Package ‘RTopper’
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Type Package

Title This package is designed to perform Gene Set Analysis across multiple genomic platforms

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Author Luigi Marchionni <marchion@jhu.edu>, Svitlana Tyekucheva <svitlana@jimmy.harvard.edu>

Maintainer Luigi Marchionni <marchion@jhu.edu>

Depends R (>= 2.11.0), Biobase

Imports limma, multtest

Suggests limma, org.Hs.eg.db, KEGG.db, GO.db

Description the RTopper package is designed to perform and integrate gene set enrichment results across multiple genomic platforms.

License GPL (>= 3)

biocViews Microarray

LazyLoad yes

NeedsCompilation no

R topics documented:

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Description

Gene sets analysis considers whether genes sharing a biological property also behave in a related way in experimental data. This technique is commonly used in high throughput genomic analyses to assist results interpretation, and has been successfully applied in cancer genome projects for integrating information from multiple genome-wide assays. The RTopper package uses gene sets analysis to overcome the diversity of genomic data providing the statistical framework for integration across data types.

Details

- Package: RTopper
- Type: Package
- Version: 0.1
- Date: 2010-11-12
- License: GPL version 3 or newer

RTopper package features

Rtopper enables two gene set-based data integration approaches:

1. Integration+GSA: computing integrated gene-to-phenotype association scores, followed by conventional gene sets analysis;
2. GSA+Integration: computing consensus significance score after all data types are analyzed individually;
3. Use of alternative enrichment test: RTopper uses the Wilcoxon rank-sum test for enrichment testing, however alternative tests can be defined and used;
4. Multiple testing correction: RTopper enables adjustment of p-values obtained from enrichment analysis;

Author(s)

Luigi Marchionni <marchion@jhu.edu>

References

adjustPvalGSE

Description
This function is an interface to the \texttt{mt.rawp2adjp} function contained in the multtest package. adjustPvalGSE works on outputs from \texttt{runBatchGSE} and \texttt{combineGSE} returning adjusted p-values

Usage
adjustPvalGSE(gseOut, proc = "BH", alpha = 0.05 , na.rm = FALSE)

Arguments
- \texttt{gseOut}: list of lists, either the output from \texttt{runBatchGSE}, or the output from \texttt{combineGSE}
- \texttt{proc}: character, the method to be used for p-values adjusting. This parameter will be passed to the \texttt{mt.rawp2adjp} function from the multtest package. The available options include: "Bonferroni", "Holm", "Hochberg", "SidakSS", code"SidakSD", "BH" (the default), "BY", "ABH", and "TSBH"
- \texttt{alpha}: numeric, the nominal type I error rate
- \texttt{na.rm}: logical, the option for handling NA values in the list of raw p-values

Details
The adjustPvalGSE function performs p-value adjusting for multiple testing correction on the list of lists resulting from enrichment analysis obtained using the \texttt{runBatchGSE} and \texttt{combineGSE} functions. This functions is based on the \texttt{mt.rawp2adjp} function contained in the multtest package.

Value
For each vector of p-value contained in the gseOut input object a data.frame is returned, containing original p-value and corrected p-values

Author(s)
Luigi Marchionni <marchion@jh.edu>

References

Examples
```r
###load gse analysis results for separate gene-to-phenotype score
data(gseResultsSep)

###adjust for multiple testing using the Benjamini and Hochberg method
gseABS.int.BH <- adjustPvalGSE(gseResultsSep)

###adjust for multiple testing using the Holm method
```
combineGSE

Description

Combines GSE results obtained from a separate set of gene-to-phenotypes scores

Usage

combineGSE(gseOut, method)

Arguments

- **gseOut**: a list of lists containing the enrichment results to be combined. This is usually the output of `runBatchGSE` obtained from a set of distinct gene-to-phenotype scores (usually one per genomic platform)
- **method**: character, this argument specifies the method used to combine the enrichment results obtained from distinct gene-to-phenotype scores (usually one per genomic platform). Available options are the computation of the geometric or arithmetic means, the use of the median, the selection of the minimum or the maximum enrichment score, and the random selection of a score (respectively "geometricMean", "mean", "median", "min", "max", and "random")

Details

This function summarizes enrichment results obtained from distinct gene-to-phenotypes scores (usually one per genomic platform) by one of several alternative methods.

Value

The output is a list of lists containing integrated enrichment results for all FGS collections

Author(s)

Luigi Marchionni <marchion@jhu.edu>

References


Examples

```r
# load gse analysis results for separate gene-to-phenotype score data(gseResultsSep)

# combine enrichment score results using geometric mean
gseABS.sep.geoMean <- combineGSE(gseResultsSep, method="geometricMean")
```
computeDrStat

```r
gseABS.sep.max <- combineGSE(gseResultsSep, method="max")
```

### Description

`computeDrStat` computes gene-to-phenotype association scores, using as input the output from `convertToDr`.

#### Usage

```r
computeDrStat(data, columns = c(1:(ncol(data)-1)), method = "dev", integrate = TRUE)
```

#### Arguments

- **data**: a list of data.frames containing genomic measurements. Each element of `dataIntersection` must account for the same set of patients(columns) and genes (rows).
- **columns**: a data.frame indicating patients' phenotypic class.
- **method**: character, the number of genomic platforms.
- **integrate**: logical, whether to integrate the gene-to-phenotype scores across platform or return separates scores for each platform.

#### Details

This function allows computing gene-to-phenotype association scores, using as input the gene-centered list produced by `computeDr`. The `computeDrStat` function works separately on each gene-centered data.frame created by the `convertToDr` function, assuming that the phenotype information is stored in the last column named "response". It is possible computing both separate association scores for each platform, as well as an integrated score, as specified by the `integrate` arguments. There are currently three methods available for obtaining the scores (see Tyekucheva et al, manuscript under review), as specified by the `methods` argument:

- "dev": this approach computes the score as the difference of deviances;
- "aic": this approach computes the score as the Akaike information criterion for model selection;
- "bic": this approach computes the score as the penalized likelihood ratio;

#### Value

A list of named vectors containing separate or integrated gene-to-phenotype association scores.

#### Author(s)

Luigi Marchionni <marchion@jhu.edu>

#### References

### load data
data(exampleData)

### convert
dataDr <- convertToDr(dat, pheno, 4)

### compute the integrated score
bicStatIntegrated <- computeDrStat(dataDr, columns = c(1:4), method="bic", integrate = TRUE)

### compute separate scores for each genomic platform
bicStatSeparate <- computeDrStat(dataDr, columns = c(1:4), method="bic", integrate = FALSE)

---

**convertToDr**

**Converts genomic data to a list suitable for computing gene-to-phenotype scores**

**Description**

convertToDr converts genomic data into a list further used for computing gene-to-phenotype association scores.

**Usage**

```r
convertToDr(dataIntersection, response, nPlatforms = length(data))
```

**Arguments**

- `dataIntersection`  
a list of data.frames containing genomic measurements. Each element of dataIntersection must account for the same set of patients(columns) and genes (rows)
- `response`  
a data.frame indicating patients’ phenotypic class
- `nPlatforms`  
umeric, the number of genomic platforms

**Details**

This function converts a list of data.frames containing distinct genomic measurements performed on the same patients into a gene-centered used in further analyses for computing gene-to-phenotype scores. Data.frame in the input list (dataIntersection) must have the same dimensions, with columns being patients, and rows being genes. Column names identify the patients, while rownames identify the genes. The argument response is used to pass phenotypic information about samples to be analyzed. This is a simple two columns data.frame in which the first column correspond to patients identifiers, and the second column to the phenotypic response encoded as binary class (using the integers 0 and 1). The nPlatforms argument specifies the number of platforms that will be analyzed.

**Value**

A list of data.frames, one for each analyzed gene, summarizing all genomic measurements and phenotypic information across patients and platforms.
dat

Author(s)
Luigi Marchionni <marchion@jhu.edu>

References
Svitlana Tyekucheva, Luigi Marchionni, Rachel Karchin, and Giovanni Parmigiani "Integrating diverse genomic data using gene sets." Manuscript submitted.

Examples

### load data
data(exampleData)

### convert
dataDr <- convertToDr(dat, pheno, 4)

dat A test dataset for the RTopper package

Description
A small subset of pre-processed Glioblasoma Multiforme (GBM) genomic data from The Cancer Genome Atlas (TCGA) project, encompassing Differential Gene Expression (DGE), and Copy Number Variation (CNV). Can be used as input to convertToDr.

Usage
data(exampleData)

Format
This object is a list of 4 data.frames containing genomic measurements obtained across distinct genomic scopes (copy number variation and gene expression), platforms (Affymetrix and Agiles), and laboratories. In particular each data.frame consist of 500 gene measurements (by rows), for 95 distinct patients (by columns) from the following 4 distinct platforms:
"dat.affy": DGE obtained using Affymetrix microarrays;
"dat.agilent": DGE obtained using Agilent microarrays;
"dat.cnvHarvard": CNV data obtained at Harvard;
"dat.cnvMskcc": CNV data obtained at Memorial Sloan Kettering Cancer Center;

Source
The Cancer Genome Atlas (TCGA) project http://cancergenome.nih.gov/

References
fgsList

A list of Functional Gene Set (FGS) to be used to run the examples in the RTopper package

Description

A list containing distinct types of FGS (i.e. Gene Ontology, KEGG pathways). Each FGS is type is a list of named character vectors, one for each FGS, containing the gene identifiers. Vectors names describe the FGS. Can be used as input to runBatchGSE.

Usage

data(fgsList)

Format

This object is a list of length two:

"go": this is a list of 5 character vectors, corresponding to 5 distinct Gene Ontology (GO) terms. Genes annotated to each GO term are identified by their gene symbol;

"kegg": this is a list of 5 character vectors, corresponding to 5 distinct KEGG pathways. Genes annotated to each KEGG pathway are identified by their gene symbol;

Source

The FGS were obtained from the org.Hs.eg.db package, (use the org.Hs.eg function to see the content); These FGS were annotated using data from GO.db and KEGG.db packages (use the GO and KEGG functions to see the content).

Examples

data(fgsList)
class(fgsList)
names(fgsList)
str(fgsList)
**gseResultsSep**

*A list of separated gene set enrichment p-values to be used to run the examples in the RTopper package*

---

**Description**

A list containing distinct named numeric vectors corresponding to the gene set enrichment p-value separately computed with `runBatchGSE` for each distinct data set. Note that for each data set there are two set of p-values, one for GO and one for KEGG. These separate p-values can be combined across data sets by the `combineGSE` function. Can be used as input to `adjustPvalGSE`.

**Usage**

```r
data(gseResultsSep)
```

**Format**

This object is a list of length four:

- "dat.affy": a list of length two: "go" is a numeric vector of length 5, containing the p-values resulting from gene set enrichment analysis of 5 GO terms on Affymetrix gene expression data;  
  "keg" is a numeric vector of length 5, containing the p-values resulting from gene set enrichment analysis of 5 KEGG pathways on Affymetrix gene expression data;

- "dat.agilent": a list of length two: "go" is a numeric vector of length 5, containing the p-values resulting from gene set enrichment analysis of 5 GO terms on Agilent gene expression data;  
  "keg" is a numeric vector of length 5, containing the p-values resulting from gene set enrichment analysis of 5 KEGG pathways on Agilent gene expression data;

- "dat.cnvHarvard": a list of length two: "go" is a numeric vector of length 5, containing the p-values resulting from gene set enrichment analysis of 5 GO terms on Harvard CNV data;  
  "keg" is a numeric vector of length 5, containing the p-values resulting from gene set enrichment analysis of 5 KEGG pathways on Harvard CNV data;

- "dat.cnvMskcc": a list of length two: "go" is a numeric vector of length 5, containing the p-values resulting from gene set enrichment analysis of 5 GO terms on MSKCC CNV data;  
  "keg" is a numeric vector of length 5, containing the p-values resulting from gene set enrichment analysis of 5 KEGG pathways on MSKCC CNV data;

**Source**

Computed using the `runBatchGSE` function from the TCGA data contained in `sepScores` and `fgsList`.

**Examples**

```r
data(gseResultsSep)
class(gseResultsSep)
names(gseResultsSep)
str(gseResultsSep)
```
intScores

A list of genomic scores integrated across distinct data sets to be used to run the examples in the RTopper package

Description

A list containing a named numeric vector corresponding to the genomic score resulting from the integration across distinct data set. These integrated gene-to-phenotype scores are computed by the `computeDrStat` function. Can be used as input to `runBatchGSE`.

Usage

data(intScores)

Format

This object is a list of length one:

"integrated": a numeric vector of length 500, corresponding to the integrated phenotype association scores computed for each of the 500 genes used in the examples;

Source

Computed using the `computeDrStat` function from the TCGA data contained in `dat` and `pheno`.

Examples

data(intScores)
class(intScores)
names(intScores)
str(intScores)

pheno

A test dataset for the RTopper package

Description

A data.frame with 2 columns containing the phenotypic class indicator for the 95 patients analyzed and used in the examples. Can be used as input to `convertToDr`.

Usage

data(exampleData)

Format

This object is a data.frame with two columns:

"Sample": the first column contains the patients identifiers;

"Class": the second columns contain a numeric indicator (0 or 1) corresponding to the phenotypic class of each patient;
runBatchGSE

Source


References


Examples

data(exampleData)
class(pheno)
colnames(pheno)
str(pheno)

```
runBatchGSE(dataList, fgsList, ...)
```

Arguments

dataList  a list containing the gene-to-phenotype scores to be used as ranking statistics in the GSE analysis. This list is usually produced by running `computeDrStat`
fgsList  a list of FGS collection, in which each element is a list of character vectors, one for each gene set
...

Details

This function performs enrichment analysis for all the gene-to-phenotype scores (argument `dataList`) passed to it over a list of Functional Gene Set (FGS) (argument `fgsList`), returning a p-value for each FGS. Additional arguments can be passed to this function to modify the way the enrichment test is performed, as follows:

* `absolute` logical, this specifies whether the absolute values of the ranking statistics should be used in the test (the default being TRUE)
gseFunc: a function to perform GSE analysis. If not specified the default is the `geneSetTest` function from the limma package. If a function is specified by the user, the membership of the analyzed genes to a FGS, and the ranking statistics must be defined in the same way this is done for `geneSetTest`, and the new function must return an integer (usually a p-value) (see the help for `geneSetTest`.

The following main arguments are used by `geneSetTest`:

- **type** character, specifies the type of statistics used to rank the genes by `geneSetTest`: 'f' for F-like statistics (default), 't' for t-like statistics, or 'auto' for an educated guess
- **alternative** character, defines the alternative with the following possible options: 'mixed' (default), 'either', 'up' or 'down', 'two.sided', 'greater', or 'less'
- **ranks.only** logical, if TRUE (default) only ranks will be used by `geneSetTest`
- **nsim** numeric, the number of randomly selected sets of genes to be used in simulations to compute the p-value

**Value**

The output is a list of lists containing the set of enrichment results for all gene-to-phenotype scores and FGS collections used as input.

**Author(s)**

Luigi Marchionni <marchion@jhu.edu>

**References**


**Examples**

```r
### require limma to run the example
require(limma)

### load integrated gene-to-phenotype scores
data(intScores)

### load separate gene-to-phenotype scores
data(sepScores)

### load list of functional gene sets
data(fgsList)

### run GSE analysis in batch with default parameters
gseABS.int <- runBatchGSE(dataList=intScores, fgsList=fgsList)

### run GSE analysis in batch with alternative parameters
gseABS.sep <- runBatchGSE(dataList=sepScores, fgsList=fgsList, absolute=FALSE, type="t", alternative="up")

### run GSE analysis in batch passing an enrichment function
gseUP.int.2 <- runBatchGSE(dataList=intScores, fgsList=fgsList, absolute=FALSE, gseFunc=wilcoxGST, alternative="up")

### define and use a new enrichment function
```
sepScores <- function (selected, statistics, threshold) {
  diffExpGenes <- statistics > threshold
  tab <- table(diffExpGenes, selected)
  pVal <- fisher.test(tab)["p.value"]
}
gseUP.sep.2 <- runBatchGSE(dataList=sepScores, fgsList=fgsList,
                           absolute=FALSE, gseFunc=gseFunc, threshold=7.5)

Description

A list containing distinct named numeric vectors corresponding to the gene-to-phenotype association scores resulting from the separate analysis of each data set. These separate gene-to-phenotype scores are computed by computeDrStat function. Can be used as input to runBatchGSE.

Usage

data(sepScores)

Format

This object is a list of length four, one element for data set:
"dat.affy": a numeric vector of length 500, corresponding to the separate phenotype association scores computed for Affymetrix gene expression data;
"dat.agilent": a numeric vector of length 500, corresponding to the separate phenotype association scores computed for Agilent gene expression data;
"dat.cnvHarvard": a numeric vector of length 500, corresponding to the separate phenotype association scores computed for Harvard CNV data;
"dat.cnvMskcc": a numeric vector of length 500, corresponding to the separate phenotype association scores computed for the MSKCC CNV data;

Source

Computed using the computeDrStat function from the TCGA data contained in data.

Examples

data(sepScores)
class(sepScores)
names(sepScores)
str(sepScores)
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