

Package ‘projectR’

October 16, 2019

Type Package

Title Functions for the projection of weights from PCA, CoGAPS, NMF, correlation, and clustering

Version 1.0.0

Date 2016-11-22

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Description Functions for the projection of data into the spaces defined by PCA, CoGAPS, NMF, correlation, and clustering.

License GPL (==2)

Imports methods, cluster, stats, limma, CoGAPS, NMF, ROCR

Suggests BiocStyle, gridExtra, grid, testthat, devtools, knitr, rmarkdown, ComplexHeatmap, viridis, ggplot2

LazyData TRUE

RoxygenNote 6.1.1

Encoding UTF-8

VignetteBuilder knitr

biocViews FunctionalPrediction, GeneRegulation, BiologicalQuestion, Software

BugReports <https://support.bioconductor.org/t/projectR/>

URL <https://github.com/genesofeve/projectR/>

git_url <https://git.bioconductor.org/packages/projectR>

git_branch RELEASE_3_9

git_last_commit 69a67dd

git_last_commit_date 2019-05-02

Date/Publication 2019-10-15

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AP.RNAseq6l3c3t	<i>CoGAPS patterns and genes weights for p.RNAseq6l3c3t</i>
-----------------	---

Description

AP.RNAseq6l3c3t contains the output of the gapsRun function in the CoGAPS package for data = p.RNAseq6l3c3t

Usage

```
AP.RNAseq6l3c3t
```

Format

A list of 12 items

auc_mat	<i>auc_mat</i>
---------	----------------

Description

Calculates AUC values for each set of weights for each label and outputs the results as a matrix

Usage

```
auc_mat(labels, weights)
```

Arguments

labels	a vector of labels whose length is equal to the number of columns in the weight matrix
weights	a matrix of weights from projection analysis

Value

A matrix of AUC values for each set of weights classifying each label.

Examples

```
projectR(data=p.ESepiGen4c11$mRNA.Seq,loadings=AP.RNAseq6l3c3t$Amean,
dataNames = map.ESepiGen4c11[["GeneSymbols"]]) -> projection
auc_mat(pd.ESepiGen4c11$Condition,projection)
```

cluster2pattern	<i>Generic cluster2pattern function</i>
-----------------	---

Description

Function to make patterns of continuous weights from clusters.

Usage

```
cluster2pattern(clusters, NP, Data, ...)

## S4 method for signature 'kmeans'
cluster2pattern(clusters, Data)

## S4 method for signature 'hclust'
cluster2pattern(clusters, NP, Data = NA)
```

Arguments

clusters	a cluster object which could be either an hclust or a kmeans object
NP	number of desired patterns
Data	data used to make clusters object
...	Additional arguments to cluster2pattern

Value

An object of class pclus containing pattern weights corresponding for each cluster.

Examples

```
k.RNAseq6l3c3t<-kmeans(p.RNAseq6l3c3t,22)
cluster2pattern(clusters=k.RNAseq6l3c3t,NP=22,Data=p.RNAseq6l3c3t)

distp <- dist(p.RNAseq6l3c3t)
hc.RNAseq6l3c3t <- hclust(distp)
cluster2pattern(clusters=hc.RNAseq6l3c3t,NP=22,Data=p.RNAseq6l3c3t)
```

clusterPlotR *Generic clusterPlotR function*

Description

plotting function for clustering objects

Usage

```
clusterPlotR(cData, cls, x, NC, ...)  
  
## S4 method for signature 'ANY,kmeans'  
clusterPlotR(cData = NA, cls = NA, x = NA,  
             NC = NA, annoIndx = NA, label = NULL, ...)  
  
## S4 method for signature 'ANY,hclust'  
clusterPlotR(cData = NA, cls = NA, x = NA,  
             NC = NA, annoIndx = NA, label = NULL, ...)
```

Arguments

cData	data used to get clusters
cls	a cluster (kmeans or hclust) object
x	a vector of length equal to number of samples to use for plotting
NC	vector of integers indicating which clusters to use
...	additional parameters for plotting. ex. pch,cex,col,labels, xlab, etc.
annoIndx	vector indexing into subsets for plotting
label	character vector to use for plotting text, defaults is NULL

Value

A plot of the mean behavior for each cluster

Examples

```
## Not run:  
k.RNAseq613c3t<-kmeans(p.RNAseq613c3t,22)  
clusterPlotR(p.RNAseq613c3t, cls=k.RNAseq613c3t,NC=1,x=pd.RNAseq613c3t$days,  
col=pd.RNAseq613c3t$color)  
  
## End(Not run)
```

correlateR	<i>correlateR</i>
------------	-------------------

Description

Function to extract genes highly correlated with a gene or reference expression pattern.

Usage

```
correlateR(genes, dat, threshtype = "R", threshold = 0.7,
  absR = FALSE, ...)
```

Arguments

genes	gene or character vector of genes for reference expression pattern
dat	matrix or data frame with genes to be used for to calculate correlation
threshtype	Default "R" indicates thresholding by R value or equivalent. Alternatively, "N" indicates a numerical cut off.
threshold	numeric indicating value at which to make threshold.
absR	logical indicating where to include both positive and negatively correlated genes
...	addition inputs to cor, such as method

Details

If threshtype is "R" than threshold must be between -1 and 1. Otherwise if top N correlated genes are required, set threshtype as "N" and set threshold = N, i.e, the number of correlated genes required.

Value

A correlation matrix

Examples

```
cor2T<-correlateR(genes="T", dat=p.RNAseq613c3t, threshtype="N", threshold=10, absR=TRUE)
```

correlateR-class	<i>correlateR</i>
------------------	-------------------

Description

class of correlateR output.

Slots

corM correlation matrix obtained from correlateR

geneMatchR *Generic geneMatchR function*

Description

Matches genes accross datasets

Usage

```
geneMatchR(data1, data2, data1Names = NULL, data2Names = NULL,
            merge = FALSE)
```

Arguments

data1	a data matrix, typically genes by samples
data2	an amplitude matrix, typically genes by factors
data1Names	rownames of data matrix, for eg genenames
data2Names	rownames of amplitude matrix to be matched to rownames of datamatrix
merge	logical indicating wether or not to merged data sets
...	Additional arguments to geneMatchR

Value

A list of genes (intersection) in both datasets. (if merge = TRUE, Also returns merged data.)

Examples

```
geneMatchR(data1=p.ESepiGen4c11$mRNA.Seq,data2=p.RNAseq6l3c3t,
            data1Names=map.ESepiGen4c11[["GeneSymbols"]])
```

initialize,correlateR-method
Constructor for correlateR

Description

Constructor for correlateR

Usage

```
## S4 method for signature 'correlateR'
initialize(.Object, corM, ...)
```

Arguments

.Object	correlateR object
corM	correlation matrix obtained from correlateR
...	additional arguments to intialize correlateR

Value

initialized correlateR object

initialize,rotatoR-method
Constructor for rotatoR

Description

Constructor for rotatoR

Usage

```
## S4 method for signature 'rotatoR'
initialize(.Object, rotatedM, ...)
```

Arguments

.Object	rotatoR object
rotatedM	rotated matrix from rotatoR function
...	additional arguments to initialize rotatoR

Value

initialized rotatoR object

intersectoR *Generic intersectoR function*

Description

A function to find and test the intersecting values of two sets of objects, presumably the genes associated with patterns in two different datasets. Both the input objects need to be of the same type either kmeans or hclust.

Usage

```
intersectoR(pSet1, pSet2, pval, ...)

## S4 method for signature 'kmeans,kmeans'
intersectoR(pSet1 = NA, pSet2 = NA,
  pval = 0.05, full = FALSE)

## S4 method for signature 'hclust,hclust'
intersectoR(pSet1 = NA, pSet2 = NA,
  pval = 0.05, full = FALSE, k = NULL)
```

Arguments

pSet1	an object for a set of patterns where each entry is a set of genes associated with a single pattern
pSet2	an object for a second set of patterns where each entry is a set of genes associated with a single pattern
pval	the maximum p-value considered significant
...	additional parameters depending on input object
full	logical indicating whether to return full data frame of significantly overlapping sets. Default is false will return summary matrix.
k	Numeric giving cut height for hclust objects, if a vector is given arguments will be applied to pSet1 and pSet2 in that order

Value

A list containing: Overlap matrix, overlap index, and overlapping sets.

Examples

```
ESepiGen4c11mRNASeq <- p.ESepiGen4c11$mRNA.Seq
rownames(ESepiGen4c11mRNASeq) <- map.ESepiGen4c11$GeneSymbols

k.RNAseq6l3c3t<-kmeans(p.RNAseq6l3c3t,22)
k.ESepiGen4c11<-kmeans(ESepiGen4c11mRNASeq,10)
intersectoR(k.RNAseq6l3c3t, k.ESepiGen4c11, pval=.05)

h.RNAseq6l3c3t<-hclust(as.dist(1-(cor(t(p.RNAseq6l3c3t)))))
h.ESepiGen4c11<-hclust(as.dist(1-(cor(t(ESepiGen4c11mRNASeq)))))
intersectoR(pSet1=h.ESepiGen4c11, pSet2=h.RNAseq6l3c3t, pval=.05, k=c(3,4))
```

map.ESepiGen4c11	<i>RNAseqing and ChIPSeq of matched genes in differentiated human iPS cells</i>
------------------	---

Description

map.ESepiGen4c11 contains gene annotations

Usage

```
map.ESepiGen4c11
```

Format

A data frames with 93 rows and 9 variables:

References

1. Gifford, C. A. et al. Transcriptional and epigenetic dynamics during specification of human embryonic stem cells. *Cell* 153, 1149-1163 (2013).

map.RNAseq613c3t	<i>RNAseqing from human 3 iPS & 3 ES cell lines in 3 experimental condition at 3 time points</i>
------------------	--

Description

map.RNAseq613c3 contains gene annotations for polyA bulk sequencing of 6 cell lines in 3 experimental condition at 3 time points.

Usage

map.RNAseq613c3t

Format

A data frames with 108 rows and 54 variables:

p.ESepiGen4c11	<i>RNAseqing and ChIPSeq of matched genes in differentiated human iPS cells</i>
----------------	---

Description

p.ESepiGen4c11 contains $\log_2(\text{RPKM} + 1)$ values for polyA bulk sequencing and \log_2 counts of normalized ChIPSeq reads of 1 cell lines with 2 replicates in 4 experimental conditions at a single time point.

Usage

p.ESepiGen4c11

Format

p.ESepiGen4c11 is a list of 6 data frames each with with 93 rows and between 4 and 9 variables:

References

1. Gifford, C. A. et al. Transcriptional and epigenetic dynamics during specification of human embryonic stem cells. *Cell* 153, 1149-1163 (2013).

p.RNAseq613c3t	<i>RNAseqing from human 3 iPS & 3 ES cell lines in 3 experimental condition at 3 time points</i>
----------------	--

Description

p.RNAseq613c3 contains $\log_2(\text{RPKM} + 1)$ values for polyA bulk sequencing of 6 cell lines in 3 experimental condition at 3 time points.

Usage

p.RNAseq613c3t

Format

A data frames with 108 rows and 54 variables:

pd.ESepiGen4c11	<i>RNAseqing and ChIPSeq of matched genes in differentiated human iPS cells</i>
-----------------	---

Description

pd.ESepiGen4c11.4cond contains sample phenotype and experimental information

Usage

pd.ESepiGen4c11

Format

A data frames with 9 rows and 2 variables:

References

1. Gifford, C. A. et al. Transcriptional and epigenetic dynamics during specification of human embryonic stem cells. *Cell* 153, 1149-1163 (2013).

pd.RNAseq6l3c3t	<i>RNAseqing from human 3 iPS & 3 ES cell lines in 3 experimental condition at 3 time points</i>
-----------------	--

Description

pd.RNAseq6l3c3t contains sample phenotype and experimental information for polyA bulk sequencing of 6 cell lines in 3 experimental condition at 3 time points.

Usage

```
pd.RNAseq6l3c3t
```

Format

A data frames with 54 rows and 38 variables:

projectR	<i>Generic projectR function</i>
----------	----------------------------------

Description

A function for the projection of new data into a previously defined feature space.

Usage

```
projectR(data, loadings, dataNames = NULL, loadingsNames = NULL, ...)
```

```
## S4 method for signature 'matrix,matrix'
projectR(data, loadings, dataNames = NULL,
  loadingsNames = NULL, NP = NA, full = FALSE,
  family = "gaussianff")
```

```
## S4 method for signature 'matrix,LinearEmbeddingMatrix'
projectR(data, loadings,
  dataNames = NULL, loadingsNames = NULL, NP = NA, full = FALSE,
  model = NA, family = "gaussianff")
```

```
## S4 method for signature 'matrix,prcomp'
projectR(data, loadings, dataNames = NULL,
  loadingsNames = NULL, NP = NA, full = FALSE)
```

```
## S4 method for signature 'matrix,rotator'
projectR(data, loadings, dataNames = NULL,
  loadingsNames = NULL, NP = NA, full = FALSE)
```

```
## S4 method for signature 'matrix,correlateR'
projectR(data, loadings, dataNames = NULL,
  loadingsNames = NULL, NP = NA, full = FALSE)
```

```
## S4 method for signature 'matrix,hclust'
projectR(data, loadings, dataNames = NULL,
         loadingsNames = NULL, full = FALSE, targetNumPatterns, sourceData)

## S4 method for signature 'matrix,kmeans'
projectR(data, loadings, dataNames = NULL,
         loadingsNames = NULL, full = FALSE, sourceData)
```

Arguments

<code>data</code>	Target dataset into which you will project. It must of type matrix.
<code>loadings</code>	loadings learned from source dataset.
<code>dataNames</code>	a vector containing unique name, i.e. gene names, for the rows of the target dataset to be used to match features with the loadings, if not provided by <code>rownames(data)</code> . Order of names in vector must match order of rows in data.
<code>loadingsNames</code>	a vector containing unique names, i.e. gene names, for the rows of loadings to be used to match features with the data, if not provided by <code>rownames(loadings)</code> . Order of names in vector must match order of rows in loadings.
<code>...</code>	Additional arguments to <code>projectR</code>
<code>NP</code>	vector of integers indicating which columns of loadings object to use. The default of <code>NP=NA</code> will use entire matrix.
<code>full</code>	logical indicating whether to return the full model solution. By default only the new pattern object is returned.
<code>family</code>	VGAM family function for model fitting (default: "gaussianff")
<code>model</code>	Optional arguments to choose method for projection
<code>targetNumPatterns</code>	desired number of patterns
<code>sourceData</code>	data used to create cluster object

Details

loadings can belong to one of several classes depending on upstream analysis. Currently permitted classes are `matrix`, `CogapsResult`, `CoGAPS`, `pclust`, `prcomp`, `rotator`, and `correlater`.

Value

A matrix of sample weights for each input basis in the loadings matrix (if `full=TRUE`, full model solution is returned).

Examples

```
projectR(data=p.ESepiGen4c11$mRNA.Seq, loadings=AP.RNAseq613c3t$Amean,
         dataNames = map.ESepiGen4c11[["GeneSymbols"]])

library("CoGAPS")
CR.RNAseq613c3t <- CoGAPS(p.RNAseq613c3t, params = new("CogapsParams",
nPatterns=5))
projectR(data=p.ESepiGen4c11$mRNA.Seq, loadings=CR.RNAseq613c3t,
         dataNames = map.ESepiGen4c11[["GeneSymbols"]])
```

```

pca.RNAseq613c3t<-prcomp(t(p.RNAseq613c3t))
pca.ESepiGen4c11<-projectR(data=p.ESepiGen4c11$mRNA.Seq,
loadings=pca.RNAseq613c3t, dataNames = map.ESepiGen4c11[["GeneSymbols"]])

pca.RNAseq613c3t<-prcomp(t(p.RNAseq613c3t))
r.RNAseq613c3t<-rotatoR(1,1,-1,-1,pca.RNAseq613c3t$rotation[,1:2])
pca.ESepiGen4c11<-projectR(data=p.ESepiGen4c11$mRNA.Seq,
loadings=r.RNAseq613c3t, dataNames = map.ESepiGen4c11[["GeneSymbols"]])

c.RNAseq613c3t<-correlateR(genes="T", dat=p.RNAseq613c3t, threshtype="N",
threshold=10, absR=TRUE)
cor.ESepiGen4c11<-projectR(data=p.ESepiGen4c11$mRNA.Seq, loadings=c.RNAseq613c3t,
NP="PositiveCOR", dataNames = map.ESepiGen4c11[["GeneSymbols"]])

```

rotatoR

rotatoR

Description

a function for rotating two basis about a point or line in that plane

Usage

```
rotatoR(x1, y1, x2, y2, basisSET)
```

Arguments

x1	a value describing a the coordinate of a point in the first basis. If no values are provided for x2
y1	a value describing a the coordinate of a point in the second basis
x2	a value describing a the coordinate of the second point in the second basis
y2	a value describing a the coordinate of the second point in the second basis
basisSET	the basis to be rotated

Value

An object of class rotatoR.

Examples

```

pca.RNAseq613c3t<-prcomp(t(p.RNAseq613c3t))
r.RNAseq613c3t<-rotatoR(1,1,-1,-1,pca.RNAseq613c3t$rotation[,1:2])

```

rotatoR-class	<i>rotatoR</i>
---------------	----------------

Description

class of rotatoR output.

Slots

rotatedM rotated basis set (matrix) that is output of rotatoR function

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