

The RTopper package: perform run Gene Set Enrichment across genomic platforms

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1 Overview

Gene Set Enrichment (GSE) analysis has been widely use to assist the interpretation of gene expression data. We propose here to apply GSE for the integration of genomic data obtained from distinct analytical platform.

In the present implementation of the **RTopper** GSE analysis is performed using the `geneSetTest` function from the **limma** package [6, 5, 7]. This function enables testing the hypothesis that a specific set of genes (a Functional Gene Set, FGS) is more highly ranked on a given statistics. In

particular this functions computes a p-value for each FGS by one or two-sided Wilcoxon rank-sum test. Alternative user-defined functions can also be used.

Furthermore multiple hypothesis testing correction is achieved by applying the Benjamini and Hochberg method [2] as implemented in the `multtest` R/Bioconductor package. Overall, this approach is conceptually analogous to Gene Set Enrichment Analysis (GSEA), as proposed by Mootha and colleagues [4, 8].

The integration can be achieved through two distinct approaches:

1. **GSE + INTEGRATION**: Separate GSE analysis on the individual genomic platforms followed by GSE results integration;
2. **INTEGRATION + GSE**: Integration of genomic data measurement using a logistic model followed by GSE analysis;

2 RTopper data structure

In this tutorial we demonstrate the functionality of `RTopper` package. To this end we will make use of simplified data generated within The Cancer Genome Atlas (TCGA) project, using Glioblastoma Multiforme (GBM) genomics data obtained from the same patients' cohort using distinct platforms, including Differential Gene Expression (DGE), Copy Number Variation (CNV), and Differential Methylation (DM). This data is included with the `RTopper` package as the dataset `exampleData`, which consists of genomic measurements (the list `dat`) for 500 genes (in rows) and 95 patients (in columns) from 4 distinct platforms:

1. DGE obtained using Affymetrix;
2. DGE obtained using Agilent;
3. CNV data generated at Harvard;
4. CNV data generated at the MSKCC;

The phenotypic class for each patient is defined in the a data.frame `pheno` consisting of 95 rows (patients, `pheno$Sample`) and 2 columns, the first being patients identifiers, and the second variable giving the group indicator (`pheno$Class`).

To load the data set type `data(exampleData)`, and to view a description of this data type `?exampleData`. The structure of the data is shown below:

```
> library(RTopper)
> data(exampleData)
> ls()

[1] "dat"      "pheno"

> class(dat)

[1] "list"

> names(dat)

[1] "dat.affy"      "dat.agilent"
[3] "dat.cnvHarvard" "dat.cnvMskcc"
```

```

> sapply(dat,class)

      dat.affy      dat.agilent dat.cnvHarvard
"data.frame"  "data.frame"    "data.frame"
dat.cnvMskcc
"data.frame"

> sapply(dat,dim)

      dat.affy dat.agilent dat.cnvHarvard
[1,]      500      500      500
[2,]      95      95      95
      dat.cnvMskcc
[1,]      500
[2,]      95

> dim(pheno)

[1] 95  2

> str(pheno)

'data.frame':      95 obs. of  2 variables:
 $ Sample: chr   "TCGA.02.0003" "TCGA.02.0007" "TCGA.02.0011" "TCGA.02.0021" ...
 $ Class : int    0 0 1 1 0 0 0 0 0 0 ...

```

In summary to perform the analysis with functions from **RTopper** the genomic data used as input must be in the following format:

1. **Genomic measurements:** a list of data.frames, in which each list item corresponds to a genomic platform, and comprises a data.frame with rows being genes and columns patients;
2. **Phenotype data:** a data.frame with 2 columns: patients and their phenotypes;
3. The number of columns of the *Genomic measurements* data.frames must match the number of rows of the *Phenotype data*;
4. The same set of genes must be measured in each platform and gene labels must be stored as rownames;

Below are shown the first 6 rows and 4 columns of each data.frame contained in **dat**, which share the same genes (shown for some of the possible combinations). Similarly column names in the **dat** data.frames correspond to rownames of **pheno**.

```

> ###data structure
> lapply(dat,function(x) head(x)[,1:3])

$dat.affy
      TCGA.02.0003 TCGA.02.0007 TCGA.02.0011
AACS      7.747995      7.685409      7.535661
AARS      9.381544      9.930156     10.197194
ABI1      8.173255      8.962803      9.895811
ACHE      5.127197      4.547297      5.146552
ACTC1     6.612645      5.825879      8.067945
ACTN2     6.257383      5.330557      5.842319

```

```

$dat.agilent
      TCGA.02.0003 TCGA.02.0007 TCGA.02.0011
AACS      -1.0070000    -1.1164000    -0.913000
AARS      -1.2665000    -0.8981250     0.263500
ABI1      -0.2765000     0.3356250     1.027250
ACHE       0.4403750    -0.0222500     0.115000
ACTC1      0.3641538     0.1234615     1.046692
ACTN2      4.3348000     2.2278000     3.330600

$dat.cnvHarvard
      TCGA.02.0003 TCGA.02.0007 TCGA.02.0011
AACS      -0.08273213  -0.08917331  -0.02075644
AARS      -0.10233281  -0.20620608  -0.05157664
ABI1      -0.86886659  -0.01214599   0.59307754
ACHE       0.31560002  -1.00166150  -0.14519639
ACTC1     -1.17495078  -0.26698279  -0.95662761
ACTN2     -0.11319016  -0.09657971   0.02582138

$dat.cnvMskcc
      TCGA.02.0003 TCGA.02.0007 TCGA.02.0011
AACS      -0.0383875   -0.09140000   0.008233333
AARS       0.0075600    0.02801667   0.104850000
ABI1      -0.7006900    0.21270000   0.499472727
ACHE       0.8676000   -0.23970000   0.075000000
ACTC1     -0.9779500   -0.11625000  -0.692950000
ACTN2     -0.1258571   -0.05394444   0.010200000

> sum(rownames(dat[[1]])%in%rownames(dat[[2]]))

[1] 500

> sum(rownames(dat[[2]])%in%rownames(dat[[3]]))

[1] 500

```

2.1 Creation of Functional Gene Sets

Functional Gene Sets (FGS) are list of genes that share a specific biological function. Examples of FGS are genes that operate in the same signaling pathway (*i.e.* Notch signaling genes), or that share the same biological function (*i.e.* Cell adhesion genes). FGS can be retrieved from various database, or can be constructed *ad hoc*. A convenient source of FGS are the R-Bioconductor metaData packages, and S4 classes and methods for handling FGS are provided by the **GSEABase** package. Below is shown a simple way to extract FGS from the human genome metaData package **org.Hs.eg.db**. As a general rule the name of the metaData package, without the **.db** extension, can be used a function to see the content of the package, as shown below:

```

> library(org.Hs.eg.db)
> org.Hs.eg()

```

Quality control information for org.Hs.eg:

This package has the following mappings:

org.Hs.egACCCNUM has 41961 mapped keys (of 56340 keys)
org.Hs.egACCCNUM2EG has 750930 mapped keys (of 750930 keys)
org.Hs.egALIAS2EG has 113648 mapped keys (of 113648 keys)
org.Hs.egCHR has 55858 mapped keys (of 56340 keys)
org.Hs.egCHRLNGTHS has 93 mapped keys (of 93 keys)
org.Hs.egCHRLLOC has 23584 mapped keys (of 56340 keys)
org.Hs.egCHRLLOCEND has 23584 mapped keys (of 56340 keys)
org.Hs.egENSEMBL has 26404 mapped keys (of 56340 keys)
org.Hs.egENSEMBL2EG has 28423 mapped keys (of 28423 keys)
org.Hs.egENSEMBLPROT has 19616 mapped keys (of 56340 keys)
org.Hs.egENSEMBLPROT2EG has 98264 mapped keys (of 98264 keys)
org.Hs.egENSEMBLTRANS has 20464 mapped keys (of 56340 keys)
org.Hs.egENSEMBLTRANS2EG has 154732 mapped keys (of 154732 keys)
org.Hs.egENZYME has 2230 mapped keys (of 56340 keys)
org.Hs.egENZYME2EG has 975 mapped keys (of 975 keys)
org.Hs.egGENENAME has 56340 mapped keys (of 56340 keys)
org.Hs.egGO has 18585 mapped keys (of 56340 keys)
org.Hs.egGO2ALLEGS has 19541 mapped keys (of 19541 keys)
org.Hs.egGO2EG has 15283 mapped keys (of 15283 keys)
org.Hs.egMAP has 36078 mapped keys (of 56340 keys)
org.Hs.egMAP2EG has 2367 mapped keys (of 2367 keys)
org.Hs.egOMIM has 16072 mapped keys (of 56340 keys)
org.Hs.egOMIM2EG has 20279 mapped keys (of 20279 keys)
org.Hs.egPATH has 5869 mapped keys (of 56340 keys)
org.Hs.egPATH2EG has 229 mapped keys (of 229 keys)
org.Hs.egPMID has 33435 mapped keys (of 56340 keys)
org.Hs.egPMID2EG has 452385 mapped keys (of 452385 keys)
org.Hs.egREFSEQ has 40744 mapped keys (of 56340 keys)
org.Hs.egREFSEQ2EG has 268751 mapped keys (of 268751 keys)
org.Hs.egSYMBOL has 56340 mapped keys (of 56340 keys)
org.Hs.egSYMBOL2EG has 56332 mapped keys (of 56332 keys)
org.Hs.egUCSCCKG has 22864 mapped keys (of 56340 keys)
org.Hs.egUNIGENE has 25400 mapped keys (of 56340 keys)
org.Hs.egUNIGENE2EG has 27472 mapped keys (of 27472 keys)
org.Hs.egUNIPROT has 19170 mapped keys (of 56340 keys)

Additional Information about this package:

DB schema: HUMAN_DB
DB schema version: 2.1
Organism: Homo sapiens
Date for NCBI data: 2015-Mar17

```

Date for GO data: 20150314
Date for KEGG data: 2011-Mar15
Date for Golden Path data: 2010-Mar22
Date for Ensembl data: 2015-Mar13

```

For instance the `org.Hs.egG02ALLEGS` environment contains the mapping of all ENTREZ Gene identifiers to the **Gene Ontology Terms** [1], while `org.Hs.egPATH2EG` maps the identifiers to **KEGG** pathways [3]. The corresponding lists of FGS can be retrieve from the corresponding environments using the the R command `as.list()`, as shown below for KEGG and GO:

```

> kegg <- as.list(org.Hs.egPATH2EG)
> go <- as.list(org.Hs.egG02ALLEGS)
> length(kegg)

[1] 229

> length(go)

[1] 19541

> str(kegg[1:5])

List of 5
 $ 04610: chr [1:69] "2" "462" "623" "624" ...
 $ 00232: chr [1:7] "9" "10" "1544" "1548" ...
 $ 00983: chr [1:52] "9" "10" "978" "1066" ...
 $ 01100: chr [1:1130] "9" "10" "15" "18" ...
 $ 00380: chr [1:42] "15" "26" "38" "39" ...

> names(kegg)[1:5]

[1] "04610" "00232" "00983" "01100" "00380"

> str(go[1:5])

List of 5
 $ G0:0000002: Named chr [1:32] "291" "1763" "1890" "3980" ...
 ..- attr(*, "names")= chr [1:32] "TAS" "IDA" "TAS" "IEA" ...
 $ G0:0000003: Named chr [1:1029] "18" "49" "49" "49" ...
 ..- attr(*, "names")= chr [1:1029] "IEA" "IEA" "IMP" "ISS" ...
 $ G0:0000011: Named chr "64145"
 ..- attr(*, "names")= chr "IBA"
 $ G0:0000012: Named chr [1:9] "3981" "7141" "7515" "23411" ...
 ..- attr(*, "names")= chr [1:9] "IDA" "IDA" "IEA" "IMP" ...
 $ G0:0000018: Named chr [1:59] "604" "641" "641" "940" ...
 ..- attr(*, "names")= chr [1:59] "IEA" "IEA" "IMP" "IEA" ...

> names(go)[1:5]

[1] "G0:0000002" "G0:0000003" "G0:0000011"
[4] "G0:0000012" "G0:0000018"

```

In the `kegg` list genes are identified by their ENTREZ Gene identifiers, while in the `dat` genes are identified by their Gene Symbol. Below is an example of the code that can be used to perform the identifiers conversion, using only a subset of KEGG and GO FGS:

```
> kegg <- lapply(kegg[sample(1:length(kegg),5)],function(x) unique(unlist(mget(x,org.Hs.egSYMBOL))))
> go <- lapply(go[sample(1:length(go),5)],function(x) unique(unlist(mget(x,org.Hs.egSYMBOL))))
> str(kegg)
```

List of 5

```
$ 04920: chr [1:68] "ACACB" "AGRP" "AKT1" "AKT2" ...
$ 05131: chr [1:61] "ABL1" "ACTB" "ACTG1" "RHOG" ...
$ 04012: chr [1:87] "ABL1" "ABL2" "AKT1" "AKT2" ...
$ 00450: chr [1:17] "CCBL1" "CTH" "MARS" "MTR" ...
$ 00240: chr [1:99] "NUDT2" "POLR3D" "CAD" "ENTPD1" ...
```

```
> str(go)
```

List of 5

```
$ GO:0005968: chr [1:4] "CHM" "CHML" "RABGGTA" "RABGGTB"
$ GO:0089712: chr [1:4] "SLC1A6" "SLC25A12" "SLC25A13" "SLC38A7"
$ GO:0000184: chr [1:119] "EIF4G1" "ETF1" "FAU" "GSPT1" ...
$ GO:0035090: chr [1:7] "ANK1" "LLGL1" "LIN7A" "PDZD11" ...
$ GO:0030522: chr [1:289] "ACTN4" "NROB1" "AHR" "ALOX15" ...
```

Finally, it is also possible to annotate FGS, mapping pathways identifiers to pathway names, as shown below for KEGG, using the KEGG.db.

```
> library(KEGG.db)
> KEGG()
```

Quality control information for KEGG:

This package has the following mappings:

```
KEGGZYMEID2GO has 4172 mapped keys (of 4172 keys)
KEGGEXTID2PATHID has 75100 mapped keys (of 75100 keys)
KEGGGO2ENZYMEID has 5224 mapped keys (of 5224 keys)
KEGGPATHID2EXTID has 3152 mapped keys (of 3152 keys)
KEGGPATHID2NAME has 390 mapped keys (of 390 keys)
KEGGPATHNAME2ID has 390 mapped keys (of 390 keys)
```

Additional Information about this package:

```
DB schema: KEGG_DB
DB schema version: 2.1
Date for KEGG data: 2011-Mar15
```

```
> names(kegg) <- paste(names(kegg),unlist(mget(names(kegg),KEGGPATHID2NAME)),sep=".")
> names(kegg)
```

```
[1] "04920.Adipocytokine signaling pathway"
[2] "05131.Shigellosis"
[3] "04012.ErbB signaling pathway"
[4] "00450.Selenocompound metabolism"
```

```
[5] "00240.Pyrimidine metabolism"
```

Similarly GO Terms can be retrieved from the `GO.db` (please refer to the vignettes of the corresponding packages for details).

```
> library(GO.db)
> GO()
```

Quality control information for GO:

This package has the following mappings:

```
GOBPANCESTOR has 27199 mapped keys (of 27199 keys)
GOBPCHILDREN has 16020 mapped keys (of 27199 keys)
GOBP OFFSPRING has 16020 mapped keys (of 27199 keys)
GOBPPARENTS has 27199 mapped keys (of 27199 keys)
GOCCANCESTOR has 3722 mapped keys (of 3722 keys)
GOCCCHILDREN has 1237 mapped keys (of 3722 keys)
GOCCOFFSPRING has 1237 mapped keys (of 3722 keys)
GOCCPARENTS has 3722 mapped keys (of 3722 keys)
GOMFANCESTOR has 9889 mapped keys (of 9889 keys)
GOMFCHILDREN has 2012 mapped keys (of 9889 keys)
GOMFOFFSPRING has 2012 mapped keys (of 9889 keys)
GOMFPARENTS has 9889 mapped keys (of 9889 keys)
GOOBSOLETE has 1950 mapped keys (of 1950 keys)
GOTERM has 40811 mapped keys (of 40811 keys)
```

Additional Information about this package:

DB schema: `GO_DB`

DB schema version: 2.1

Date for GO data: 20150314

```
> names(go) <- paste(names(go), Term(names(go)), sep=".")
> names(go)
```

```
[1] "GO:0005968.Rab-protein geranylgeranyltransferase complex"
[2] "GO:0089712.L-aspartate transmembrane transport"
[3] "GO:0000184.nuclear-transcribed mRNA catabolic process, nonsense-mediated decay"
[4] "GO:0035090.maintenance of apical/basal cell polarity"
[5] "GO:0030522.intracellular receptor signaling pathway"
```

Finally we can combine the two FGS collections into a named list for further used in GSE analysis (see below).

```
> fgsList <- list(go=go,kegg=kegg)
```

3 Data analysis with RTopper

To compute gene-to-phenotype association scores the first step required is the conversion of the data into a list, where each list item corresponds to a gene, and comprises a data.frame with the rows being patients, and columns being measurements for each data type, along with the class phenotype (*the response*). Importantly each element of the list with the data should have the same genes and patients.

The `convertToDr` function is used to make such conversion. Below is a short description of the arguments to this function:

- **dataIntersection**: a list of data.frames containing the same set of patients(columns) and genes (rows)
- **response**: a data.frame indicating patients' phenotypic class;
- **nPlatforms**: the number of platforms;

This can be achieved as follows using our examples data:

```
> dataDr <- convertToDr(dat, pheno, 4)
> class(dataDr)

[1] "list"

> length(dataDr)

[1] 500

> names(dataDr)[1:5]

[1] "AACS" "AARS" "ABI1" "ACHE" "ACTC1"

> str(dataDr[1:2])
```

List of 2

```
$ AACS:'data.frame':      95 obs. of  5 variables:
 ..$ dat.affy      : num [1:95] 7.75 7.69 7.54 7.3 7.01 ...
 ..$ dat.agilent   : num [1:95] -1.007 -1.116 -0.913 -1.061 -1.775 ...
 ..$ dat.cnvHarvard: num [1:95] -0.0827 -0.0892 -0.0208 -0.1811 -0.0625 ...
 ..$ dat.cnvMskcc  : num [1:95] -0.03839 -0.0914 0.00823 0.03456 0.0573 ...
 ..$ response      : int [1:95] 0 0 1 1 0 0 0 0 0 0 ...

$ AARS:'data.frame':      95 obs. of  5 variables:
 ..$ dat.affy      : num [1:95] 9.38 9.93 10.2 9.54 9.37 ...
 ..$ dat.agilent   : num [1:95] -1.266 -0.898 0.264 -0.599 -1.437 ...
 ..$ dat.cnvHarvard: num [1:95] -0.1023 -0.2062 -0.0516 -0.0923 -0.1199 ...
 ..$ dat.cnvMskcc  : num [1:95] 0.00756 0.02802 0.10485 0.0841 0.12262 ...
 ..$ response      : int [1:95] 0 0 1 1 0 0 0 0 0 0 ...
```

It is now possible to compute gene-to-phenotype association scores, using as input the gene-centered list produced by `convertToDr`. Therefore the `computeDrStat` function assumes that each gene-centered data.frame contains a column (the last one) called **'response'**, as created by the `convertToDr`. Below is a short description of the arguments to this function:

- **data**: a list of data.frames, one for each gene analyzed, containing the genomic measurements from all platforms (by column) for all the patients (by row), along with the phenotypic response;
- **columns**: a numeric vector indicating column indexes corresponding the genomic measurements to be used for computing the gene-to-phenotype association scores; the default is `columns = c(1:(ncol(data) - 1))`, assuming the phenotypic response to be the last column;
- **method**: the method used to compute the association score;
- **integrate**: logical, whether an integrated gene-to-phenotype score should be computed, or separate scores for each platform/data sets specified by **columns**;

In the current implementation of the **RTopper** there are three methods for computing gene-to-phenotype association scores:

1. **dev**: this approach computes the score as the difference of deviances (as described in Tyekucheva et al, manuscript under review [9]);
2. **aic**: this approach computes the score as the Akaike information criterion for model selection;
3. **bic**: this approach computes the score as the penalized likelihood ratio;

3.1 Integrated Gene-to-Phenotype score computation

This approach first integrates genomic data across platform, and subsequently perform GSE to identify the FGS most strongly associated with the integrated score. Below is an example of application to compute the gene-to-phenotype association scores for 4 data type simultaneously:

```
> bicStatInt <- computeDrStat(dataDr, columns = c(1:4), method="bic", integrate = TRUE)
> names(bicStatInt)

[1] "integrated"

> str(bicStatInt)

List of 1
 $ integrated: Named num [1:500] -11.43 -15.93 -8.85 -13.52 -7.26 ...
 ..- attr(*, "names")= chr [1:500] "AACS" "AARS" "ABI1" "ACHE" ...
```

3.2 Separate Gene-to-Phenotype score computation

This approach first computes gene-to-phenotype score separately for each platform, uses the scores to perform separate GSE analysis in each platform for identifying the FGS most strongly associated with the score, and finally integrates the results from GSE analysis, Below is an example of this approach:

```
> bicStatSep <- computeDrStat(dataDr, columns = c(1:4), method="bic", integrate = FALSE)
> names(bicStatSep)

[1] "dat.affy"          "dat.agilent"
[3] "dat.cnvHarvard"    "dat.cnvMskcc"
```

```
> str(bicStatSep)
```

```
List of 4
```

```
$ dat.affy      : Named num [1:500] 0.545 -4.269 -2.334 -4.471 -3.625 ...
..- attr(*, "names")= chr [1:500] "AACS" "AARS" "ABI1" "ACHE" ...
$ dat.agilent   : Named num [1:500] -3.57 -4.5 -3.66 -4.52 -1.05 ...
..- attr(*, "names")= chr [1:500] "AACS" "AARS" "ABI1" "ACHE" ...
$ dat.cnvHarvard: Named num [1:500] -4.49 -3.64 3.13 -3.26 -2.57 ...
..- attr(*, "names")= chr [1:500] "AACS" "AARS" "ABI1" "ACHE" ...
$ dat.cnvMskcc  : Named num [1:500] -4.53 -4.48 2.1 -2.55 -4.25 ...
..- attr(*, "names")= chr [1:500] "AACS" "AARS" "ABI1" "ACHE" ...
```

3.3 Gene Set Enrichment using integrated and separate score

After the gene-to-phenotype scores have been obtained it is possible to perform a GSE analysis. To this end we will use the `runBatchGSE` function, as shown below. This function enables to perform GSE analysis over multiple collections of FGS, and over multiple ranking statistics. In the current implementation of the `runBatchGSE` the default is performing the enrichment analysis using the `geneSetTest` function from the `limma` package, and most of the arguments passed to `runBatchGSE` are indeed passed to `geneSetTest` (see the relative help for the details).

As an alternative the user can also define his own function to test for FGS enrichment, passing the selection of genes within the FGS and the ranking statistics in the same way as done for `geneSetTest`. In this tutorial we apply `geneSetTest` in order to perform a Wilcoxon rank-sum test, using the absolute value of the gene-to-phenotype scores as the ranking statistics.

```
> args(runBatchGSE)
```

```
function (dataList, fgsList, ...)
NULL
```

Below a short description of the arguments that can be passed to this function:

- **dataList**: a list containing gene-to-phenotype scores to be used as ranking statistics in the GSE analysis;
- **fgsList**: a list of FGS collection, in which each element is a list of character vectors, one for each gene set;
- **...**: any other argument to be passed to lower level functions, including the lower level enrichment function to be used (like the `geneSetTest` function from the `limma` package, which is used as the default);
- **absolute**: logical specifying 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, when not specified (the default) the `geneSetTest` from the `limma` package is used. When a function is specified, 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` in the `limma` package)

Below are few examples to perform Wilcoxon rank-sum test over multiple FGS collections, and over multiple ranking statistics, using the `runBatchGSE`. To this end we will use the **KEGG** and **GO** collections created above, and the separate and integrated gene-to-phenotype scores computed using the `computeDrStat`. The output of this function is a named list of lists, containing an element for each ranking statistics considered in the input. Each one of these elements, in turn, is another list, containing the GSE results for each collection sets. In the examples below we will therefore obtain a list of length one in the case of the integrated gene-to-phenotype score, and a list of length four (one element for each genomic platform) in the case of the separate scores. For all the rankings we will obtain GSE result for both the collections of FGS.

3.4 INTEGRATION + GSE

The integrated gene-to-phenotype scores we have computed can be used to perform a GSE analysis. Below are reported few examples, using the default options, as well as passing several specific arguments to `geneSetTest` (see the relative help for details).

3.4.1 One-sided Wilcoxon rank-sum test using absolute ranking statistics

This can be accomplished by calling the `runBatchGSE` with default values, or by specifying each argument, as shown below:

```
> gseABS.int <- runBatchGSE(dataList=bicStatInt, fgsList=fgsList)
> gseABS.int <- runBatchGSE(dataList=bicStatInt, fgsList=fgsList,
+                           absolute=TRUE, type="f", alternative="mixed")
```

3.4.2 One-sided Wilcoxon rank-sum test using signed ranking statistics

When the signed ranking statistics has a sign, it is possible to perform a one-sided test assessing both tails separately, as well as a two-sided test. This can be accomplished by passing the corresponding arguments to `runBatchGSE`, as shown below:

```
> gseUP.int <- runBatchGSE(dataList=bicStatInt, fgsList=fgsList,
+                           absolute=FALSE, type="t", alternative="up")
> gseDW.int <- runBatchGSE(dataList=bicStatInt, fgsList=fgsList,
+                           absolute=FALSE, type="t", alternative="down")
> gseBOTH.int <- runBatchGSE(dataList=bicStatInt, fgsList=fgsList,
+                             absolute=FALSE, type="t", alternative="either")
```

3.4.3 Performing a simulation-based GSE test

It is also possible to perform an enrichment analysis comparing each FGS to randomly selected gene lists of the same size of the FGS. In this case the p-value is computed by simulation as the proportion of times the mean of the statistics in the FGS is smaller (or larger) than in the `nsim` random simulated sets of genes.

```
> gseABSsim.int <- runBatchGSE(dataList=bicStatInt, fgsList=fgsList,
+                              absolute=TRUE, type="f", alternative="mixed",
```

```

+                               ranks.only=FALSE, nsim=1000)
> gseUPsim.int <- runBatchGSE(dataList=bicStatInt, fgsList=fgsList,
+                               absolute=FALSE, type="t", alternative="up",
+                               ranks.only=FALSE, nsim=1000)

```

Results from this analysis are named lists of lists, as shown below:

```
> str(gseUP.int)
```

```
List of 1
```

```
$ integrated:List of 2
```

```
..$ go : Named num [1:5] NA 0.29 0.561 NA 0.041
```

```
.. ..- attr(*, "names")= chr [1:5] "GO:0005968.Rab-protein geranylgeranyltransferase complex"
```

```
..$ kegg: Named num [1:5] 0.449 NA 0.8444 NA 0.0477
```

```
.. ..- attr(*, "names")= chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis"
```

```
> gseABSSsim.int
```

```
$integrated
```

```
$integrated$go
```

```
GO:0005968.Rab-protein geranylgeranyltransferase complex
```

```
NA
```

```
GO:0089712.L-aspartate transmembrane transport
```

```
0.6523477
```

```
GO:0000184.nuclear-transcribed mRNA catabolic process, nonsense-mediated decay
```

```
0.3926074
```

```
GO:0035090.maintenance of apical/basal cell polarity
```

```
NA
```

```
GO:0030522.intracellular receptor signaling pathway
```

```
0.9760240
```

```
$integrated$kegg
```

```
04920.Adipocytokine signaling pathway
```

```
0.6113886
```

```
05131.Shigellosis
```

```
NA
```

```
04012.ErbB signaling pathway
```

```
0.1788212
```

```
00450.Selenocompound metabolism
```

```
NA
```

```
00240.Pyrimidine metabolism
```

```
0.8091908
```

3.4.4 Passing alternative enrichment functions to runBatchGSE

Below is show how to define and pass alternative enrichment functions to runBatchGSE. We will first show how to use the limma wilcoxGST function, which is a synonym for geneSetTest using ranks.only=TRUE and type="t".

```
> library(limma)
> gseUP.int.2 <- runBatchGSE(dataList=bicStatInt, fgsList=fgsList,
+                             absolute=FALSE, gseFunc=wilcoxGST, alternative="up")
```

As shown below this approach will return the same results obtained with `geneSetTest` passing appropriate arguments.

```
> str(gseUP.int.2)
```

```
List of 1
```

```
 $ integrated:List of 2
```

```
 ..$ go : Named num [1:5] NA 0.29 0.561 NA 0.041
```

```
 .. ..- attr(*, "names")= chr [1:5] "G0:0005968.Rab-protein geranylgeranyltransferase complex"
```

```
 ..$ kegg: Named num [1:5] 0.449 NA 0.8444 NA 0.0477
```

```
 .. ..- attr(*, "names")= chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis"
```

```
> all(gseUP.int.2$go==gseUP.int$go)
```

```
[1] TRUE
```

We can finally also pass any new user-defined enrichment function, provided that the arguments are passed in the same way as with `geneSetTest`, as shown below using the Fisher's exact test, and a threshold for defining the list of differentially expressed genes.

```
> gseFunc <- function(selected, statistics, threshold) {
+   diffExpGenes <- statistics > threshold
+   tab <- table(diffExpGenes, selected)
+   pVal <- fisher.test(tab)[["p.value"]]
+ }
```

```
> gseUP.int.3 <- runBatchGSE(dataList=bicStatInt, fgsList=fgsList,
+                             absolute=FALSE, gseFunc=gseFunc, threshold=7.5)
```

As shown below this approach will test for over-representation of the a specific gene set within the genes defined as differentially expressed (in our example the genes showing an integrated association score larger than 7.5). Results are somewhat comparable to what obtained using the Wilcoxon rank-sum test.

```
> str(gseUP.int.3)
```

```
List of 1
```

```
 $ integrated:List of 2
```

```
 ..$ go : Named num [1:5] NA 1 1 NA 1
```

```
 .. ..- attr(*, "names")= chr [1:5] "G0:0005968.Rab-protein geranylgeranyltransferase complex"
```

```
 ..$ kegg: Named num [1:5] 1 NA 1 NA 0.0298
```

```
 .. ..- attr(*, "names")= chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis"
```

```
> data.frame(fisher=gseUP.int.3$integrated$kegg,wilcoxon=gseUP.int$integrated$kegg)
```

	fisher
04920.Adipocytokine signaling pathway	1.00000
05131.Shigellosis	NA
04012.ErbB signaling pathway	1.00000
00450.Selenocompound metabolism	NA
00240.Pyrimidine metabolism	0.02976

	wilcoxon
04920.Adipocytokine signaling pathway	0.44897258
05131.Shigellosis	NA
04012.ErbB signaling pathway	0.84438786
00450.Selenocompound metabolism	NA
00240.Pyrimidine metabolism	0.04771396

3.5 GSE + INTEGRATION

The individual gene-to-phenotype scores computed for each platform can be similarly used to perform separate GSE analyses for each considered genomic platform, applying the same code and functions used to perform GSE analysis in the **INTEGRATION + GSE** approach above.

```
> gseABS.sep <- runBatchGSE(dataList=bicStatSep, fgsList=fgsList)
```

This step of GSE analysis on separate platform is then followed by GSE results integration, which is achieved using the `combineGSE` function, which summarizes the individual p-values from the tests. To this end different methods are available, including the computation of the geometric or arithmetic means, the use of the median, the selection of the minimum or the maximum p-value, and the random selection (respectively `geometricMean`, `mean`, `median`, `min`, `max`, and `random`). Few examples are shown below:

```
> gseABS.geoMean.sep <- combineGSE(gseABS.sep, method="geometricMean")
> gseABS.max.sep <- combineGSE(gseABS.sep, method="max")
```

Also in this case the results from the combination are named lists of lists, as shown below:

```
> names(gseABS.sep)

[1] "dat.affy"      "dat.agilent"
[3] "dat.cnvHarvard" "dat.cnvMskcc"
```

```
> str(gseABS.sep)
```

```
List of 4
```

```
$ dat.affy      :List of 2
..$ go : Named num [1:5] NA 0.696 0.473 NA 0.924
.. ..- attr(*, "names")= chr [1:5] "G0:0005968.Rab-protein geranylgeranyltransferase complex"
..$ kegg: Named num [1:5] 0.554 NA 0.499 NA 0.986
.. ..- attr(*, "names")= chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis"

$ dat.agilent   :List of 2
..$ go : Named num [1:5] NA 0.423 0.921 NA 0.569
.. ..- attr(*, "names")= chr [1:5] "G0:0005968.Rab-protein geranylgeranyltransferase complex"
..$ kegg: Named num [1:5] 0.321 NA 0.787 NA 0.825
.. ..- attr(*, "names")= chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis"

$ dat.cnvHarvard:List of 2
..$ go : Named num [1:5] NA 0.9532 0.0558 NA 0.373
.. ..- attr(*, "names")= chr [1:5] "G0:0005968.Rab-protein geranylgeranyltransferase complex"
..$ kegg: Named num [1:5] 0.4537 NA 0.142 NA 0.0785
.. ..- attr(*, "names")= chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis"

$ dat.cnvMskcc  :List of 2
```

```

..$ go : Named num [1:5] NA 0.483 0.977 NA 0.381
.. ..- attr(*, "names")= chr [1:5] "G0:0005968.Rab-protein geranylgeranyltransferase complex"
..$ kegg: Named num [1:5] 0.618 NA 0.26 NA 0.137
.. ..- attr(*, "names")= chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis"

> str(gseABS.geoMean.sep)

List of 1
 $ combinedScore:List of 2
  ..$ go : Named num [1:5] NA 0.607 0.393 NA 0.523
  .. ..- attr(*, "names")= chr [1:5] "G0:0005968.Rab-protein geranylgeranyltransferase complex"
  ..$ kegg: Named num [1:5] 0.473 NA 0.347 NA 0.306
  .. ..- attr(*, "names")= chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis"

> gseABS.geoMean.sep

$combinedScore
$combinedScore$go
      G0:0005968.Rab-protein geranylgeranyltransferase complex
                                     NA
      G0:0089712.L-aspartate transmembrane transport
                                     0.6069100
G0:0000184.nuclear-transcribed mRNA catabolic process, nonsense-mediated decay
                                     0.3925083
      G0:0035090.maintenance of apical/basal cell polarity
                                     NA
      G0:0030522.intracellular receptor signaling pathway
                                     0.5228761

$combinedScore$kegg
04920.Adipocytokine signaling pathway
      0.4726189
      05131.Shigellosis
      NA
      04012.ErbB signaling pathway
      0.3468525
      00450.Selenocompound metabolism
      NA
      00240.Pyrimidine metabolism
      0.3056998

```

3.6 Multiple testing correction

Finally the `adjustPvalGSE` enables to adjust the p-values computed by the `runBatchGSE`. This functions is an interface to the `mt.rawp2adjp` function from the `multtest` package.

```

> gseABS.int.BH <- adjustPvalGSE(gseABS.int)
> gseABS.int.holm <- adjustPvalGSE(gseABS.int, proc = "Holm")

```

Also in this case the results after the adjustment are named lists of lists, as shown below:

```

> names(gseABS.int.BH)
[1] "integrated"
> names(gseABS.int.holm)
[1] "integrated"
> str(gseABS.int.BH)
List of 1
 $ integrated:List of 2
  ..$ go : num [1:5, 1:2] NA 0.715 0.444 NA 0.963 ...
  .. ..- attr(*, "dimnames")=List of 2
  .. .. ..$ : chr [1:5] "G0:0005968.Rab-protein geranylgeranyltransferase complex" "G0:0089712"
  .. .. ..$ : chr [1:2] "rawp" "BH"
  ..$ kegg: num [1:5, 1:2] 0.559 NA 0.158 NA 0.765 ...
  .. ..- attr(*, "dimnames")=List of 2
  .. .. ..$ : chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis" "04012.ErbB"
  .. .. ..$ : chr [1:2] "rawp" "BH"
> str(gseABS.int.holm)
List of 1
 $ integrated:List of 2
  ..$ go : num [1:5, 1:2] NA 0.715 0.444 NA 0.963 ...
  .. ..- attr(*, "dimnames")=List of 2
  .. .. ..$ : chr [1:5] "G0:0005968.Rab-protein geranylgeranyltransferase complex" "G0:0089712"
  .. .. ..$ : chr [1:2] "rawp" "Holm"
  ..$ kegg: num [1:5, 1:2] 0.559 NA 0.158 NA 0.765 ...
  .. ..- attr(*, "dimnames")=List of 2
  .. .. ..$ : chr [1:5] "04920.Adipocytokine signaling pathway" "05131.Shigellosis" "04012.ErbB"
  .. .. ..$ : chr [1:2] "rawp" "Holm"

```

4 System Information

Session information:

```

> sessionInfo()

R version 3.2.0 RC (2015-04-08 r68161)
Platform: x86_64-apple-darwin10.8.0 (64-bit)
Running under: OS X 10.6.8 (Snow Leopard)

locale:
[1] en_US.UTF-8/en_US.UTF-8/en_US.UTF-8/C/en_US.UTF-8/en_US.UTF-8

attached base packages:
[1] stats4      parallel  stats      graphics
[5] grDevices  utils      datasets  methods
[9] base

```

other attached packages:

```
[1] limma_3.24.0      GO.db_3.1.2
[3] KEGG.db_3.1.2     org.Hs.eg.db_3.1.2
[5] RSQLite_1.0.0     DBI_0.3.1
[7] AnnotationDbi_1.30.0 GenomeInfoDb_1.4.0
[9] IRanges_2.2.0     S4Vectors_0.6.0
[11] RTopper_1.14.0    Biobase_2.28.0
[13] BiocGenerics_0.14.0
```

loaded via a namespace (and not attached):

```
[1] MASS_7.3-40      splines_3.2.0
[3] tools_3.2.0      survival_2.38-1
[5] multtest_2.24.0
```

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