Package ‘GeneExpressionSignature’

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Title Gene Expression Signature based Similarity Metric
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Description This package gives the implementations of the gene expression signature and its distance to each. Gene expression signature is represented as a list of genes whose expression is correlated with a biological state of interest. And its distance is defined using a nonparametric, rank-based pattern-matching strategy based on the Kolmogorov-Smirnov statistic. Gene expression signature and its distance can be used to detect similarities among the signatures of drugs, diseases, and biological states of interest.

Depends R (>= 2.13), Biobase, PGSEA
Suggests apcluster,GEOquery
License GPL-2
LazyLoad yes
biocViews GeneExpression
NeedsCompilation no

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exampleSet

sample data, a subset of the C-MAP

Description

sample data, a subset of the C-MAP as , which is a collection of 50 genome-wide transcriptional expression data from cultured human cells treated with 15 different small molecules.

Usage
data(exampleSet)

Format

A ExpressionSet: assay data represents the 50 genome-wide transcriptional expression data, phenotypic data describes 15 different small molecules corresponds to the expression data (assay data).

References

http://www.sciencemag.org/content/313/5795/1929.short Lamb et al., The Connectivity Map: Using Gene-Expression Signatures to Connect Small Molecules, Genes, and Disease, science 2006

Examples
data(exampleSet)
getRLs

Convert gene expression profiles to a ranked list (data preprocessing)

**Description**

Sorting the microarray probe-set identifiers according to the differential expression values with respect to the untreated hybridization to obtain a ranked list. Gene-expression profiles in are represented in a nonparametric fashion.

**Usage**

```r
getRLs(control, treatment)
```

**Arguments**

- `control` a matrix, including the vehicle control gene expression profiles corresponding to the treatment gene expression profiles.
- `treatment` a matrix, is composed of gene expression profiles.

**Details**

The genes on the array are rank-ordered according to their differential expression relative to the control. First, control and treatment values less than a primary threshold value (quartile) were set to that threshold value. Finally, probe sets were ranked in descending order of d, where d is the ratio of the corresponding treatment-to-control values. For probe sets where d=1, a lower threshold was applied to the original difference values and a new treatment to control ratio (d') calculated. These probe sets were then sub-sorted in descending order of d.

**Value**

A matrix is composed of ranked lists, a ranked list represents the corresponding gene expression profiles.

**Examples**

```r
if (require(GEOquery)){
  #treatment gene-expression profiles
  GSM118720 <- getGEO(filename=system.file("extdata/GSM118720.soft",package="GeneExpressionSignature"))
  #control gene-expression profiles
  GSM118721 <- getGEO(filename=system.file("extdata/GSM118721.soft",package="GeneExpressionSignature"))
  #data ranking according to the different expression values
  control <- as.matrix(as.numeric(Table(GSM118721)[,2]))
  treatment <- as.matrix(as.numeric(Table(GSM118720)[,2]))
  ranked_list <- getRLs(control,treatment)
}
```
RankMerging

*Merging the ranker lists with the same labels of the biological states into a single list with the Iorio’s method.*

**Description**

Merging the assay data according to phenotypic data of the input ExpressionSet. Each group of the ranked lists with the same phenotypic data is aggregated into a single list, return it as an ExpressionSet object.

**Usage**

```
RankMerging(exprSet, MergingDistance = c("Spearman", "Kendall"), weighted = TRUE)
```

**Arguments**

- `exprSet`: an ExpressionSet object, each column of assay data represents a ranked list obtained by preprocessing the corresponding gene expression profile, and phenotypic data represents the short description (characteristics of gene expression profile, such as the drug type, the disease state) about the assay data.
- `MergingDistance`: distance to be used which "measures" the similarity of ordered lists, the default is "Spearman"
- `weighted`: there are two rank merging approaches for two cases: if `weighted=FALSE`, all ranked list with the same biological state are treated equally important, a simple but useful method average ranking technique is selected; otherwise, `weighted=TRUE`, each individual ranked lists has its own ranked weights, this takes the iterative rank-aggregating algorithm, default is TRUE.

**Details**

The krubor function is used in the aggregating procedure. And the following methods are used in the implementation: a measure of the distance between two ranked lists (Spearman’s Footrule), a method to merge two or more ranked lists the (Borda Merging Method), and a algorithm to obtain a single ranked list from a set of them in a hierarchical way (the Kruskal Algorithm). If choose Kendall as distance, the effectiveness of this function is certainly limited by the size of the merging problem.

**See Also**

`SignatureDistance`

**Examples**

```r
#load the sample expressionSet
data(exampleSet)

## Merging each group of the ranked lists in the exampleSet with the same phenotypic data into a single PRL
MergingSet=RankMerging(exampleSet,"Spearman",weighted=TRUE)
```
ScoreGSEA

Compute pairwise distances between samples with method in package GSEA

Description

Compute pairwise distances between sample according to their (Prototype Ranked List) PRL, a N x N distance matrix is generated by calling this function, N is the length of PRL.

Usage

ScoreGSEA(MergingSet, SignatureLength, ScoringDistance = c("avg", "max"), p.value = F)

Arguments

MergingSet an ExpressionSet object. The assay data represents the PRLs of the samples, each column represents one PRL. The number of sample of this argument must be greater than 1, otherwise, this function is not meaningful.

SignatureLength the length of "gene signature". In order to compute pairwise distances among samples, genes lists are ranked according to the gene expression ratio (fold change). And the "gene signature" includes the most up-regulated genes (near the top of the list) and the most down-regulated genes (near the bottom of the list).

ScoringDistance the distance measurements between PRLs: the Average Enrichment Score Distance $D_{avg} = \frac{(TES_{x,y} + TES_{y,x})}{2}$, and the Maximum Enrichment Score Distance $D_{max} = \min(TES_{x,y}, TES_{y,x})/2$. The $D_{avg}$ is more stringent than $D_{max}$, where $D_{max}$ is more sensitive to weak similarities, with lower precision but large recall.

p.value logical, if TRUE return a matrix of p.values of the distance matrix, default FALSE

Details

Once the PRL obtained for each sample, the distances between samples are calculated based on gene signature, including the expression of genes that seemed to consistently vary in response to the across different experimental conditions (e.g., different cell lines and different dosages). We take two distance measurements between PRLs: the Average Enrichment Score Distance $D_{avg} = \frac{(TES_{x,y} + TES_{y,x})}{2}$, and the Maximum Enrichment Score Distance $D_{max} = \min(TES_{x,y}, TES_{y,x})/2$. The $D_{avg}$ is more stringent than $D_{max}$, where $D_{max}$ is more sensitive to weak similarities, with lower precision but large recall.

Value

an distance-matrix, the max distance is more sensitive to weak similarities, providing a lower precision but a larger recall.

If p.value is set to TRUE, then a list is returned that consists of the distance matrix as well as their p.values, otherwise, without p.values in the result.

See Also

ScorePGSEA, SignatureDistance
Examples

```r
# load the sample expressionSet
data(exampleSet)

# Merging each group of the ranked lists in the exampleSet with the same phenotypic data into a single PRL
MergingSet=RankMerging(exampleSet,"Spearman")

# get the distance matrix
ds=ScoreGSEA(MergingSet,250,"avg")
```

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**ScorePGSEA**

*Compute pairwise distances between samples with method in package PGSEA*

**Description**

Compute pairwise distances between sample according to their (Prototype Ranked List) PRL, get a N x N distance matrix is generated by calling this function, N is the length of PRL.

**Usage**

```r
ScorePGSEA(MergingSet, SignatureLength, ScoringDistance = c("avg", "max"), p.value = F)
```

**Arguments**

- **MergingSet**: an ExpressionSet object. The assay data represents the PRLs of the samples, each column represents one PRL. The number of sample must be greater than 1, otherwise, this function is not meaningful.
- **SignatureLength**: the length of "gene signature". In order to compute pairwise distances among samples, genes lists are ranked according to the gene expression ratio (fold change). And the "gene signature" includes the most up-regulated genes (near the top of the list) and the most down-regulated genes (near the bottom of the list).
- **ScoringDistance**: the distance measurements between PRLs: the Average Enrichment Score Distance (avg), or the Maximum Enrichment Score Distance (max).
- **p.value**: logical, if TRUE return a matrix of p.values of the distance matrix, default FALSE

**Details**

This function has the same function with ScoreGSEA, just with different methods.

**See Also**

- ScoreGSEA
- SignatureDistance
**SignatureDistance**

**Examples**

```r
# load the sample expressionSet
data(exampleSet)

# Merging each group of the ranked lists in the exampleSet with the same phenotypic data into a single PRL
MergingSet=RankMerging(exampleSet,"Spearman")

# get the distance matrix
ds=ScorePGSEA(MergingSet,250, ScoringDistance="avg")
```

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**SignatureDistance**  
Compute pairwise distances comprehensively.

**Description**

This function integrated the function for rank merging and distance scoring, we can do the rank merging and distance scoring simply with it.

**Usage**

```r
SignatureDistance(exprSet, SignatureLength, MergingDistance = c("Spearman", "Kendall"), ScoringMethod = c("GSEA", "PGSEA"), ScoringDistance = c("avg", "max"), weighted = TRUE, ...)
```

**Arguments**

- `exprSet`  
an ExpressionSet object, each column of assay data represents a ranked list obtained by preprocessing the corresponding gene expression profile, and phenotypic data represents the short description (characteristics of gene expression profile, such as the drug type, the disease state) about the assay data.

- `SignatureLength`  
the length of "gene signature". In order to compute pairwise distances among samples, genes lists are ranked according to the gene expression ratio (fold change). And the "gene signature" includes the most up-regulated genes (near the top of the list) and the most down-regulated genes (near the bottom of the list).

- `MergingDistance`  
distance to be used which "measures" the similarity of ordered lists, Spearman or Kendall

- `ScoringMethod`  
method to be used to perform distance scoring, GSEA or PGSEA

- `ScoringDistance`  
the distance measurements between PRLs: the Average Enrichment Score Distance (avg), and the Maximum Enrichment Score Distance (max).

- `weighted`  
there are tow rank merging approaches for two cases: if weighted=FALSE, all ranked list with the same biological state are treated equally important, a simple but useful method average ranking technique is selected; otherwise, weighted=TRUE, each individual ranked lists has its own ranked weights, this takes the iterative rank-aggregating algorithm, default is TRUE.

- `...`  
additional arguments can be passed to the internal procedures

**See Also**

`RankMerging, ScoreGSEA, ScorePGSEA`
Examples

#load the sample expressionSet
data(exampleSet)

#distance scoring
SignatureDistance(exampleSet, SignatureLength=250, MergingDistance="Spearman", ScoringMethod="GSEA", ScoringDistance="avg", weighted=TRUE)
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