

Package ‘GOSim’

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Title Computation of functional similarities between GO terms and gene products; GO enrichment analysis

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Suggests igraph

LazyLoad Yes

Description This package implements several functions useful for computing similarities between GO terms and gene products based on their GO annotation. Moreover it allows for computing a GO enrichment analysis

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calc.diffusion.kernel

Calculation and loading of diffusion kernel matrices

Description

Manifold embeddings of gene ontology terms via diffusion kernel techniques. Diffusion kernels are positive semidefinite similarity measures calculated from the graph Laplacian. They are interpreted as the result of a local heat diffusion process along the graph structure.

Usage

```
calc.diffusion.kernel(method="diffKernelLapl", m=7)
```

```
load.diffusion.kernel(method="diffKernelLapl")
```

Arguments

method	one of "diffKernelLapl", "diffKernelpower", "diffKernelLLE", "diffKernelexpm"
m	(1) Half the power of the transition probability matrix (an integer > 0). (2) an arbitrary positive time constant for the exponential diffusion kernel

Details

"diffKernelLapl" pseudo inverse of the (unnormalized) graph Laplacian: Takes into account all powers of diffusion and incorporates all paths from one node to another one.

"diffKernelpower" even power of the transition probability matrix: Takes into account local transitions of path length m

"diffKernelLLE" local linear embedding into an Euclidean space: The focus is to preserve local distances to nearest neighbors. The LLE kernel emphasizes short-range interactions between GO terms.

diffKernelexpm" $\exp(-mL)$, where t is a positive constant, L is the (unnormalized) graph Laplacian and expm denotes the matrix exponential. This kernel takes into account all positive integer powers of diffusion, but with an exponential decay of the influence of long-range interactions.

Value

`calc.diffusion.kernel` puts a kernel matrix / similarity matrix named "`<method><ontology><organism><evidence levels>.rda`" in the current directory. This file has then to be moved into the data directory of GOSim. It can be used afterwards by calling `load.diffusion.kernel`. Sometimes it is necessary to switch to the GOSim directory for this purpose.

References

Lerman G. & Shaknovich B., Defining Functional Distance using Manifold Embeddings of Gene Ontology Annotations, PNAS, 104(27): 11334 - 11339, 2007

See Also

[load.diffusion.kernel](#)

Examples

```
## Not run:
  calc.diffusion.kernel(method="diffKernelpower", m=5)
  load.diffusion.kernel("diffKernelpower")
  getTermSim(c("GO:0007166", "GO:0007267", "GO:0007584", "GO:0007165", "GO:0007186"), method="diffKernelpower")
## End(Not run) # --> this may take some time ...
```

calcICs

Calculate information contents of GO terms.

Description

Recalculates the information content of all GO terms.

Usage

```
calcICs()
```

Arguments

none

Details

This functions should only be invoked, if one wants to calculate the information content for GO terms with respect to combinations of evidence codes other than the precomputed ones or, if a new version of the organism annotation packages has been installed. By default the information contents are precomputed using all evidence codes and evidence codes "IMP, IGI, IDA, IEP, IPI" together.

Value

Puts a file named "ICs<ontology><organism><evidence levels>.rda" in the current directoy. This file has then to be moved into the data directory of GOSim. It can be used afterwards by calling [setOntology](#). Sometimes it is necessary to switch to the GOSim directory for this purpose.

See Also

[setEvidenceLevel](#)

Examples

```
## Not run:

    setEvidenceLevel("IMP")
    setOntology("CC", loadIC=FALSE) # important: setOntology assumes that the IC file al
    calcICs()
## End(Not run) # --> this may take some time ...
```

clusterEvaluation *Evaluate a given grouping of genes or GO terms.*

Description

Evaluate a given grouping of genes or terms with respect to their GO similarity.

Usage

```
evaluateClustering(clust, Sim)
```

Arguments

clust	vector of cluster labels (integer or character) for each gene
Sim	similarity matrix

Details

If necessary, more details than the description above

Value

evaluateClustering returns a list with two items:

```
clusterstats      matrix (ncluster x 2) of median within cluster similarities and median absolute
                  deviations
clustersil        cluster silhouette values
```

Author(s)

Holger Froehlich

References

Rousseeuw, P., Silhouettes: a graphical aid to the interpretation and validation of cluster analysis, J. Comp. and Applied Mathematics, 1987, 20, 53-6

See Also

[getGeneSimPrototypes](#), [getGeneSim](#), [getTermSim](#), [GOenrichment](#)

Examples

```
## Not run:
setOntology("BP")
gomap <- get("gomap", env=GOSimEnv)
allgenes = sample(names(gomap), 1000) # suppose these are all genes
genesOfInterest = sample(allgenes, 20) # suppose these are all genes of interest

sim = getGeneSim(genesOfInterest, verbose=FALSE) # and these are their similarities
hc = hclust(as.dist(1-sim), method="ward") # use them to perform a clustering
plot(hc) # plot the cluster tree
cl = cutree(hc, k=3) # take 3 clusters

if(require(cluster)){
  ev = evaluateClustering(cl, sim) # evaluate the clustering
  print(ev$clusterstats) # print out some statistics
  plot(ev$clustersil, main="") # plot the cluster silhouettes
}

# investigate cluster 1 further
if(require(topGO))
  GOenrichment(genesOfInterest[cl == 1], allgenes, cutoff=0.05) # print out wh
## End(Not run)
```

`filterGO`*Filter GO.*

Description

Filter out genes from a list not mapping to the actual ontology. Genes not mapping to the currently set ontology ("BP","MF","CC") and not having one of the predefined evidence codes (default is to use all evidence codes) are removed.

Usage

```
filterGO(genelist)
```

Arguments

```
genelist      character vector of Entrez gene IDs
```

Value

List with items

```
"genename"   gene ID
```

```
"annotation" character vector of GO terms mapping to the gene within the actual ontology
```

Note

The result depends on the currently set ontology. **IMPORTANT:** The result refers to the GO library that was used to precompute the information content of GO terms.

Author(s)

Holger Froehlich

See Also

[setOntology](#), [setEvidenceLevel](#), [getGOInfo](#), [calcICs](#)

Examples

```
filterGO(c("12345", "4559"))  
# --> 4559 is removed
```

getAncestors *get list of ALL ancestors associated to each GO term*

Description

Returns the list of all (also indirect) ancestors (= less specific terms) associated to each GO term. The type of relationship between GO terms ("is a" or "part of") is ignored.

Usage

```
getAncestors ()
```

Value

List with entry names for each GO term. Each entry contains a character vector with the ancestor GO terms.

Note

The result is computed within the currently set ontology ("BP","MF","CC"). It directly uses the "GO" library to compute the result.

Author(s)

Holger Froehlich

See Also

[getOffsprings](#), [getChildren](#), [getParents](#), [setOntology](#)

Examples

```
## Not run: getAncestors ()
```

getChildren *Get a list of all direct children of each GO term.*

Description

Returns the list of all direct children (= more specific terms one hierarchy level down) associated to each GO term. The type of relationship between GO terms ("is a" or "part of") is ignored.

Usage

```
getChildren ()
```

Value

List with entry names for each GO term. Each entry contains a character vector with the direct children GO terms.

Note

The result is computed within the currently set ontology ("BP","MF","CC"). It directly uses the "GO" library to compute the result.

Author(s)

Holger Froehlich

See Also

[getOffsprings](#), [getParents](#), [getAncestors](#), [setOntology](#)

Examples

```
## Not run: getChildren()
```

getDisjCommAnc *Get disjoint common ancestors.*

Description

Returns the GO terms representing the disjoint common ancestors of two GO terms.

Usage

```
getDisjCommAnc(term1, term2)
```

Arguments

term1	GO term 1
term2	GO term 2

Details

The result is computed within the currently set ontology ("BP","MF","CC").

Value

Character vector of GO terms

Author(s)

Holger Froehlich

References

Couto, F.; Silva, M. & Coutinho, P., Semantic Similarity over the Gene Ontology: Family Correlation and Selecting Disjunctive Ancestors, Conference in Information and Knowledge Management, 2005

See Also

[getTermSim](#), [getGOGraph](#), [setOntology](#)

Examples

```
if(require(graph))
  getDisjCommAnc("GO:0006955", "GO:0007584")
```

`getGeneFeatures` *Get simple feature vector representation of genes*

Description

Computes feature vectors for list of genes: Each gene is represented by a vector describing the presence/absence of all GO terms. The absence of each GO term is additionally weighted by its information content.

Usage

```
getGeneFeatures(genelist, pca=FALSE, normalization=FALSE, verbose=TRUE)
```

Arguments

<code>genelist</code>	character vector of Entrez gene IDs
<code>pca</code>	perform PCA on feature vectors to reduce dimensionality
<code>normalization</code>	scale the feature vectors to norm 1
<code>verbose</code>	print out additional information

Details

The PCA postprocessing determines the principal components that can explain at least 95% of the total variance in the feature space.

Value

matrix with rows being genes and columns being GO terms.

Note

The result depends on the currently set ontology ("BP", "MF", "CC").

Author(s)

Holger Froehlich

References

M. Mistry, P Pavlidis, Gene Ontology term overlap as a measure of gene functional similarity, BMC Bioinformatics, 9:327, 2008.

See Also

[getGeneSimPrototypes](#), [selectPrototypes](#), [getGeneSim](#), [getTermSim](#), [setOntology](#)

Examples

```
# see selectPrototypes
```

```
getGeneFeaturesPrototypes
```

Get feature vector representation of genes relative to prototype genes

Description

Computes feature vectors for list of genes: Each gene is represented by its similarities to predefined prototype genes.

Usage

```
getGeneFeaturesPrototypes(genelist, prototypes = NULL,
                          similarity = "max", similarityTerm = "JiangConrath",
                          pca = TRUE, normalization = TRUE, verbose = TRUE)
```

Arguments

<code>genelist</code>	character vector of Entrez gene IDs
<code>prototypes</code>	character vector of Entrez gene IDs representing the prototypes
<code>similarity</code>	method to calculate the similarity to prototypes
<code>similarityTerm</code>	method to compute the GO term similarity
<code>pca</code>	perform PCA on feature vectors to reduce dimensionality
<code>normalization</code>	scale the feature vectors to norm 1
<code>verbose</code>	print out additional information

Details

If no prototypes are passed, the method calls the [selectPrototypes](#) function with no arguments. Hence, the prototypes in this case are the 250 genes with most known annotations.

The PCA postprocessing determines the principal components that can explain at least 95% of the total variance in the feature space.

The method to calculate the functional similarity of a gene to a certain prototype can be any of those described in [getGeneSim](#).

Value

List with items

"features" feature vectors for each gene: n x d data matrix

"prototypes" prototypes (= principal components, if PCA has been performed)

Note

The result depends on the currently set ontology ("BP", "MF", "CC").

Author(s)

Holger Froehlich

References

[1] H. Froehlich, N. Speer, C. Spieth, and A. Zell, Kernel Based Functional Gene Grouping, Proc. Int. Joint Conf. on Neural Networks (IJCNN), 6886 - 6891, 2006

[2] N. Speer, H. Froehlich, A. Zell, Functional Grouping of Genes Using Spectral Clustering and Gene Ontology, Proc. Int. Joint Conf. on Neural Networks (IJCNN), pp. 298 - 303, 2005

See Also

[getGeneSimPrototypes](#), [selectPrototypes](#), [getGeneSim](#), [getTermSim](#), [setOntology](#)

Examples

```
# see selectPrototypes
```

getGeneSim *Compute functional similarity for genes*

Description

Calculate the pairwise functional similarities for a list of genes using different strategies.

Usage

```
getGeneSim(genelist, similarity="funSimMax", similarityTerm="relevance", normalization
```

Arguments

genelist	character vector of Entrez gene IDs
similarity	method to calculate the functional similarity between gene products
similarityTerm	method to compute the similarity of GO terms
normalization	normalize similarities yes/no
method	"sqrt": normalize $\text{sim}(x,y) \leftarrow \text{sim}(x,y)/\sqrt{\text{sim}(x,x)*\text{sim}(y,y)}$; "Lin": normalize $\text{sim}(x,y) \leftarrow 2*\text{sim}(x,y)/(\text{sim}(x,x) + \text{sim}(y,y))$; "Tanimoto": normalize $\text{sim}(x,y) \leftarrow \text{sim}(x,y)/(\text{sim}(x,x) + \text{sim}(y,y) - \text{sim}(x,y))$. NOTE: normalization does not have any effect, if similarity = "funSimMax", "funSimAvg" or similarity = "OA" and avg=TRUE
avg	standardize the OA kernel by the maximum number of GO terms for both genes
verbose	print out some information

Details

The method to calculate the pairwise functional similarity between gene products can either be:

"max" the maximum similarity between any two GO terms

"mean" the average similarity between any two GO terms

funSimMax the average of best matching GO term similarities. Take the maximum of the scores achieved by assignments of GO terms from gene 1 to gene 2 and vice versa. [2]

funSimAvg the average of best matching GO term similarities. Take the average of the scores achieved by assignments of GO terms from gene 1 to gene 2 and vice versa. [2]

"OA" the optimal assignment (maximally weighted bipartite matching) of GO terms associated to the gene having fewer annotation to the GO terms of the other gene. [1]

"hausdorff" the translation of the Hausdorff distance between two sets: Let A and B be the two sets of GO terms associated to two genes. Then $\text{sim}(A, B) = \min(\min(\max_{x \in A}(x, y)), \min(\max_{y \in B}(x, y)))$. [3]

"dot" the dot product between feature vectors describing the absence/presence of each GO term. The absence of each GO term is weighted by its information content. Depending on the type of later normalization one can arrive at the cosine similarity (method="sqrt") or at the Tanimoto coefficient (method="Tanimoto").[4]

Arguments

genelist	character vector of Entrez gene IDs
prototypes	character vector of Entrez gene IDs representing the prototypes
similarity	method to calculate the similarity to prototypes
similarityTerm	method to compute the GO term similarity
pca	perform PCA on feature vectors to reduce dimensionality
normalization	normalize similarities to [0,1]: $\text{sim}(x,y) \leftarrow 0.5 * (\text{sim}(x,y) / \sqrt{(\text{sim}(x,x) * \text{sim}(y,y)) + 1})$
verbose	print additional information

Details

The method calls [getGeneFeaturesPrototypes](#) to calculate the feature vectors. The functional similarity between two genes is essentially given by the dot product between their feature vectors.

Value

List with items

"similarity" n x n similarity matrix (n = number of genes)

"prototypes" prototypes (= principal components, if PCA has been performed)

"features" feature vectors for each gene: n x d data matrix

Note

The result depends on the currently set ontology ("BP", "MF", "CC").

Author(s)

Holger Froehlich

References

- [1] H. Froehlich, N. Speer, C. Spieth, and A. Zell, Kernel Based Functional Gene Grouping, Proc. Int. Joint Conf. on Neural Networks (IJCNN), 6886 - 6891, 2006
- [2] N. Speer, H. Froehlich, A. Zell, Functional Grouping of Genes Using Spectral Clustering and Gene Ontology, Proc. Int. Joint Conf. on Neural Networks (IJCNN), pp. 298 - 303, 2005

See Also

[getGeneFeaturesPrototypes](#), [selectPrototypes](#), [getGeneSim](#), [getTermSim](#), [setOntology](#)

Examples

```
## Not run:
may take some time ...
proto=selectPrototypes(n=50) # --> returns a character vector of 50 genes with the highest
getGeneSimPrototypes(c("207", "208", "360"), prototypes=proto, similarityTerm="Resnik")
## End(Not run)
```

getGOGraph	<i>(1) Get GO graph with specified GO terms at its leave; (2) Get GO Graph with GO terms at leaves associated to one or several genes of interest.</i>
------------	--

Description

The function `getGOGraph` returns a `graphNEL` object representing the GO graph with leaves specified in the argument. The function `getGOGraphsGenes` returns a set of `graphNEL` objects. The *i*th graph object is created by call to `getGOGraph` with the GO terms associated to gene *i*. It hence shows for each gene, where its GO terms are located within the GO structure.

Usage

```
getGOGraph(term, prune=Inf)

getGOGraphsGenes(genelist, prune=Inf)
```

Arguments

<code>term</code>	character vector of GO terms
<code>genelist</code>	character vector of Entrez gene IDs
<code>prune</code>	do not show the complete graph, but prune it after the specified number of ancestors

Details

The result is computed within the currently set ontology ("BP", "MF", "CC").

Value

`graphNEL` object(s)

Note

directly calls the function `GOMGraph` in the "GOstats" library

Author(s)

Holger Froehlich

Examples

```
if(require(graph)){
  G=getGOGraph(c("GO:0006955","GO:0007584"))
  if(require(Rgraphviz)){
    plot(G)
    Gs = getGOGraphsGenes(c("207","7494"))
    plot(Gs[[1]]) # plot the first of both GO graphs
  }
}
```

getGOInfo

Obtain GO terms and their description for a list of genes.

Description

Obtain the GO terms and their description for a list of genes.

Usage

```
getGOInfo(geneIDs)
```

Arguments

geneIDs character vector of Entrez gene IDs

Value

List with entry names equal to the gene IDs. Each list contains a sublist with entry names equal to the GO terms associated to the corresponding gene ID. Each entry also contains a description of the GO term, its definition and the ontology ("BP","CC","MF") it belongs to.

Note

The corresponding information is directly extracted from the "GO" library. The result depends on the currently set ontology ("BP","MF","CC"), i.e. only GO terms within the actual ontology are considered. The shown GO information refers to the actually installed GO library.

Author(s)

Holger Froehlich

See Also

[setOntology](#)

Examples

```
## Not run:
if(require(annotate)){
  setOntology("BP")
  getGOInfo(c("207", "7494"))
}
## End(Not run)
```

getMinimumSubsumer *Compute minimum subsumer of two GO terms.*

Description

Returns the minimum subsumer (i.e. the common ancestor having the maximal information content) of two GO terms

Usage

```
getMinimumSubsumer(term1, term2)
```

Arguments

term1	GO term 1
term2	GO term 2

Details

The result is computed within the currently set ontology ("BP", "MF", "CC").

Value

GO term representing the minimum subsumer. If there is no minimum subsumer within the currently set GO category (e.g. because one of the GO terms does not exist), the result is the string "NA".

Author(s)

Holger Froehlich

References

P. Resnik, Using Information Content to evaluate semantic similarity in a taxonomy, Proc. 14th Int. Conf. Artificial Intel., 1995

See Also

[getTermSim](#), [getGOGraph](#), [setOntology](#)

Examples

```
# setOntology("BP")
getMinimumSubsumer("GO:0006955", "GO:0007584")
# returns GO:0050896
```

getOffsprings	<i>Get all offspring associated with one or more GO term</i>
---------------	--

Description

Returns the list of all (also indirect) offspring (= more specific terms) associated to each GO term. The type of relationship between GO terms ("is a" or "part of") is ignored.

Usage

```
getOffsprings()
```

Value

List with entry names for each GO term. Each entry contains a character vector with the offspring GO terms.

Note

The result is computed within the currently set ontology ("BP", "MF", "CC"). It directly uses the "GO" library to compute the result.

Author(s)

Holger Froehlich

See Also

[getChildren](#), [getParents](#), [getAncestors](#), [setOntology](#)

Examples

```
## Not run: getOffsprings()
```

getParents *Get direct parents for each GO term.*

Description

Returns the list of all direct parents (= less specific terms one hierarchy level up) associated to each GO term. The type of relationship between GO terms ("is a" or "part of") is ignored.

Usage

```
getParents()
```

Value

List with entry names for each GO term. Each entry contains a character vector with the direct parent GO terms.

Note

The result is computed within the currently set ontology ("BP","MF","CC"). It directly uses the "GO" library to compute the result.

Author(s)

Holger Froehlich

See Also

[getOffsprings](#), [getChildren](#), [getAncestors](#), [setOntology](#)

Examples

```
## Not run: getParents()
```

getTermSim *Get pairwise GO term similarities.*

Description

Returns the pairwise similarities between GO terms. Different calculation method are implemented.

Usage

```
getTermSim(termlist, method = "relevance", verbose = TRUE)
```

Arguments

termList	character vector of GO terms
method	one of the supported methods for GO term similarity (see below)
verbose	print out various information or not

Details

Currently the following methods for computing GO term similarities are implemented:

"Resnik" information content of minimum subsumer (ICms) [1]

"JiangConrath" $1 - \min(1, IC(term1) - 2ICms + IC(term2))$ [2]

"Lin" $\frac{2ICms}{(IC(term1)+IC(term2))}$ [3]

"CoutoEnriched" FuSSiMeg enriched term similarity by Couto et al. [4]. Requires enrichment factors to be set by `setEnrichmentFactors`.

"CoutoResnik" average information content of common disjunctive ancestors of term1 and term2 (ICshare) [5]

"CoutoJiangConrath" $1 - \min(1, IC(term1) - 2ICshare + IC(term2))$ [5]

"CoutoLin" $\frac{2ICshare}{(IC(term1)+IC(term2))}$ [5]

"diffKernel" diffusion kernel similarity from a pre-loaded kernel matrix (see `load.diffusion.kernel`). The diffusion kernel is calculated using one of the methods described in [6].

"relevance" `sim_Lin * (1 - exp(-ICms))` [7]

"GIC" summed information content of common ancestors divided by summed information content of all ancestors of term1 and term2 [8]

Value

n x n matrix (n = number of GO terms) with similarities between GO terms scaled to [0,1]. If a GO term does not exist for the currently set ontology, the similarity is set to "NA".

Note

All calculations use normalized information contents for each GO term. Normalization is achieved by dividing each information content by the maximum information content within the currently set ontology ("BP", "MF", "CC")

Author(s)

Holger Froehlich

References

- [1] P. Resnik, Using Information Content to evaluate semantic similarity in a taxonomy, Proc. 14th Int. Conf. Artificial Intel., 1995
- [2] J. Jiang, D. Conrath, Semantic Similarity based on Corpus Statistics and Lexical Taxonomy, Proc. Int. Conf. Research in Comp. Ling., 1998

- [3] D. Lin, An Information-Theoretic Definition of Similarity, Proc. 15th Int. Conf. Machine Learning, 1998
- [4] F. Couto, M. Silva, P. Coutinho, Implementation of a Functional Semantic Similarity Measure between Gene-Products, DI/FCUL TR 03-29, Department of Informatics, University of Lisbon, 2003
- [5] Couto, F.; Silva, M. & Coutinho, P., Semantic Similarity over the Gene Ontology: Family Correlation and Selecting Disjunctive Ancestors, Conference in Information and Knowledge Management, 2005
- [6] Lerman G. & Shakhovich B., Defining Functional Distance using Manifold Embeddings of Gene Ontology Annotations, PNAS, 104(27): 11334 - 11339, 2007
- [7] A. Schlicker, F. Domingues, J. Rahnenfuehrer, T. Lengauer, A new measure for functional similarity of gene products based on Gene Ontology, BMC Bioinformatics, 7, 302, 2006.
- [8] C. Pesquita, D. Faria, H. Bastos, A. Falcao, F. Couto, Evaluating GO-based Semantic Similarity Measures, In: Proc. 10th Annual Bio-Ontologies Meeting 2007, 37 - 40, 2007

See Also

[getMinimumSubsumer](#), [getDisjCommAnc](#), [setEnrichmentFactors](#), [setOntology](#), [load.diffusion.kernel](#)

Examples

```
# setOntology ("BP")

# Lin's method
getTermSim(c("GO:0006955", "GO:0007584"), method="Lin")
if(require(graph)){
  # Couto's method combined with Jiang-Conrath distance
  getTermSim(c("GO:0006955", "GO:0007584"), method="CoutoJiangConrath")

  # set enrichment factors
  setEnrichmentFactors(alpha=0.1, beta=0.5)
  getTermSim(c("GO:0006955", "GO:0007584"), method="CoutoEnriched")
}
```

GOenrichment

GO enrichment analysis

Description

This function performs a GO enrichment analysis using topGO. It combines the two former functions "GOenrichment" and "analyzeCluster".

Usage

```
GOenrichment(genesOfInterest, allgenes, cutoff=0.01)
```

Arguments

<code>genesOfInterest</code>	character vector of Entrez gene IDs or vector of statistics (p-values, t-statistics, ...) named with entrez gene IDs
<code>allgenes</code>	character vector of Entrez gene IDs or vector of statistics named with entrez gene IDs
<code>cutoff</code>	significance cutoff for GO enrichment analysis

Details

If the parameters 'genesOfInterest' and 'allgenes' are both character vectors of Entrez gene IDs, the 'elim' method is used, otherwise the 'weight' method. For more details please refer to the topGO vignette.

Value

<code>GOterms</code>	list of significant GO terms and their description
<code>p.values</code>	vector of p-values for significant GO terms
<code>genes</code>	list of genes associated to each GO term

Author(s)

Holger Froehlich

References

Adrian Alexa, Jörg Rahnenführer, Thomas Lengauer: Improved scoring of functional groups from gene expression data by decorrelating GO graph structure, *Bioinformatics*, 2006, 22(13):1600-1607

See Also

[evaluateClustering](#)

Examples

```
## Not run:

setOntology("BP")
gomap <- get("gomap",env=GOSimEnv)
allgenes = sample(names(gomap), 1000) # suppose these are all genes
allpvalues = runif(1000) # an these are their pvalues
names(allpvalues) = allgenes
if(require(topGO) & require(annotate))
  GOenrichment(allpvalues[allpvalues<0.05], allpvalues) # GO enrichment analysis

## End(Not run)
```

Description

- "ICshumanBPall"** Information content of GO terms in "biological process" using all evidence codes for human
- "ICshumanCCall"** Information content of GO terms in "cellular component" using all evidence codes for human
- "ICshumanMFall"** Information content of GO terms in "molecular function" using all evidence codes for human
- "ICsyeastBPall"** Information content of GO terms in "biological process" using all evidence codes for yeast
- "ICsyeastCCall"** Information content of GO terms in "cellular component" using all evidence codes for yeast
- "ICsyeastMFall"** Information content of GO terms in "molecular function" using all evidence codes for yeast
- "ICsflyBPall"** Information content of GO terms in "biological process" using all evidence codes for fly
- "ICsflyCCall"** Information content of GO terms in "cellular component" using all evidence codes for fly
- "ICsflyMFall"** Information content of GO terms in "molecular function" using all evidence codes for fly
- "ICsmalariaBPall"** Information content of GO terms in "biological process" using all evidence codes for malaria
- "ICsmalariaCCall"** Information content of GO terms in "cellular component" using all evidence codes for malaria
- "ICsmalariaMFall"** Information content of GO terms in "molecular function" using all evidence codes for malaria
- "ICsratBPall"** Information content of GO terms in "biological process" using all evidence codes for rat
- "ICsratCCall"** Information content of GO terms in "cellular component" using all evidence codes for rat
- "ICsratMFall"** Information content of GO terms in "molecular function" using all evidence codes for rat
- "ICsmouseBPall"** Information content of GO terms in "biological process" using all evidence codes for mouse
- "ICsmouseCCall"** Information content of GO terms in "cellular component" using all evidence codes for mouse
- "ICsmouseMFall"** Information content of GO terms in "molecular function" using all evidence codes for mouse

Format

A vector of double values

Note

The currently used IC values can be accessed within the GOSimEnv environment.

internal	<i>internal functions</i>
----------	---------------------------

Description

internal functions: do not call these functions directly.

Usage

various

Arguments

various

Value

various

Author(s)

Holger Froehlich

selectPrototypes	<i>Heuristic selection of prototypes and dimensionality reduction of feature vectors.</i>
------------------	---

Description

- Heuristic selection of prototypes
- Dimensionality reduction of feature vectors

Usage

```
selectPrototypes(n = 250, method = "frequency", data = NULL, verbose = TRUE)
```

Arguments

n	number of prototypes or maximum number of clusters
method	method to select prototypes or to perform subset selection
data	data matrix (l x d) of feature vectors (l = number of genes)
verbose	print out information

Details

The following heuristics to perform automatic selection of prototypes are implemented:

"frequency" select n genes with highest number of GO annotations in the currently selected ontology

"random" select n genes uniform randomly over all genes with annotations in the currently selected ontology

To perform dimensionality reduction implemented methods are:

"PCA" dimensionality reduction via principal component analysis; the number of principal components is determined such that at least 95% of total variance in feature space can be explained

"clustering" EM-clustering in feature space

Value

If the function is called to automatically select prototypes, a character vector of gene IDs is returned.

If the function is called to perform dimensionality via PCA, the result is a list with items

If the function is called to perform clustering in feature space, the cluster centers are returned in a l x k matrix (each column is one cluster center). The "Mclust" function in the package "mclust" is called to perform the clustering. The BIC is used to calculate the optimal number of clusters in the range 2,...,n.

Note

The result depends on the currently set ontology ("BP","MF","CC").

Author(s)

Holger Froehlich

References

- [1] H. Froehlich, N. Speer, C. Spieth, and A. Zell, Kernel Based Functional Gene Grouping, Proc. Int. Joint Conf. on Neural Networks (IJCNN), pp. 6886 - 6891, 2006
- [2] N. Speer, H. Froehlich, A. Zell, Functional Grouping of Genes Using Spectral Clustering and Gene Ontology, Proc. Int. Joint Conf. on Neural Networks (IJCNN), pp. 298 - 303, 2005

See Also

[getGeneFeaturesPrototypes](#), [getGeneSimPrototypes](#), [setOntology](#)

Examples

```
## Not run:
# takes too much time in the R CMD check
proto=selectPrototypes(n=50) # --> returns a character vector of 50 genes with the highest
feat=getGeneFeaturesPrototypes(c("207", "208", "7494"), prototypes=proto, pca=FALSE) # --> comp
selectPrototypes(data=feat$features, method="pca") # ... and PCA projection

## End(Not run)
```

```
setEnrichmentFactors
```

Set the depth and density enrichment factors for GO term similarity.

Description

Sets the depth and density enrichment factors for the enriched FuSSiMeg GO term similarity measure by Couto et al.

Usage

```
setEnrichmentFactors(alpha = 0.5, beta = 0.5)
```

Arguments

alpha	depth factor
beta	density factor

Value

none

Note

The enrichment factors are stored internally and are used by the function [getTermSim](#), if one uses the method "CoutoEnriched" to calculate GO term similarities

Author(s)

Holger Froehlich

References

F.Couto, M. Silva, P. Coutinho, Implementation of a Functional Semantic Similarity Measure between Gene-Products, DI/FCUL TR 03-29, Department of Informatics, University of Lisbon, 2003

See Also

[getTermSim](#)

Examples

```
if(require(graph)){
  setEnrichmentFactors(alpha=0.1,beta=0.5)
  getTermSim(c("GO:0006955","GO:0007584"),method="CoutoEnriched")
}
```

setEvidenceLevel *Specifies to use only GO terms with given evidence codes.*

Description

Specifies to use only GO terms with given evidence codes. This, in combination with the specified GO ontology ("BP", "MF", "CC"), influences, how the information content for individual GO terms is calculated.

Usage

```
setEvidenceLevel(evidences = "all", organism=org.Hs.egORGANISM, gomap=org.Hs.egGO)
```

Arguments

evidences	character vector of evidence codes
organism	organism, for which to load a mapping of Entrez gene IDs to GO terms (see details)
gomap	mapping of Entrez gene IDs to GO terms to be used (see details)

Details

Each evidence code can be one of:

- "IMP"** inferred from mutant phenotype
- "IGI"** inferred from genetic interaction
- "IPI"** inferred from physical interaction
- "ISS"** inferred from sequence similarity
- "IDA"** inferred from direct assay
- "IEP"** inferred from expression pattern
- "IEA"** inferred from electronic annotation
- "TAS"** traceable author statement
- "NAS"** non-traceable author statement
- "ND"** no biological data available
- "IC"** inferred by curator


```

$'11306'$'GO:0006810'$Ontology
1
"BP"
$'11306'$'GO:0006879'
$'11306'$'GO:0006879'$GOID
1
"GO:0006879"
$'11306'$'GO:0006879'$Evidence
1
"IMP"
$'11306'$'GO:0006879'$Ontology
1
"BP"

```

Value

The mapping is stored in the GOSimEnv environment.

Note

By default all evidence codes are used. If another behavior is wanted, one has to recalculate the information content of all GO terms via `calcICs`. The evidence level influences the behavior of all other functions, especially `filterGO` and `getGOInfo`.

Author(s)

Holger Froehlich

References

<www.geneontology.org>

See Also

`setOntology`, `calcICs`, `filterGO`, `getGOInfo`

Examples

```
## Not run: setEvidenceLevel("all") # the default behavior
```

setOntology *Set an ontology as base for subsequent computations.*

Description

Sets the ontology that all subsequent computations are based on and loads the information content of all GO terms within this ontology. At load time of the library the default ontology is "BP". Further, on running this function the environment GOSimEnv is reinitialized, i.e. all global settings or parameters used in the library are reset to their default values.

Usage

```
setOntology(ont = "BP", loadIC=TRUE)
```

Arguments

ont	the ontology to use ("BP","MF","CC")
loadIC	Should the corresponding file with precomputed IC-values be loaded? Default: TRUE. WARNING: If the file is not loaded, no calculations can be performed! This might only be useful, if you want to recalculate IC values.

Details

The following ontologies can be used:

- "BP" biological process
- "MF" molecular function
- "CC" cellular component

Value

none.

Author(s)

Holger Froehlich

Examples

```
# set ontology to "molecular function"
## Not run:
setOntology("MF")
# calculate Resnik similarity of two GO terms within this ontology
getTermSim(c("GO:0004060", "GO:0003867"), method="Resnik")
## End(Not run)
```

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