

# Package 'RDAVIDWebService'

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**Type** Package

**Title** An R Package for retrieving data from DAVID into R objects using Web Services API.

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**Description** Tools for retrieving data from the Database for Annotation, Visualization and Integrated Discovery (DAVID) using Web Services into R objects. This package offers the main functionalities of DAVID website including: i) user friendly connectivity to upload gene/background list/s, change gene/background position, select current specie/s, select annotations, etc. ii) Reports of the submitted Gene List, Annotation Category Summary, Gene/Term Clusters, Functional Annotation Chart, Functional Annotation Table

**License** GPL (>=2)

**URL** <http://www.bdmg.com.ar>, <http://david.abcc.ncifcrf.gov/>

**Imports** Category, GO.db, RBGL, rJava

**Depends** R (>= 2.14.1), methods, graph, GStats, ggplot2

**Collate** 'DAVIDdemo-ids.R' 'DAVIDdemo-geneList.R'  
'DAVIDdemo-annotationSummary.R'  
'DAVIDdemo-functionalAnnotationChart.R'  
'DAVIDdemo-annotationTable.R' 'DAVIDdemo-clusterReport.R'  
'DAVIDResult-class.R' 'DAVIDGenes-class.R'  
'DAVIDFunctionalAnnotationChart-class.R' 'DAVIDCluster-class.R'  
'DAVIDGeneCluster-class.R' 'DAVIDTermCluster-class.R'  
'DAVIDFunctionalAnnotationTable-class.R' 'DAVIDGODag-class.R'  
'DAVIDWebService-class.R' 'DAVIDClasses-show.R'  
'DAVIDClasses-ids.R' 'DAVIDClasses-plot2D.R'  
'DAVIDClasses-summary.R' 'DAVIDClasses-genes.R'  
'DAVIDClasses-categories.R' 'DAVIDResult-getters.R'  
'DAVIDGenes-methods.R' 'DAVIDCluster-methods.R'  
'DAVIDFunctionalAnnotationTable-methods.R'  
'DAVIDGODag-methods.R' 'DAVIDWebService-accessors.R'  
'DAVIDWebService-methods.R' 'DAVIDWebService-reports.R'  
'DAVIDClasses-constructor.R' 'RDAVIDWebService-package.R'

**Suggests** Rgraphviz

**InstallableEverywhere** yes

**biocViews** Visualization, DifferentialExpression, GraphAndNetwork

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DAVIDWebService-package

*An R Package for retrieving data from DAVID into R objects using Web Services*

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## Description

Tools for retrieving data from the Database for Annotation, Visualization and Integrated Discovery (DAVID) using Web Services into R objects. This package offers the main functionalities of DAVID website including:

**Connectivity:** upload gene/background list/s, change gene/background position, select current specie/s, select annotations, etc. from R.

**Exploration:** native R objects of submitted Gene List, Annotation Category Summary, Gene/Term Clusters, Functional Annotation Chart and Functional Annotation Tables. In addition it offers the usual many-genes-to-many-terms visualization and induced Gene Ontology direct acyclic graph GOstats-based conversion method, in order to visualize GO structure.

## Author(s)

Cristobal Fresno <crisobalfresno@gmail.com> and Elmer A. Fernandez <elmerfer@gmail.com>

## References

1. The Database for Annotation, Visualization and Integrated Discovery (david.abcc.ncifcrf.gov)
2. Xiaoli Jiao, Brad T. Sherman, Da Wei Huang, Robert Stephens, Michael W. Baseler, H. Clifford Lane, Richard A. Lempicki, DAVID-WS: A Stateful Web Service to Facilitate Gene/Protein List Analysis *Bioinformatics* 2012 doi:10.1093/bioinformatics/bts251
3. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C. & Lempicki, R. A DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. *Nucleic Acids Res*, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175
4. Cristobal Fresno, Elmer A. Fernandez (2013) RDAVIDWebService: a versatile R interface to DAVID, *Bioinformatics*, 29(21), 2810-2811., <http://bioinformatics.oxfordjournals.org/content/29/21/2810>.

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annotationSummary1

*DAVID's website annotation summary example files*

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## Description

These datasets correspond to the unfolded main summary categories data obtained in the Annotation Summary Results page in the Database for Annotation, Visualization and Integrated Discovery (DAVID) website, using as input file, the ones provided for demo purposes (demoList1 or demoList2) with default options. No statistical analysis is performed on these results.

**Usage**

```
data(annotationSummary1)
```

```
data(annotationSummary2)
```

**Format**

annotationSummary1/2 are data.frame for demoList1/2 input ids, respectively, with the following columns.

**Main.Category** factor with the main categories used in the present analysis.

**ID** integer to identify the annotation category.

**Name** character with the name of category (the ones available in getAllAnnotationCategoryNames function).

**X.** numeric with the percentage of the gene list ids present in the term.

**Count** integer with the number of ids of the gene list that belong to this term.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

**References**

1. The Database for Annotation, Visualization and Integrated Discovery ([davidgenelist.abcc.ncifcrf.gov](http://davidgenelist.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C.; Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. *Nucleic Acids Res*, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175
3. DAVID Help page [http://david.abcc.ncifcrf.gov/helps/functional\\_annotation.html#summary](http://david.abcc.ncifcrf.gov/helps/functional_annotation.html#summary)

**See Also**

Other DataExamples: [demoList1](#), [demoList2](#), [geneList1](#), [geneList2](#)

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annotationTable1

*DAVID's website functional annotation table example files*

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**Description**

These datasets correspond to the Functional Annotation Table report obtained in the Database for Annotation, Visualization and Integrated Discovery (DAVID) website, using as input file, the ones provided for demo purposes (demoList1 or demoList2) for GOTERM\_BP\_ALL, GOTERM\_MF\_ALL and GOTERM\_CC\_ALL categories. No statistical analysis is performed on these results.

**Usage**

```
data(annotationTable1)
```

```
data(annotationTable2)
```

**Format**

annotationTable1/2 are data.frame for demoList1/2 input ids, respectively, with the following columns.

**Gene** Three Columns with the same data included in Gene List Report (ID, Gene.Name and Species) but coding for DAVID ID, i. e., comma separated character with input ids if, two or more stands for the same gene.

**Annotation** As many columns as Annotation Categories were used. In each column, a comma separated style is use to delimitate the different terms where is evidence reported for DAVID ID record.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

**References**

1. The Database for Annotation, Visualization and Integrated Discovery ([davidgenelist.abcc.ncifcrf.gov](http://davidgenelist.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C.; Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. Nucleic Acids Res, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175
3. DAVID Help page [http://david.abcc.ncifcrf.gov/helps/functional\\_annotation.html#EXP2](http://david.abcc.ncifcrf.gov/helps/functional_annotation.html#EXP2)

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categories

*categories for the different DAVIDWebService package class objects*

---

**Description**

Obtain ids related information, according to the given function call (see Values).

**Usage**

```
categories(object)
```

```
## S4 method for signature 'DAVIDFunctionalAnnotationChart'
categories(object)
```

```
## S4 method for signature 'DAVIDFunctionalAnnotationTable'
categories(object)
```

**Arguments**

object DAVIDWebService class object. Possible values are: DAVIDFunctionalAnnotationChart or DAVIDFunctionalAnnotationTable.

**Value**

according to the call, one of the following objects can be returned:

DAVIDFunctionalAnnotationChart

factor vector of the "Category" column.

DAVIDFunctionalAnnotationTable

character vector with the name of available main categories in the dictionary/membership.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

**See Also**

Other DAVIDFunctionalAnnotationChart: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart-class](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [ids](#), [ids](#), [ids](#), [ids](#), [ids](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#)

Other DAVIDFunctionalAnnotationTable: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable-class](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [dictionary](#), [dictionary](#), [genes](#), [genes](#), [genes](#), [genes](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [membership](#), [membership](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [subset](#), [subset](#)

**Examples**

```
{
##DAVIDFunctionalAnnotationChart example
##Load the Functional Annotation Chart file report for the input demo
##file 2, using data function. Then, create a DAVIDFunctionalAnnotationChart
##object using the loaded data.frame funChart2.
data(funChart2)
davidFunChart2<-DAVIDFunctionalAnnotationChart(funChart2)

##In Addition to the usual data.frame accessors, the user can inspect the
##main categories used in the analysis.
categories(davidFunChart2)

##DAVIDFunctionalAnnotationTable example
##Load the Functional Annotation Table file report for the input demo
##file 1, using data function. Then, create a DAVIDFunctionalAnnotationTable
##object using the loaded data.frame annotationTable1.
data(annotationTable1)
```

```
davidFunTable1<-DAVIDFunctionalAnnotationTable(annotationTable1)

##Now, the user can inspect the main categories used in the analysis.
categories(davidFunTable1)
}
```

---

cluster

*Methods for DAVIDCluster class object*

---

## Description

Obtain DAVIDCluster related information, according to the given function call (see Values).

## Usage

```
cluster(object)

## S4 method for signature 'DAVIDCluster'
cluster(object)

enrichment(object)

## S4 method for signature 'DAVIDCluster'
enrichment(object)

members(object)

## S4 method for signature 'DAVIDCluster'
members(object)
```

## Arguments

object            DAVIDCluster class object.

## Value

according to the call, one of the following objects can be returned:

cluster	list with DAVIDCluster object slot.
enrichment	numeric vector with DAVID cluster's enrichment score.
members	list with DAVID Cluster's members.

## Author(s)

Cristobal Fresno and Elmer A Fernandez

**See Also**

Other DAVIDCluster: [DAVIDCluster-class](#), [dictionary](#), [dictionary](#), [membership](#), [membership](#), [subset](#), [subset](#), [summary](#), [summary](#), [summary](#), [summary](#)

Other DAVIDCluster: [DAVIDCluster-class](#), [dictionary](#), [dictionary](#), [membership](#), [membership](#), [subset](#), [subset](#), [summary](#), [summary](#), [summary](#), [summary](#)

Other DAVIDCluster: [DAVIDCluster-class](#), [dictionary](#), [dictionary](#), [membership](#), [membership](#), [subset](#), [subset](#), [summary](#), [summary](#), [summary](#), [summary](#)

**Examples**

```
{
##DAVIDGeneCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo list 1 file to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/geneClusterReport1.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidGeneCluster1<-DAVIDGeneCluster(untar(fileName, list=TRUE))
davidGeneCluster1

##Now we can invoke DAVIDCluster ancestor functions to inspect the report
##data of each cluster. For example, we can call summary to get a general
##idea, and then inspect the cluster with the higher Enrichment Score, to see
##which members belong to it, etc. or simply, returning the whole cluster as
##a list with EnrichmentScore and Members.
summary(davidGeneCluster1)
higherEnrichment<-which.max(enrichment(davidGeneCluster1))
clusterGenes<-members(davidGeneCluster1)[[higherEnrichment]]
wholeCluster<-cluster(davidGeneCluster1)[[higherEnrichment]]

##DAVIDTermCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo file 2 to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/termClusterReport2.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidTermCluster2<-DAVIDTermCluster(untar(fileName, list=TRUE))
davidTermCluster2

##Now we can invoke DAVIDCluster ancestor functions to inspect the report
##data of each cluster. For example, we can call summary to get a general
##idea, and then inspect the cluster with the higher Enrichment Score, to see
##which members belong to it, etc. Or simply returning the whole cluster as a
##list with EnrichmentScore and Members.
summary(davidTermCluster2)
higherEnrichment<-which.max(enrichment(davidTermCluster2))
clusterGenes<-members(davidTermCluster2)[[higherEnrichment]]
wholeCluster<-cluster(davidTermCluster2)[[higherEnrichment]]
}
```



---

DAVIDCluster-class     *class "DAVIDCluster"*

---

### Description

This virtual class represents the output of a DAVID "Cluster" report, with "DAVIDTermCluster" and "DAVIDGeneCluster" as possible heirs, according to the report used.

### Type

This class is a "Virtual" one.

### Extends

- *DAVIDResult* in the conceptual way.

### Heirs

- DAVIDTermCluster: DAVID's Functional Annotation Clustering report.
- DAVIDGeneCluster: DAVID's Functional Classification Tool report.

### Slots

cluster    named list with the different clustered terms/genes: Members, represented as DAVID-Genes object; and EnrichmentScore, a numeric with DAVID cluster enrichment score.

### Methods

show    signature(object="DAVIDCluster"): basic console output.

summary    signature(object="DAVIDCluster"): basic summary console output.

initialize    signature(object="DAVIDCluster",    fileName="character"): basic cluster report file parser.

cluster    signature(object="DAVIDCluster"): getter for the corresponding slot.

enrichment    signature(object="DAVIDCluster"): obtain the enrichment score of each cluster.

members    signature(object="DAVIDCluster"): obtain the corresponding cluster members.

### Author(s)

Cristobal Fresno and Elmer A Fernandez

### References

1. The Database for Annotation, Visualization and Integrated Discovery ([david.abcc.ncifcrf.gov](http://david.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C.; Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. *Nucleic Acids Res*, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175

**See Also**

Other DAVIDCluster: [cluster](#), [cluster](#), [dictionary](#), [dictionary](#), [enrichment](#), [enrichment](#), [members](#), [members](#), [membership](#), [membership](#), [subset](#), [subset](#), [summary](#), [summary](#), [summary](#), [summary](#)

---

DAVIDFunctionalAnnotationChart-class

*class "DAVIDFunctionalAnnotationChart"*

---

**Description**

This class represents the output of "Functional Annotation Chart" of DAVID. It is an heir of DAVIDResult in the conceptual way, and also a data.frame with additional features, such as identifying the unique and duplicate ids, searching for genes with a given id, etc.

**Type**

This class is a "Concrete" one.

**Extends**

- *DAVIDResult* in the conceptual way.
- *data.frame* in order to extend the basic features.

**Slots**

no additional to the ones inherited from DAVIDResult and data.frame classes.

**Methods**

`show` signature(object="DAVIDFunctionalAnnotationChart"): returns a basic console output.

`valid` signature(object="DAVIDFunctionalAnnotationChart"): logical which checks DAVID's file output name ("Category", "Term", "Count", etc.) presence.

`DAVIDFunctionalAnnotationChart` signature( object="character"): constructor with the name of the .tab file report to load.

`DAVIDFunctionalAnnotationChart` signature( object="data.frame"): data.frame already loaded to use when constructing the object.

`as` signature(object="DAVIDFunctionalAnnotationChart"): coerce a data.frame into a DAVID-FunctionalAnnotationChart object.

`categories` signature( object="DAVIDFunctionalAnnotationChart"): obtain the factor vector of the "Category" column.

`ids` signature(object="DAVIDFunctionalAnnotationChart"): obtain a list with character/integer vector with the ids of the corresponding term.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

## References

1. The Database for Annotation, Visualization and Integrated Discovery (david.abcc.ncifcrf.gov)
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C. & Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. Nucleic Acids Res, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175

## See Also

Other DAVIDFunctionalAnnotationChart: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [categories](#), [categories](#), [categories](#), [ids](#), [ids](#), [ids](#), [ids](#), [ids](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#)

## Examples

```
{
##Load the Functional Annotation Chart file report for the input demo
##file 2, using data function. Then, create a DAVIDFunctionalAnnotationChart
## object using the loaded data.frame funChart2. In addition, the user can
##use the file name of the downloaded file report.
data(funChart2)
davidFunChart2<-DAVIDFunctionalAnnotationChart(funChart2)

##In Addition to the usual data.frame accessors, the user can inspect the
##main categories used in the analysis.
categories(davidFunChart2)

##Obtain the ids of the genes present in each Term, as a list of character
##vector
ids(davidFunChart2)

##Or plot a 2D tile matrix with the reported evidence (green) or not (black).
##Just to keep it simple, for the first five terms present in funChart2
##object.
plot2D(DAVIDFunctionalAnnotationChart(funChart2[1:5, ]),
color=c("FALSE"="black", "TRUE"="green"))
}
```

---

DAVIDFunctionalAnnotationTable-class

*class "DAVIDFunctionalAnnotationTable"*

---

## Description

This class represents the output of a DAVID Functional Annotation Table report. In this class no statistical analysis is carried out.

**Type**

This class is a "Concrete" one.

**Extends**

- *DAVIDResult* in the conceptual way, and to reuse some functionalities such as plot2D, type and so on.

**Slots**

**Genes** a DAVIDGenes object with the submitted genes.

**Dictionary** a look up list of data.frame of each main annotation category, where the specified IDs and Terms used can be found.

**Membership** list with logical membership matrix, where gene ids are coded by rows and the respective annotation category ids as columns.

**Methods**

`initialize` signature(.Object= "DAVIDFunctionalAnnotationTable", fileName="character"): basic Functional Annotation Table report file parser.

`DAVIDFunctionalAnnotationTable` signature(fileName= "character"): high level Functional Annotation Table report file parser.

`valid` signature(object= "DAVIDFunctionalAnnotationTable"): logical which checks for Membership, Dictionary and Genes cohesion.

`show` signature(object="DAVIDFunctionalAnnotationTable"): returns a basic console output.

`genes` signature(object="DAVIDFunctionalAnnotationTable"): returns a DAVIDGenes object.

`subset` signature(object= "DAVIDFunctionalAnnotationTable", selection=c("Membership", "Dictionary"): returns a subset list using the selection slot, looking up the category parameter if provided. Otherwise, it returns all the available main categories. Drop parameter indicates whether to drop list structure or not, if a list of length==1 is to be returned.

`dictionary` signature(object= "DAVIDFunctionalAnnotationTable", category, drop=TRUE): returns subset using selection="Dictionary" and category and drop parameters.

`membership` signature(object= "DAVIDFunctionalAnnotationTable", category="character", drop=TRUE): returns subset using selection="Membership" and category and drop parameters.

`genes` signature(object= "DAVIDFunctionalAnnotationTable", ...): returns a DAVIDGenes object slot, according to additional ... parameters.

`categories` signature(object= "DAVIDFunctionalAnnotationTable"): returns a character vector with the main annotation categories available..

`plot2D` signature(object="DAVIDFunctionalAnnotationTable", category, id, names.genes=FALSE, names.ggplot2): tile plot of genes id vs functional annotation category membership. If missing, all available data is used. In addition, names.genes and names.category parameters indicate whether to use or not, genes and category names respectively. Default value is FALSE.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

## References

1. The Database for Annotation, Visualization and Integrated Discovery (david.abcc.ncifcrf.gov)
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C. & Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. Nucleic Acids Res, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175

## See Also

Other DAVIDFunctionalAnnotationTable: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [categories](#), [categories](#), [categories](#), [dictionary](#), [dictionary](#), [genes](#), [genes](#), [genes](#), [genes](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [membership](#), [membership](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [subset](#), [subset](#)

## Examples

```
{
##Load the Functional Annotation Table file report for the input demo
##file 1, using data function. Then, create a DAVIDFunctionalAnnotationTable
##object using the loaded data.frame annotationTable1. In addition, the user
##can use the file name of the downloaded file report.
data(annotationTable1)
davidFunTable1<-DAVIDFunctionalAnnotationTable(annotationTable1)

##Now we can obtain the genes for the given ids, or the complete list if the
##parameter is omitted.
genes(davidFunTable1, id=c("37166_at", "41703_r_at"))

##Or the main categories used on the analysis, in order to get the
##dictionary for a specific category (ID and Term fields), for the head of
##the data.frame.
categories(davidFunTable1)
head(dictionary(davidFunTable1, categories(davidFunTable1)[1]))

##And what about the membership of the genes in these terms? Just for the
##first six ids we can use:
head(membership(davidFunTable1, categories(davidFunTable1)[1]))

##Or simply plot the membership of only for the first six terms in this
##category, with only the genes of the first six terms with at least one
##evidence code.
##Category filtering...
categorySelection<-list(head(dictionary(davidFunTable1,
categories(davidFunTable1)[1])$ID))
names(categorySelection)<-categories(davidFunTable1)[1]

##Gene filter...
id<-membership(davidFunTable1, categories(davidFunTable1)[1])[,1:6]
id<-ids(genes(davidFunTable1))[rowSums(id)>0]

##Finally the membership tile plot
```

```
plot2D(davidFunTable1, category=categorySelection, id=id,
names.category=TRUE)
}
```

---

DAVIDGeneCluster-class

*class "DAVIDGeneCluster"*

---

### Description

This class represents the output of a DAVID Gene Functional Classification Tool report.

### Type

This class is a "Concrete" one.

### Extends

- *DAVIDCluster* and uses its constructor to parse the report.

### Slots

the ones inherited from *DAVIDCluster*.

### Methods

`initialize` signature(.Object="DAVIDGeneCluster", fileName="character"): basic cluster report file parser.

`DAVIDGeneCluster` signature(fileName="character"): high level gene cluster report file parser.

`ids` signature(object="DAVIDGeneCluster"): list with the member ids within each cluster.

`genes` signature(object="DAVIDGeneCluster"): list with the DAVIDGenes members within each cluster.

`plot2D` signature(object="DAVIDGeneCluster", color=c("FALSE"="black", "TRUE"="green"), names=FALSE): ggplot2 tile plot with gene membership to each cluster.

### Author(s)

Cristobal Fresno and Elmer A Fernandez

### References

1. The Database for Annotation, Visualization and Integrated Discovery ([david.abcc.ncifcrf.gov](http://david.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C. & Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. *Nucleic Acids Res*, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175

**See Also**

Other DAVIDGeneCluster: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [genes](#), [genes](#), [genes](#), [genes](#), [ids](#), [ids](#), [ids](#), [ids](#), [ids](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#)

**Examples**

```
{
##Load the Gene Functional Classification Tool file report for the
##input demo list 1 file to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/geneClusterReport1.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidGeneCluster1<-DAVIDGeneCluster(untar(fileName, list=TRUE))
davidGeneCluster1

##Now we can invoke DAVIDCluster ancestor functions to inspect the report
##data, of each cluster. For example, we can call summary to get a general
##idea, and the inspect the cluster with higher Enrichment Score, to see
##which members belong to it, etc. Or simply returning the whole cluster as
##a list with EnrichmentScore and Members.
summary(davidGeneCluster1)
higherEnrichment<-which.max(enrichment(davidGeneCluster1))
clusterGenes<-members(davidGeneCluster1)[[higherEnrichment]]
wholeCluster<-cluster(davidGeneCluster1)[[higherEnrichment]]

##Then, we can obtain the ids of the members calling clusterGenes object
##which is a DAVIDGenes class or directly using ids on davidGeneCluster1.
ids(clusterGenes)
ids(davidGeneCluster1)[[higherEnrichment]]

##Obtain the genes of the first cluster using davidGeneCluster1 object.
##Or, using genes on DAVIDGenes class once we get the members of the cluster.
genes(davidGeneCluster1)[[1]]
genes(members(davidGeneCluster1)[[1]])

##Finally, we can inspect a 2D tile membership plot, to visually inspect for
##overlapping of genes across the clusters. Or use a scaled version of gene
##names to see the association of gene cluster, e.g., cluster 3 is related to
##ATP genes.
plot2D(davidGeneCluster1)
plot2D(davidGeneCluster1,names=TRUE)+
theme(axis.text.y=element_text(size=rel(0.9)))
}
```

---

DAVIDGenes

*High level constructors for DAVIDWebService package's classes.*


---

**Description**

Different ways to build the different DAVIDWebService's object according to the signature in use.

**Usage**

```
DAVIDGenes(object)

## S4 method for signature 'character'
DAVIDGenes(object)

## S4 method for signature 'data.frame'
DAVIDGenes(object)

## S4 method for signature 'DAVIDGenes'
initialize(.Object, fileName)

as(object, Class, strict=TRUE,
  ext=possibleExtends(thisClass, Class))

DAVIDFunctionalAnnotationChart(object)

## S4 method for signature 'character'
DAVIDFunctionalAnnotationChart(object)

## S4 method for signature 'data.frame'
DAVIDFunctionalAnnotationChart(object)

## S4 method for signature 'DAVIDFunctionalAnnotationChart'
initialize(.Object,
  fileName)

as(object, Class, strict=TRUE,
  ext=possibleExtends(thisClass, Class))

## S4 method for signature 'DAVIDCluster'
initialize(.Object, fileName)

## S4 method for signature 'DAVIDGeneCluster'
initialize(.Object,
  fileName)

DAVIDGeneCluster(object)

## S4 method for signature 'character'
DAVIDGeneCluster(object)

## S4 method for signature 'DAVIDTermCluster'
initialize(.Object,
  fileName)

DAVIDTermCluster(object)

## S4 method for signature 'character'
DAVIDTermCluster(object)

## S4 method for signature 'DAVIDFunctionalAnnotationTable'
```



```

initialize(.Object,
  fileName)

as(object, Class, strict=TRUE,
  ext=possibleExtends(thisClass, Class))

DAVIDFunctionalAnnotationTable(object)

## S4 method for signature 'character'
DAVIDFunctionalAnnotationTable(object)

## S4 method for signature 'data.frame'
DAVIDFunctionalAnnotationTable(object)

## S4 method for signature 'DAVIDGODag'
initialize(.Object, funChart, type=c("BP", "MF", "CC"), pvalueCutoff=0.1, removeUnattached=FALSE, ...)

DAVIDGODag(funChart, ...)

## S4 method for signature 'DAVIDFunctionalAnnotationChart'
DAVIDGODag(funChart,
  ...)

```

### Arguments

object	could be a character with the file name of the .tab report or data.frame already loaded.
fileName	character with the file name of the .tab report to load.
.Object	character to use in new function call. Possible values are: "DAVIDGenes", "DAVIDFunctionalAnnotationChart" or "DAVIDCluster".
Class	character to use in the <code>as</code> function call. Possible values are: "DAVIDGenes" and "DAVIDFunctionalAnnotationChart".
strict, ext	see <code>as</code> function.
funChart	DAVIDFunctionalAnnotationChart object.
type	character to indicate Gene Ontology main category: "BP", "MF" or "CC".
pvalueCutoff	numeric >0 <=1 to indicate the p-value to use as the threshold for enrichment. Default value is 0.1
removeUnattached	Should unattached nodes be removed from GO DAG? Default value is FALSE.
...	Additional parameters for lower level constructors (initialize).

### Value

a DAVIDWebService object according to function call:

DAVIDGenes	object with genes description related data.
DAVIDFunctionalAnnotationChart	object with the respective report.
DAVIDFunctionalAnnotationTable	object with the respective report.



```
## object using the loaded data.frame funChart2.
data(funChart2)
davidFunChart2<-DAVIDFunctionalAnnotationChart(funChart2)

##In addition, the user can use the file name of the downloaded file report.
##Here, we need to first uncompressed the report included in the package, in
##order to load it.
setwd(tempdir())
fileName<-system.file("files/functionalAnnotationChartReport2.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidFunChart2<-DAVIDFunctionalAnnotationChart(untar(fileName, list=TRUE))

##DAVIDFunctionalAnnotationTable example
##Load the Functional Annotation Table file report for the input demo
##file 1, using data function. Then, create a DAVIDFunctionalAnnotationTable
##object using the loaded data.frame annotationTable1.
data(annotationTable1)
davidFunTable1<-DAVIDFunctionalAnnotationTable(annotationTable1)

##In addition, the user can use the file name of the downloaded file report.
##Here, we need to first uncompressed the report included in the package, in
##order to load it.
setwd(tempdir())
fileName<-system.file("files/annotationTableReport1.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidFunTable1<-DAVIDFunctionalAnnotationTable(untar(fileName, list=TRUE))

##Example DAVIDGODag
##Load the Functional Annotation Chart file report for the input demo
##file 2, using data function. Then, create a DAVIDGODag object using
##Molecular Function main category of DAVIDFunctionalAnnotationChart object,
##obtained from the loaded data.frame funChart2. In addition, we have
##selected a threshold pvalue of 0.001 and removed unattached nodes, in case
##DAVID/GO.db database are not using the same version.
data(funChart2)
davidGODag<-DAVIDGODag(DAVIDFunctionalAnnotationChart(funChart2), type="MF",
pvalueCutoff=0.001, removeUnattached=TRUE)

##DAVIDGeneCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo list 1 file to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/geneClusterReport1.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidGeneCluster1<-DAVIDGeneCluster(untar(fileName, list=TRUE))

##DAVIDTermCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo file 2 to create a DAVIDGeneCluster object.
setwd(tempdir())
```

```

fileName<-system.file("files/termClusterReport2.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidTermCluster2<-DAVIDTermCluster(untar(fileName, list=TRUE))
}

```

---

DAVIDGenes-class      *class "DAVIDGenes"*

---

### Description

This class represents the output of "Show Genes Result" of DAVID. It is an heir of DAVIDResult in the conceptual way, and also a data.frame with additional features, such as identifying the unique and duplicate ids, searching for genes with a given id, etc.

### Type

This class is a "Concrete" one.

### Extends

- DAVIDResult in the conceptual way.
- data.frame in order to extend the basic features.

### Slots

none additional to the ones inherited from DAVIDResult and data.frame classes.

### Methods

valid signature(object="DAVIDGenes"): logical which checks for data.frame name (ID, Name) presence.

DAVIDGenes signature(object="character"): constructor with the name of the .tab file report to load.

DAVIDGenes signature(object="data.frame"): data.frame already loaded to use when constructing the object.

ids signature(object="DAVIDGenes"): character vector with gene submitted ids.

### Author(s)

Cristobal Fresno and Elmer A Fernandez

### References

1. The Database for Annotation, Visualization and Integrated Discovery ([david.abcc.ncifcrf.gov](http://david.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C. & Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. Nucleic Acids Res, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175

**See Also**

Other DAVIDGenes: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [genes](#), [genes](#), [genes](#), [ids](#), [ids](#), [ids](#), [ids](#), [ids](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#)

**Examples**

```
{
##Load Show Gene List file report for the input demo file 1, using data
##function. Then, create a DAVIDGenes object using the loaded data.frame
##geneList1. In addition, the user can use the file name of the downloaded
##file report.
data(geneList1)
davidGenes1<-DAVIDGenes(geneList1)

##Now we can inspect davidGenes1 as it was an common data.frame
head(davidGenes1)

##Additional getters for this object are also available, to obtain the
##different columns: ids, genes and species.
ids(davidGenes1)
genes(davidGenes1)
species(davidGenes1)

##Or even look up for a particular gene id, which will return only the
##matched ones.
genes(davidGenes1, ids=c("38926_at", "35367_at", "no match"))

##Obtain the genes with duplicate manufacturer ids or just the genes that
##do not have duplicate ids (uniqueIds).
duplicateIds(davidGenes1)
uniqueIds(davidGenes1)
}
```

---

DAVIDGODag-class	<i>class "DAVIDGODag"</i>
------------------	---------------------------

---

**Description**

This concrete class represents an induced GO DAG generated by the DAVID Functional Annotation Chart report a.k.a a `DAVIDFunctionalAnnotationChart` object.

**Type**

This class is a "Concrete" one.

**Extends**

- *GOHyperGResult* directly, in order to reuse *GOSTats* functionalities.

**Slots**

the ones inherited from GOHyperGResult

**Methods**

show signature(object="DAVIDGODag"): basic console output.

summary signature(object="DAVIDGODag", ...): basic summary console output.

initialize signature(object="DAVIDGODag", fileName="character"): basic cluster report file parser.

DAVIDGODag signature(object="DAVIDGODag", fileName="character"): high level constructor to parse the file report.

universeMappedCount, universeCounts, counts signature( object="DAVIDGODag"): modifications to the corresponding GOstats/Category library functions, to keep the same behavior, for DAVIDGODag object.

fdrs, benjaminis, bonferronis signature( object="DAVIDGODag"): Adjusted method specific p-values for the corresponding nodes/terms.

terms signature(object="DAVIDGODag"): character vector with GO node names.

popTotals, popHits, listTotals signature( object="DAVIDGODag"): integer vector with the number of ids, to use in the EASE score calculations, when building the 2x2 contingency table.

percentages signature(object="DAVIDGODag"): numeric vector with the percentage of the gene list ids present in the term.

foldEnrichments signature(object="DAVIDGODag"):numeric vector with the ratio of the two proportions for each node/term. For example, if 40/400 (i.e. 10%) of your input genes involved in "kinase activity" and the background information is 300/30000 genes (i.e. 1%) associating with "kinase activity", roughly  $10\%/1\%=10$  fold enrichment.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

**References**

1. The Database for Annotation, Visualization and Integrated Discovery (david.abcc.ncifcrf.gov)
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C. & Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. Nucleic Acids Res, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175
3. Falcon, S; Gentleman, R.; Using GOstats to test gene lists for GO term association, Bioinformatics 23 (2007) 257-258.

**See Also**

Other DAVIDGODag: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [benjaminis](#), [benjaminis](#), [bonferronis](#), [bonferronis](#), [counts](#), [counts](#), [fdrs](#), [fdrs](#), [foldEnrichments](#), [foldEnrichments](#),

`initialize`, `initialize`, `initialize`, `initialize`, `initialize`, `initialize`, `initialize`,  
`listTotals`, `listTotals`, `percentages`, `percentages`, `popHits`, `popHits`, `popTotals`, `popTotals`,  
`summary`, `summary`, `summary`, `summary`, `terms`, `terms`, `universeCounts`, `universeMappedCount`,  
`upsideDown`, `upsideDown`

## Examples

```
{
##Load the Functional Annotation Chart file report for the input demo
##file 2, using data function. Then, create a DAVIDGODag object using
##Molecular Function main category of DAVIDFunctionalAnnotationChart object,
##obtained from the loaded data.frame funChart2. In addition, we have
##selected a threshold pvalue of 0.001 and removed unattached nodes, in case
##DAVID/GO.db database are not using the same version.
data(funChart2)
davidGODag<-DAVIDGODag(DAVIDFunctionalAnnotationChart(funChart2), type="MF",
pvalueCutoff=0.001, removeUnattached=TRUE)

##Now, we can inspect the enrichment GO DAG using G0stats functionalities:
##counts, pvalues, sigCategories, universeCounts, geneMappedCount, etc.
##However, oddsRatios, expectedCounts and universeMappedCount are not
##available because these results are not available on DAVID's Functional
##Annotation Chart report. In addition geneIdUniverse are not the ones of
##the universe but the ids on the category (geneIdsByCategory).
davidGODag
counts(davidGODag)
pvalues(davidGODag)
sigCategories(davidGODag, p=0.0001)
universeCounts(davidGODag)
geneMappedCount(davidGODag)
geneIdsByCategory(davidGODag)
summary(davidGODag)

##In addition, the new nodeData attributes (term, listTotal, popHit,
##popTotal, foldEnrichment, bonferroni, benjamini, fdr) can be retrieved.
terms(davidGODag)
listTotals(davidGODag)
popHits(davidGODag)
popTotals(davidGODag)
foldEnrichments(davidGODag)
bonferronis(davidGODag)
benjamins(davidGODag)
fdrs(davidGODag)

##The user can even plot the enrichment GO DAG if Rgraphviz package is
##available.
plotGOTermGraph(g=goDag(davidGODag), r=davidGODag, max.nchar=30,
node.shape="ellipse")
}
```

---

DAVIDResult-class      *class "DAVIDResult"*

---





---

DAVIDTermCluster-class  
*class "DAVIDTermCluster"*

---

### Description

This class represents the output of a DAVID Functional Annotation Clustering report.

### Type

This class is a "Concrete" one.

### Extends

- *DAVIDCluster* and uses its constructor to parse the report.

### Slots

the ones inherited from DAVIDCluster.

### Methods

`initialize` signature(.Object="DAVIDTermCluster", fileName="character"): basic cluster report file parser.

`DAVIDTermCluster` signature(fileName="character"): high level gene cluster report file parser.

`ids` signature(object="DAVIDTermCluster"): list with the member ids within each cluster.

`plot2D` signature(object="DAVIDTermCluster", number=1, color=c("FALSE"="black", "TRUE"="green")): ggplot2 tile plot of genes vs functional annotation category membership of the given cluster number.

### Author(s)

Cristobal Fresno and Elmer A Fernandez

### References

1. The Database for Annotation, Visualization and Integrated Discovery ([david.abcc.ncifcrf.gov](http://david.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C. & Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. *Nucleic Acids Res*, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175

### See Also

Other DAVIDTermCluster: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [ids](#), [ids](#), [ids](#), [ids](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#)

**Examples**

```

{
##Load the Gene Functional Classification Tool file report for the
##input demo file 2 to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/termClusterReport2.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidTermCluster2<-DAVIDTermCluster(untar(fileName, list=TRUE))
davidTermCluster2

##Now we can invoke DAVIDCluster ancestor functions to inspect the report
##data, of each cluster. For example, we can call summary to get a general
##idea, and the inspect the cluster with higher Enrichment Score, to see
##which members belong to it, etc. Or simply returning the whole cluster as a
##list with EnrichmentScore and Members.
summary(davidTermCluster2)
higherEnrichment<-which.max(enrichment(davidTermCluster2))
clusterGenes<-members(davidTermCluster2)[[higherEnrichment]]
wholeCluster<-cluster(davidTermCluster2)[[higherEnrichment]]

##Then, we can obtain the ids of the term members calling clusterGenes object
##which is a DAVIDFunctionalAnnotationChart class or directly using ids on
##davidTermCluster2 for the higherEnrichment cluster.
ids(clusterGenes)
ids(davidTermCluster2)[[higherEnrichment]]

##Finally, we can inspect a 2D tile membership plot, to visual inspect for
##overlapping of genes across the term members of the selected cluster.
plot2D(davidTermCluster2, number=higherEnrichment)
}

```

---

DAVIDWebService-class *Main class to connect to DAVID Web Service*

---

**Description**

A reference class to manage DAVID's Web Service connectivity, to run Set Enrichment Analysis (SEA) or Modular Enrichment Analysis (MEA) on a candidate list of gene/protein(s) with respect to a background list (the genome of the organism by default).

**Usage**

```
DAVIDWebService(...)
```

**Arguments**

... additional parameters. See Methods section.

**Details**

DAVIDWebService class is implemented as a reference class, to manage a single instance connection to DAVID's server by means of web services using a registered e-mail. For user registration, go to

<http://david.abcc.ncifcrf.gov/webservice/register.html>. The implementation uses Java Remote Method Implementation (RMI) to connect the client and server side of DAVID. The main functionalities include:

1. Connectivity: upload gene/background list/s, change gene/background position, select current specie/s, select annotations, etc. from R.
2. Reports: Submitted Gene List, Annotation Category Summary, Gene/Term Clusters, Functional Annotation Chart and Functional Annotation Table as native R objects.

### Fields

**stub:** Java jobjRef which corresponds to a sample/session/client/stub/DAVIDWebServiceStub object for the client side of DAVID.

**email:** character.

### Methods

**show():** prints DAVIDWebService object.

**summary():** return a data.frame with a summary of all available annotations in DAVID in terms of percentage of gene list ids present in the category and numbers of terms where they can be found (see `getAnnotationSummary`)

**initialize(email="", ..., url):** constructor for DAVIDWebService object, which includes: Java Virtual Machine initialization (... if required), and stub initialization with the provided email (if present) and using the url parameter for the API website.

**setEmail(mail):** Set the email field with the given registered user email parameter for authentication purposes.

**getEmail():** Returns the current authentication email in use.

**getStub:** Returns jobjRef object with the current stub field in use.

**is.connected():** Check if connected to the DAVID server.

**connect():** Try to establish a connection with the DAVID server using the provided email.

**getIdTypes():** Returns all acceptable DAVID idTypes.

**getAllAnnotationCategoryNames():** Returns all available annotation category names.

**getDefaultCategoryNames():** Returns all default category names.

**getGeneListNames():** Returns submitted gene list names.

**getBackgroundListNames():** Returns submitted background names.

**getListName(listType=c("Gene", "Background"), position=1L):** Get the name of the selected list type at a given position.

**getSpecieNames():** Return specie/s of the current gene list.

**getCurrentGeneListPosition():** Return the position of current gene list.

**getCurrentBackgroundListPosition():** Return the position of current background list.

**getCurrentSpeciesPosition():** Return current specie/s used positions for the uploaded gene list.

**setCurrentGeneListPosition(position):** Use the gene list of the given position.

**setCurrentBackgroundPosition(position):** Use the gene list of the given position.

**setCurrentSpecies(species):** Select the specie/s of the submitted gene list to use in the analysis.

`setAnnotationCategories(categories)`: Select the specie/s of the submitted gene list to use in the analysis.

`addList(inputIds, idType, listName, listType=c("Gene", "Background"))`: Add a gene or background to the current session.

`getGeneCategoriesReport()`: Get the gene report categories.

`getAnnotationSummary()`: Generate the summary of all available annotation in DAVID in terms of percentage of gene list ids present in the category and numbers of terms where the can be found.

`getGeneListReportFile(fileName)`: Generate the Gene List Report a.k.a Show Gene List in DAVID website and save it into a file.

`getGeneListReport()`: `getGeneListReport` but as an R object.

`getFunctionalAnnotationChartFile(fileName, threshold=0.1, count=2L)`: Generate the Functional Annotation Chart Report for the selected functional categories, for the given EASE threshold and number of genes and save it to a file.

`getFunctionalAnnotationChart(...)`: `getFunctionalAnnotationChart` but as an R object.

`getClusterReportFile(fileName, type=c("Term", "Gene"), overlap=4L, initialSeed=4L, finalSeed=4L)`: Generate the Term/Gene Cluster Report for the given configuration.

`getClusterReport(type=c("Term", "Gene"), ...)`: Wrapper for `getClusterReportFile` function.

`getFunctionalAnnotationTableFile(fileName)`: Generate Functional Annotation Table Report File, which is a gene-centric view of the genes and their associated annotation terms (selected only). There is no statistics applied in this report.

`getFunctionalAnnotationTable()`: `getFunctionalAnnotationTable` but as an R object.

### Limitations

1. A job with more than 3000 genes to generate gene or term cluster report will not be handled by DAVID due to resource limit.
2. No more than 200 jobs in a day from one user or computer.
3. DAVID Team reserves right to suspend any improper uses of the web service without notice.

### Author(s)

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### References

1. The Database for Annotation, Visualization and Integrated Discovery ([david.abcc.ncifcrf.gov](http://david.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C. & Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. *Nucleic Acids Res*, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175
3. Huang, D. W.; Sherman, B. T. & Lempicki, R. A. Bioinformatics enrichment tools: paths toward the comprehensive functional analysis of large gene lists. *Nucleic Acids Res*, Laboratory of Immunopathogenesis and Bioinformatics, Clinical Services Program, SAIC-Frederick, Inc., National Cancer Institute at Frederick, Frederick, MD 21702, USA., 2009, 37, 1-13

4. Xiaoli Jiao, Brad T. Sherman, Da Wei Huang, Robert Stephens, Michael W. Baseler, H. Clifford Lane, Richard A. Lempicki, DAVID-WS: A Stateful Web Service to Facilitate Gene/Protein List Analysis *Bioinformatics* 2012 doi:10.1093/bioinformatics/bts251
5. Cristobal Fresno, Elmer A. Fernandez (2013) RDAVIDWebService: a versatile R interface to DAVID, *Bioinformatics*, 29(21), 2810-2811., <http://bioinformatics.oxfordjournals.org/content/29/21/2810>.

### See Also

Other DAVIDWebService: `addList`, `addList`, `connect`, `connect`, `getAllAnnotationCategoryNames`, `getAllAnnotationCategoryNames`, `getAnnotationSummary`, `getAnnotationSummary`, `getBackgroundListNames`, `getBackgroundListNames`, `getClusterReport`, `getClusterReport`, `getClusterReportFile`, `getClusterReportFile`, `getCurrentBackgroundListPosition`, `getCurrentBackgroundListPosition`, `getCurrentGeneListPosition`, `getCurrentGeneListPosition`, `getCurrentSpeciesPosition`, `getCurrentSpeciesPosition`, `getDefaultCategoryNames`, `getDefaultCategoryNames`, `getEmail`, `getEmail`, `getFunctionalAnnotationChart`, `getFunctionalAnnotationChart`, `getFunctionalAnnotationChartFile`, `getFunctionalAnnotationChartFile`, `getFunctionalAnnotationTable`, `getFunctionalAnnotationTable`, `getFunctionalAnnotationTableFile`, `getFunctionalAnnotationTableFile`, `getGeneCategoriesReport`, `getGeneCategoriesReport`, `getGeneListNames`, `getGeneListNames`, `getGeneListReport`, `getGeneListReport`, `getGeneListReportFile`, `getGeneListReportFile`, `getIdTypes`, `getIdTypes`, `getListName`, `getListName`, `getSpecieNames`, `getSpecieNames`, `getStub`, `getStub`, `is.connected`, `is.connected`, `setAnnotationCategories`, `setAnnotationCategories`, `setCurrentBackgroundPosition`, `setCurrentBackgroundPosition(position)`, `setCurrentGeneListPosition`, `setCurrentGeneListPosition`, `setCurrentSpecies`, `setCurrentSpecies`, `setEmail`, `setEmail`, `setEmail`, `DAVIDWebService-method`, `summary`, `summary`, `summary`, `summary`

---

demoList1

*DAVID's website demoList1 example id files*

---

### Description

This datasets are the same example input id files present in the Database for Annotation, Visualization and Integrated Discovery.

### Usage

```
data(demoList1)
```

```
data(demoList2)
```

### Format

character vector with AFFYMETRIX\_3PRIME\_IVT\_ID manufacturer identification codes (ids)

**demoList1** 164 ids in total.

**demoList2** 403 ids in total.

### Author(s)

Cristobal Fresno and Elmer A Fernandez

## References

1. The Database for Annotation, Visualization and Integrated Discovery ([david.abcc.ncifcrf.gov](http://david.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C.; Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. *Nucleic Acids Res*, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175

## See Also

Other DataExamples: [annotationSummary1](#), [annotationSummary2](#), [geneList1](#), [geneList2](#)

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funChart1

*DAVID's website Functional Annotation Chart example files*

---

## Description

These datasets correspond to the reports obtained using Functional Annotation Chart Reports in the Database for Annotation, Visualization and Integrated Discovery (DAVID) website, using as input file the ones provided for demo purposes (demoList1 or demoList2) with GOTERM\_BP\_ALL, GOTERM\_MF\_ALL and GOTERM\_CC\_ALL categories.

## Usage

```
data(funChart1)
```

```
data(funChart2)
```

## Format

funChart1/2 are data.frame for demoList1/2 input ids, respectively, with the following columns.

**Category** factor with the main categories under used in the present analysis.

**Term** character with the name of the term in format id~name (if available).

**Count** integer with the number of ids of the gene list that belong to this term.

**X.** after converting user input gene IDs to corresponding DAVID gene ID, it refers to the percentage of DAVID genes in the list associated with particular annotation term. Since DAVID gene ID is unique per gene, it is more accurate to use DAVID ID percentage to present the gene-annotation association by removing any redundancy in user gene list, i.e. two user IDs represent same gene.

**PValue** numeric with the EASE Score of the term (see DAVID Help page).

**Genes** character in comma separated style with the genes present in the term.

**List.Total, Pop.Hits, Pop.Total** integers (in addition to Count) to build the 2x2 contingency table in order to compute the EASE Score (see DAVID Help page).

**Fold.Enrichment** numeric with the ratio of the two proportions. For example, if 40/400 (i.e. 10%) of your input genes involved in "kinase activity" and the background information is 300/30000 genes (i.e. 1%) associating with "kinase activity", roughly  $10\%/1\%=10$  fold enrichment.

**Bonferroni, Benjamini, FDR** numerics with p-value adjust different criterias (see p.adjust)

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

**References**

1. The Database for Annotation, Visualization and Integrated Discovery ([david.abcc.ncifcrf.gov](http://david.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C.; Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. *Nucleic Acids Res*, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175
3. DAVID Help page [http://david.abcc.ncifcrf.gov/helps/functional\\_annotation.html#E3](http://david.abcc.ncifcrf.gov/helps/functional_annotation.html#E3)

---

geneCluster1

*DAVID's website gene/term cluster report example files*

---

**Description**

These datasets correspond to the Functional Annotation Clustering or Gene Functional Classification report obtained in the Database for Annotation, Visualization and Integrated Discovery (DAVID) website, using as input file the ones provided for demo purposes (demoList1 or demoList2) with GOTERM\_BP\_ALL, GOTERM\_MF\_ALL and GOTERM\_CC\_ALL categories.

**Format**

geneCluster1/2 or termCluster1/2 are tab delimitate unstructured files with DAVID format where:

**Cluster header** 1. TypeGene Cluster or Annotation Cluster.

2. Numberinteger to indicate the cluster label.
3. Enrichment Scorenumeric with the geometric mean (in -log scale) of members p-values in a corresponding annotation cluster, is used to rank their biological significance. Thus, the top ranked annotation groups most likely have consistent lower p-values for their annotation members.

**Members Header** according to the type of cluster it can be:

1. Genethe character vector with "ID", "Gene" and "Name".
2. Annotationthe same columns of a Functional Annotation Chart (see `getFunctionalAnnotationChart`).

**Members Body** member data per line according to the respective type of cluster.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

## References

1. The Database for Annotation, Visualization and Integrated Discovery ([davidgenelist.abcc.ncifcrf.gov](http://davidgenelist.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C.; Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. Nucleic Acids Res, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175
3. DAVID Help page [http://david.abcc.ncifcrf.gov/helps/functional\\_classification.html#textmode](http://david.abcc.ncifcrf.gov/helps/functional_classification.html#textmode)

---

geneList1

*DAVID's website gene list example files*

---

## Description

These datasets correspond to the reports obtained using Show Gene List in the Database for Annotation, Visualization and Integrated Discovery (DAVID) website, using as input file the ones provided for demo purposes (demoList1 or demoList2) with default options.

## Usage

```
data(geneList1)
```

```
data(geneList2)
```

## Format

geneList1/2 are data.frame for demoList1/2 input ids, respectively, with the following columns.

**ID** character with the Gene List ID present in DAVID knowledge base, in the submitted type. If more than one ids map to the same DAVID ID, the record is a comma separated character.

**Name** character with the name of the gene as seen in DAVID knowledge base, in a comma separated fashion (if more than one ID maps to the same DAVID ID).

**Species** factor with the name of the Specie.

## Author(s)

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## References

1. The Database for Annotation, Visualization and Integrated Discovery ([david.abcc.ncifcrf.gov](http://david.abcc.ncifcrf.gov))
2. Huang, D. W.; Sherman, B. T.; Tan, Q.; Kir, J.; Liu, D.; Bryant, D.; Guo, Y.; Stephens, R.; Baseler, M. W.; Lane, H. C.; Lempicki, R. A. DAVID Bioinformatics Resources: expanded annotation database and novel algorithms to better extract biology from large gene lists. Nucleic Acids Res, Laboratory of Immunopathogenesis and Bioinformatics, SAIC-Frederick, Inc., National Cancer Institute at Frederick, MD 21702, USA., 2007, 35, W169-W175
3. DAVID Help page [http://david.abcc.ncifcrf.gov/helps/functional\\_annotation.html#E3](http://david.abcc.ncifcrf.gov/helps/functional_annotation.html#E3)



**See Also**

Other DataExamples: [annotationSummary1](#), [annotationSummary2](#), [demoList1](#), [demoList2](#)

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genes	<i>genes for the different DAVIDWebService package class objects.</i>
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**Description**

Obtain genes related information, according to the given function call (see Values).

**Usage**

```
genes(object, ...)

## S4 method for signature 'DAVIDGenes'
genes(object, ids)

## S4 method for signature 'DAVIDGeneCluster'
genes(object)

## S4 method for signature 'DAVIDFunctionalAnnotationTable'
genes(object,
      ...)
```

**Arguments**

object	DAVIDGenes or DAVIDGeneCluster class object.
ids	character vector with the ids to fetch.
...	Additional parameters for internal functions (if applicable).

**Value**

according to the call one of the following objects can be returned

DAVIDGenes	a DAVIDGenes object with the matched genes of ids parameter. If missing, returns all the genes.
DAVIDGeneCluster	list with DAVIDGenes objects for each cluster.
DAVIDFunctionalAnnotationTable	a DAVIDGenes objects, according to ... parameter used internally on genes(DAVIDGenes, ...).

**Author(s)**

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**See Also**

Other DAVIDFunctionalAnnotationTable: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable-class](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [categories](#), [categories](#), [categories](#), [dictionary](#), [dictionary](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [membership](#), [membership](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [subset](#), [subset](#)

Other DAVIDGeneCluster: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster-class](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [ids](#), [ids](#), [ids](#), [ids](#), [ids](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#)

Other DAVIDGenes: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes-class](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [ids](#), [ids](#), [ids](#), [ids](#), [ids](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#)

**Examples**

```
{
##DAVIDGenes example:
##Load Show Gene List file report for the input demo file 1, using data
##function. Then, create a DAVIDGenes object using the loaded data.frame
##geneList1.
data(geneList1)
davidGenes1<-DAVIDGenes(geneList1)

##Now, get the genes using the ids look up parameter with the first
##six ids. If ids omitted, all the available are returned.
genes(davidGenes1, ids=head(ids(davidGenes1)))

##DAVIDFunctionalAnnotationTable example:
##Load the Functional Annotation Table file report for the input demo
##file 1, using data function. Then, create a DAVIDFunctionalAnnotationTable
##object using the loaded data.frame annotationTable1.
data(annotationTable1)
davidFunTable1<-DAVIDFunctionalAnnotationTable(annotationTable1)

##Now we can obtain the genes for the given ids, or the complete list if the
##parameter is omitted.
genes(davidFunTable1, id=c("37166_at", "41703_r_at"))

##DAVIDGeneCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo list 1 file to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/geneClusterReport1.tab.tar.gz",
package="RDAVIDWebService")
```

```

untar(fileName)
davidGeneCluster1<-DAVIDGeneCluster(untar(fileName, list=TRUE))

##Then, we can obtain the genes of the first cluster using davidGeneCluster1
##object. Or, using genes on DAVIDGenes class once we get the members of the
##cluster
genes(davidGeneCluster1)[[1]]
genes(members(davidGeneCluster1)[[1]])
}

```

---

getGeneCategoriesReport

*Obtain DAVID website reports*


---

### Description

DAVIDWebService class methods to obtain DAVID website reports from R. This includes the different functionalities starting from the basic "Show Gene List" or "Annotation Summary", to Set Enrichment Analysis using "Functional Annotation Chart" or Modular Enrichment Analysis using "Functional Annotation Clustering" or "Gene Functional Classification Tool". Note that DAVIDWebService is a Reference class, hence invoke it using object\_name\$method\_name(parameters). In addition, the user can use the S4 version style function call (see Details).

### Usage

```

getGeneCategoriesReport(object)

## S4 method for signature 'DAVIDWebService'
getGeneCategoriesReport(object)

getAnnotationSummary(object)

## S4 method for signature 'DAVIDWebService'
getAnnotationSummary(object)

getGeneListReportFile(object, fileName)

## S4 method for signature 'DAVIDWebService'
getGeneListReportFile(object,
  fileName)

getGeneListReport(object)

## S4 method for signature 'DAVIDWebService'
getGeneListReport(object)

getFunctionalAnnotationChartFile(object, fileName,
  threshold=0.1, count=2L)

## S4 method for signature 'DAVIDWebService'
getFunctionalAnnotationChartFile(object,
  fileName, threshold=0.1, count=2L)

```

```

getFunctionalAnnotationChart(object, ...)

## S4 method for signature 'DAVIDWebService'
getFunctionalAnnotationChart(object,
...)

getClusterReportFile(object, fileName, type=c("Term",
"Gene"), overlap=4L, initialSeed=4L, finalSeed=4L,
linkage=0.5, kappa=35L)

## S4 method for signature 'DAVIDWebService'
getClusterReportFile(object,
fileName, type=c("Term", "Gene"), overlap=4L,
initialSeed=4L, finalSeed=4L, linkage=0.5, kappa=35L)

getClusterReport(object, type=c("Term", "Gene"), ...)

## S4 method for signature 'DAVIDWebService'
getClusterReport(object,
type=c("Term", "Gene"), ...)

getFunctionalAnnotationTableFile(object, fileName)

## S4 method for signature 'DAVIDWebService'
getFunctionalAnnotationTableFile(object,
fileName)

getFunctionalAnnotationTable(object)

## S4 method for signature 'DAVIDWebService'
getFunctionalAnnotationTable(object)

```

### Arguments

object	DAVIDWebService class object.
fileName	character with the name of the file to store the Report.
threshold	numeric with the EASE score (at most equal) that must be present in the category to be included in the report. Default value is 0.1.
count	integer with the number of genes (greater equal) that must be present in the category to be included in the report. Default value is 2.
type	character with the type of cluster to obtain Term/Genes. Default value "Term".
overlap	integer with the minimum number of annotation terms overlapped between two genes in order to be qualified for kappa calculation. This parameter is to maintain necessary statistical power to make kappa value more meaningful. The higher value, the more meaningful the result is. Default value is 4L.
initialSeed, finalSeed	integer with the number of genes in the initial (seeding) and final (filtering) cluster criteria. Default value is 4L for both.
linkage	numeric with the percentage of genes that two clusters share in order to become one.

kappa integer (kappa \* 100), with the minimum kappa value to be considered biological significant. The higher setting, the more genes will be put into unclustered group, which lead to higher quality of functional classification result with a fewer groups and a fewer gene members. Kappa value 0.3 starts giving meaningful biology based on our genome-wide distribution study. Anything below 0.3 have great chance to be noise.

... additional parameters for getXXFile functions.

## Details

Available functions include:

**getGeneCategoriesReport:** Get the gene categories report.

**getAnnotationSummary:** Generate the summary of all available annotation in DAVID in terms of percentage of gene list ids present in the category and numbers of terms where the can be found.

**getGeneListReportFile:** Generate the Gene List Report a.k.a Show Gene List in DAVID website and save it into a file.

**getGeneListReport:** Generate Gene List Report a.k.a Show Gene List in DAVID website and import it as a DAVIDGenes object into R.

**getFunctionalAnnotationChartFile:** Generate the Functional Annotation Chart Report for the selected functional categories, for the given EASE threshold and number of genes and save it to a file.

**getFunctionalAnnotationChart:** Generate the Functional Annotation Chart Report for the selected functional categories, for the given EASE threshold and number of genes, and import it as a DAVIDFunctionalAnnotationChart object in R.

**getClusterReportFile:** Generate the Term/Gene Cluster Report for the given configuration.

**getClusterReport:** Generate the Term/Gene Cluster Report for the given configuration, and import it as a DAVIDGeneCluster or DAVIDTermCluster object, according to function call.

**getFunctionalAnnotationTableFile:** Generate Functional Annotation Table Report File, which is a gene-centric view of the genes and their associated annotation terms (selected only). There is no statistics applied in this report.

**getFunctionalAnnotationTable:** Generate Functional Annotation Table Report and import it as a DAVIDFunctionalAnnotationTable object in R.

## Value

according to the call one of the following objects can be returned

**getGeneCategoriesReport**

integer vector with the IDs of the categories.

**getAnnotationSummary**

data.frame with the annotation summary report with the following columns:

1. **Main.Category:** factor with the main categories under used in the present analysis.
2. **ID:** integer to identify the annotation category.
3. **Name:** character with the name of category (the available ones in getAlAnnotationCategoryNames function).
4. **X.:** numeric with the percentage of the gene list ids present in the term.

5. **Count:** integer with the number of ids of the gene list that belong to this term.

`getGeneListReportFile`

data.frame with the Gene List Report with the following columns:

1. **ID:** character with the Gene List ID present in DAVID knowledge base, in the submitted type. If more than one ids map to the same DAVID ID, the record is a comma separated character.
2. **Name:** character with the name of the gene as seen in DAVID knowledge base, in a comma separated fashion (if more than one ID maps to the same DAVID ID).
3. **Species:** factor with the name of the Specie.

`getGeneListReport`

Generate Gene List Report a.k.a Show Gene List in DAVID website and import it as a DAVIDGenes object in R.

`getFunctionalAnnotationChartFile`

file with the following columns:

1. **Category:** factor with the main categories under used in the present analysis.
2. **Term:** character with the name of the term in format id~name (if available).
3. **Count:** integer with the number of ids of the gene list that belong to this term.
4. **X.:** after converting user input gene IDs to corresponding DAVID gene ID, it refers to the percentage of DAVID genes in the list associated with a particular annotation term. Since DAVID gene ID is unique per gene, it is more accurate to use DAVID ID percentage to present the gene-annotation association by removing any redundancy in user gene list, i.e. two user IDs represent same gene.
5. **PValue:** numeric with the EASE Score of the term (see DAVID Help page).
6. **Genes:** character in comma separated style with the genes present in the term.
7. **List.Total, Pop.Hits, Pop.Total:** integers (in addition to Count) to build the 2x2 contingency table in order to compute the EASE Score (see DAVID Help page).
8. **Fold.Enrichment:** numeric with the ratio of the two proportions. For example, if 40/400 (i.e. 10%) of your input genes involved in "kinase activity" and the background information is 300/30000 genes (i.e. 1%) associating with "kinase activity", roughly  $10\% / 1\% = 10$  fold enrichment.
9. **Bonferroni, Benjamini, FDR:** numerics with p-value adjust different criteria (see p.adjust).

`getFunctionalAnnotationChart`

Generate the Functional Annotation Chart Report for the selected functional categories, for the given EASE threshold and number of genes, and import it as a DAVIDFunctionalAnnotationChart object in R.

`getClusterReportFile`

file with the following columns:

1. **Annotation/Gene Cluster:** integer with the number of cluster.
2. **EnrichmentScore:** numeric with the geometric mean (in -log scale) of members p-values in a corresponding annotation cluster, is used to rank their biological significance. Thus, the top ranked annotation groups most likely have consistent lower p-values for their annotation members.

3. **Members:** according to the type of cluster, changes the associated data to include Gene List or Functional Chart Report (see `getGeneListReport` and `getFunctionalAnnotationChart`).

`getClusterReport`

Generate the Term/Gene Cluster Report for the given configuration, and import it as a `DAVIDGeneCluster` or `DAVIDTermCluster` according to function call.

`getFunctionalAnnotationTableFile`

file with the following columns:

1. **Gene:** Three Columns with the same data included in Gene List Report (ID, Gene.Name and Species) but coding for DAVID ID, i. e., comma separated character with input ids if two or more stands for the same gene.
2. **Annotation:** as many columns as Annotation Categories were in used. In each column, a comma separated style is use to delimitate the different terms where is reported evidence for DAVID ID record.

`getFunctionalAnnotationTable:`

Generate Functional Annotation Table Report, which is a gene-centric view of the genes and their associated annotation terms (selected only), and import it as a `DAVIDFunctionalAnnotationTable` object in R.

## References

1. [http://david.abcc.ncifcrf.gov/helps/functional\\_annotation.html#E3](http://david.abcc.ncifcrf.gov/helps/functional_annotation.html#E3)
2. [http://david.abcc.ncifcrf.gov/helps/functional\\_classification.html#clustering](http://david.abcc.ncifcrf.gov/helps/functional_classification.html#clustering)
3. Cohen, J: A coefficient of agreement for nominal scales, Educational and Psychological Measurement, 1960, 20, 37-46.

## See Also

`p.adjust` and `fisher.test`

Other `DAVIDWebService`: `DAVIDWebService-class`, `addList`, `addList`, `connect`, `connect`, `getAllAnnotationCategories`, `getAllAnnotationCategoryNames`, `getBackgroundListNames`, `getBackgroundListNames`, `getCurrentBackgroundListPosition`, `getCurrentBackgroundListPosition`, `getCurrentGeneListPosition`, `getCurrentGeneListPosition`, `getCurrentSpeciesPosition`, `getCurrentSpeciesPosition`, `getDefaultCategoryNames`, `getDefaultCategoryNames`, `getEmail`, `getEmail`, `getGeneListNames`, `getGeneListNames`, `getIdTypes`, `getIdTypes`, `getListName`, `getListName`, `getSpecieNames`, `getSpecieNames`, `getStub`, `getStub`, `is.connected`, `is.connected`, `setAnnotationCategories`, `setAnnotationCategories`, `setCurrentBackgroundPosition`, `setCurrentBackgroundPosition`, `setCurrentGeneListPosition`, `setCurrentGeneListPosition`, `setCurrentSpecies`, `setCurrentSpecies`, `setEmail`, `setEmail`, `setEmail`, `DAVIDWebService-method`, `summary`, `summary`, `summary`, `summary`

---

ids

*ids for the different DAVIDWebService package class objects*

---

## Description

Obtain ids related information, according to the given function call (see Values).

**Usage**

```

ids(object)

## S4 method for signature 'DAVIDGenes'
ids(object)

## S4 method for signature 'DAVIDFunctionalAnnotationChart'
ids(object)

## S4 method for signature 'DAVIDGeneCluster'
ids(object)

## S4 method for signature 'DAVIDTermCluster'
ids(object)

```

**Arguments**

**object** DAVIDWebService class object. Possible values are: DAVIDGenes, DAVID-FunctionalAnnotationChart, DAVIDGeneCluster or DAVIDTermCluster.

**Value**

according to the call one of the following objects can be returned

DAVIDGenes            character vector with gene submitted ids.  
DAVIDFunctionalAnnotationChart  
                         list with character/integer vector of ids of the corresponding "Category".  
DAVIDGeneCluster, DAVIDTermCluster  
                         list with character/integer vector of ids of the members of each cluster.

**Author(s)**

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**See Also**

Other DAVIDFunctionalAnnotationChart: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart-class](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [categories](#), [categories](#), [categories](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#)

Other DAVIDGeneCluster: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster-class](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [genes](#), [genes](#), [genes](#), [genes](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#), [plot2D](#)

Other DAVIDGenes: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#),



DAVIDGenes, DAVIDGenes, DAVIDGenes, DAVIDGenes-class, DAVIDTermCluster, DAVIDTermCluster, as, as, as, genes, genes, genes, genes, initialize, initialize, initialize, initialize, initialize, initialize, initialize, initialize

Other DAVIDTermCluster: DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationTable, DAVIDFunctionalAnnotationTable, DAVIDFunctionalAnnotationTable, DAVIDGODag, DAVIDGODag, DAVIDGeneCluster, DAVIDGeneCluster, DAVIDGenes, DAVIDGenes, DAVIDGenes, DAVIDTermCluster, DAVIDTermCluster, DAVIDTermCluster-class, as, as, as, initialize, initialize, initialize, initialize, initialize, initialize, initialize, initialize, plot2D, plot2D, plot2D, plot2D, plot2D, plot2D

## Examples

```
{
##DAVIDGenes example:
##Load Show Gene List file report for the input demo file 1, using data
##function. Then, create a DAVIDGenes object using the loaded data.frame
##geneList1. Once, the report is loaded, we can retrieve the ids.
data(geneList1)
davidGenes1<-DAVIDGenes(geneList1)
ids(davidGenes1)

##DAVIDFunctionalAnnotationChart example:
##Load the Functional Annotation Chart file report for the input demo
##file 2, using data function. Then, create a DAVIDFunctionalAnnotationChart
##object using the loaded data.frame funChart2. Once the report is loaded,
##the user can obtain the ids of the genes present in each Term, as a list of
##character vector.
data(funChart2)
davidFunChart2<-DAVIDFunctionalAnnotationChart(funChart2)
ids(davidFunChart2)

##DAVIDGeneCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo list 1 file to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/geneClusterReport1.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidGeneCluster1<-DAVIDGeneCluster(untar(fileName, list=TRUE))
davidGeneCluster1

##Now we can invoke DAVIDCluster ancestor functions to inspect the report
##data, of each cluster. For example, we can call summary to get a general
##idea, and the inspect the cluster with higher Enrichment Score, to see
##which members belong to it, etc. Or simply returning the whole cluster as
##a list with EnrichmentScore and Members.
summary(davidGeneCluster1)
higherEnrichment<-which.max(enrichment(davidGeneCluster1))
clusterGenes<-members(davidGeneCluster1)[[higherEnrichment]]
wholeCluster<-cluster(davidGeneCluster1)[[higherEnrichment]]

##Now, we can obtain the ids of the first cluster directly using
##davidGeneCluster1 or by using DAVIDGenes class on the same cluster.
ids(davidGeneCluster1)[[1]]
ids(members(davidGeneCluster1)[[1]])
```

```

##DAVIDTermCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo file 2 to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/termClusterReport2.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidTermCluster2<-DAVIDTermCluster(untar(fileName, list=TRUE))
davidTermCluster2

##Now we can invoke DAVIDCluster ancestor functions to inspect the report
##data, of each cluster. For example, we can call summary to get a general
##idea, and the inspect the cluster with higher Enrichment Score, to see
##which members belong to it, etc. Or simply returning the whole cluster as a
##list with EnrichmentScore and Members.
summary(davidTermCluster2)
higherEnrichment<-which.max(enrichment(davidTermCluster2))
clusterGenes<-members(davidTermCluster2)[[higherEnrichment]]
wholeCluster<-cluster(davidTermCluster2)[[higherEnrichment]]

##Then, we can obtain the ids of the term members calling clusterGenes object
##which is a DAVIDFunctionalAnnotationChart class or directly using ids on
##davidTermCluster2 for the higherEnrichment cluster.
ids(clusterGenes)
ids(davidTermCluster2)[[higherEnrichment]]
}

```

---

is.connected

*Methods to manipulate DAVID website*


---

## Description

DAVIDWebService class methods to manipulate DAVID website status from R. This includes different functionalities to set up and track the connexion, upload a Gene/Background list, check the species, etc. Note that DAVIDWebService is a Reference class, hence invoke it using object\_name\$method\_name(parameters). In addition, the user can use the S4 version style function call (see Details).

## Usage

```

is.connected(object)

## S4 method for signature 'DAVIDWebService'
is.connected(object)

connect(object)

## S4 method for signature 'DAVIDWebService'
connect(object)

getIdTypes(object)

```

```
## S4 method for signature 'DAVIDWebService'  
getIdTypes(object)  
  
addList(object, inputIds, idType, listName,  
listType=c("Gene", "Background"))  
  
## S4 method for signature 'DAVIDWebService'  
addList(object, inputIds,  
idType, listName, listType=c("Gene", "Background"))  
  
getAllAnnotationCategoryNames(object)  
  
## S4 method for signature 'DAVIDWebService'  
getAllAnnotationCategoryNames(object)  
  
getDefaultCategoryNames(object)  
  
## S4 method for signature 'DAVIDWebService'  
getDefaultCategoryNames(object)  
  
getGeneListNames(object)  
  
## S4 method for signature 'DAVIDWebService'  
getGeneListNames(object)  
  
getBackgroundListNames(object)  
  
## S4 method for signature 'DAVIDWebService'  
getBackgroundListNames(object)  
  
getListName(object, listType=c("Gene", "Background"),  
position=1L)  
  
## S4 method for signature 'DAVIDWebService'  
getListName(object,  
listType=c("Gene", "Background"), position=1L)  
  
getSpecieNames(object)  
  
## S4 method for signature 'DAVIDWebService'  
getSpecieNames(object)  
  
getCurrentGeneListPosition(object)  
  
## S4 method for signature 'DAVIDWebService'  
getCurrentGeneListPosition(object)  
  
getCurrentBackgroundListPosition(object)  
  
## S4 method for signature 'DAVIDWebService'  
getCurrentBackgroundListPosition(object)
```

```
    getCurrentSpeciesPosition(object)

    ## S4 method for signature 'DAVIDWebService'
    getCurrentSpeciesPosition(object)

    getTimeout(object)

    ## S4 method for signature 'DAVIDWebService'
    getTimeout(object)

    getHttpProtocolVersion(object)

    ## S4 method for signature 'DAVIDWebService'
    getHttpProtocolVersion(object)

    setCurrentGeneListPosition(object, position)

    ## S4 method for signature 'DAVIDWebService'
    setCurrentGeneListPosition(object,
    position)

    setCurrentBackgroundPosition(object, position)

    ## S4 method for signature 'DAVIDWebService'
    setCurrentBackgroundPosition(object,
    position)

    setCurrentSpecies(object, species)

    ## S4 method for signature 'DAVIDWebService'
    setCurrentSpecies(object,
    species)

    setAnnotationCategories(object, categories)

    ## S4 method for signature 'DAVIDWebService'
    setAnnotationCategories(object,
    categories)

    setTimeout(object, milliSeconds)

    ## S4 method for signature 'DAVIDWebService'
    setTimeout(object, milliSeconds)

    setHttpProtocolVersion(object, version)

    ## S4 method for signature 'DAVIDWebService'
    setHttpProtocolVersion(object, version)
```

### Arguments

object            DAVIDWebService class object.

inputIds	character vector with the associated ids.
idType	character with the type of submitted ids.
listName	character to identify the submitted list.
listType	character with the type of list (Gene, Background). Default value is "Gene".
position	integer with the position of the gene/background list to set.
species	numeric vector with the specie/s to use.
categories	character vector with the category name/s to use in the analysis.
milliSeconds	integer with time defined in milli seconds.
version	character with HTTP_PROTOCOL_VERSION to use. At present available strings are: "1.1", "1.0", "HTTP/1.1" and "HTTP/1.0"

## Details

Available functions include:

**connect:** Try to establish a connection with DAVID server using the provided email.

**is.connected:** Check if connected to DAVID server.

**getIdTypes:** Returns all acceptable DAVID idTypes.

**addList:** Add a gene or background to the current session.

**getAllAnnotationCategoryNames:** Returns all available annotation category names.

**getDefaultCategoryNames:** Returns all default category names.

**getGeneListNames:** Returns all list names

**getBackgroundListNames:** Returns background names.

**getListName:** Get the name of the selected list type at a given position.

**getSpecieNames:** Return specie/s of the current gene list.

**getCurrentGeneListPosition:** Return the position of current gene list.

**getCurrentBackgroundListPosition:** Return the position of current background list.

**getCurrentSpeciesPosition:** Return current specie/s used positions for the uploaded gene list.

**setCurrentGeneListPosition:** Use the gene list of the given position.

**setCurrentBackgroundPosition:** Use the background list of the given position.

**setCurrentSpecies:** Select the specie/s of the submitted gene list to use in the analysis.

**setAnnotationCategories:** Let the user to select specific annotation categories.

**getTimeout:** Get apache Axis time out in milliSeconds.

**setTimeout:** Set apache Axis time out in milliSeconds.

**getHttpProtocolVersion:** Get apache Axis HTTP\_PROTOCOL\_VERSION.

**setHttpProtocolVersion:** Set apache Axis HTTP\_PROTOCOL\_VERSION. possible values are defined in org.apache.axis2.transport.http.HTTPConstants class with HEADER\_PROTOCOL\_XX property. At present available strings are: "1.1", "1.0", "HTTP/1.1" and "HTTP/1.0".

**Value**

according to the call one of the following objects can be returned

is.connected	TRUE if user has registered email with DAVID knowledge base, FALSE otherwise.
getIdTypes	character vector with the available DAVID input ID type.
addList	list with two items: i)inDavid, a numeric with the percentage of the inputIds in DAVID knowledge database, ii)unmappedIds, a character vector with the unmapped ids if listType is "Gene", NA_character_ otherwise.
getAllAnnotationCategoryNames	character vector with the available DAVID annotation categories.
getDefaultCategoryNames	character vector with a subset of the available DAVID annotation categories, chosen by default.
getGeneListNames	return a character vector with the name of the submitted gene list/s.
getBackgroundListNames	character vector with the name of the available background gene list/s for the submitted gene list/s.
getListName	character with the name of the list.
getSpeciesNames	character vector with the specie/s and in brackets the number of DAVID Ids of the current gene list, e.g. Homo sapiens(155).
getCurrentGeneListPosition	integer with the position of current gene list if available, NA_integer_ otherwise.
getCurrentBackgroundListPosition	integer with the position of current background list if available, NA_integer_ otherwise.
getCurrentSpeciesPosition	integer vector with the specie/s position under use for the gene list under use if available, NA_character_ otherwise.

**See Also**

Other DAVIDWebService: [DAVIDWebService-class](#), [getAnnotationSummary](#), [getAnnotationSummary](#), [getClusterReport](#), [getClusterReport](#), [getClusterReportFile](#), [getClusterReportFile](#), [getEmail](#), [getEmail](#), [getFunctionalAnnotationChart](#), [getFunctionalAnnotationChart](#), [getFunctionalAnnotationChartFile](#), [getFunctionalAnnotationTable](#), [getFunctionalAnnotationTable](#), [getFunctionalAnnotationTableFile](#), [getFunctionalAnnotationTableFile](#), [getGeneCategoriesReport](#), [getGeneCategoriesReport](#), [getGeneListReport](#), [getGeneListReport](#), [getGeneListReportFile](#), [getGeneListReportFile](#), [getStub](#), [getStub](#), [setEmail](#), [setEmail](#), [setEmail](#), [DAVIDWebService-method](#), [summary](#), [summary](#), [summary](#), [summary](#)

Other DAVIDWebService: [DAVIDWebService-class](#), [getAnnotationSummary](#), [getAnnotationSummary](#), [getClusterReport](#), [getClusterReport](#), [getClusterReportFile](#), [getClusterReportFile](#), [getEmail](#), [getEmail](#), [getFunctionalAnnotationChart](#), [getFunctionalAnnotationChart](#), [getFunctionalAnnotationChartFile](#), [getFunctionalAnnotationTable](#), [getFunctionalAnnotationTable](#), [getFunctionalAnnotationTableFile](#), [getFunctionalAnnotationTableFile](#), [getGeneCategoriesReport](#), [getGeneCategoriesReport](#), [getGeneListReport](#), [getGeneListReport](#), [getGeneListReportFile](#), [getGeneListReportFile](#), [getStub](#), [getStub](#), [setEmail](#), [setEmail](#), [setEmail](#), [DAVIDWebService-method](#), [summary](#), [summary](#), [summary](#), [summary](#)







getGeneListReportFile, getStub, getStub, setEmail, setEmail, setEmail, DAVIDWebService-method, summary, summary, summary, summary

Other DAVIDWebService: DAVIDWebService-class, getAnnotationSummary, getAnnotationSummary, getClusterReport, getClusterReport, getClusterReportFile, getClusterReportFile, getEmail, getEmail, getFunctionalAnnotationChart, getFunctionalAnnotationChart, getFunctionalAnnotationChartFile, getFunctionalAnnotationTable, getFunctionalAnnotationTable, getFunctionalAnnotationTableFile, getFunctionalAnnotationTableFile, getGeneCategoriesReport, getGeneCategoriesReport, getGeneListReport, getGeneListReport, getGeneListReportFile, getGeneListReportFile, getStub, getStub, setEmail, setEmail, setEmail, DAVIDWebService-method, summary, summary, summary, summary

Other DAVIDWebService: DAVIDWebService-class, getAnnotationSummary, getAnnotationSummary, getClusterReport, getClusterReport, getClusterReportFile, getClusterReportFile, getEmail, getEmail, getFunctionalAnnotationChart, getFunctionalAnnotationChart, getFunctionalAnnotationChartFile, getFunctionalAnnotationTable, getFunctionalAnnotationTable, getFunctionalAnnotationTableFile, getFunctionalAnnotationTableFile, getGeneCategoriesReport, getGeneCategoriesReport, getGeneListReport, getGeneListReport, getGeneListReportFile, getGeneListReportFile, getStub, getStub, setEmail, setEmail, setEmail, DAVIDWebService-method, summary, summary, summary, summary

## Examples

```
david <- DAVIDWebService$new()
david$is.connected()
##Or the equivalent S4 style function call
is.connected(david)
```

---

plot2D

*Visualization of biological relationships*

---

## Description

plot2D uses a 2D tile ggplot to explore biological relationships between two variables such as annotation category and genes, for Functional Annotation Chart/Table or Term cluster results. For Gene cluster, the cluster number vs genes membership is plotted.

## Usage

```
plot2D(object,...)

## S4 method for signature 'DAVIDResult'
plot2D(object, dataFrame)

## S4 method for signature 'DAVIDFunctionalAnnotationChart'
plot2D(object,color=c("FALSE"="black",
"TRUE"="green"))

## S4 method for signature 'DAVIDGeneCluster'
plot2D(object,color=c("FALSE"="black", "TRUE"="green"),names=FALSE)

## S4 method for signature 'DAVIDTermCluster'
plot2D(object,number=1,color=c("FALSE"="black", "TRUE"="
```

```

"green"))

## S4 method for signature 'DAVIDFunctionalAnnotationTable'
plot2D(object,
  category, id, names.genes=FALSE,
  names.category=FALSE,color=c("FALSE"="black", "TRUE"="green"))

```

### Arguments

object	DAVIDResult heirs (DAVIDFunctionalAnnotationChart/Table or DAVIDGeneCluster/TermCluster)
dataFrame	data.frame with three columns (x, y and fill) to be used in ggplot. X(Y) is a character/factor with the X(Y)-axis labels and "fill" is the color to be used for x-y labels.
color	named character vector to indicate tile color. Default value is c("FALSE"="black", "TRUE"="green").
names	should gene names be plotted? Default value is FALSE, i.e, use ids.
number	integer to indicate which cluster to plot. Default value is 1.
category	character vector to select the main annotation categories. By default is missing in order to use all the available ones.
id	character vector to indicate which gene ids to use. By default is missing in order to use all the available ones.
names.genes, names.category	Should genes and/or category names used? Default value is FALSE, i.e., use both ids.
...	Additional parameters for heirs functions.

### Value

a ggplot object if the object is not empty.

### Author(s)

Cristobal Fresno and Elmer A Fernandez

### See Also

Other DAVIDFunctionalAnnotationChart: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart-class](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [categories](#), [categories](#), [categories](#), [ids](#), [ids](#), [ids](#), [ids](#), [ids](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#)

Other DAVIDFunctionalAnnotationTable: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable-class](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [categories](#), [categories](#), [categories](#), [dictionary](#), [dictionary](#), [genes](#), [genes](#), [genes](#), [genes](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [membership](#), [membership](#), [subset](#), [subset](#)

Other DAVIDGeneCluster: DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationTable, DAVIDFunctionalAnnotationTable, DAVIDFunctionalAnnotationTable, DAVIDGODag, DAVIDGODag, DAVIDGeneCluster, DAVIDGeneCluster, DAVIDGeneCluster-class, DAVIDGenes, DAVIDGenes, DAVIDGenes, DAVIDTermCluster, DAVIDTermCluster, as, as, as, genes, genes, genes, genes, ids, ids, ids, ids, ids, initialize, initialize, initialize, initialize, initialize, initialize, initialize

Other DAVIDResult: DAVIDResult-class, type, type

Other DAVIDTermCluster: DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationTable, DAVIDFunctionalAnnotationTable, DAVIDFunctionalAnnotationTable, DAVIDGODag, DAVIDGODag, DAVIDGeneCluster, DAVIDGeneCluster, DAVIDGenes, DAVIDGenes, DAVIDGenes, DAVIDTermCluster, DAVIDTermCluster, DAVIDTermCluster-class, as, as, as, ids, ids, ids, ids, ids, initialize, initialize, initialize, initialize, initialize, initialize, initialize, initialize

## Examples

```
{
##DAVIDFunctionalAnnotationChart example:
##Load the Functional Annotation Chart file report for the input demo
##file 2, using data function. Just to keep it simple, for the first five
##terms present in funChart2 object, create a DAVIDFunctionalAnnotationChart
##object and plot a 2D tile matrix with the reported evidence (green) or not
##(black).
data(funChart2)
plot2D(DAVIDFunctionalAnnotationChart(funChart2[1:5, ]),
color=c("FALSE"="black", "TRUE"="green"))

##DAVIDFunctionalAnnotationTable example
##Load the Functional Annotation Table file report for the input demo
##file 1, using data function. Then, create a DAVIDFunctionalAnnotationTable
##object using the loaded data.frame annotationTable1.
data(annotationTable1)
davidFunTable1<-DAVIDFunctionalAnnotationTable(annotationTable1)

##Plot the membership of only for the first six terms in this
##category, with only the genes of the first six terms with at least one
##evidence code.
##Category filtering...
categorySelection<-list(head(dictionary(davidFunTable1,
categories(davidFunTable1)[1])$ID))
names(categorySelection)<-categories(davidFunTable1)[1]

##Gene filter...
id<-membership(davidFunTable1, categories(davidFunTable1)[1])[,1:6]
id<-ids(genes(davidFunTable1))[rowSums(id)>0]

##Finally the membership tile plot
plot2D(davidFunTable1, category=categorySelection, id=id,
names.category=TRUE)

##DAVIDGeneCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo list 1 file to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/geneClusterReport1.tab.tar.gz",
```

```

package="RDAVIDWebService")
untar(fileName)
davidGeneCluster1<-DAVIDGeneCluster(untar(fileName, list=TRUE))

##We can inspect a 2D tile membership plot, to visual inspect for
##overlapping of genes across the clusters. Or use an scaled version of gene
##names to see the association of gene cluster, e.g., cluster 3 is related to
##ATP genes.
plot2D(davidGeneCluster1)
plot2D(davidGeneCluster1,names=TRUE)+
theme(axis.text.y=element_text(size=rel(0.9)))

##DAVIDTermCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo file 2 to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/termClusterReport2.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidTermCluster2<-DAVIDTermCluster(untar(fileName, list=TRUE))

##Finally, we can inspect a 2D tile membership plot, to visual inspect for
##overlapping of genes across the term members of the selected cluster,
##e.g., the first cluster .
plot2D(davidTermCluster2, number=1)
}

```

---

setEmail

*Accessor methods for DAVIDWebService class*


---

## Description

Setter/getters for DAVIDWebService class fields.

## Usage

```

setEmail(object, mail)

## S4 method for signature 'DAVIDWebService'
setEmail(object, mail)

## S4 method for signature 'character'
setEmail(mail)

getEmail(object)

## S4 method for signature 'DAVIDWebService'
getEmail(object)

getStub(object)

## S4 method for signature 'DAVIDWebService'
getStub(object)

```

**Arguments**

object	DAVIDWebService class object.
mail	character with a registered e-mail account at DAVID's website.

**Details**

Note that DAVIDWebService is a Reference class, hence invoke it using object\_name\$setter/getter(parameters). In addition, the user can use the S4 version style function call.

**Value**

according to the call one of the following objects can be returned

setEmail	character with the given e-mail to set.
getEmail	character with the e-mail under use.
getstub	jobjRef object with the stub java object to interface with DAVID API.

**References**

1. DAVID web <http://david.abcc.ncifcrf.gov>
2. DAVID API <http://david.abcc.ncifcrf.gov/content.jsp?file=WS.html>

**See Also**

Other DAVIDWebService: [DAVIDWebService-class](#), [addList](#), [addList](#), [connect](#), [connect](#), [getAllAnnotationCate](#), [getAllAnnotationCategoryNames](#), [getAnnotationSummary](#), [getAnnotationSummary](#), [getBackgroundListNames](#), [getBackgroundListNames](#), [getClusterReport](#), [getClusterReport](#), [getClusterReportFile](#), [getClusterReportFi](#), [getCurrentBackgroundListPosition](#), [getCurrentBackgroundListPosition](#), [getCurrentGeneListPosition](#), [getCurrentGeneListPosition](#), [getCurrentSpeciesPosition](#), [getCurrentSpeciesPosition](#), [getDefaultCategoryNames](#), [getDefaultCategoryNames](#), [getFunctionalAnnotationChart](#), [getFunctionalAnnota](#), [getFunctionalAnnotationChartFile](#), [getFunctionalAnnotationChartFile](#), [getFunctionalAnnotationTable](#), [getFunctionalAnnotationTable](#), [getFunctionalAnnotationTableFile](#), [getFunctionalAnnotationTableFile](#), [getGeneCategoriesReport](#), [getGeneCategoriesReport](#), [getGeneListNames](#), [getGeneListNames](#), [getGeneListReport](#), [getGeneListReport](#), [getGeneListReportFile](#), [getGeneListReportFile](#), [getIdTypes](#), [getIdTypes](#), [getListName](#), [getListName](#), [getSpecieNames](#), [getSpecieNames](#), [is.connected](#), [is.connected](#), [setAnnotationCategories](#), [setAnnotationCategories](#), [setCurrentBackgroundPosition](#), [setCurrentBackgroundPosition\(position\)](#), [setCurrentGeneListPosition](#), [setCurrentGeneListPosition](#), [setCurrentSpecies](#), [setCurrentSpecies](#), [summary](#), [summary](#), [summary](#), [summary](#)

**Examples**

```
{
##Create a DAVIDWebService object
david<-DAVIDWebService$new()

##Invoke Reference class style function setter/getters
david$setEmail("valid_mail@david.org")
david$getEmail()
stub<-david$getStub()

##Or the equivalent S4 style function call setter/getters
setEmail(david, "valid_mail@david.org")
getEmail(david)
```

```
stub<-getStub(david)
}
```

---

show

*Basic console output*

---

## Description

The different implementations of show function for the DAVIDWebService package classes.

## Usage

```
## S4 method for signature 'DAVIDResult'
show(object)

## S4 method for signature 'DAVIDGenes'
show(object)

## S4 method for signature 'DAVIDFunctionalAnnotationChart'
show(object)

## S4 method for signature 'DAVIDCluster'
show(object)

## S4 method for signature 'DAVIDFunctionalAnnotationTable'
show(object)

## S4 method for signature 'DAVIDWebService'
show(object)
```

## Arguments

object            DAVIDXX class members (where XX stands for Result, Genes, Term/GeneCluster, FunctionalAnnotationChart/Table or DAVIDWebService).

## Value

Basic console output.

## Author(s)

Cristobal Fresno and Elmer A Fernandez

## Examples

```
{
##DAVIDGenes example:
##Load Show Gene List file report for the input demo file 1, using data
##function. Then, create a DAVIDGenes object using only the head of the
##loaded data.frame geneList1 (just to keep it simple).
data(geneList1)
davidGenes1<-DAVIDGenes(head(geneList1))
davidGenes1
```

```

##DAVIDFunctionalAnnotationChart example
##Load the Functional Annotation Chart file report for the input demo
##file 2, using data function. Then, create a DAVIDFunctionalAnnotationChart
##object using the head of the loaded data.frame funChart2 (just to keep
##it simple).
data(funChart2)
davidFunChart2<-DAVIDFunctionalAnnotationChart(head(funChart2))
davidFunChart2

##DAVIDFunctionalAnnotationTable example:
##Load the Functional Annotation Table file report for the input demo
##file 1, using data function. Then, create a DAVIDFunctionalAnnotationTable
##object using the loaded data.frame annotationTable1.
data(annotationTable1)
davidFunTable1<-DAVIDFunctionalAnnotationTable(annotationTable1)
davidFunTable1
}

```

---

species

*Methods for DAVIDGenes class object*


---

### Description

Obtain DAVIDGenes related information, according to the given function call (see Values).

### Usage

```

species(object)

## S4 method for signature 'DAVIDGenes'
species(object)

duplicateIds(object, collapse = FALSE)

## S4 method for signature 'DAVIDGenes'
duplicateIds(object,
collapse=FALSE)

uniqueIds(object)

## S4 method for signature 'DAVIDGenes'
uniqueIds(object)

```

### Arguments

object	DAVIDGenes class object.
collapse	logical indicating if duplicate ids should be grouped as a comma separated id. Default value is FALSE.
...	Additional parameters for internal functions (if applicable).

**Value**

according to the call one of the following objects can be returned

show	console output of the class and associated data.
species	character vector with the levels of Species if available.
uniqueIds	a DAVIDGenes object with only the gene names with a unique id.
duplicateIds	a DAVIDGenes object with only the gene names with at least two ids. If collapse is TRUE, a data.frame in where all the ids that matched the same gene name, are coded in comma separated style.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

**Examples**

```
{
##Load Show Gene List file report for the input demo file 1, using data
##function. Then, create a DAVIDGenes object using the loaded data.frame
##geneList1. In addition, the user can use the file name of the downloaded
##file report.
data(geneList1)
davidGenes1<-DAVIDGenes(geneList1)

##Now we can inspect davidGenes1 as it was an common data.frame
head(davidGenes1)

##Additional getters for this object are also available, to obtain the
##different columns: ids, genes and species.
ids(davidGenes1)
genes(davidGenes1)
species(davidGenes1)

##Or even look up for a particular gene id, which will return only the
##matched ones.
genes(davidGenes1, ids=c("38926_at", "35367_at", "no match"))

##Obtain the genes with duplicate manufacturer ids or just the genes that
##do not have duplicate ids (uniqueIds).
duplicateIds(davidGenes1)
uniqueIds(davidGenes1)
}
```

---

subset

*Methods for DAVIDFunctionalAnnotationTable class object*


---

**Description**

Obtain DAVIDFunctionalAnnotationTable related information, according to the given function call (see Values).



**Usage**

```
subset(x, ...)

## S4 method for signature 'DAVIDFunctionalAnnotationTable'
subset(x,selection=c("Membership",
"Dictionary"), category, drop=TRUE)

dictionary(object, ...)

## S4 method for signature 'DAVIDFunctionalAnnotationTable'
dictionary(object,
...)

membership(object, ...)

## S4 method for signature 'DAVIDFunctionalAnnotationTable'
membership(object,
...)
```

**Arguments**

object, x	DAVIDFunctionalAnnotationTable class object.
selection	which slot to use to obtain the subset. Possible values are "Membership" or "Dictionary".
category	named list with main annotation category, which contains a character vector with the ids to use. Default value is missing in order to use all available categories of the report.
drop	Should list structure be drop if length==1? Default value TRUE.
...	Additional parameters for subset function call.

**Value**

according to the call one of the following objects can be returned

subset	list with filtered categories/ids according to function call.
enrichment	numeric vector with DAVID cluster's enrichment score.
members	list with DAVID Cluster's members.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

**See Also**

Other DAVIDCluster: [DAVIDCluster-class](#), [cluster](#), [cluster](#), [enrichment](#), [enrichment](#), [members](#), [members](#), [summary](#), [summary](#), [summary](#), [summary](#)

Other DAVIDFunctionalAnnotationTable: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable-class](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [categories](#), [categories](#), [categories](#), [genes](#)

genes, genes, genes, initialize, initialize, initialize, initialize, initialize, initialize, initialize, initialize, plot2D, plot2D, plot2D, plot2D, plot2D, plot2D

Other DAVIDFunctionalAnnotationTable: DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationChart, DAVIDFunctionalAnnotationTable, DAVIDFunctionalAnnotationTable, DAVIDFunctionalAnnotationTable, DAVIDFunctionalAnnotationTable-class, DAVIDGODag, DAVIDGODag, DAVIDGeneCluster, DAVIDGeneCluster, DAVIDGenes, DAVIDGenes, DAVIDGenes, DAVIDTermCluster, DAVIDTermCluster, as, as, as, categories, categories, categories, genes, genes, genes, initialize, initialize, initialize, initialize, initialize, initialize, initialize, initialize, plot2D, plot2D, plot2D, plot2D, plot2D, plot2D

## Examples

```
{
##Load the Functional Annotation Table file report for the input demo
##file 1, using data function. Then, create a DAVIDFunctionalAnnotationTable
##object using the loaded data.frame annotationTable1.
data(annotationTable1)
davidFunTable1<-DAVIDFunctionalAnnotationTable(annotationTable1)

##Obtain the head of the dictionary and the membership matrix for the first
##annotated genes used in davidFunTable1 object.
head(membership(davidFunTable1, categories(davidFunTable1)[1]))
head(dictionary(davidFunTable1, categories(davidFunTable1)[1]))
head(genes(davidFunTable1))
}
```

---

summary

*Basic summary for DAVIDWebService package classes.*

---

## Description

The different implementations of summary function for the DAVIDWebService package classes.

## Usage

```
summary(object, ...)

## S4 method for signature 'DAVIDCluster'
summary(object)

## S4 method for signature 'DAVIDGODag'
summary(object, ...)

## S4 method for signature 'DAVIDWebService'
summary(object)
```

## Arguments

object	DAVIDXX class members (where XX stands for Term/GeneCluster, GODag or DAVIDWebService).
...	Additional parameters.

**Value**

data.frame with summary output.

**Author(s)**

Cristobal Fresno and Elmer A Fernandez

**See Also**

Other DAVIDCluster: [DAVIDCluster-class](#), [cluster](#), [cluster](#), [dictionary](#), [dictionary](#), [enrichment](#), [enrichment](#), [members](#), [members](#), [membership](#), [membership](#), [subset](#), [subset](#)

Other DAVIDGODag: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGODag-class](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [benjamins](#), [benjamins](#), [bonferronis](#), [bonferronis](#), [counts](#), [counts](#), [fdrs](#), [fdrs](#), [foldEnrichments](#), [foldEnrichments](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [listTotals](#), [listTotals](#), [percentages](#), [percentages](#), [popHits](#), [popHits](#), [popTotals](#), [popTotals](#), [terms](#), [terms](#), [universeCounts](#), [universeMappedCount](#), [upsideDown](#), [upsideDown](#)

Other DAVIDWebService: [DAVIDWebService-class](#), [addList](#), [addList](#), [connect](#), [connect](#), [getAllAnnotationCategoryNames](#), [getAllAnnotationCategoryNames](#), [getAnnotationSummary](#), [getAnnotationSummary](#), [getBackgroundListNames](#), [getBackgroundListNames](#), [getClusterReport](#), [getClusterReport](#), [getClusterReportFile](#), [getClusterReportFile](#), [getCurrentBackgroundListPosition](#), [getCurrentBackgroundListPosition](#), [getCurrentGeneListPosition](#), [getCurrentGeneListPosition](#), [getCurrentSpeciesPosition](#), [getCurrentSpeciesPosition](#), [getDefaultCategoryNames](#), [getDefaultCategoryNames](#), [getEmail](#), [getEmail](#), [getFunctionalAnnotationChart](#), [getFunctionalAnnotationChart](#), [getFunctionalAnnotationChartFile](#), [getFunctionalAnnotationChartFile](#), [getFunctionalAnnotationTable](#), [getFunctionalAnnotationTable](#), [getFunctionalAnnotationTableFile](#), [getFunctionalAnnotationTableFile](#), [getGeneCategoriesReport](#), [getGeneCategoriesReport](#), [getGeneListNames](#), [getGeneListNames](#), [getGeneListReport](#), [getGeneListReport](#), [getGeneListReportFile](#), [getGeneListReportFile](#), [getIdTypes](#), [getIdTypes](#), [getListName](#), [getListName](#), [getSpecieNames](#), [getSpecieNames](#), [getStub](#), [getStub](#), [is.connected](#), [is.connected](#), [setAnnotationCategories](#), [setAnnotationCategories](#), [setCurrentBackgroundPosition](#), [setCurrentBackgroundPosition\(position\)](#), [setCurrentGeneListPosition](#), [setCurrentGeneListPosition](#), [setCurrentSpecies](#), [setCurrentSpecies](#), [setEmail](#), [setEmail](#), [setEmail](#), [DAVIDWebService-method](#)

**Examples**

```
{
##DAVIDGODag example:
##Load the Functional Annotation Chart file report for the input demo
##file 2, using data function. Then, create a DAVIDGODag object using
##Molecular Function main category of DAVIDFunctionalAnnotationChart object,
##obtained from the loaded data.frame funChart2. In addition, we have
##selected a threshold pvalue of 0.001 and removed unattached nodes, in case
##DAVID/GO.db database are not using the same version.
data(funChart2)
davidGODag<-DAVIDGODag(DAVIDFunctionalAnnotationChart(funChart2), type="MF",
pvalueCutoff=0.001, removeUnattached=TRUE)
summary(davidGODag)
```

```
##DAVIDGeneCluster example:
```

```

##Load the Gene Functional Classification Tool file report for the
##input demo list 1 file to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/geneClusterReport1.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidGeneCluster1<-DAVIDGeneCluster(untar(fileName, list=TRUE))
davidGeneCluster1

##Now we can invoke DAVIDCluster ancestor functions to inspect the report
##data, of each cluster. For example, we can call summary to get a general
##idea
summary(davidGeneCluster1)

##DAVIDTermCluster example:
##Load the Gene Functional Classification Tool file report for the
##input demo file 2 to create a DAVIDGeneCluster object.
setwd(tempdir())
fileName<-system.file("files/termClusterReport2.tab.tar.gz",
package="RDAVIDWebService")
untar(fileName)
davidTermCluster2<-DAVIDTermCluster(untar(fileName, list=TRUE))
davidTermCluster2

##Now we can invoke DAVIDCluster ancestor functions to inspect the report
##data, of each cluster. For example, we can call summary to get a general
##idea
summary(davidTermCluster2)
}

```

---

terms

*Methods for DAVIDGODag class object*


---

## Description

Obtain DAVIDGODag related information, according to the given function call (see Values).

## Usage

```

terms(x, ...)

## S4 method for signature 'DAVIDGODag'
terms(x, ...)

percentages(object)

## S4 method for signature 'DAVIDGODag'
percentages(object)

listTotals(object)

## S4 method for signature 'DAVIDGODag'

```

```
listTotals(object)

  popHits(object)

  ## S4 method for signature 'DAVIDGODag'
popHits(object)

  popTotals(object)

  ## S4 method for signature 'DAVIDGODag'
popTotals(object)

  foldEnrichments(object)

  ## S4 method for signature 'DAVIDGODag'
foldEnrichments(object)

  bonferronis(object)

  ## S4 method for signature 'DAVIDGODag'
bonferronis(object)

  benjaminis(object)

  ## S4 method for signature 'DAVIDGODag'
benjaminis(object)

  fdrs(object)

  ## S4 method for signature 'DAVIDGODag'
fdrs(object)

  counts(object, ...)

  ## S4 method for signature 'DAVIDGODag'
counts(object, ...)

  upsideDown(graph)

  ## S4 method for signature 'graph'
upsideDown(graph)

  ## S4 method for signature 'DAVIDGODag'
universeCounts(r)

  ## S4 method for signature 'DAVIDGODag'
universeMappedCount(r)
```

### Arguments

object, x, r	DAVIDGODag class object.
graph	a graph object with the GO DAG structure.

... Additional parameters (if required).

### Value

according to the call one of the following objects can be returned

`upsideDown` the same graph but the arcs with its directions in the other way around. Hence, plot layout would make upside down the graph.

`universeMappedCount`, `universeCounts`, `counts`  
modifications to the corresponding `GOstats/Category` library functions, to keep the same behavior for `DAVIDGODag` objects.

`fdrs`, `benjamini`, `bonferroni`  
Adjusted method specific p-values for the corresponding nodes/terms.

`terms` character vector with GO node names.

`popTotals`, `popHits`, `listTotals`  
integer vector with the number of ids, to use in the EASE score calculations, when building the 2x2 contingency table.

`percentages` numeric vector with the percentage of the gene list ids present in the term.

`foldEnrichments`  
numeric vector with the ratio of the two proportions for each node/term. For example, if 40/400 (i.e. 10%) of your input genes involved in "kinase activity" and the background information is 300/30000 genes (i.e. 1%) associating with "kinase activity", roughly  $10\%/1\%=10$  fold enrichment.

### Author(s)

Cristobal Fresno and Elmer A Fernandez

### See Also

Other `DAVIDGODag`: [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationChart](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDFunctionalAnnotationTable](#), [DAVIDGODag](#), [DAVIDGODag](#), [DAVIDGODag-class](#), [DAVIDGeneCluster](#), [DAVIDGeneCluster](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDGenes](#), [DAVIDTermCluster](#), [DAVIDTermCluster](#), [as](#), [as](#), [as](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [initialize](#), [summary](#), [summary](#), [summary](#), [summary](#)

### Examples

```
{
##Load the Functional Annotation Chart file report for the input demo
##file 2, using data function. Then, create a DAVIDGODag object using
##Molecular Function main category of DAVIDFunctionalAnnotationChart object,
##obtained from the loaded data.frame funChart2. In addition, we have
##selected a threshold pvalue of 0.001 and removed unattached nodes, in case
##DAVID/GO.db database are not using the same version.
data(funChart2)
davidGODag<-DAVIDGODag(DAVIDFunctionalAnnotationChart(funChart2), type="MF",
pvalueCutoff=0.001, removeUnattached=TRUE)

##Now, we can inspect the enrichment GO DAG using GOstats functionalities:
##counts, pvalues, sigCategories, universeCounts, geneMappedCount, etc.
##However, oddsRatios, expectedCounts and universeMappedCount are not
##available because these results are not available on DAVID's Functional
```



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