

# Package ‘categoryCompare’

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**Version** 1.0.2

**Title** Meta-analysis of high-throughput experiments using feature annotations

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**License** GPL-2

**Depends** R (>= 2.10), Biobase (>= 1.15.29), AnnotationDbi (>= 0.1.15),Category

**Suggests** methods, GSEABase, hwriter, colorspace, graph, GO.db,KEGG.db, estrogen, org.Hs.eg.db, hgu95av2.db, limma, affy,genefilter

**Imports** Biobase (>= 1.15.29), AnnotationDbi (>= 0.1.15), hwriter,GSEABase, Category (>= 2.21.2), GOstats, annotate, colorspace,graph, RCytoscape (>= 1.5.11)

**LazyLoad** yes

## Description

Calculates significant annotations (categories) in each of two (or more) feature (i.e. gene) lists, determines the overlap between the annotations, and returns graphical and tabular data about the significant annotations and which combinations of feature lists the annotations were found to be significant. Interactive exploration is facilitated through the use of RCytoscape (heavily suggested).

**biocViews** Bioinformatics, Annotation, GO, MultipleComparisons,Pathways, GeneExpression

**SystemRequirements** Cytoscape (>= 2.8.0) (if used for visualization of results, heavily suggested), CytoscapeRPC plugin (>= 1.8)

**TODO** Text and HTML output without graphs.

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categoryCompare-package

*Meta-analysis of high-throughput experiments using feature annotations*

---

## Description

Calculates significant annotations (categories) in each of two (or more) feature (i.e. gene) lists, determines the overlap between the annotations, and returns graphical and tabular data about the significant annotations and which combinations of feature lists the annotations were found to be significant. Interactive exploration is facilitated through the use of RCytoscape (heavily suggested).

## Details

Package:	categoryCompare
Version:	0.99.1
License:	GPL-2
Depends:	Biobase (>= 1.15.29), AnnotationDbi (>= 0.1.15), Category
Suggests:	methods, GSEABase, hwriter, colorspace, graph, GO.db, KEGG.db, estrogen, org.Hs.eg.db, hgu9
Imports:	Biobase (>= 1.15.29), AnnotationDbi (>= 0.1.15), hwriter, GSEABase, Category (>= 2.21.2), GO
LazyLoad:	yes
biocViews:	Bioinformatics, Annotation, GO, MultipleComparisons, Pathways, GeneExpression
SystemRequirements:	Cytoscape (>= 2.8.0) (if used for visualization of results, heavily suggested), CytoscapeRPC plug
TODO:	Text and HTML output without graphs.

Built: R 2.15.0; ; 2012-03-15 18:42:40 UTC; windows

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```

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Further information is available in the following vignettes:

categoryCompare\_vignette    categoryCompare: High-throughput data meta-analysis using gene annotations (source)

### Author(s)

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breakEdges

*Break Cytoscape (or graphNEL) Network Edges*

---

### Description

Removes those edges in a graph network with the edge attribute that is "under" or "over" the cutoff value supplied.

### Usage

```
breakEdges(cwObject, cutoff, edgeAtt='weight', valDir='under', layout='force-directed')
```

### Arguments

cwObject	The object returned by <a href="#">ccOutCyt</a>
cutoff	What is the cutoff to select edges
edgeAtt	Which edge attribute should be used for deciding which edges to select
valDir	Select the edge attribute with values "under" (default) or "over" the input value
layout	What type of layout should be used after the edges are broken

### Details

When viewing annotations returned by [ccCompare](#), unless a hierarchical view is used, the annotations are linked by edges weighted by how many genes are shared between each annotation. The default cutoff is 0.1, any edges above that are kept. This can result in a big hairy mess when viewing the annotation network in Cytoscape. To help clean up the display and encourage finding functionally related sets of annotations, it is very useful to remove edges with low values of overlap, and re-layout the network. `breakEdges` does just that.

The other option is to break the highly overlapping edges prior to sending the network to Cytoscape, thereby speeding up the process, as the fewer edges one has to "push" to Cytoscape the faster it will go.

### Value

The first returns nothing, rather it modifies the network in Cytoscape itself.

The second returns a new `ccCompareResult` object for use by [ccOutCyt](#)

**Author(s)**

Robert M Flight

**See Also**[ccOutCyt breakEdges-methods](#)**Examples**

```
## Not run:
require(RCytoscape)
g <- makeSimpleGraph()
cw <- CytoscapeWindow('breakEdges',g)
displayGraph(cw)
layout(cw)
redraw(cw)
breakEdges(cw, 'score', 0, layout=NULL)

## End(Not run)
```

---

breakEdges-methods      *Methods for Function breakEdges in Package **categoryCompare***

---

**Description**

Methods for function breakEdges in package **categoryCompare**

**Methods**

signature(cwObject = "ccCompareResult", cutoff = "numeric") Allows one to remove edges in the ccCompareResult mainGraph slot prior to passing it into Cytoscape for visualization. Given that the number of edges can be rather large (especially for Gene Ontology) this can easily speed up the transfer, without actually losing any information.

signature(cwObject = "CytoscapeWindowClass", cutoff = "numeric") Once an annotation graph is in Cytoscape, remove edges above or below the cutoff. Note that this does not affect the original graph in the ccCompareResult object.

**Author(s)**

Robert M Flight

**See Also**[CytoscapeWindowClass breakEdges ccCompareResult ccOutCyt](#)

**Examples**

```

data(ccData)

# breaking the edges in a ccCompareResult
ccResults$BP <- breakEdges(ccResults$BP, 0.8)
## Not run:
hasCy <- (if (.Platform$OS.type %in% "windows") { (length(grep("Cytoscape", system("tasklist", intern=TRUE))) > 0)}

if hasCy {

cwObj <- ccOutCyt(ccResults$BP, ccOpts)
# now breaking them in the CytoscapeWindow object
breakEdges(cwObj, 0.85)
Sys.sleep(10)
RCytoscape::deleteWindow(cwObj)
}

## End(Not run)

```

---

ccCompare-methods

*Comparison of enriched annotations*


---

**Description**

Takes the results from [ccEnrich](#) and compares the enriched annotations based on the settings previously set in [ccOptions](#). Returns a `ccCompareResult` or `ccCompareCollection` object, see [Details](#).

**Usage**

```
ccCompare(ccEnrichResult, ccOptions)
```

**Arguments**

`ccEnrichResult` The enriched annotations collection returned from [ccEnrich](#). This can be the `ccEnrichCollection`, `GOccEnrichResult`, or `KEGGccEnrichResult`

`ccOptions` A [ccOptions](#) object that will determine which lists to actually compare against each other. See details below.

**Details**

Based on the enrichments found for each gene list, we now want to compare the annotations between lists. `ccCompare` accesses the annotations for each enrichment performed for each list, and makes the comparisons defined in `ccOptions`.

**Value**

`ccCompare` generates both a graph of the comparisons (to show how the categories are linked to each list and each other) and tabular output. The tabular output is a data frame, with ID for each term that was considered as a candidate annotation for each list, as well as a long description (`Desc`) of what the term is, and then membership and statistics from each gene list.

For each type of comparison (GO, KEGG, etc) a `ccCompareResult` is generated, with the following slots:

mainGraph	Annotations arranged as a graph
mainTable	The tabular results from all enrichment calculations combined into one
allAnnotation	A list of lists, where each entry is the annotation identifier, then a list for each comparison, with the genes that are annotated to that term that also belong to each list

The default is to generate an overlap graph for GO and KEGG, where the overlap is a measure of the similarity of the features (genes) annotated to each annotation term (based on a formula from `EnrichmentMap`). Optionally for GO, one can generate a hierarchical layout where the parent GO terms of the significant terms will also be included in the graph, with term origin saved in the node annotation (see example below to do this).

Only those terms with more than 10 and less than 500 annotated genes (according to the GO annotation file) are included.

When using weighted overlap graphs and **RCytoscape** for viewing, it is recommended to use `breakEdges` and `minNodes` to remove edges with low weights and nodes with only a few genes from the dataset annotated to them.

### Author(s)

Robert M Flight

### See Also

[ccCompareResult](#) [ccCompareCollection](#) [ccOutCyt](#) [breakEdges](#) [outType](#) [ccEnrich](#)

### Examples

```
## Not run:
require(GO.db)
require(KEGG.db)
require(org.Hs.eg.db)

## End(Not run)
data(ccData)

# note that enrichLists is generated from ccEnrich
# ccResults <- ccCompare(enrichLists,ccOpts)
ccResults

# use the GO hierarchy tree
graphType(enrichLists$BP) <- "hierarchical"
# ccResultsBPHier <- ccCompare(enrichLists$BP,ccOpts)
ccResultsBPHier
```

---

ccCompareCollection-class

*Class "ccCompareCollection"*

---

### Description

Holds multiple `ccCompareResult` objects.

### Objects from the Class

Objects can be created by calls of the form `new("ccCompareCollection", ...)`.

These are not normally created by the user, but rather by `ccCompare` while performing the categorical comparisons for each type of category

### Slots

`.Data`: Object of class "list"

`names`: Object of class "character"

### Extends

Class "`namedList`", directly. Class "`list`", by class "namedList", distance 2. Class "`vector`", by class "namedList", distance 3. Class "`AssayData`", by class "namedList", distance 3.

### Methods

No methods defined with class "ccCompareCollection" in the signature.

### Author(s)

Robert M Flight

### See Also

`ccCompareResult` `ccCompare`

### Examples

```
showClass("ccCompareCollection")
```

---

ccCompareGeneric-methods

*Methods for Function* `ccCompareGeneric` in Package **categoryCompare**

---

### Description

Methods for function `ccCompareGeneric` in package **categoryCompare**

### Methods

```
signature(gccResult = "GENccEnrichResult", ccOptions = "ccOptions")
```



---

```
ccCompareResult-class  Class "ccCompareResult"
```

---

### Description

Holds the results from a single category comparison

### Objects from the Class

Objects can be created by calls of the form `new("ccCompareResult", ...)`.

### Slots

**mainGraph:** Object of class "graph". Holds the graph describing the relationships between the annotations

**subGraph:** Object of class "list". Not currently used

**mainTable:** Object of class "data.frame". Table of results, with all the various statistics for each annotation in the category

**allAnnotation:** Object of class "list". For each annotation, which genes from which comparison are annotated to that particular annotation

**categoryName:** Object of class "character". Which category (e.g. GO, KEGG, etc) was used

**ontology:** Object of class "character". If GO, which ontology was used

### Methods

**allAnnotation** signature(object = "ccCompareResult"): ...

**mainGraph** signature(object = "ccCompareResult"): ...

**mainTable<-** signature(object = "ccCompareResult"): ...

### Author(s)

Robert M Flight

### See Also

[ccCompare](#) [ccCompareCollection](#)

### Examples

```
showClass("ccCompareResult")
```

---

ccData

*Test data for categoryCompare*

---

### Description

Processed data from the estrogen example data set

### Usage

```
data(ccData)
```

### Format

table10: Log-ratio output from **limma** for the comparison of presence-absence of estrogen at 10 hours

table48: Log-ratio output from **limma** for the comparison of presence-absence of estrogen at 48 hours

glUniverse: All of the genes measured on the chip

gseaRes: Toy results of GSEA analysis of 3 different tissues

enrichLists: Apply [ccEnrich](#) to a ccGeneList from table10 and table48

ccResults: Apply [ccCompare](#) to enrichLists

ccResultsBPHier: Modify enrichLists\$BP to use a "hierarchical" layout

geneLists: a ccGeneList generated from genes in table10 and table48

ccOpts: a ccOptions object describing what we are going to do as far as feature list comparisons

### Author(s)

Robert M Flight

### Source

Taken from the **estrogen** package in Bioconductor, and then processed using the normal **affy** and **limma** tools.

### See Also

[ccGeneList](#) [ccEnrichCollection](#) [ccCompareCollection](#) [ccEnrich](#) [ccCompare](#)

### Examples

```
data(ccData)
```

---

`ccEnrich-method`*Perform annotation enrichment for multiple gene lists*

---

## Description

Takes a `ccGeneList` object containing all the information needed to perform enrichment calculations for Gene Ontology.

## Usage

```
ccEnrich(ccGeneList)
```

## Arguments

`ccGeneList` A `ccGeneList` object, which is really just a list of lists, with some extra slots to tell us how to examine results. Each entry in the list should be named to allow identification later on. Each sub list should contain a vector `genes` denoting the genes of interest, a vector `universe` denoting the gene background (i.e. all genes on the chip), and an entry `annotation` denoting an organism database package (such as `org.Hs.eg.db`). See `ccGeneList` for more details regarding this object.

## Details

This function is essentially a wrapper for `hyperGTestCC` that performs all of the calculations for the many gene lists in one go, returning a list of `HyperGResultCC` objects, one for each of the `ccTypes` and each gene list. These various `HyperGResultCC` objects can then be accessed and results compared among the lists for each of the ontologies

## Value

A list of `HyperGResultCC` objects, one for each `ccType` and gene list, returned as `ccEnrichResult` objects for each `ccType`. This can be passed with a `ccOptions` object to `ccCompare` to generate actual annotation comparisons.

## Author(s)

Robert M Flight

## See Also

`ccGeneList`, `hyperGTestCC`, `ccEnrichResult`

## Examples

```
## Not run:
require(GO.db)
require(KEGG.db)
require(org.Hs.eg.db)

## End(Not run)
data(ccData)
```

```

g10 <- (unique(table10$Entrez[1:100]))
g48 <- (unique(table48$Entrez[1:100]))

list10 <- list(genes=g10, universe=gUniverse, annotation="org.Hs.eg.db")
list48 <- list(genes=g48, universe=gUniverse, annotation="org.Hs.eg.db")

geneLists <- list(T10=list10, T48=list48)
geneLists <- new("ccGeneList", geneLists, ccType=c("BP","KEGG"))
geneLists <- new("ccGeneList", geneLists, ccType=c("CC","KEGG"))

# set number of fdr runs to 0 to speed up runtime, not generally recommended.
geneLists <- new("ccGeneList", geneLists, ccType = c('BP','KEGG'), pvalueCutoff=0.01, fdr=0)
# enrichLists <- ccEnrich(geneLists)

```

---

ccEnrichCollection-class

*Class "ccEnrichCollection"*

---

### Description

Holds multiple classes of `ccEnrichResult` in one object to allow `ccCompare` to work on only the one object and generate all of the results of a comparison.

### Objects from the Class

Objects can be created by calls of the form `new("ccEnrichCollection", ...)`.

### Slots

**.Data:** Object of class "list"

**names:** Object of class "character" The names (generally GO ontologies or KEGG, but can be changed) of each set of results

### Extends

Class "`namedList`", directly. Class "`list`", by class "`namedList`", distance 2. Class "`vector`", by class "`namedList`", distance 3. Class "`AssayData`", by class "`namedList`", distance 3.

### Methods

**pvalueCutoff<-** signature(`r = "ccEnrichCollection"`): Changes the `pvalueCutoff` to be used to decide significant annotations for all of the contained `ccEnrichResult` objects

**pvalueType<-** signature(`object = "ccEnrichCollection"`): Changes whether to use p-values or fdr values to determine those annotations that are significant in all of the contained `ccEnrichResult` objects

**minCount<-** signature(`object = "ccEnrichCollection"`): how many features have to be annotated to a term to be reported as significant

**graphType** signature(`object = "ccEnrichCollection"`): Gets the type of graph that should be output for this collection

**Author(s)**

Robert M Flight

**See Also**[ccEnrich](#) [hyperGTestCC](#) [ccCompare](#) [ccEnrichResult](#)**Examples**

```
data(ccData)
enrichLists
```

---

 ccEnrichResult-class    *Class "ccEnrichResult"*


---

**Description**

Acts as a container object for multiple [HyperGResultCC](#) objects.

**Objects from the Class**

Objects can be created by calls of the form `new("ccEnrichResult", ...)`.

**Extends**

Class "[namedList](#)", directly. Class "[list](#)", by class "[namedList](#)", distance 2. Class "[vector](#)", by class "[namedList](#)", distance 3. Class "[AssayData](#)", by class "[namedList](#)", distance 3.

**Methods**

**fdR** signature(object = "ccEnrichResult"): get the number of runs using random feature lists were performed

**pvalueCutoff** signature(r = "ccEnrichResult"): what is the pvalueCutoff to determine significant annotations

**pvalueCutoff<-** signature(r = "ccEnrichResult"): change the pvalueCutoff for an annotation to be considered significant

**pvalueType<-** signature(object = "ccEnrichResult"): change whether p-values used are from "FDR" or raw p-values

**minCount** signature(object = "ccEnrichResult"): how many features need to belong to an annotation to be reported

**minCount<-** signature(object = "ccEnrichResult"): adjust the minCount

**graphType** signature(object = "ccEnrichResult"): what type of graph should be generated (generally set by the class of object)

**graphType<-** signature(object = "ccEnrichResult"): change the type of graph to generate by [ccCompare](#)

**Author(s)**

Robert M Flight

**Examples**

```

data(ccData)
enrichRes <- enrichLists[[1]]
fdr(enrichRes)
pvalueType(enrichRes)
enrichRes
pvalueType(enrichRes) <- 'pval'
enrichRes

pvalueCutoff(enrichRes)
pvalueCutoff(enrichRes) <- 0.01
enrichRes

```

---

ccGeneList-class	<i>Class "ccGeneList"</i>
------------------	---------------------------

---

**Description**

This stores the actual gene lists and related information that will be used in categoryCompare.

**Objects from the Class**

Objects can be created by calls of the form `new("ccGeneList", list)`. `ccGeneList` is actually just an extension of R list objects. The input `list` should be a list of lists. See [Details](#) for more information.

**Slots**

**fdr**: Object of class "numeric" The number of fdr runs to perform to account for different list sizes and term dependence

**pvalueCutoff**: Object of class "numeric" Value used to determine whether or not a particular term is significant or not

**ccType**: Object of class "character" What types of annotations to use. Currently supported ones include "BP", "MF", "CC" (from Gene Ontology) and "KEGG"

**testDirection**: Object of class "character" Are you interested in "over" or "under" represented annotations

**Methods**

**fdr** signature(object = "ccGeneList"): how many random runs to perform

**fdr<-** signature(object = "ccGeneList"): change the number of random runs

**pvalueCutoff** signature(object = "ccGeneList"): what is the pvalue to consider significant

**pvalueCutoff<-** signature(object = "ccGeneList"): change the cutoff for significance

**ccType** signature(object = "ccGeneList"): what type of annotations are going to be examined

**ccType<-** signature(object = "ccGeneList"): change the type of annotations to examine

**testDirection** signature(object = "ccGeneList"): query for "over" or "under" represented annotations

**testDirection<-** signature(object = "ccGeneList"): change the type of representation ("over" or "under")

**listNames** signature(object = "ccGeneList"): what are the names of the lists contained

**Details**

The input list should be a list of lists, with at least three sub-lists.

```
testList <- list(list1=list(genes='...',universe='...',annotation='...'), list2=list(...))
```

**genes** : These are the gene identifiers of the genes that are of interest (differentially expressed genes)

**universe** : All of the genes that were measured in this particular experiments (i.e. all the genes on the chip)

**annotation** : What organism or chip do these ID's come from (e.g. "org.Hs.eg.db" for Human Entrez gene ID's, "hgu133a.db" for probe ID's from the Affymetrix U133A chip)

**data** : A data-frame that contains extra information about the genes of interest. At the very least, the data-frame must have a column ID that matches the ID's contained in genes

What actually happens when running ccEnrich is that the appropriate HyperGParamsCC objects are generated for each geneList and each type of annotation (e.g. BP, CC, KEGG), and then the calculations performed on each one.

**Note**

The ccGeneList object is what will undergo all of the enrichment calculations. When the results are combined with the [ccOptions](#) object, we can get our results of actual comparisons between experiments.

**Author(s)**

Robert M Flight

**See Also**

[ccOptions](#)

**Examples**

```
data(ccData)
g10 <- (unique(table10$Entrez[1:100]))
g48 <- (unique(table48$Entrez[1:100]))

list10 <- list(genes=g10, universe=gUniverse, annotation="org.Hs.eg.db")
list48 <- list(genes=g48, universe=gUniverse, annotation="org.Hs.eg.db")

geneLists <- list(T10=list10, T48=list48)
geneLists <- new("ccGeneList", geneLists, ccType=c("BP","KEGG"))
geneLists
```

---

ccOptions-class      *Class "ccOptions"*

---

### Description

These objects store the various options required by categoryCompare for actually making comparisons and generating output.

### Objects from the Class

Objects can be created by calls of the form `new("ccOptions", listNames=c('list1','list2',etc))`. This is the minimum call required, and will generate a ccOptions object where comparisons are assumed between all the lists supplied. See the examples section for more examples of how to initialize new objects.

### Slots

**listNames:** Object of class "character" The actual names of the various datasets defined in the ccData object

**compareNames:** Object of class "character" Which lists to compare, each entry should be a comma separated list

**compareIndx:** Object of class "list" List indices for each of the comparison, not usually set by the user. Generated automatically.

**compareColors:** Object of class "character" For graphical and tabular output each comparison can be colored. Should be one color for each comparison. Can be either an n by 3 matrix of rgb triples, or a character vector of hexadecimal color codes, or character vector of color names ('red','green','blue', etc)

**cssClass:** Object of class "character" Classnames used when generating HTML tables to color entries. Generated automatically upon initialization, or modifying compareNames

**outType:** Object of class "character" Sets the type of output generated by ccTables. Valid types are "html", "text", "rcytoscape" or "none", default is "text" when the ccOptions object is initialized without an outType specified.

### Methods

**compareColors** signature(object = "ccOptions"): ...

**compareColors<-** signature(object = "ccOptions"): ...

**compareIndx** signature(object = "ccOptions"): ...

**compareNames** signature(object = "ccOptions"): ...

**compareNames<-** signature(object = "ccOptions"): ...

**cssClass** signature(object = "ccOptions"): ...

**listNames** signature(object = "ccOptions"): ...

**listNames<-** signature(object = "ccOptions"): ...

**outType** signature(object = "ccOptions"): ...

**outType<-** signature(object = "ccOptions"): ...



**Author(s)**

Robert M Flight

**Examples**

```

showClass("ccOptions")
## A very basic "ccOptions" for a comparison of two sets of data, "list1" and "list2"
c1 <- new("ccOptions", listNames=c('list1','list2'))
c1

## Now lets get a little more complicated
c1 <- new("ccOptions", listNames=c('list1','list2'), compareNames=c('list1,list2','list1,list3'), compareC
c1

# set the type of output you want to eventually produce
c1 <- new("ccOptions", listNames=c('list1','list2'), outType='html')
c1

c1 <- new("ccOptions", listNames=c('list1','list2'), outType=c('html','text','none'))
c1

## Using RGB colors
ccCols <- matrix(c(255,0,0, 0,0,255), nrow=2, ncol=3)
ccCols <- rgb(ccCols, maxColorValue=255)
c1 <- new("ccOptions", listNames=c('list1','list2','list3'), compareNames=c('list1,list2','list1,list3'), c
c1

## Using Hex colors
c1 <- new("ccOptions", listNames=c('list1','list2','list3'), compareNames=c('list1,list2','list1,list3'), c
c1

## or even using a color palette from R. Note that you need at least enough colors to cover all of individu
c1 <- new("ccOptions", listNames=c('list1','list2','list3'), compareNames=c('list1,list2','list1,list3'), c
c1

```

---

ccOutCyt-methods

*Methods for Function ccOutCyt in Package categoryCompare*


---

**Description**

Passes a ccCompareResult object to Cytoscape for interactive visualization of ccCompare results.

**Details**

Note that only some basic, required methods have been imported from RCytoscape for use with categoryCompare, and these are hidden in the functions within categoryCompare and are not visible to the user. If access to all the functionality of RCytoscape is desired (and trust me, there is a lot of useful stuff in there), then the user should use library(RCytoscape) directly.

**Methods**

signature(ccCompRes = "ccCompareResult", ccOpts = "ccOptions", ...) At a minimum, this method requires a ccCompareResult and a ccOptions to work.

... may include:

**layout = "character"** to override the default layout set by ccCompare, as well as options

**postText = "character"** to add a user set string to the Cytoscape window

In addition, any of the arguments to CytoscapeWindow may also be set, such as host or port.

**See Also**

[ccCompareResult](#) [ccOptions](#) [ccCompare](#) [CytoscapeWindowClass](#)

**Examples**

```
## Not run:
hasCy <- (if (.Platform$OS.type %in% "windows") { (length(grep("Cytoscape", system("tasklist", intern=TRUE))) > 0)}

if hasCy {

ccResults$BP <- breakEdges(ccResults$BP, 0.8)
cwObj <- ccOutCyt(ccResults$BP, ccOpts)
Sys.sleep(10)
RCytoscape::deleteWindow(cwObj)
}

## End(Not run)
```

---

ccSigList-class	<i>Class "ccSigList"</i>
-----------------	--------------------------

---

**Description**

Holds a generic list of significant annotations. Allows one to use Bioconductor annotation packages, or when combined into a GENccEnrichResult, use custom annotation / gene mappings.

**Objects from the Class**

Objects can be created by calls of the form new("ccSigList", ...).

**Slots**

**sigID:** Object of class "character"  
**categoryName:** Object of class "character"  
**ontology:** Object of class "character"  
**annotation:** Object of class "character"

**Methods**

**annotation** signature(object = "ccSigList"): ...  
**category** signature(object = "ccSigList"): ...  
**ontology** signature(object = "ccSigList"): ...  
**sigID** signature(object = "ccSigList"): ...

**Author(s)**

Robert M Flight

**See Also**

[GENccEnrichResult ccCompareGeneric](#)

**Examples**

```
showClass("ccSigList")
```

---

cwReload-methods

*Methods for Function cwReload in Package categoryCompare*


---

**Description**

Methods for function cwReload in package **categoryCompare**

**Methods**

```
signature(oldCW = "CytoscapeWindowClass", windowName = "character", ccOpts = "ccOptions")
```

---

cytOutData-methods

*Methods for Function cytOutData*


---

**Description**

Takes the saveObj generated by cytOutNodes and writes the data to a file

**Value**

A text file with the annotations previously saved using cytOutNodes

**Methods**

```
signature(saveObj = "list", compareResult = "ccCompareResult", mergedData = "mergedData")
```

saveObj is the list object generated by cytOutNodes, compareResult is the object from ccCompare, and mergedData is created using mergeLists, but is optional.

... : optional arguments also include: orgType, default is "header" where each group is separate, "annotate" pushes all the data into one table with a new column that designates which groups the annotation was found in; fileName, the name of a text file to output the results to; displayFile, whether or not to display the file (default is "FALSE")

**Examples**

```
## Not run:
hasCy <- (if (.Platform$OS.type %in% "windows") { (length(grep("Cytoscape", system("tasklist", intern=TRUE)))

if hasCy {
data(ccData)
ccResults$BP <- breakEdges(ccResults$BP, 0.8)
cwObj <- ccOutCyt(ccResults$BP, ccOpts)
# user selects some nodes in Cytoscape
RCytoscape::selectNodes(cwObj, c("GO:0007017", "GO:0000226", "GO:0007051", "GO:0007052"))
savedNodes <- cytOutNodes("random1", cwObj) # save them
# and selects some other nodes
RCytoscape::selectNodes(cwObj, c("GO:0071103", "GO:0034728", "GO:0006323", "GO:0030261", "GO:0006334"), pr
savedNodes <- cytOutNodes("random2", cwObj, savedNodes)

# now spit results out to a file
cytOutData(savedNodes, ccResults$BP)
}
## End(Not run)
```

---

cytOutNodes-methods      *Methods for Function cytOutNodes*

---

**Description**

Allows export of currently selected nodes in the Cytoscape window for data export

**Methods**

signature(descStr = "character", cwObj = "CytoscapeWindowClass", saveObj = "list")  
descStr is a string describing the nodes that are currently selected, cwObj is the CytoscapeWindow that the nodes are in, and then saveObj is a previously generated cytOutNodes list, and is optional.

**Examples**

```
## Not run:
hasCy <- (if (.Platform$OS.type %in% "windows") { (length(grep("Cytoscape", system("tasklist", intern=TRUE)))

if hasCy {
ccResults$BP <- breakEdges(ccResults$BP, 0.8)
cwObj <- ccOutCyt(ccResults$BP, ccOpts)
# user selects some nodes in Cytoscape
RCytoscape::selectNodes(cwObj, c("GO:0007017", "GO:0000226", "GO:0007051", "GO:0007052"))
savedNodes <- cytOutNodes("random1", cwObj) # save them
# and selects some other nodes
RCytoscape::selectNodes(cwObj, c("GO:0071103", "GO:0034728", "GO:0006323", "GO:0030261", "GO:0006334"), pr
savedNodes <- cytOutNodes("random2", cwObj, savedNodes)

}

## End(Not run)
```

---

fdr	<i>Number of FDR runs to perform</i>
-----	--------------------------------------

---

**Description**

Queries or sets the number of random runs to perform to generate an estimate of the false discovery rate. Defaults to 50

**Usage**

```
fdr(object)
```

**Arguments**

object	Can be <code>ccGeneList</code> , <code>HyperGParamsCC</code> , <code>HyperGResultCC</code> , <code>ccEnrichResult</code> See Details for more information.
--------	--

**Details**

`fdr(object)` gets the number of fdr runs for `ccGeneList`, `HyperGParamsCC`, `HyperGResultCC`, `ccEnrichResult`

`fdr(object)<-` will set the number of fdr runs to be used by `ccEnrich` and `HyperGTestCC` when performing calculations on either a `ccGeneList` or `HyperGParamsCC`, respectively

**Author(s)**

Robert M Flight

**See Also**

[HyperGResultCC](#) [ccEnrichResult](#) [ccGeneList](#) [HyperGParamsCC](#)

---

GENccEnrichResult-class

*Class "GENccEnrichResult"*

---

**Description**

Holds generic `ccEnrich` type results

**Objects from the Class**

Objects can be created by calls of the form `new("GENccEnrichResult", ...)`.

**Slots**

.Data: Object of class "list" The actual list containing the ccEnrichResults  
 categoryName: Object of class "character"  
 ontology: Object of class "character"  
 geneAnnMapping: Object of class "namedList"  
 graphType: Object of class "character"  
 names: Object of class "character"

**Extends**

Class "namedList", directly. Class "list", by class "namedList", distance 2. Class "vector", by class "namedList", distance 3. Class "AssayData", by class "namedList", distance 3.

**Methods**

[ signature(x = "GENccEnrichResult", i = "ANY", j = "ANY"): Subsets the object to just those lists that are desired  
**categoryName** signature(object = "GENccEnrichResult"):  
**ccCompareGeneric** signature(gccResult = "GENccEnrichResult", ccOptions = "ccOptions"):  
 ...  
**geneAnnMapping** signature(object = "GENccEnrichResult"): ...  
**graphType** signature(object = "GENccEnrichResult"): ...  
**graphType<-** signature(object = "GENccEnrichResult"): ...  
**ontology** signature(object = "GENccEnrichResult"): ...

**Author(s)**

Robert M Flight

**See Also**

[ccCompareGeneric](#) [ccSigList](#)

**Examples**

```
data(ccData)
locA <- grep("A", gseaRes$Tissues)
locL <- grep("L", gseaRes$Tissues)
locM <- grep("M", gseaRes$Tissues)

A <- new("ccSigList", sigID=gseaRes$KEGGID[locA], categoryName="KEGG", annotation="org.Mm.eg")
L <- new("ccSigList", sigID=gseaRes$KEGGID[locL], categoryName="KEGG", annotation="org.Mm.eg")
M <- new("ccSigList", sigID=gseaRes$KEGGID[locM], categoryName="KEGG", annotation="org.Mm.eg")
ccEnrichCol <- list(A=A, L=L, M=M)
ccEnrichCol <- new("GENccEnrichResult", ccEnrichCol, categoryName="KEGG")
```

---

getGeneSymbol	<i>Entrez to name, symbol, GO and path conversion, as well as general ID to ID conversion.</i>
---------------	--

---

**Description**

Get different attributes for the Entrez gene Ids

**Usage**

```
getGeneSymbol(id, annPackage)
getGeneName(id, annPackage)
getG02ALLEGS(id, annPackage)
getPath2EG(id, annPackage)
getAnnotation(id, annPackage, mapID, doUnlist=TRUE)
```

**Arguments**

id	The IDs one wants to get information for.
annPackage	Which annotation package to use.
mapID	Which mapping to use
doUnlist	should the results be unlisted or not?

**Details**

The type of ID will change depending on the function. For `getGene...` the ID should be Entrez IDs. For `getG02ALLEGS` Gene Ontology IDs should be used, and for `getPath2EG` KEGG pathways IDs should be used. For `getAnnotation`, any ID can be used.

**Value**

Returns the requested information.

**Note**

These functions are generally called internally for mapping between genes and various objects.

**Author(s)**

Robert M Flight

---

graphType-methods      *graphType*

---

### Description

Gets and sets the graphType for a couple of different ccEnrichResults objects

### Methods

signature(object = "ccEnrichResult")  
signature(object = "GENccEnrichResult")

### See Also

[ccEnrichResult](#) [GENccEnrichResult](#)

---

HyperGParamsCC-class      *Class "HyperGParamsCC"*

---

### Description

This class extends the HyperGParams class in Category by providing options for multiple testing and the storing of extra data in addition to the gene list of interest (not currently used, but might be in the future).

### Objects from the Class

Objects can be created by calls of the form `new("HyperGParamsCC", ...)`. In general the user will not create these directly, but they are created and used by to carry out the enrichment calculations.

### Slots

**fdr:** Object of class "numeric" The number of FDR runs to perform  
**data:** Object of class "data.frame" Extra data stored in the object  
**geneIds:** Object of class "ANY" The genes of interest  
**universeGeneIds:** Object of class "ANY" The gene universe or background used (all the genes on the chip)  
**annotation:** Object of class "character" The annotation package used to get information about the geneIds  
**datPkg:** Object of class "DatPkg" Generated automatically from the annotation slot  
**categorySubsetIds:** Object of class "ANY" A specific set of category IDs that one wants to restrict the testing to  
**categoryName:** Object of class "character" What type of category to use, currently either "GO" or "KEGG"  
**pvalueCutoff:** Object of class "numeric" What should be the p-value to decide significance  
**testDirection:** Object of class "character" "over" or "under" represented annotation terms



**Extends**

Class "[GOHyperGParams](#)", directly.

**Methods**

No methods defined with class "HyperGParamsCC" in the signature.

**Author(s)**

Robert M Flight

**See Also**

[HyperGResultCC ccEnrich](#) Category-package

**Examples**

```
showClass("HyperGParamsCC")
```

---

HyperGResultCC-class    *Class "HyperGResultCC"*

---

**Description**

Contains the results of performing a hypergeometric test on a [HyperGParams](#) object.

**Objects from the Class**

Objects can be created by calls of the form `new("HyperGResultCC", ...)`.

**Slots**

**fdr:** Object of class "numeric" The number of FDR runs performed  
**fdrvalues:** Object of class "numeric" The FDR values generated  
**pvalueType:** Object of class "character" Whether to use p-values or FDR values in determining the significant terms returned  
**data:** Object of class "data.frame" Extra data  
**pvalues:** Object of class "numeric" P-values calculated for each term  
**oddsRatios:** Object of class "numeric"  
**expectedCounts:** Object of class "numeric"  
**catToGeneId:** Object of class "list"  
**organism:** Object of class "character"  
**annotation:** Object of class "character"  
**geneIds:** Object of class "ANY"  
**testName:** Object of class "character"  
**pvalueCutoff:** Object of class "numeric"  
**testDirection:** Object of class "character"

**Extends**

Class "[HyperGResult](#)", directly. Class "[HyperGResultBase](#)", by class "[HyperGResult](#)", distance 2.

**Methods**

**fdR** signature(object = "HyperGResultCC"): ...  
**fdRvalues** signature(object = "HyperGResultCC"): ...  
**pCC** signature(object = "HyperGResultCC"): ...  
**pvalueCutoff<-** signature(r = "HyperGResultCC"): ...  
**pvalueType** signature(object = "HyperGResultCC"): ...  
**pvalueType<-** signature(object = "HyperGResultCC"): ...  
**minCount** signature(object = "HyperGResultCC"): ...  
**minCount<-** signature(object = "HyperGResultCC"): ...

**Author(s)**

Robert M Flight

**See Also**

[hyperGTestCC](#)

**Examples**

```
showClass("HyperGResultCC")
```

---

hyperGTestCC

*Hypergeometric testing with false discovery rate*

---

**Description**

Performs the hypergeometric testing for [HyperGParamsCC](#) objects.

**Usage**

```
hyperGTestCC(p)
```

**Arguments**

p                    A [HyperGParamsCC](#) object

**Details**

This is the heart of `categoryCompare`, the function that calculates the HyperGeometric statistics for the given categories of annotation for each gene list.

**Value**

Returns a [HyperGResultCC](#) object

**Author(s)**

Robert M Flight

**See Also**[HyperGParamsCC](#) [HyperGResultCC](#) [GOHyperGParamsCC](#) [KEGGHyperGParamsCC](#) [GOHyperGResultCC](#) [KEGGHyperGResultCC](#)**Examples**

```
require(GO.db)
require(org.Hs.eg.db)
data(ccData)
g10 <- unique(table10$Entrez)
testGO <- new("GOHyperGParamsCC", geneIds=g10, universeGeneIds=gUniverse,
annotation="org.Hs.eg.db", ontology="CC", conditional=FALSE,
testDirection="over",fdr=0, pvalueCutoff = 0.01)
# ccHypRes <- hyperGTestCC(testGO)
# summary(ccHypRes)
```

---

listNames
-----------

<i>listNames</i>
------------------

---

**Description**

Extracts the listNames from [ccGeneList](#) or [ccOptions](#) objects.

**Usage**

```
listNames(object)
```

**Arguments**

object	This will be either a <a href="#">ccGeneList</a> or <a href="#">ccOptions</a> object
--------	--

**Author(s)**

Robert M Flight

**See Also**[ccGeneList](#) [ccOptions](#)

---

mergedData-class      *Class "mergedData"*

---

### Description

Stores merged data tables from the "data" entry in a ccGeneList. This is useful for output later.

### Objects from the Class

Objects can be created by calls of the form `new("mergedData", ...)`.

### Slots

.Data: Object of class "list"  
useIDName: Object of class "character"  
names: Object of class "character"  
row.names: Object of class "data.frameRowLabels"  
.S3Class: Object of class "character"

### Extends

Class "[data.frame](#)", directly. Class "[list](#)", by class "data.frame", distance 2. Class "[oldClass](#)", by class "data.frame", distance 2. Class "[data.frameOrNULL](#)", by class "data.frame", distance 2. Class "[vector](#)", by class "data.frame", distance 3.

### Methods

`signature(saveObj = "list", compareResult = "ccCompareResult", mergedData = "mergedData")`

### Author(s)

Robert M. Flight

### See Also

[mergeLists](#) [cytOutData](#)

### Examples

```
showClass("mergedData")

data(ccData)
mergeDat <- mergeLists(geneLists, ccOpts)
```

---

mergeLists-methods      *Function mergeLists in Package categoryCompare*

---

## Description

Merges the gene lists or the data tables from a ccGeneList object, providing a single table with all the input data, that can then be queried later, using cytTableOut

## Usage

```
mergeLists(ccGeneList, ccOptions, isGene=TRUE)
```

## Arguments

ccGeneList	a ccGeneList object
ccOptions	a ccOptions object
isGene	are the identifiers genes, or something else (metabolites, etc)

## Value

A mergedData object which is really just a glorified data frame. If the ccGeneList input had a data list, then these are all merged into a single table. Otherwise, it contains just the gene names and which list they were present in.

## Methods

```
signature(ccGeneList = "ccGeneList", ccOptions = "ccOptions")
```

## See Also

[ccGeneList ccOptions mergedData](#)

## Examples

```
data(ccData)
g10 <- (unique(table10$Entrez[1:100]))
g48 <- (unique(table48$Entrez[1:100]))

list10 <- list(genes=g10, universe=gUniverse, annotation="org.Hs.eg.db", data=table10[1:100,])
list48 <- list(genes=g48, universe=gUniverse, annotation="org.Hs.eg.db", data=table48[1:100,])

geneLists <- list(T10=list10, T48=list48)
geneLists <- new("ccGeneList", geneLists, ccType=c("BP", "KEGG"))
ccOpts <- new("ccOptions", listNames = names(geneLists))
mergedDat <- mergeLists(geneLists, ccOpts)

list10 <- list(genes=g10, universe=gUniverse, annotation="org.Hs.eg.db")
list48 <- list(genes=g48, universe=gUniverse, annotation="org.Hs.eg.db")
geneLists <- list(T10=list10, T48=list48)
geneLists <- new("ccGeneList", geneLists, ccType=c("BP", "KEGG"))
ccOpts <- new("ccOptions", listNames = names(geneLists))
mergedDat <- mergeLists(geneLists, ccOpts)
```

---

 minCount

*minCount*


---

**Description**

Extracts and sets the minimum number of genes that an annotation must have to be considered in subsequent steps.

**Usage**

```
minCount(object)
```

**Arguments**

object            This will be either a `HyperGResultCC`, `ccEnrichResult`, or `ccEnrichCollection` object. See Details for more information.

**Details**

`minCount(object)` fetches the set `minCount` for `HyperGResultCC` and `ccEnrichResult` objects  
`minCount(object)<-` will set the `minCount` for `HyperGResultCC` objects, and when applied to `ccEnrichResult` and `ccEnrichCollection` sets the `minCount` for all of the contained objects, so be careful if you want to use different `minCounts` for different results

**Author(s)**

Robert M Flight

**See Also**

[HyperGResultCC](#) [ccEnrichResult](#) [ccEnrichCollection](#)

**Examples**

```
data(ccData)
enrichLists
minCount(enrichLists) <- 5
enrichLists
```

---

 minNodes

*Delete nodes with less than a certain number of genes annotated*


---

**Description**

Deletes from the graph those annotations with less than a certain number of genes

**Usage**

```
minNodes(cwObj, cutoff)
```

**Arguments**

cwObj            a CytoscapeWindowClass object returned from ccOutCyt  
 cutoff            the minimum number of genes that an annotation must have

**Author(s)**

Robert M Flight

**See Also**

CytoscapeWindowClass [ccOutCyt](#)

**Examples**

```

## Not run:
hasCy <- (if (.Platform$OS.type %in% "windows") { (length(grep("Cytoscape", system("tasklist", intern=TRUE))) > 0)}

if hasCy {
  data(ccData)
  ccResults$BP <- breakEdges(ccResults$BP, 0.8)
  cwObj <- ccOutCyt(ccResults$BP, ccOpts)

  minNodes(cwObj, 5)

}
## End(Not run)

```

---

pvalueType

*Type of p-values to return from object*

---

**Description**

Queries or sets the type of p-values to return from objects, either base calculated (pvals) or from fdr calculations (fdr)

**Usage**

```
pvalueType(object)
```

**Arguments**

object            Can be HyperGResultCC, ccEnrichResult, ccEnrichCollection. See Details for more information

**Details**

pvalueType(object) gets the type of p-values to be returned from HyperGResultCC and ccEnrichResult objects

pvalueType(object)<- will set the type of p-values to be returned from HyperGResultCC, ccEnrichResult, ccEnrichCollection. Note that for a ccEnrichCollection, the type is changed for all contained ccEnrichResults

**Author(s)**

Robert M Flight

**See Also**[HyperGResultCC ccEnrichResult ccEnrichCollection](#)**Examples**

```
# pvalueType-Methods
data(ccData)

## Not run: pvalueType(enrichLists) # this returns an error
pvalueType(enrichLists[[1]])
pvalueType(enrichLists[[1]][[1]])

# change the type for one of the results
pvalueType(enrichLists[[1]]) <- 'pval' # note, I do not recommend changing it for a single result in a call
enrichLists

# change for all of the results
pvalueType(enrichLists) <- 'pval'
enrichLists
```

---

resetColors-methods     *resetColors*

---

**Description**

If the color of particular nodes have been modified from the original color scheme in ccOptions, this will reset them

**Methods**

signature(cwObj = "CytoscapeWindowClass", ccOpts = "ccOptions") What CytoscapeWindow to apply this to, and what ccOptions to use for the color scheme.

**Optional Arguments:** Note that optional arguments include node.attribute.name (default is 'fillcolor') and mode (default is 'lookup')

**Note**

This is most commonly used with the cwReload function, as the color scheme of the network does not get saved in the CYS file.

**Author(s)**

Robert M Flight

**See Also**[CytoscapeWindowClass ccOptions setNodeColorRule cwReload](#)



---

`show-methods`*Methods for Function show in Package 'categoryCompare'*

---

**Description**

The show and summary methods for [HyperGResultCC](#) objects generated using [hyperGTestCC](#)

**Methods**

```
show, signature(object = "HyperGResultCC")
summary, signature(object = "HyperGResultCC")
```

**Author(s)**

Robert M Flight

**Examples**

```
## Not run:
data(ccData)
show(enrichLists)
summary(enrichLists[[1]][[1]])

## End(Not run)
```

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