

# Package ‘GOGANPA’

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**Type** Package

**Title** GO-Functional-Network-based Gene-Set-Analysis

**Version** 1.0

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**Description**

Accounting for genes' functional-non-equivalence within pathways in classical Gene-set-analysis.

**Depends** R (>= 2.10), GANPA, WGCNA

**Suggests** GANPA, WGCNA, GANPAdata

**License** GPL-2

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GOGANPA-package

*GO-Functional-Network-based Gene-Set-Analysis*

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Accounting for genes' functional-non-equivalence within pathways in classical Gene-set-analysis.

**Details**

Package: GOGANPA  
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Date: 2011-12-30  
License: GPL-2

### Author(s)

Billy Chang

Maintainer: Billy Chang <billy.chang@utoronto.ca>

### References

Chang, B, Kustra, R, and Tian, WD, (2012) Functional-Network-Based Gene-Set-Analysis using Gene Ontology. Submitted.

Fang, ZY, Tian, WD, and Ji, HB. (September 6, 2011) A network-based gene-weighting approach for pathway analysis. Cell Research. Advanced Publication.

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getGNET

*Gene Network Construction by Similarity-Thresholding*

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### Description

Construct a gene network by linking gene-pairs with GO similarity above a chosen threshold.

### Usage

```
getGNET(simMat, rho)
```

### Arguments

simMat	The GO-similarity matrix. Missing and negative entries are not allowed. The gene names should be assigned to the row and column names.
rho	The threshold, chosen e.g. by <code>selectRho</code> . Gene-pairs with similarity above the threshold will be linked.

### Value

A list, where each element contains the names of the genes connected to the corresponding gene indicated by the element-header.

**Note**

Note that certain GO-similarity measures are unbounded (e.g. the Resnik similarity). This code will not normalize the similarity matrix, and rho should therefore be chosen according to the range of the GO-similarity values inside simMat.

**Author(s)**

Billy Chang

**References**

Chang, B., Kustra, R. and Tian, WD (2012) Functional-Network-based Gene Set Analysis using Gene Ontology. Submitted.

Zhang, B. and Horvath, S. (2005) A General Framework for Weighted Gene Co-Expression Network Analysis. *Statistical Applications in Genetics and Molecular Biology*. 4:1:A17.

**See Also**

[selectRho](#)

**Examples**

```
#Not to Run
data("simMatSmall", package="GOGANPA")
gNET <- getGNET(simMatSmall, rho=0.7)
hist(sapply(gNET, length)) # network connectivities (excluding unconnected genes)
```

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GOGANPA

*GO-based Gene-Set-Analysis*

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**Description**

A Gene-Set-Analysis method that accounts of functional-linkages among Genes, as indicated by a GO-functional network constructed from a GO-similarity matrix.

**Usage**

```
GOGANPA(gExprs.obj, gsets, gNET = NULL, simMat = NULL, rho = NULL, msp.groups,
check.exprs = TRUE, msp.correction = TRUE, size.min = 15, size.max = 500,
permN = 2000, randN = 30, permFDR.cutoff = 0.15, output.label = "GOGANPAResult")
```

**Arguments**

<code>gExprs.obj</code>	Gene expression experiment data object
<code>gsets</code>	A list of gene sets.
<code>gNET</code>	A gene association network stored in a list.
<code>simMat</code>	The GO-similarity matrix. Missing and negative entries are not allowed. The gene names should be assigned to the row and column names.
<code>rho</code>	The threshold, chosen e.g. by <code>selectRho</code> . If NULL, then chosen automatically by <code>selectRho</code> .
<code>msp.groups</code>	A list of multi-subunit-proteins-coding genes.
<code>check.exprs</code>	Logical (TRUE by default). Check and correct the missing values and scaling in the <code>gExprs.obj</code> . If the scale is natural, it will be converted to log2.
<code>msp.correction</code>	Logical (TRUE). Whether to do a correction for multi-subunit proteins in gene weighting.
<code>size.min</code>	Minimum size of gene sets used for analysis. By default 15 genes.
<code>size.max</code>	Maximum size of gene sets used for analysis. By default 500 genes.
<code>permN</code>	Sample permutation times. By default 2000 times.
<code>randN</code>	Gene randomization times. Can be set smaller (say, 30) if you do not care randomization-based significance so as to be faster.
<code>permFDR.cutoff</code>	Sample permutation FDR cutoff. A number between 0 and 1. Set it larger if wish to see the significance of more gene sets.
<code>output.label</code>	A label to name output files.

**Details**

Exactly one of `gNET` and `simMat` must be NULL. If `simMat` and `rho` are provided, `getGNET` will be called to obtain the gene network. If `simMat` is provided but `rho` is missing, then `selectRho` will also be called to provide an automatic choice of `rho`. This code is based on GANPA (Fang et. al. 2011), the gene network, `gNET`, whether supplied or derived from `simMat`, will be fed into `GSE.Test.Main` in the package GANPA for weighted Gene-Set-Analysis.

**Value**

A .csv file containing various statistics.

**Author(s)**

Billy Chang

**References**

- Chang, B., Kustra, R. and Tian, WD (2012) Functional-Network-based Gene Set Analysis using Gene Ontology. Submitted.
- Fang, ZY, Tian, WD, and Ji, HB. (September 6, 2011) A network-based gene-weighting approach for pathway analysis. Cell Research. Advanced Publication.

**See Also**

[getGNET](#), [selectRho](#)

**Examples**

```
#Not to Run
require(GANPA)
data("simMatSmall", package="GOGANPA")
data("gExprs.p53", "gsets.msigdb.pnas", "msp.groups", package="GANPAdata")
set.seed(1000)
GOGANPA(gExprs.obj=gExprs.p53, gsets=gsets.msigdb.pnas, gNET=NULL, simMat=simMatSmall, rho=NULL,
        msp.groups=msp.groups, check.exprs=TRUE, msp.correction=TRUE,
        size.min=15, size.max=500, permN=2000, randN=30,
        permFDR.cutoff=0.15, output.label="GOGANPAResult")
```

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selectRho

*Choosing a threshold based on the Scale-Free-Topology-Criterion*

---

**Description**

Determine the threshold parameter which will result in a network with optimal scale-free fitness.

**Usage**

```
selectRho(simMat, rhovec = NULL)
```

**Arguments**

simMat	The GO-similarity matrix. Missing and negative entries are not allowed. The gene names should be assigned to the row and column names.
rhovec	a vector of candidate thresholds, or if NULL, a set of thresholds chosen according to the range of the similarity matrix.

**Details**

The scale-free fitness measure is based on linear-regression-based R-squared goodness-of-fit measure.

**Value**

A list, with elements:

criterion	a summary table of the candidate thresholds' resulting fits.
bestrho	The candidate threshold with the highest R-squared.

**Note**

Note that certain GO-similarity measures are unbounded (e.g. the Resnik similarity). This code will not normalize the similarity matrix, and `rhoVec`, if supplied, should be chosen according to the range of the GO-similarity values inside `simMat`.

**Author(s)**

Billy Chang

**References**

Chang, B., Kustra, R. and Tian, WD (2012) Functional-Network-based Gene Set Analysis using Gene Ontology. Submitted.

Zhang, B. and Horvath, S. (2005) A General Framework for Weighted Gene Co-Expression Network Analysis. *Statistical Applications in Genetics and Molecular Biology*. 4:1:A17.

**See Also**

[getGNET](#)

**Examples**

```
#Not to Run
data("simMatSmall", package="GOGANPA")
fit <- selectRho(simMatSmall)
plot(fit$criterion[,1], fit$criterion[,2])
abline(v=fit$bestrho, col=2)
```

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simMatSmall

*A Resnik Similarity Matrix.*

---

**Description**

A Resnik Similarity Matrix (normalized) for 2000 human genes sampled from 14173 annotated human genes.

**Details**

The similarity matrix was computed using the R package `csbl.go` (<http://csbi.ltdk.helsinki.fi/csbl.go>), using a GO term specificity table computed using GO BP annotations for all human Entrez Genes available in the Bioconductor package `org.Hs.eg.db`, version 2.6.4 (not the default table provided `csbl.go`).

**Note**

This matrix is provided for test-running GOGANPA only. Although it is sampled from the similarity matrix used in Chang et. al. (2012), it cannot be used to reproduce the results presented in Chang et. al. (2012).

**Author(s)**

Billy Chang

**References**

Ovaska, K, Laakso M, and Hautaniemi, S. (2008) Fast Gene Ontology based clustering for microarray experiments. *BioData Mining*. 1:11.

Chang, B., Kustra, R. and Tian, WD (2012) Functional-Network-based Gene Set Analysis using Gene Ontology. Submitted.

**Examples**

```
#Not to Run  
data("simMatSmall", package='GOGANPA')
```



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