Package 'ORCME'

October 12, 2022

Type Package				
Title Order Restricted Clustering for Microarray Experiments				
Version 2.0.2				
Date 2015-07-15				
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Description Provides clustering of genes with similar dose response (or time course) profiles. It implements the method described by Lin et al. (2012).				
Imports Iso				
License GPL-3				
LazyLoad yes				
Repository CRAN				
Repository/R-Forge/Project orcme				
Repository/R-Forge/Revision 65				
Repository/R-Forge/DateTimeStamp 2015-07-23 12:31:52				
Date/Publication 2015-07-31 12:12:01				
NeedsCompilation no				
Depends R (>= 2.10)				
R topics documented:				
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doseData

Dose Data Example

Description

Dose data; a vector of length 12 with 3 observations for each of 4 doses.

Usage

```
data(doseData)
```

Format

The format is: num [1:12] 1 1 1 2 2 2 3 3 ...

Examples

```
data(doseData)
doseData
```

geneData

Gene Expression Data Example

Description

This dose-response microarray data contains 1000 genes and 4 doses (one control dose (zero dose) and three increasing dose) with 3 arrays at each dose level. Due to confidetiality, it is only part of the real data set.

Usage

```
data(geneData)
```

Format

A data frame with 1000 observations on the following 12 variables.

- X1 Sample one with zero dose
- X1.1 Sample two with zero dose
- X1.2 Sample three with zero dose
- X2 Sample one with second dose
- X2.1 Sample two with second dose

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- X2.2 Sample three with second dose
- X3 Sample one with third dose
- X3.1 Sample two with third dose
- X3.2 Sample three with third dose
- X4 Sample one with fourth dose
- X4.1 Sample two with fourth dose
- X4.2 Sample three with fourth dose

References

Testing for Trend in Dose-Response Microarray Experiments: a Comparison of Testing Procedures, Multiplicity, and Resampling-Based Inference, Lin et al. 2007, Stat. App. in Gen. & Mol. Bio., 6(1), article 26.

Examples

data(geneData)

head(geneData)

monotoneDirection

The monotone means under increasing/decreasing trend

Description

The function calculates the likelihood for the increasing and decreasing trend in the dose response for all the given genes separately gene-by-gene. The trend with the higher likelihood is chosen and the isotonic regression is applied on the means.

Usage

```
monotoneDirection(geneData, doseData)
```

Arguments

geneData gene expression matrix for all genes

doseData indicates the dose levels

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Value

A list with components

direction the direction with the higher likelihood of increasing (indicated by "up") or de-

creasing (indicated by "dn") trend.

incData isotonic means with respect to dose for those genes that were classified as fol-

lowing the increasing trend.

decData isotonic means with respect to dose for those genes that were classified as fol-

lowing the decreasing trend.

obsincData observed gene expression matrix for those genes that were classified as follow-

ing the increasing trend.

observed gene expression matrix for those genes that were classified as follow-

ing the decreasing trend.

arrayMean isotonic means with respect to dose for all genes.

Author(s)

Adetayo Kasim, Martin Otava and Tobias Verbeke

References

Lin D., Shkedy Z., Yekutieli D., Amaratunga D., and Bijnens, L. (editors). (2012) Modeling Doseresponse Microarray Data in Early Drug Development Experiments Using R. Springer.

Cheng, Y. and Church, G. M. (2000). Biclustering of expression data. In: Proceedings of the Eighth International Conference on Intelligent Systems for Molecular Biology, 1, 93-103.

See Also

ORCME, plotIsomeans

Examples

```
data(doseData)
data(geneData)

dirData <- monotoneDirection(geneData = geneData,doseData = doseData)

## direction of monotone trend
Direction <- dirData$direction

## Isotonic means for upward genes
incData <- as.data.frame(dirData$incData)

##Isotonic means for downward genes
decData <- as.data.frame(dirData$decData)

## observd data upward genes
obsIncData <- as.data.frame(dirData$obsincData)

## observed data for downward genes
obsDecData <- as.data.frame(dirData$obsdecData)

## isotonic means for all genes
isoMeans <- as.data.frame(dirData$arrayMean)</pre>
```

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ORCME	Order restricted clustering for dose-response trends in microarray experiments

Description

The function performs delta-clustering of a microarray data. It can be used for clustering of both the time-course or dose-response microarray data.

Usage

```
ORCME(DRdata, lambda, phi, robust=FALSE)
```

Arguments

phi

DRdata	matrix of a microarray data with rows corresponding to genes and columns corresponding to time points or different doses
lambda	assumed proportion of coherence relative to the observed data, it ranges between 0 and 1. A lambda value of 1 considers the observed data as a cluster and lambda value of 0 finds every possible pattern within the data.

minimum number of genes in a cluster

robust logical variable that determines, if algorithm uses robust version based on me-

dian polish and absolute values, instead of mean square error. Default is FALSE.

Value

The matrix of classification into clusters: each row represents one gene and columns found clusters. The matrix consist of the Booleans values, in each row there is only one of them TRUE which means that the gene was classified into the respective cluster.

Author(s)

Adetayo Kasim, Martin Otava and Tobias Verbeke

References

Lin D., Shkedy Z., Yekutieli D., Amaratunga D., and Bijnens, L. (editors). (2012) Modeling Doseresponse Microarray Data in EarlyDrug Development Experiments Using R. Springer.

Cheng, Y. and Church, G. M. (2000). Biclustering of expression data. In: Proceedings of the Eighth International Conference on Intelligent Systems for Molecular Biology, 1, 93-103.

See Also

monotoneDirection, plotIsomeans

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Examples

```
data(doseData)
data(geneData)

dirData <- monotoneDirection(geneData = geneData,doseData = doseData)
incData <- as.data.frame(dirData$incData)

print(orcme <- ORCME(DRdata=incData,lambda=0.15,phi=2))
orcmeRobust <- ORCME(DRdata=incData,lambda=0.15,phi=2, robust=TRUE)

# number of genes within cluster
colSums(orcme)
colSums(orcmeRobust)</pre>
```

plotCluster

Plotting the gene specific profiles for one given cluster of genes

Description

The function is plotting the profiles of the genes that belongs to the same cluster. It is not providing the clustering itself, just plotting the results of clustering from input. Optionally, the function can center the profiles around the gene-specific means.

Usage

```
plotCluster(DRdata, doseData, ORCMEoutput, clusterID,
zeroMean=FALSE, xlabel, ylabel, main="")
```

Arguments

DRdata the microarray data with rows corresponding to genes and columns correspond-

ing to time points or different doses

doseData indicates the dose levels

ORCMEoutput the matrix of classification into clusters: each row represents one gene and

columns found clusters. The matrix consist of the Booleans values, in each row there is only one of them TRUE which means that the gene was classified

into the respective gene

clusterID id of the cluster to be plotted

zeroMean if TRUE, it centers the gene profiles around the gene-specific means, default is

FALSE

xlabel a title for the x axis ylabel a title for the y axis

main an overall title for the plot

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Value

Plot of the gene specific profiles dependent one the dose level (or time point) that are classified into the given cluster.

Author(s)

Adetayo Kasim, Martin Otava and Tobias Verbeke

References

Lin D., Shkedy Z., Yekutieli D., Amaratunga D., and Bijnens, L. (editors). (2012) Modeling Doseresponse Microarray Data in Early Drug Development Experiments Using R. Springer.

Cheng, Y. and Church, G. M. (2000). Biclustering of expression data. In: Proceedings of the Eighth International Conference on Intelligent Systems for Molecular Biology, 1, 93-103.

See Also

```
ORCME, plotIsomeans
```

Examples

```
data(doseData)
data(geneData)

dirData <- monotoneDirection(geneData = geneData,doseData = doseData)
incData <- as.data.frame(dirData$incData)
ORCMEoutput <- ORCME(DRdata=incData,lambda=0.15,phi=2)

plotCluster(DRdata=incData,doseData=doseData, ORCMEoutput=ORCMEoutput,clusterID=4,zeroMean=FALSE, xlabel="Dose",ylabel="Gene Expression")</pre>
```

plotIsomeans

Plot of the observed gene expression and the isotonic means with respect to dose

Description

The function is plotting the observed data points of the gene expression and isotonic means with respect to dose for one particular gene.

Usage

```
plotIsomeans(monoData, obsData, doseData, geneIndex)
```

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Arguments

monoData isotonic means with respect to dose for all genes

obsData observed gene expression for all genes

doseData indicates the dose levels

geneIndex index of the gene to be plotted

Value

Plot of the data points and the isotonic means for each dose with the isotonic regression curve.

Author(s)

Adetayo Kasim, Martin Otava and Tobias Verbeke

References

Lin D., Shkedy Z., Yekutieli D., Amaratunga D., and Bijnens, L. (editors). (2012) Modeling Doseresponse Microarray Data in Early Drug Development Experiments Using R. Springer.

Cheng, Y. and Church, G. M. (2000). Biclustering of expression data. In: Proceedings of the Eighth International Conference on Intelligent Systems for Molecular Biology, 1, 93-103.

See Also

ORCME, monotoneDirection

Examples

```
data(doseData)
data(geneData)

dirData <- monotoneDirection(geneData = geneData,doseData = doseData)
incData <- as.data.frame(dirData$incData)
obsIncData <- as.data.frame(dirData$obsincData)

## gene-specific profile plot
plotIsomeans(monoData=incData,obsData=obsIncData,doseData=doseData,geneIndex=10)</pre>
```

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plotLambda	Plot the variaty of the properties dependent on the proportion of heterogeneity in observed data set
	•

Description

This function provides the plots of the dependency of the variety of properties on the proportion of heterogeneity in observed data set. It is not using the clustering as simple input, but it is also computing additional properties. The function can plot within cluster sum of squares, number of cluster, penalized within cluster sum of squares, Calsanzik and Harabasx index and Hartigan index.

Usage

plotLambda(lambdaChoiceOutput,output)

Arguments

lambdaChoiceOutput

the output of the function resampleORCME

output

the variable that determines which output would be plotted, the values are "wss" for the cluster sum of squares, "ncluster" for the number of cluster, "pwss" for the penalized within cluster sum of squares, "ch" for the Calsanzik and Harabasx index and "h" for the Llustines index.

index and "h" for the Hartigan index

Value

A plot of one of the properties mentioned above dependent on the proportion of heterogeneity. The confidence intervals are plotted instead of the point estimates.

Author(s)

Adetayo Kasim, Martin Otava and Tobias Verbeke

References

Lin D., Shkedy Z., Yekutieli D., Amaratunga D., and Bijnens, L. (editors). (2012) Modeling Doseresponse Microarray Data in Early Drug Development Experiments Using R. Springer.

Cheng, Y. and Church, G. M. (2000). Biclustering of expression data. In: Proceedings of the Eighth International Conference on Intelligent Systems for Molecular Biology, 1, 93-103.

See Also

ORCME, resampleORCME

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Examples

```
data(doseData)
data(geneData)

dirData <- monotoneDirection(geneData = geneData,doseData = doseData)
incData <- as.data.frame(dirData$incData)

lambdaVector <- c(0.05,0.50,0.95)

lambdaChoiceOutput <- resampleORCME(clusteringData=incData, lambdaVector=lambdaVector)
plotLambda(lambdaChoiceOutput,output="wss")
plotLambda(lambdaChoiceOutput,output="ncluster")
plotLambda(lambdaChoiceOutput,output="pwss")
plotLambda(lambdaChoiceOutput,output="ch")
plotLambda(lambdaChoiceOutput,output="ch")</pre>
```

resampleORCME

Estimation of the proportion of the heterogeneity in the observed data for clustering

Description

The function is computing within cluster sum of squares for given proportion of heterogeneity. Minimal number of genes per cluster is fixed as 2. The sum of squares is computed through resampling the 100 data sets with 100 genes randomly sampled with replacement from the reduced expression data.

Usage

```
resampleORCME(clusteringData, lambdaVector, robust=FALSE)
```

Arguments

clusteringData the microarray data with rows corresponding to genes and columns corresponding to time points or different doses

lambdaVector vector of assumed proportions of of heterogeneity of the observed data, it ranges

between 0 and 1. A lambda value of 1 considers the observed data as a cluster

and lambda value of 0 finds every possible pattern within the data

robust logical variable that determines, if algorithm uses robust version based on me-

dian polish and absolute values, instead of mean square error. Default is FALSE.

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Value

A list of matrices that represent one of the 100 iterations. Every matrix consist of the columns

lambda vector of the proportions of heterogeneity given as input

within clusters sum of squares for given proportion of heterogeneity
total clusters sum of squares for given proportions of heterogeneity
nc number of clusters as a function for given proportions of heterogeneity

Author(s)

Adetayo Kasim, Martin Otava and Tobias Verbeke

References

Lin D., Shkedy Z., Yekutieli D., Amaratunga D., and Bijnens, L. (editors). (2012) Modeling Doseresponse Microarray Data in Early Drug Development Experiments Using R. Springer.

Cheng, Y. and Church, G. M. (2000). Biclustering of expression data. In: Proceedings of the Eighth International Conference on Intelligent Systems for Molecular Biology, 1, 93-103.

See Also

```
ORCME, plotLambda
```

Examples

```
data(doseData)
data(geneData)

dirData <- monotoneDirection(geneData = geneData,doseData = doseData)
incData <- as.data.frame(dirData$incData)

lambdaVector <- c(0.05,0.50,0.95)

resampleORCME(clusteringData=incData, lambdaVector=lambdaVector, robust=FALSE)</pre>
```

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