending on which method is used: \texttt{mb.1D}, \texttt{mb.paired}, and \texttt{mb.2D}.

The arguments \texttt{condition.grp}, \texttt{rep.grp}, and \texttt{time.grp}, if specified, should have lengths equal to the number of arrays. The \texttt{i\_th} elements of these three arguments should correspond to the biological condition, replicate, and time for the \texttt{i\_th} column (array) in the expression value matrix of the input object, respectively. The default assumes the columns of \( M \) are in the ascending order of \texttt{condition.grp} first, and then \texttt{rep.grp}, and finally \texttt{time.grp}.

Arguments \texttt{one.sample} and \texttt{ref} are for \texttt{method=paired} only.

When \texttt{type=robust}, the numerator of the \( T^2 \) statistic is calculated using the weighted average time course vector(s), where the weight at each data point is determined using Huber’s weight function with the default tuning constant 1.345.

Warning: When there are only 2 replicates within conditions, \texttt{type="robust"} produces the same rankings as \texttt{type="none"} since there is no consensus on gene expression values. Check the output weights for these outliers.

\textbf{Value}

Object of \texttt{MArrayTC}.

\textbf{Author(s)}

Yu Chuan Tai <yuchuan@stat.berkeley.edu>

\textbf{References}


\textbf{See Also}

timecourse Vignette.

\textbf{Examples}

data(fruitfly)
colnames(fruitfly) ## check if arrays are arranged in the default order
gnames <- rownames(fruitfly)
assay <- rep(c("A", "B", "C"), each = 12)
time.grp <- rep(c(1:12), 3)
size <- rep(3, nrow(fruitfly))

out1 <- mb.long(fruitfly, times=12, reps=size, rep.grp = assay, time.grp = time.grp)
summary(out1)
plotProfile(out1, type="b", gnames=gnames, legloc=c(2,15), pch=c("A","B","C"), xlab="Hour")

## Simulate gene expression data
## Note: this simulation is for demonstration purpose only,
## and does not necessarily reflect the real
## features of longitudinal time course data
## one biological condition, 5 time points, 3 replicates
## 500 genes, 10 genes change over time

\[
SS <- \begin{bmatrix}
0.01 & -0.0008 & -0.003 & 0.007 & 0.002 \\
-0.0008 & 0.02 & 0.002 & -0.0004 & -0.001 \\
-0.003 & 0.002 & 0.03 & -0.0054 & -0.009 \\
0.007 & -0.0004 & -0.00538 & 0.02 & 0.0008 \\
0.002 & -0.001 & -0.009 & 0.0008 & 0.07
\end{bmatrix}
\]

sim.Sigma <- function()
{
S <- matrix(rep(0,25),ncol=5)
x <- mvrnorm(n=10, mu=rep(0,5), Sigma=10*SS)
for(i in 1:10)
S <- S+crossprod(t(x[i,]))
solve(S)
}

sim.data1 <- function(x, indx=1)
{
mu <- rep(runif(1,8,x[1]),5)
if(indx==1) res <- as.numeric(t(mvrnorm(n=3, mu=mu+rnorm(5,sd=4), Sigma=sim.Sigma())))
if(indx==0) res <- as.numeric(t(mvrnorm(n=3, mu=mu, Sigma=sim.Sigma())))
res
} # Not Run

M1 <- matrix(rep(14,500*15), ncol=15)
M1[1:10,] <- t(apply(M1[1:10,],1,sim.data1))
M1[11:500,] <- t(apply(M1[11:500,],1,sim.data1, 0))

## Which genes are nonconstant?
MB.1D1 <- mb.long(M1, times=5, reps=rep(3, 500))
MB.1D1$percent # check the percent of moderation

plotProfile(MB.1D1,type="b") # plots the no. 1 gene

plotProfile(MB.1D1,type="b",ranking=10) # plots the no. 10 gene

plotProfile(MB.1D1, type="b", gid="8", gnames=genenames) #plots the gene with ID "8"

# assign the following labellings to columns of M1
# which is actually the same as the default

trt <- rep("wildtype", 15)
assay <- rep(c("A","B","C"), rep(5,3))
time.grp <- rep(c(0, 1, 3, 4, 6), 3)

# MB.1D2 should give the same results as MB.1D1
#MB.1D2 <- mb.long(M1, times=5, reps=rep(3, 500), conditiongrp = trt, rep.grp = assay, 
#time.grp=time.grp)
## suppose now the replicates are in this order instead
assay <- rep(c("A","C","B"), rep(5,3))

## then
MB.1D3 <- mb.long(M1, times=5, reps=rep(3, 500), condition.grp = trt, rep.grp = assay, time.grp=time.grp)
MB.1D3$rep.group # check the replicate and time group
MB.1D3$time.group

## Now let's simulate another dataset with two biological conditions
## 500 genes also, 10 of them have different expected time course profiles
## between these two biological conditions
## 3 replicates, 5 time points for each condition

sim.data2 <- function(x, indx=1)
{
  mu <- rep(runif(1,8,x[1]),5)
  if(indx==1)
    res <- c(as.numeric(t(mvrnorm(n=3, mu=mu+rnorm(5,sd=5), Sigma=sim.Sigma()))),
             as.numeric(t(mvrnorm(n=3, mu=mu+rnorm(5,sd=3.2), Sigma=sim.Sigma()))))
  if(indx==0) res <- as.numeric(t(mvrnorm(n=6, mu=mu+rnorm(5,sd=3), Sigma=sim.Sigma())))
  res
}

M2 <- matrix(rep(14,500*30), ncol=30)
M2[1:10,] <- t(apply(M2[1:10,],1,sim.data2))
M2[11:500,] <- t(apply(M2[11:500,],1,sim.data2, 0))

## assume it is a paired two-sample problem
trt <- rep(c("wt","mt"),each=15)
assay <- rep(rep(c("1.2.04","2.4.04","3.5.04"),each=5),2)
size <- matrix(3, nrow=500, ncol=2)
MB.paired <- mb.long(M2, method="paired", times=5, reps=size, condition.grp=trt, rep.grp=assay)
MB.paired$con.group # check the condition, replicate and time groups
MB.paired$rep.group
MB.paired$time.group

plotProfile(MB.paired, type="b")
genenames <- as.character(1:500)
plotProfile(MB.paired, gid="12", type="b", gnames=genenames) # plots the gene with ID "12"

## assume it is an unpaired two-sample problem
assay <- rep(rep(c("1.2.04","2.4.04","3.5.04","5.21.04","7.17.04","8.4.04"),each=5))
MB.2D <- mb.long(M2, method="2", times=5, reps=size, condition.grp=trt, rep.grp=assay)
MB.2D$con.group # check the condition, replicate and time groups
MB.2D$rep.group
MB.2D$time.group

plotProfile(MB.2D, type="b", gnames=genenames) # plot the no. 1 gene

## Now let's simulate another dataset with two biological conditions
## 500 genes also, 10 of them have different expected time course profiles
## between these two biological conditions
## the first condition has 3 replicates, while the second condition has 4 replicates,
## 5 time points for each condition

```r
sim.data3 <- function(x, indx=1)
{
    mu <- rep(runif(1,8,x[1]),5)
    if(indx==1)
        res <- c(as.numeric(t(mvrnorm(n=3, mu=mu+rnorm(5,sd=5), Sigma=sim.Sigma()))),
                  as.numeric(t(mvrnorm(n=4, mu=mu+rnorm(5,sd=3.2), Sigma=sim.Sigma()))))
    if(indx==0) res <- as.numeric(t(mvrnorm(n=7, mu=mu+rnorm(5,sd=3), Sigma=sim.Sigma())))
    res
}

M3 <- matrix(rep(14,500*35), ncol=35)
M3[1:10,] <- t(apply(M3[1:10,],1,sim.data3))
M3[11:500,] <- t(apply(M3[11:500,],1,sim.data3, 0))

assay <- rep(c("1.2.04","2.4.04","3.5.04","5.21.04","7.17.04","9.10.04","12.1.04"),each=5)
trt <- c(rep(c("wildtype","mutant"),each=15),rep("mutant",5))
# Note that "mutant" < "wildtype", the sample sizes are (4, 3)
size <- matrix(c(4,3), nrow=500, ncol=2, byrow=TRUE)
MB.2D.2 <- mb.long(M3, method="2", times=5, reps=size, rep.grp=assay, condition.grp=trt)
MB.2D.2$con.group # check the condition, replicate and time groups
MB.2D.2$rep.group
MB.2D.2$time.group

plotProfile(MB.2D.2, type="b") # plot the no. 1 gene
```

---

**mb.MANOVA**

Multivariate Empirical Bayes Analysis of Variance for Longitudinal Replicated Developmental Microarray Time Course Data

### Description

Computes the MB-statistics for longitudinal replicated developmental microarray time course data with multiple biological conditions.

### Usage

```r
mb.MANOVA(object, times, D, size, nu = NULL, Lambda = NULL,
          beta.d = NULL, beta = NULL, alpha.d = NULL, alpha = NULL,
          condition.grp, time.grp = NULL, rep.grp = NULL, p = 0.02)
```

### Arguments

- **object**: Required. An object of class matrix, MAList, marrayNorm, or ExpressionSet containing log-ratios or log-values of expression for a series of microarrays.
- **times**: Required. A positive integer giving the number of time points.
- **D**: Required. A positive integer giving the number of biological conditions. D>1
- **size**: Required. A numeric matrix corresponding to the sample sizes for all genes across different biological conditions, when biological conditions are sorted in ascending order. Rows represent genes while columns represent biological conditions.
**mb.MANOVA**

- **nu**
  an optional positive value giving the degrees of moderation for the fully moderated Wilks' Lambda.

- **Lambda**
  an optional numeric matrix giving the common covariance matrix for the fully moderated Wilks' Lambda.

- **beta.d**
  an optional numeric vector of length D giving the condition-specific scale parameters for the common covariance matrix of the expected time course vectors under the alternative.

- **beta**
  an optional numeric value giving the scale parameter for the common covariance matrix of the common expected time course vector under the null.

- **alpha.d**
  an optional numeric matrix giving the condition-specific means of the expected time course vectors under the alternative.

- **alpha**
  an optional numeric vector of length times giving the common mean of the expected time course vector under the null.

- **condition.grp**
  Required. A numeric or character vector with length equals to the number of arrays, assigning the biological condition group to each array.

- **rep.grp**
  an optional numeric or character vector with length equals to the number of arrays, assigning the replicate group to each array.

- **time.grp**
  an optional numeric vector with length equals to the number of arrays, assigning the time point group to each array.

- **p**
  a numeric value between 0 and 1, assumed proportion of genes which are differentially expressed.

**Details**

This function implements the multivariate empirical Bayes Analysis of Variance model for identifying genes with different temporal profiles across multiple biological conditions, as described in Tai (2005).

**Value**

Object of `MArrayTC`.

**Author(s)**

Yu Chuan Tai <yuchuan@stat.berkeley.edu>

**References**


**See Also**

timecourse Vignette.
Examples

SS <- matrix(c( 0.01, -0.0008, -0.003, 0.007, 0.002, 
              -0.0008, 0.02, 0.002, -0.0004, -0.001, 
              -0.003, 0.002, 0.03, -0.0054, -0.009, 
              0.007, -0.0004, -0.00538, 0.02, 0.0008, 
              0.002, -0.001, -0.009, 0.0008, 0.07), ncol=5)

sim.Sigma <- function()
{
  S <- matrix(rep(0,25),ncol=5)
  x <- mvrnorm(n=10, mu=rep(0,5), Sigma=10*SS)
  for(i in 1:10)
    S <- S + crossprod(t(x[i,]))
  solve(S)
}

## Now let's simulate a dataset with three biological conditions
## 500 genes in total, 10 of them have different expected time course profiles
## across biological conditions
## the first condition has 3 replicates, while the second condition has 4 replicates,
## and the third condition has 2 replicates. 5 time points for each condition.

sim.data <- function(x, indx=1)
{
  mu <- rep(runif(1,8,x[1]),5)
  if(indx==1)
    res <- c(as.numeric(t(mvrnorm(n=3, mu=mu+rnorm(5,sd=5), Sigma=sim.Sigma()))),
             as.numeric(t(mvrnorm(n=4, mu=mu+rnorm(5,sd=3.2), Sigma=sim.Sigma()))),
             as.numeric(t(mvrnorm(n=2, mu=mu+rnorm(5,sd=2), Sigma=sim.Sigma()))))
  if(indx==0) res <- as.numeric(t(mvrnorm(n=9, mu=mu+rnorm(5,sd=3), Sigma=sim.Sigma())))
  res
}

M <- matrix(rep(14,500*45), ncol=45)
M[1:10,] <- t(apply(M[1:10,],1,sim.data))
M[11:500,] <- t(apply(M[11:500,],1,sim.data, 0))

assay <- rep(c("1.2.04","2.4.04","3.5.04","5.21.04","7.17.04","9.10.04","12.1.04","1.2.05","4.1.05"),each=5)
trt <- c(rep("wildtype","mutant1"),each=15),rep("mutant1",5), rep("mutant2", 10))

# Caution: since "mutant1" < "mutant2" < "wildtype", the sample sizes should be in the order of 4,2,3,
# but NOT 3,4,2.
size <- matrix(c(4,2,3), byrow=TRUE, nrow=500, ncol=3)
MB.multi <- mb.MANOVA(M, times=5, D=3, size=size, rep.grp=assay, condition.grp=trt)

plotProfile(MB.multi, stats="MB", type="b") # plots the no. 1 gene
plotProfile

Description

Computes the Helmert orthogonal transformation matrix.

Usage

ot.helmert(k)

Arguments

k

a positive integer giving the number of time points.

Details

This function is for internal use only and is not to be called by the user.

Value

a numeric matrix.

Author(s)

Yu Chuan Tai <yuchuan@stat.berkeley.edu>

plotProfile

Description

Plots the longitudinal temporal profile of a gene.

Usage

plotProfile(object, stats=c("HotellingT2", "MB"), ranking=1, gid=NULL, gnames=NULL, desc=NULL, type=c("p","l","b"), col=2:100, lty=1:100, pch=1:100, lwd=2, xlab="Time", ylab="Expression", legloc=NULL, xlim=NULL, ylim=NULL, cex.main=1,...)

Arguments

object

a MArrayTC object.

stats

a character indicating which statistic the ranking is based on.

ranking

a numeric value giving the ranking of the gene to be plotted.

gid

an optional character giving the ID of the gene to be plotted.

gnames

an optional character vector with the i_th element corresponds to the gene ID of the i_th gene in object$M.

desc

an optional character vector with the i_th element corresponds to the gene description of the i_th gene in object$M.

type

a character indicating the plot type, "p" for points, "l" for lines, and "b" for both.

col

a character or numeric vector giving the colors for different biological conditions. Default is 2:100.
univ.func

1ty a character or numeric vector giving the line types for different replicates. Default is 1:100.
pch a character or numeric vector giving the point types for different replicates. Default is 1:100.
lwd optional. The default sets to 2.
xlab character. The label for the x-axis.
ylab character. The label for the y-axis.
legloc an optional vector giving the location of the legend.
xlim an optional vector giving the upper- and lower- limits of x-axis.
ylim an optional vector giving the upper- and lower- limits of y-axis.
cex.main optional. The default sets to 1
... any other arguments passed onto plot

Details

This function takes an object of MArrayTC as the input and plots the temporal profile of a single gene. The user can specify either the ranking based on stats or the gene ID of the gene to be plotted.

See points for possible values for pch, col and cex.
See mb.long for examples.

Author(s)

Yu Chuan Tai <yuchuan@stat.berkeley.edu>

---

univ.func     Univariate Data

Description

Transforms multivariate vectors into univariate values using the Helmert matrix.

Usage

univ.func(dummy, M, k, n, indx = 1)

Arguments

dummy a numeric gene index.
M a numeric matrix containing the log-values or log-ratios of a gene.
k a positive integer giving the number of time points.
n a positive integer giving the number of replicates.
indx a positive integer between 1 and k, indicating which row of the Helmert matrix to transform the vectors.

Details

This function is for internal use only and is not to be called by the user.
**Value**

A numeric vector with length equals to n.

**Author(s)**

Yu Chuan Tai <yuchuan@stat.berkeley.edu>

**See Also**

*ot.helmert*.
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