# Package 'psygenet2r'

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**Title** psygenet2r - An R package for querying PsyGeNET and to perform comorbidity studies in psychiatric disorders

**Version** 1.36.0

**Description** Package to retrieve data from PsyGeNET database (www.psygenet.org) and to perform comorbidity studies with PsyGeNET's and user's data.

**Depends** R (>= 3.4)

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LazyData true

**Imports** stringr, RCurl, igraph, ggplot2, reshape2, grid, parallel, biomaRt, BgeeDB, topGO, Biobase, labeling, GO.db

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DataGeNET.Psy-class Class DataGeNET.Psy

# Description

Class DataGeNET.Psy is the basic object use in psygenet2r package. It is the main data container to using the different functions to query PsyGeNET database and generate teir output. The constructors of this class are the functions psygenetGene and psygenetDisease.

#### **Slots**

type Character containing 'gene' of 'disease'. It is used to eprform the correct query to Psy-GeNET.

search Character containing 'single' of 'list'. It is used to eprform the correct query to Psy-GeNET.

database Character containing the name of the database that will be queried. It can take the values 'MODELS' to use Comparative Toxigenomics Database, data from mouse and rat; 'GAD' to use Genetic Association Database; 'CTD' to use Comparative Toxigenomics Database, data from human; 'PsyCUR' to use Psychiatric disorders Gene association manually curated; 'CURATED' to use Human, manually curated databases (PsyCUR and CTD); or 'ALL' to use all these databases.

enrichedPD 3

```
term Charcter with the term(s) to search into the database(s). qresult data.frame with the obtained result
```

# See Also

psygenetGene, psygenetDisease, DataGeNET.Psy-methods

Enrichment of a user's input (genes) in PsyGeNET's diseases.

# Description

Test the enrichment of a given gene list on Psychiatric Disorders from PsyGeNET.

### Usage

```
enrichedPD(gene, database = "ALL", verbose = FALSE, warnings = FALSE)
```

# Arguments

gene	Name or vector of names (that can be both code or uml) to specific genes from PsyGeNET.
database	Name of the database that will be queried. It can take the values 'psycur15' to use data validated by experts for first release of PsyGeNET; 'psycur16' to use data validated by experts for second release of PsyGeNET; or 'ALL' to use both databases. Default 'ALL'.
verbose	By default FALSE. Change it to TRUE to get a on-time log from the function.
warnings	By default TRUE. Change it to FALSE to not see the warnings.

# Value

A data. frame with the enricment at each Psychiatric Disorder

```
enrichedPD(c("ADCY2", "AKAP13", "ANK3"), "ALL")
```

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extract

Raw data from DataGeNET.Psy and JaccardIndexPsy.

# **Description**

Obtain the raw data from a PsyGeNET's query stored in a DataGeNET.Psy object or the raw data with all the Jaccard Index for the disease of interest of an JaccardIndexPsy object.

# Usage

```
extract(object, ...)
## S4 method for signature 'DataGeNET.Psy'
extract(object)
## S4 method for signature 'JaccardIndexPsy'
extract(object, order.cl = "pval", ...)
```

### **Arguments**

object Object of class DataGeNET.Psy or JaccardIndexPsy
... NO USED
order.cl Order resulting data.frame by the name of this column.

#### Value

A data. frame containing the raw result from PsyGeNET or a data. frame with the result Jaccard Index for each disease.

A data. frame containing the raw result from PsyGeNET

A data. frame with the result Jaccard Index for each disease.

# Methods (by class)

- DataGeNET.Psy: Extract function for DataGeNET.Psy
- JaccardIndexPsy: Extract function for JacardIndexPsy

```
data(qr)
extract(qr)[1:2, ] # get internat data.frame
## Not run:
#Being x an JaccardIndexPsy
extract(x)
## End(Not run)
```

extractSentences 5

extractSentences	Method to obtain the sentences that support a gene-disease association from a DataGeNET. Psy object.

# Description

PsyGeNET contains a list of sentences that support a gene-disease association from public literature. The internal table of a DataGeNET.Psy object contains this information. The method extractSentences allows to extract those sentences that support a gene-disease association given a DataGeNET.Psy object and a disorder of interest.

# Usage

```
extractSentences(object, disorder, verbose)
## S4 method for signature 'DataGeNET.Psy'
extractSentences(object, disorder, verbose = FALSE)
```

# **Arguments**

object Object of class DataGeNET.Psy.

disorder The disorder of interest. Only those sentences supporting a gene-disease asso-

ciation with this specific disorder will be extracted. Disorder must be provided by using "Disease Id" or "Disease Name". The "Disease Id" can be provided with or without the "uml" tag. Example of a "Disease Name": "schizophrenia"; Example of a "Disease Id": "umls:C0036341", that is equivalent of using

"C0036341".

verbose If set to TRUE informative messages are show.

### Value

A data frame showing the sentences.

### Methods (by class)

• DataGeNET.Psy: Get sentences or evidences

```
data(qr)
extractSentences(qr, "Depression")
```

6 geneAttrPlot

geneAttrPlot Ploting the relation between genes and disease-categories
--

# Description

Given a set of genes or a result of psygenetGene creates four types of plots showing the relation of the genes with the disease's category in psyGeNET.

# Usage

```
geneAttrPlot(x, type = "pie", ..., verbose = FALSE)
```

# Arguments

x	Vector of genes of interest of DataGeNET.Psy resulting of psyegnetDisease.
type	Type of the drawn chart. By default it is "pie". It can takes "pie" to plot a pie chart with the number of genes for each psychiatric category, "disease category" for visualizing a barplot with the total and specific number of genes for each psychiatric disorder, "evidence index" for a barplot showing for each psychiatric disorder the number of gene-disease associations according to the Evidence index and "gene" for visualizing a barplot with the total and specific number of diseases associated to each gene.
	(Check NOTE section) Passed to inner functions for different plots.
verbose	By default FALSE. Change it to TRUE to get a on-time log from the function.

# Value

A plot for a DataGeNET. Psy in terms of the panther-class.

# Note

The "Evidence Index" is gotten from PsyGeNET. For more information about it and its calculation, pease visit psygenet.org. Argument . . . can be filled with specific argument depending on the type of plot:

Argument	Color	Description
cuiBarColor	Purple	Determines the color of the bar for diseases
diseaseCategoryBarColor	Yellow	Determines the color of the bar for psychiatric categories
uniqueGenesBarColor	Orange	Determines the color of the bar for unique genes for a disease category
totalGenesBarColor	Blue	Determines the color of the bar for total genes for a disease category
	cuiBarColor diseaseCategoryBarColor uniqueGenesBarColor	cuiBarColor Purple diseaseCategoryBarColor Yellow uniqueGenesBarColor Orange

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getUMLs

Query PsyGeNET for given gene(s) and generates an DataGeNET.Psy

### **Description**

Given the name of one or multiple gene and retrives their information from PsyGeNET and creates an object of type DataGeNET.Psy.

# Usage

```
getUMLs(word, database = "ALL")
```

### **Arguments**

word

Disese to convert to UMLS using PsyGeNET database.

database

Name of the database that will be queried. It can take the values 'psycur15' to use data validated by experts for first release of PsyGeNET; 'psycur16' to use data validated by experts for second release of PsyGeNET; or 'ALL' to use both databases. Default 'ALL'.

#### Value

The corresponding UMLs for the input disease/s

### **Examples**

```
umls <- getUMLs( word = "depressive", database = "ALL" )</pre>
```

jaccardEstimation

Calculation of the Jaccard Index between ideseases

# **Description**

This function is able to calculate the Jacard Index between: 1. muliple disases, 2. a set og genes and multiple diseases, 3. a set of genes and multiple main psychiatric disorders and 4. multiple diseases and multiple main psychiatric disorders.

# Usage

```
jaccardEstimation(pDisease, sDisease, database = "ALL", nboot = 100,
    ncores = 1, verbose = FALSE)
```

Argument	S
----------	---

pDisease	vector of diseases, vector of genes, vector of main psychiatric disorder.
sDisease	vector of diseases, vector of genes, vector of main psychiatric disorder. Only necessary when comparing genes vs. diseases, genes vs. main psychiatric disorders or diseases vs. main psychiatric disorders. To compare multiple diseases only use pDisease.
database	Name of the database that will be queried. It can take the values 'psycur15' to use data validated by experts for first release of PsyGeNET; 'psycur16' to use data validated by experts for second release of PsyGeNET; or 'ALL' to use both databases.
nboot	Number of iterations sued to compute the pvalue associted to the calculated Jaccard Index (default 100).
ncores	Number of cores used to calculate the pvalue associated to the computed Jaccard Index (default 1).
verbose	By default FALSE. Change it to TRUE to get a on-time log from the function.

### **Details**

Warning: The main psychiatric disorders are understood as a single set of genes composed by the genes of all the diseases that the main psychiatric disorder cotains.

#### Value

An object of class JaccardIndexPsy with the computed calculation of the JaccardIndex.

# **Examples**

```
ji <- jaccardEstimation( c( "COMT", "CLOCK", "DRD3" ), "umls:C00005586", "ALL" )

JaccardIndexPsy-class Class JaccardIndexPsy</pre>
```

# Description

Class JaccardIndexPsy is theresult of the process to look for a Jaccard Index between muliple diseases in psygenet2r package.

### **Slots**

```
nit Number of iterations to calculate the estimated Jaccard index
type Slot to save type of query (disease-disease, gene-disease)
table data.frame containing the result table of Jaccard indexes
i1 [internal use] vector with names of first component
i2 [internal use] vector with names of second component
```

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# See Also

```
psygenet Gene, psygenet Disease, Jaccard Index Psy-methods \\
```

# Examples

```
ji <- jaccardEstimation( c( "COMT", "CLOCK", "DRD3" ), "umls:C0005586", "ALL" )</pre>
```

ndisease

Getter from DataGeNET.Psy.

# Description

Obtain the number of unique diseases in a DataGeNET.Psy.

# Usage

```
ndisease(object)
## S4 method for signature 'DataGeNET.Psy'
ndisease(object)
```

# Arguments

object

Object of class DataGeNET.Psy.

# Value

The number of unique diseases

# Methods (by class)

• DataGeNET.Psy: Get number of diseases

```
data(qr)
ndisease(qr)
```

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ngene

Getter from DataGeNET.Psy.

# **Description**

Obtain the number of unique genes in a DataGeNET.Psy.

# Usage

```
ngene(object)
## S4 method for signature 'DataGeNET.Psy'
ngene(object)
```

# **Arguments**

object

Object of class DataGeNET.Psy.

# Value

The number of unique genes

The number of unique genes

# Methods (by class)

• DataGeNET.Psy: Get number of genes

# **Examples**

```
data(qr)
ngene(qr)
```

pantherGraphic

Query PsyGeNET for given genes and creates a representation in base of their panther-class

# Description

Given a vector of genes of interest (or using a DataGeNET.Psy object), this function creates a representation of a the panther-class these genes belongs to.

# Usage

```
pantherGraphic(x, database = "ALL", evidenceIndex, verbose = FALSE)
```

#### **Arguments**

x Vector of genes of interest of DataGeNET.Psy resulting of psyegnetDisease.

database Name of the database that will be queried. It can take the values 'psycur15' to

use data validated by experts for first release of PsyGeNET; 'psycur16' to use data validated by experts for second release of PsyGeNET; or 'ALL' to use both

databases. Default 'ALL'.

evidenceIndex threshold to take into account a gene in the analysis

verbose By default FALSE. Change it to TRUE to get a on-time log from the function.

#### Value

A plot for a DataGeNET. Psy in terms of the panther-class.

### **Examples**

```
d.alch <- pantherGraphic( c( "COMT", "CLOCK", "DRD3" ), "ALL" )</pre>
```

plot, DataGeNET. Psy, ANY-method

Plots the content of a DataGeNET. Psy object

### **Description**

This functions llows to create a variety of plots for DataGeNEt.Psy and JaccardIndexPsy objects.

### Usage

```
## S4 method for signature 'DataGeNET.Psy,ANY'
plot(x, y,
  layout = igraph::layout.fruchterman.reingold, type = "GDA network",
  verbose = FALSE, ...)
```

#### **Arguments**

x Object of class DataGeNET.Psy

y NOT USED

layout Function to design the location of the different nodes. By default layout.fruchterman.reingold

from igraph is used.

type Type of the drawn chart. By default it is "GDA network" but it also can be

"GDCA network", "GDCA heatmap", "GDA heatmap" and "publications". The first two are network representations of the second two. While the last one draws

a barplot with the number of PMIDs between genes and diseases.

verbose By default FALSE. If set to TRUE information on the drawing process will be

shown.

... (Check NOTE section) Passed to inner functions for different plots.

# Value

A plot for DataGeNET.Psy.

#### Note

The "Evidence Index" is gotten from PsyGeNET. For more information about it and its calculation, pease visit psygenet.org. Argument . . . can be filled with specific argument depending on the type of plot:

Туре	Argument	Color	Description
GDA network	geneColor	Yellow	Determines the color of the gene nodes
	diseaseColor	Blue	Determines the color of the disease nodes
GDA heatmap	highColor	Blue	Determines the color of the heatmap for the highest value
_	lowColor	White	Determines the color of the heatmap for the lowest value
GDCA network	ei0color	Yellow	Determines the color for those associations with evidence index 0
	eiAmbiguitcolor	Grey	Determines the color for those associations with evidence index greater that
	ei1color	Blue	Determines the color for those associations with evidence index 1
GDCA heatmap	AUDcolor	#FF3C32	
_	BDcolor	#FFC698	
	DEPcolor	#9BE75E	
	SCHZcolor	#1F6024	
	CUDcolor	#5AB69C	
	SIDEPcolor	#50B8D6	

# **Examples**

```
data(qr)
plot(qr) # for all-disease plot
plot(qr, type = 'GDCA network') # for MPI plot
```

CanUDcolor

SYPSYcolor

geneColor

```
plot, JaccardIndexPsy, ANY-method
```

Plot the content of a JaccardIndexPsy object.

# Description

This functions llows to create a variety of plots for DataGeNEt.Psy and JaccardIndexPsy objects.

# Usage

```
## S4 method for signature 'JaccardIndexPsy,ANY'
plot(x, y, cutOff, zero.remove = TRUE,
   noTitle = FALSE, lowColor = "white", highColor = "mediumorchid4",
   verbose = FALSE, ...)
```

#5467C3

#A654C3

Orange

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### **Arguments**

x Object of class JaccardIndexPsy.

y NOT USED

cutOff Number to filter the shown results.

zero.remove By deffault TRUE. It removes those relations with a Jaccard Index of 0.

noTitle By default FALSE. If set to true no title will be added to the plot.

lowColor By default "white". It can be changed to any other color.

highColor By default "mediumorchid4". It can be changed to any other color.

verbose By default FALSE. If set to TRUE information on the drawing process will be

shown.

... NOT USED

#### Value

A plot for JaccardIndexPsy.

### **Examples**

```
## Not run:
#Being x an JaccardIndexPsy
qr <- plot(x)
## End(Not run)</pre>
```

psygenet2r

psygenet2r: Package to query PsyGeNET database and to perform

comorbidity studies

# **Description**

psygenet2r has two categories of functions: querying functions and analysis and plotting functions.

# querying functions

The functions to retrieve data from PsyGeNET are psygenetDisease and psygenetGene. There are some other support functions like psygenetGeneSentences.

### analysis and plotting functions

The functions extract and extractSentences allows to retrieve the row data obtained from on-line resources. The functions plot and pantherGraphic draws a variety of charts to illustrate the obtained results. The function enrichedPD was built to perform enrichment studies on PsyGeNET data. Finally the function jaccardEstimation computes a Jaccard Index from a given input on PsyGeNET data.

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psygenetDisease	•	PsyGeNET PNET.Psy	for	given	disease(s)	and	generates	an

# **Description**

Given the name of one or multiple diseases and retrives their information from PsyGeNET and creates an object of type DataGeNET.Psy.

### Usage

```
psygenetDisease(disease, database = "ALL", evidenceIndex = c(">", 0),
  verbose = FALSE, warnings = TRUE)
```

# **Arguments**

disease	Name or vector of names (that can be both code or uml) to specific diseases from
	D. C.NET Th. 1'

PsyGeNET. The diseases non existing in PsyGeNET will be removed from the

output.

database Name of the database that will be queried. It can take the values 'psycur15' to

use data validated by experts for first release of PsyGeNET; 'psycur16' to use data validated by experts for second release of PsyGeNET; or 'ALL' to use both

databases. Default 'ALL'.

evidenceIndex A vector with two elements: 1) character with greather '>' or with lower '<'

meaing greather or equal and lower or equal; 2) the evidence index cut-off to be

compared. By default: c('>', 0).

verbose By default FALSE. Change it to TRUE to get a on-time log from the function.

warnings By default TRUE. Change it to FALSE to don't see the warnings.

#### Value

An object of class DataGeNET.Psy

#### Note

The "Evidence Index" is gotten from PsyGeNET. For more information about it and its calculation, pease visit psygenet.org.

```
d.sch <- psygenetDisease( "schizophrenia", "ALL" )</pre>
```

psygenetDiseaseSentences

Query PsyGeNET for given disease(s) and extract the pmids sentences that report a gene-disease association.

# **Description**

Given a disease or a disease list, retrives the pmids and sentences for each gene-disease association from PsyGeNET and creates an object of type DataGeNET.Psy.

# Usage

```
psygenetDiseaseSentences(diseaseList, database = "ALL", verbose = FALSE)
```

### **Arguments**

diseaseList Name or vector of names (that can be both code or uml) to specific diseases from

PsyGeNET. The diseases non existing in PsyGeNET will be removed from the

output.

database Name of the database that will be queried. It can take the values 'psycur15' to

use data validated by experts for first release of PsyGeNET; 'psycur16' to use data validated by experts for second release of PsyGeNET; or 'ALL' to use both

databases. Default 'ALL'.

verbose By default FALSE. Change it to TRUE to get a on-time log from the function.

### Value

An object of class DataGeNET. Psy

# **Examples**

psygenetGene

Query PsyGeNET for given gene(s) and generates an DataGeNET. Psy

# Description

Given the name of one or multiple gene and retrives their information from PsyGeNET and creates an object of type DataGeNET.Psy.

### Usage

```
psygenetGene(gene, database = "ALL", evidenceIndex = c(">", 0),
  verbose = FALSE, warnings = TRUE)
```

### **Arguments**

gene Name or vector of names (that can be both code or symbol) to specific genes

from PsyGeNET. The genes non existing in PsyGeNET will be removed from

the output.

database Name of the database that will be queried. It can take the values 'psycur15' to

use data validated by experts for first release of PsyGeNET; 'psycur16' to use data validated by experts for second release of PsyGeNET; or 'ALL' to use both

databases. Default 'ALL'.

evidenceIndex A vector with two elements: 1) character with greather '>' or with lower '<'

meaing greather or equal and lower or equal; 2) the evidence index cut-off to be

compared. By default: c('>', 0).

verbose By default FALSE. Change it to TRUE to get a on-time log from the function.

warnings By default TRUE. Change it to FALSE to not see the warnings.

#### Value

An object of class DataGeNET. Psy

### Note

The "Evidence Index" is gotten from PsyGeNET. For more information about it and its calculation, pease visit psygenet.org.

#### **Examples**

```
d.alch <- psygenetGene( "ALDH2", "ALL" )</pre>
```

psygenetGeneSentences Query PsyGeNET for given gene(s) and extract the pmids sentences

that report a gene-disease association.

### **Description**

Given a gene or a gene list, retrives the pmids and sentences for each gene-disease association from PsyGeNET and creates an object of type DataGeNET.Psy.

# Usage

```
psygenetGeneSentences(geneList, database = "ALL", verbose = FALSE)
```

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# **Arguments**

geneList Name or vector of names (that can be both code or symbol) to specific genes

from PsyGeNET. The genes non existing in PsyGeNET will be removed from

the output.

database Name of the database that will be queried. It can take the values 'psycur15' to

use data validated by experts for first release of PsyGeNET; 'psycur16' to use data validated by experts for second release of PsyGeNET; or 'ALL' to use both

databases. Default 'ALL'.

verbose By default FALSE. Change it to TRUE to get a on-time log from the function.

#### Value

An object of class DataGeNET.Psy

### **Examples**

qr

DataGeNET. Psy obtained from quering PsyGeNET for gene '4852'.

### **Description**

A dataset obtained from PsyGeNET after being queried with psygenetGene usig the term '4852' on "ALL" database.

#### Usage

```
data("qr")
```

# **Format**

The format is: Formal class 'DataGeNET.Psy' [package "psygenet2r"] with 5 slots .. type : chr "gene" .. search : chr "" .. database: chr "ALL" .. term : chr "4852" .. qresult :'data.frame'

#### Value

A DataGeNET.Psy object.

### Source

```
http://psygenet.org
```

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# **Examples**

```
ngene(qr)
ndisease(qr)
```

topAnatEnrichment

Enrichment of a user's input (genes) in anatomical terms (TopAnat).

# Description

Test the enrichment of a given gene list on Psychiatric Disorders from PsyGeNET.

# Usage

```
topAnatEnrichment(gene, datatype = c("rna_seq", "affymetrix", "est",
   "in_situ"), statistic = "fisher", cutOff = 1, verbose = FALSE,
   warnings = FALSE)
```

# Arguments

gene	Name or vector of names (that can be both code or uml) to specific genes from PsyGeNET.
datatype	It can take the values 'rna_seq', 'affymetrix', "est" or "in situ". Default c("rna_seq", "affymetrix", "est", "in_situ").
statistic	By default it is "fisher". But it can be changed to "ks", "t", "globaltest", "sum" or "ks.ties". All from runTest.
cut0ff	Default 1.
verbose	By default FALSE. Change it to TRUE to get a on-time log from the function.
warnings	By default TRUE. Change it to FALSE to not see the warnings.

### Value

A data. frame with the enrichment results

```
## Not run:
topAnatEnrichment(gene=c("ADCY2", "AKAP13", "ANK3"))
## End(Not run)
```

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universe

Vector with gene universe for Jaccard Index

# Description

Vector with all the gene names from DisGeNET database (http://www.disgenet.org) used as gene universe for Jaccard Index computation.

# Usage

universe

### **Format**

An object of class character of length 8947.

# **Details**

```
data("universe", package = "psygenet2r")
```

### Value

A character vector.

# Source

```
http://www.disgenet.org
```

```
length(universe)
universe[1:10]
```

# **Index**

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