Package 'TDbasedUFEadv'

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Type Package

Title Advanced package of tensor decomposition based unsupervised feature extraction

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Language en-US

Description This is an advanced version of TDbasedUFE, which is a comprehensive package to perform Tensor decomposition based unsupervised feature extraction.

In contrast to TDbasedUFE which can perform simple the feature selection and the multiomics analyses, this package can perform more complicated and advanced features, but they are not so popularly required. Only users who require more specific features can make use of its functionality.

biocViews GeneExpression, FeatureExtraction, MethylationArray, SingleCell, Software

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Imports TDbasedUFE, Biobase, GenomicRanges, utils, rTensor, methods, graphics, RTCGA, stats, enrichplot, DOSE, STRINGdb, enrichR, hash, shiny

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BugReports https://github.com/tagtag/TDbasedUFEadv/issues

URL https://github.com/tagtag/TDbasedUFEadv

Suggests knitr, rmarkdown, testthat (>= 3.0.0), RTCGA.rnaseq, RTCGA.clinical, BiocStyle, MOFAdata

VignetteBuilder knitr

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TDbasedUFEadv-package TDbasedUFEadv: Advanced package of tensor decomposition based unsupervised feature extraction

Description

This is an advanced version of TDbasedUFE, which is a comprehensive package to perform Tensor decomposition based unsupervised feature extraction. In contrast to TDbasedUFE which can perform simple the feature selection and the multiomics analyses, this package can perform more complicated and advanced features, but they are not so popularly required. Only users who require more specific features can make use of its functionality.

Author(s)

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See Also

Useful links:

- https://github.com/tagtag/TDbasedUFEadv
- Report bugs at https://github.com/tagtag/TDbasedUFEadv/issues

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computeSVD Title Perform SVD toward reduced matrix generated from a tensor with partial summation	th
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Description

Title Perform SVD toward reduced matrix generated from a tensor with partial summation

Usage

```
computeSVD(matrix1, matrix2, dim = 10L, scale = TRUE)
```

Arguments

matrix1	The first original matrix that generates a tensor
matrix2	The second original matrix that generates a tensor
dim	The number of singular value vectors to be computed
scale	If matrix should be scaled or not

Value

Singular value vectors attributed to two sets of objects associated with singular value vectors attributed to features, by multiplying

Examples

```
matrix1 <- matrix(runif(200),20)
matrix2 <- matrix(runif(400),20)
SVD <- computeSVD(matrix1,matrix2)</pre>
```

prepareCondDrugandDisease

Prepare condition matrix for expDrug

Description

Prepare condition matrix for expDrug

Usage

```
prepareCondDrugandDisease(expDrug)
```

Arguments

expDrug input gene expression profile

4 prepareCondTCGA

Value

Condition matrix for expDrug

Examples

```
library(RTCGA.rnaseq)
Cancer_cell_lines <- list(ACC.rnaseq,BLCA.rnaseq,BRCA.rnaseq)
Drug_and_Disease <- prepareexpDrugandDisease(Cancer_cell_lines)
Cond <- prepareCondDrugandDisease(Drug_and_Disease$expDrug)</pre>
```

prepareCondTCGA

Prepare Sample label for TCGA data

Description

Prepare Sample label for TCGA data

Usage

```
prepareCondTCGA(
   Multi_sample,
   Clinical,
   ID_column_of_Multi_sample,
   ID_column_of_Clinical
)
```

Arguments

```
Multi_sample list of sample ids

Clinical List of clinical data matrix from RTCGA.clinical

ID_column_of_Multi_sample

Column numbers used for conditions

ID_column_of_Clinical
```

Column numbers that include corresponding sample ids in clinical data

Value

list of sample labels

```
library(RTCGA.clinical)
library(RTCGA.rnaseq)
Clinical <- list(BLCA.clinical, BRCA.clinical, CESC.clinical, COAD.clinical)
Multi_sample <- list(
    BLCA.rnaseq[seq_len(100), 1, drop = FALSE],
    BRCA.rnaseq[seq_len(100), 1, drop = FALSE],</pre>
```

```
CESC.rnaseq[seq_len(100), 1, drop = FALSE],
  COAD.rnaseq[seq_len(100), 1, drop = FALSE]
)
ID_column_of_Multi_sample <- c(770, 1482, 773, 791)
ID_column_of_Clinical <- c(20, 20, 12, 14)
cond <- prepareCondTCGA(
  Multi_sample, Clinical,
  ID_column_of_Multi_sample, ID_column_of_Clinical
)</pre>
```

prepareexpDrugandDisease

Generating gene expression of drug treated cell lines and a disease cell line

Description

Generating gene expression of drug treated cell lines and a disease cell line

Usage

```
prepareexpDrugandDisease(Cancer_cell_lines)
```

Arguments

```
Cancer_cell_lines <- list(ACC.rnaseq,BLCA.rnaseq,BRCA.rnaseq) list that includes individual data set from RTCGA.rnaseq
```

Value

list of expDrug and expDisease

```
library(RTCGA.rnaseq)
Cancer_cell_lines <- list(ACC.rnaseq,BLCA.rnaseq,BRCA.rnaseq)
Drug_and_Disease <- prepareexpDrugandDisease(Cancer_cell_lines)</pre>
```

prepareTensorfromList Prepare tensor from a list that includes multiple profiles

Description

Prepare tensor from a list that includes multiple profiles

Usage

```
prepareTensorfromList(Multi, proj_dim)
```

Arguments

Multi a list that includes multiple profiles proj_dim the number of projection dimensions

Value

a tensor as a bundle of singular value vectors obtained by applying SVD to individual omics

Examples

```
library(MOFAdata)
data("CLL_data")
data("CLL_covariates")
Z <- prepareTensorfromList(CLL_data,10L)</pre>
```

```
prepareTensorfromMatrix
```

Generate tensor from two matrices

Description

Generate tensor from two matrices

Usage

```
prepareTensorfromMatrix(matrix1, matrix2)
```

Arguments

```
matrix1 the first input matrix
matrix2 the second input matrix
```

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Value

A tensor generated from the first and second matrices

Examples

```
Z <- prepareTensorfromMatrix(matrix(runif(100),10),matrix(runif(100),10))</pre>
```

prepareTensorRect

Prepare tensor generated from two matrices that share samples

Description

Prepare tensor generated from two matrices that share samples

Usage

```
prepareTensorRect(
  sample,
  feature,
  value,
  featureRange = GRanges(NULL),
  sampleData = list(NULL)
)
```

Arguments

sample Character vector of sample names feature list of features from two matrices

value array, contents of

featureRange Genomic Ranges to be associated with features sampleData List of conditional labeling associated with samples

Value

Tensor generated from two matrices that share samples

```
matrix1 <- matrix(runif(1000),200) #row features, column samples
matrix2 <- matrix(runif(2000),400) #row features, column samples
Z <- prepareTensorfromMatrix(t(matrix1),t(matrix2))
Z <- prepareTensorRect(sample=as.character(seq_len(50)),
feature=list(as.character(seq_len(200)),as.character(seq_len(400))),
sampleData=list(rep(seq_len(2),each=25)),value=Z)</pre>
```

8 selectFeatureProj

selectFeatureProj	Select feature when projection strategy is employed for the case where
	features are shared with multiple omics profiles

Description

Select feature when projection strategy is employed for the case where features are shared with multiple omics profiles

Usage

```
selectFeatureProj(
  HOSVD,
  Multi,
  cond,
  de = 1e-04,
  p0 = 0.01,
  breaks = 100L,
  input_all = NULL
)
```

Arguments

HOSVD	HOSVD
Multi	list of omics profiles, row: sample, column: feature
cond	list of conditions for individual omics profiles
de	initial value for optimization of standard deviation
p0	Threshold P-value
breaks	The number of bins of histogram of P-values
input_all	The number of selected feature. if null, interactive mode is activated

Value

list composed of logical vector that represent which features are selected and p-values

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Select features through the selection of singular value vectors

Description

Select features through the selection of singular value vectors

Usage

```
selectFeatureRect(
   SVD,
   cond,
   de = rep(1e-04, 2),
   p0 = 0.01,
   breaks = 100L,
   input_all = NULL
)
```

Arguments

SVD	SVD computed from matrix generated by partial summation of a tensor
cond	Condition to select singular value vectors
de	Initial values to be used for optimization of standard deviation
p0	Threshold value for the significance
breaks	Number of bins of histogram of P-values
input_all	The ID of selected singular value vectors. If it is null, interactive mode is activated.

Value

List of lists that includes P-vales as well as if individual features selected.

```
set.seed(0)
matrix1 <- matrix(runif(2000),200)
matrix2 <- matrix(runif(4000),200)
SVD <- computeSVD(matrix1,matrix2)
index_all <- selectFeatureRect(SVD,
list(NULL,rep(seq_len(2),each=5),rep(seq_len(2),each=10)),de=rep(0.5,2),input_all=1)</pre>
```

10 selectFeatureTransRect

selectFeatureTransRect

Select features for a tensor generated from two matrices that share samples.

Description

Select features for a tensor generated from two matrices that share samples.

Usage

```
selectFeatureTransRect(
  HOSVD,
  cond,
  de = rep(1e-04, 2),
  p0 = 0.01,
  breaks = 100L,
  input_all = NULL
)
```

mode

Arguments

HOSVD	HOSVD
cond	list of conditions
de	initial values for optimization of standard deviation
р0	threshold value for the significance
breaks	number of bins of the histogram of P-values
input_all	The selected singular value vectors attributed to samples. if NULL, interactive

Value

list of logical vector that represent if the individual features are selected and P-values.

```
library(TDbasedUFE)
set.seed(0)
matrix1 <- matrix(runif(1000),20) #row features, column samples
matrix2 <- matrix(runif(2000),40) #row features, column samples
Z <- prepareTensorfromMatrix(t(matrix1),t(matrix2))
Z <- prepareTensorRect(sample=as.character(seq_len(50)),
feature=list(as.character(seq_len(20)),as.character(seq_len(40))),
sampleData=list(rep(seq_len(2),each=25)),value=Z)
HOSVD <- computeHosvd(Z)
cond <- list(attr(Z,"sampleData")[[1]],NULL,NULL)
index_all <- selectFeatureTransRect(HOSVD,cond,de=c(0.1,0.1),
input_all=2,p0=1e-10)</pre>
```

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TensorRect-class

Class definitions

Description

Class definitions

Slots

```
sample character.
feature list.
value array.
featureRange GRanges.
sampleData list.
```

transSVD

Convert SVD to that for the case where samples are shared between two matrices

Description

Convert SVD to that for the case where samples are shared between two matrices

Usage

```
transSVD(SVD)
```

Arguments

SVD

input SVD object generated from computeSVD function

Value

converted SVD objects

```
matrix1 <- matrix(runif(200),20)
matrix2 <- matrix(runif(400),20)
SVD <- computeSVD(matrix1,matrix2)
SVD <- transSVD(SVD)</pre>
```

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