

Package ‘DOSE’

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Type Package

Title Disease Ontology Semantic and Enrichment analysis

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Description This package implements five methods proposed by Resnik, Schlicker, Jiang, Lin and Wang respectively for measuring semantic similarities among DO terms and gene products. Enrichment analyses including hypergeometric model and gene set enrichment analysis are also implemented for discovering disease associations of high-throughput biological data.

Depends R (>= 3.5.0)

Imports AnnotationDbi, enrichit (>= 0.0.4), ggplot2, GOSemSim (>= 2.37.1), methods, reshape2, utils, yulab.utils (> 0.2.2)

Suggests prettydoc, clusterProfiler, gson (>= 0.0.5), knitr, memoise, org.Hs.eg.db, rmarkdown, testthat

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URL <https://yulab-smu.top/contribution-knowledge-mining/>

BugReports <https://github.com/GuangchuangYu/DOSE/issues>

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DOSE-package

DOSE: Disease Ontology Semantic and Enrichment analysis

Description

This package implements five methods proposed by Resnik, Schlicker, Jiang, Lin and Wang respectively for measuring semantic similarities among DO terms and gene products. Enrichment analyses including hypergeometric model and gene set enrichment analysis are also implemented for discovering disease associations of high-throughput biological data.

Author(s)

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See Also

Useful links:

- <https://yulab-smu.top/contribution-knowledge-mining/>
- Report bugs at <https://github.com/GuangchuangYu/DOSE/issues>

`clusterSim`*clusterSim*

Description

semantic similarity between two gene clusters

Usage

```
clusterSim(  
  cluster1,  
  cluster2,  
  ont = "HDO",  
  organism = "hsa",  
  measure = "Wang",  
  combine = "BMA"  
)
```

Arguments

<code>cluster1</code>	a vector of gene IDs
<code>cluster2</code>	another vector of gene IDs
<code>ont</code>	one of "HDO", "HPO" and "MPO"
<code>organism</code>	one of "hsa" and "mmu"
<code>measure</code>	One of "Resnik", "Lin", "Rel", "Jiang" and "Wang" methods.
<code>combine</code>	One of "max", "avg", "rcmax", "BMA" methods, for combining

Details

given two gene clusters, this function calculates semantic similarity between them.

Value

similarity

Author(s)

Yu Guangchuang

Examples

```
## Not run:  
cluster1 <- c("835", "5261", "241", "994")  
cluster2 <- c("307", "308", "317", "321", "506", "540", "378", "388", "396")  
clusterSim(cluster1, cluster2)  
  
## End(Not run)
```

computeIC	<i>compute information content</i>
-----------	------------------------------------

Description

compute information content

Usage

```
computeIC(ont = "HDO")
```

Arguments

ont one of "DO", "HPO" and "MPO"

Author(s)

Guangchuang Yu <https://yulab-smu.top>

DataSet	<i>Datasets</i>
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Description

Information content and DO term to entrez gene IDs mapping

doseSim	<i>doseSim</i>
---------	----------------

Description

measuring similarities between two DO term vectors.

Usage

```
doseSim(DO1, DO2, measure = "Wang", ont = "HDO")
```

```
doSim(DO1, DO2, measure = "Wang", ont = "HDO")
```

Arguments

DO1 DO term, MPO term or HPO term vector
 DO2 DO term, MPO term or HPO term vector
 measure one of "Wang", "Resnik", "Rel", "Jiang", "Lin", and "TCSS".
 ont one of "HDO", "HPO" and "MPO"

Details

provide two term vectors, this function will calculate their similarities.

Value

score matrix

Author(s)

Guangchuang Yu <https://yulab-smu.top>

dose_params

Shared parameters for DOSE functions

Description

Shared parameters for DOSE functions

Arguments

gene	a vector of entrez gene id
organism	one of "hsa" and "mmu"
ont	one of "HDO", "HPO" or "MPO"
pvalueCutoff	pvalue cutoff
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
universe	background genes
minGSSize	minimal size of genes annotated by ontology term for testing
maxGSSize	maximal size of each geneSet for analyzing
qvalueCutoff	qvalue cutoff
readable	whether mapping gene ID to gene Name
geneList	order ranked geneList
exponent	weight of each step
nPerm	permutation numbers
verbose	print message or not
adaptive	logical, use adaptive permutation or not (default: FALSE)
minPerm	minimum number of permutations for adaptive mode (default: 1000)
maxPerm	maximum number of permutations for adaptive mode (default: 10000)
method	method of GSEA, one of "multilevel", "permute", "sample"

enrichDGN *Enrichment analysis based on the DisGeNET* (<http://www.disgenet.org/>)

Description

given a vector of genes, this function will return the enrichment NCG categories with FDR control

Usage

```
enrichDGN(
  gene,
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  universe,
  minGSSize = 10,
  maxGSSize = 500,
  qvalueCutoff = 0.2,
  readable = FALSE
)
```

Arguments

gene	a vector of entrez gene id
pvalueCutoff	pvalue cutoff
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
universe	background genes
minGSSize	minimal size of genes annotated by ontology term for testing
maxGSSize	maximal size of each geneSet for analyzing
qvalueCutoff	qvalue cutoff
readable	whether mapping gene ID to gene Name

Value

A enrichResult instance

Author(s)

Guangchuang Yu

References

Janet et al. (2015) DisGeNET: a discovery platform for the dynamical exploration of human diseases and their genes. *Database* bav028 <http://database.oxfordjournals.org/content/2015/bav028.long>

`enrichDGNv`*enrichDGN*

Description

Enrichment analysis based on the DisGeNET (<http://www.disgenet.org/>)

Usage

```
enrichDGNv(  
  snp,  
  pvalueCutoff = 0.05,  
  pAdjustMethod = "BH",  
  universe,  
  minGSSize = 10,  
  maxGSSize = 500,  
  qvalueCutoff = 0.2,  
  readable = FALSE  
)
```

Arguments

<code>snp</code>	a vector of SNP
<code>pvalueCutoff</code>	pvalue cutoff
<code>pAdjustMethod</code>	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
<code>universe</code>	background genes
<code>minGSSize</code>	minimal size of genes annotated by ontology term for testing
<code>maxGSSize</code>	maximal size of each geneSet for analyzing
<code>qvalueCutoff</code>	qvalue cutoff
<code>readable</code>	whether mapping gene ID to gene Name

Details

given a vector of genes, this function will return the enrichment NCG categories with FDR control

Value

A `enrichResult` instance

Author(s)

Guangchuang Yu

References

Janet et al. (2015) DisGeNET: a discovery platform for the dynamical exploration of human diseases and their genes. *Database* bav028 <http://database.oxfordjournals.org/content/2015/bav028.long>

`enrichDO`*DO Enrichment Analysis*

Description

Given a vector of genes, this function will return the enrichment DO categories with FDR control.

Usage

```
enrichDO(  
  gene,  
  ont = "HDO",  
  organism = "hsa",  
  pvalueCutoff = 0.05,  
  pAdjustMethod = "BH",  
  universe,  
  minGSSize = 10,  
  maxGSSize = 500,  
  qvalueCutoff = 0.2,  
  readable = FALSE  
)
```

Arguments

<code>gene</code>	a vector of entrez gene id
<code>ont</code>	one of "HDO", "HPO" or "MPO"
<code>organism</code>	one of "hsa" and "mmu"
<code>pvalueCutoff</code>	pvalue cutoff
<code>pAdjustMethod</code>	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
<code>universe</code>	background genes
<code>minGSSize</code>	minimal size of genes annotated by ontology term for testing
<code>maxGSSize</code>	maximal size of each geneSet for analyzing
<code>qvalueCutoff</code>	qvalue cutoff
<code>readable</code>	whether mapping gene ID to gene Name

Value

A `enrichResult` instance.

Author(s)

Guangchuang Yu <https://yulab-smu.top>

Examples

```
data(geneList)  
gene = names(geneList)[geneList > 1]  
yy = enrichDO(gene, pvalueCutoff=0.05)  
summary(yy)
```

`enrichNCG`*enrichNCG*

Description

Enrichment analysis based on the Network of Cancer Genes database (<http://ncg.kcl.ac.uk/>)

Usage

```
enrichNCG(  
  gene,  
  pvalueCutoff = 0.05,  
  pAdjustMethod = "BH",  
  universe,  
  minGSSize = 10,  
  maxGSSize = 500,  
  qvalueCutoff = 0.2,  
  readable = FALSE  
)
```

Arguments

<code>gene</code>	a vector of entrez gene id
<code>pvalueCutoff</code>	pvalue cutoff
<code>pAdjustMethod</code>	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
<code>universe</code>	background genes
<code>minGSSize</code>	minimal size of genes annotated by ontology term for testing
<code>maxGSSize</code>	maximal size of each geneSet for analyzing
<code>qvalueCutoff</code>	qvalue cutoff
<code>readable</code>	whether mapping gene ID to gene Name

Details

given a vector of genes, this function will return the enrichment NCG categories with FDR control

Value

A `enrichResult` instance

Author(s)

Guangchuang Yu

gene2DO *convert Gene ID to DO Terms*

Description

provide gene ID, this function will convert to the corresponding DO Terms

Usage

```
gene2DO(gene, organism = "hsa", ont = "HDO")
```

Arguments

gene	entrez gene ID
organism	organism
ont	ont

Value

DO Terms

Author(s)

Guangchuang Yu <https://yulab-smu.top>

geneSim *geneSim*

Description

measuring similarities bewteen two gene vectors.

Usage

```
geneSim(  
  geneID1,  
  geneID2 = NULL,  
  ont = "HDO",  
  organism = "hsa",  
  measure = "Wang",  
  combine = "BMA"  
)
```

Arguments

geneID1	entrez gene vector
geneID2	entrez gene vector
ont	one of "HDO" and "MPO"
organism	one of "hsa" and "mmu"
measure	one of "Wang", "Resnik", "Rel", "Jiang", and "Lin".
combine	One of "max", "avg", "rcmax", "BMA" methods, for combining semantic similarity scores of multiple DO terms associated with gene/protein.

Details

provide two entrez gene vectors, this function will calculate their similarity.

Value

score matrix

Author(s)

Guangchuang Yu <https://yulab-smu.top>

Examples

```
g <- c("835", "5261", "241", "994")
geneSim(g)
```

gseDGN

DisGeNET Gene Set Enrichment Analysis

Description

perform gsea analysis

Usage

```
gseDGN(  
  geneList,  
  exponent = 1,  
  nPerm = 1000,  
  minGSSize = 10,  
  maxGSSize = 500,  
  pvalueCutoff = 0.05,  
  pAdjustMethod = "BH",  
  verbose = TRUE,  
  method = "multilevel",  
  adaptive = FALSE,  
  minPerm = 1000,  
  maxPerm = 10000,  
  ...  
)
```

Arguments

geneList	order ranked geneList
exponent	weight of each step
nPerm	permutation numbers
minGSSize	minimal size of genes annotated by ontology term for testing
maxGSSize	maximal size of each geneSet for analyzing
pvalueCutoff	pvalue cutoff
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
verbose	print message or not
method	method of GSEA, one of "multilevel", "permute", "sample"
adaptive	logical, use adaptive permutation or not (default: FALSE)
minPerm	minimum number of permutations for adaptive mode (default: 1000)
maxPerm	maximum number of permutations for adaptive mode (default: 10000)
...	other parameter

Value

gseaResult object

Author(s)

Guangchuang Yu

gseDO

DO Gene Set Enrichment Analysis

Description

perform gsea analysis

Usage

```
gseDO(
  geneList,
  ont = "HDO",
  organism = "hsa",
  exponent = 1,
  nPerm = 1000,
  minGSSize = 10,
  maxGSSize = 500,
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  verbose = TRUE,
  method = "multilevel",
  adaptive = FALSE,
  minPerm = 1000,
  maxPerm = 10000,
  ...
)
```

Arguments

geneList	order ranked geneList
ont	one of "HDO", "HPO" or "MPO"
organism	one of "hsa" and "mmu"
exponent	weight of each step
nPerm	permutation numbers
minGSSize	minimal size of genes annotated by ontology term for testing
maxGSSize	maximal size of each geneSet for analyzing
pvalueCutoff	pvalue cutoff
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
verbose	print message or not
method	method of GSEA, one of "multilevel", "permute", "sample"
adaptive	logical, use adaptive permutation or not (default: FALSE)
minPerm	minimum number of permutations for adaptive mode (default: 1000)
maxPerm	maximum number of permutations for adaptive mode (default: 10000)
...	other parameter

Value

gseaResult object

Author(s)

Guangchuang Yu

gseNCG

NCG Gene Set Enrichment Analysis

Description

perform gsea analysis

Usage

```
gseNCG(
  geneList,
  exponent = 1,
  nPerm = 1000,
  minGSSize = 10,
  maxGSSize = 500,
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  verbose = TRUE,
  method = "multilevel",
  adaptive = FALSE,
  minPerm = 1000,
  maxPerm = 10000,
  ...
)
```

Arguments

geneList	order ranked geneList
exponent	weight of each step
nPerm	permutation numbers
minGSSize	minimal size of genes annotated by ontology term for testing
maxGSSize	maximal size of each geneSet for analyzing
pvalueCutoff	pvalue cutoff
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
verbose	print message or not
method	method of GSEA, one of "multilevel", "permute", "sample"
adaptive	logical, use adaptive permutation or not (default: FALSE)
minPerm	minimum number of permutations for adaptive mode (default: 1000)
maxPerm	maximum number of permutations for adaptive mode (default: 10000)
...	other parameter

Value

gseaResult object

Author(s)

Guangchuang Yu

mclusterSim

mclusterSim

Description

Pairwise semantic similarity for a list of gene clusters

Usage

```
mclusterSim(
  clusters,
  ont = "HDO",
  organism = "hsa",
  measure = "Wang",
  combine = "BMA"
)
```

Arguments

clusters	A list of gene clusters
ont	one of "HDO", "HPO" and "MPO"
organism	organism
measure	one of "Wang", "Resnik", "Rel", "Jiang", and "Lin".
combine	One of "max", "avg", "rcmax", "BMA" methods, for combining semantic similarity scores of multiple DO terms associated with gene/protein.

Value

similarity matrix

Author(s)

Guangchuang Yu

Examples

```
## Not run:
cluster1 <- c("835", "5261", "241")
cluster2 <- c("578", "582")
cluster3 <- c("307", "308", "317")
clusters <- list(a=cluster1, b=cluster2, c=cluster3)
mclusterSim(clusters, measure="Wang")

## End(Not run)
```

reexports

Objects exported from other packages

Description

These objects are imported from other packages. Follow the links below to see their documentation.

ggplot2 [facet_grid](#)

GOSemSim [get_organism](#)

simplot

simplot

Description

plotting similarity matrix

Usage

```
simplot(
  sim,
  xlab = "",
  ylab = "",
  color.low = "white",
  color.high = "red",
  labs = TRUE,
  digits = 2,
  labs.size = 3,
  font.size = 14
)
```

Arguments

<code>sim</code>	similarity matrix
<code>xlab</code>	xlab
<code>ylab</code>	ylab
<code>color.low</code>	color of low value
<code>color.high</code>	color of high value
<code>labs</code>	logical, add text label or not
<code>digits</code>	round digit numbers
<code>labs.size</code>	lable size
<code>font.size</code>	font size

Value

ggplot object

Author(s)

Yu Guangchuang

theme_dose

theme_dose

Description

ggplot theme of DOSE

Usage

```
theme_dose(font.size = 14)
```

Arguments

<code>font.size</code>	font size
------------------------	-----------

Value

ggplot theme

Examples

```
library(ggplot2)
qplot(1:10) + theme_dose()
```

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