Package ‘psygenet2r’

May 29, 2024

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

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Suggests testthat, knitr, rmarkdown, BiocStyle

NeedsCompilation no

VignetteBuilder knitr

biocViews Software, BiomedicalInformatics, Genetics, Infrastructure, DataImport, DataRepresentation

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DataGeNET.Psy-class

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 their output. The constructors of this class are the functions psygenetGene and psygenetDisease.

Slots

type Character containing 'gene' of 'disease'. It is used to perform the correct query to PsyGeNET.

search Character containing 'single' of 'list'. It is used to perform the correct query to PsyGeNET.

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

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

See Also
psygenetGene, psygenetDisease, DataGeNET.Psy-methods

---

enrichedPD  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

Examples
enrichedPD(c("ADCY2", "AKAP13", "ANK3"), "ALL")
**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 a JaccardIndexPsy object.

**Usage**

```r
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 JaccardIndexPsy

**Examples**

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

## End(Not run)
```
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

```r
extractSentences(object, disorder, verbose)
```

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

Arguments

- **object**: Object of class DataGeNET.Psy.
- **disorder**: The disorder of interest. Only those sentences supporting a gene-disease association 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

Examples

```r
data(qr)
eachSentences(qr, "Depression")
```
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, please visit psygenet.org. Argument ... can be filled with specific argument depending on the type of plot:

<table>
<thead>
<tr>
<th>Type</th>
<th>Argument</th>
<th>Color</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>gene</td>
<td>cuiBarColor</td>
<td>Purple</td>
<td>Determines the color of the bar for diseases</td>
</tr>
<tr>
<td></td>
<td>diseaseCategoryBarColor</td>
<td>Yellow</td>
<td>Determines the color of the bar for psychiatric categories</td>
</tr>
<tr>
<td>disease category</td>
<td>uniqueGenesBarColor</td>
<td>Orange</td>
<td>Determines the color of the bar for unique genes for a disease category</td>
</tr>
<tr>
<td></td>
<td>totalGenesBarColor</td>
<td>Blue</td>
<td>Determines the color of the bar for total genes for a disease category</td>
</tr>
</tbody>
</table>
getUMLs

---

**getUMLs**

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

**Description**

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

**Usage**

getUMLs(word, database = "ALL")

**Arguments**

<table>
<thead>
<tr>
<th>Argument</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>word</td>
<td>Disease to convert to UMLS using PsyGeNET database.</td>
</tr>
<tr>
<td>database</td>
<td>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'.</td>
</tr>
</tbody>
</table>

**Value**

The corresponding UMLs for the input disease/s

**Examples**

```r
umls <- getUMLs( word = "depressive", database = "ALL" )
```

---

**jaccardEstimation**

Calculation of the Jaccard Index between diseases

**Description**

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

**Usage**

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

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 contains.

Value

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

Examples

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

Description

Class JaccardIndexPsy is the result of the process to look for a Jaccard Index between multiple 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
See Also

psygenetGene, psygenetDisease, JaccardIndexPsy-methods

Examples

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

data(qr)
ndisease(qr)
ngene  

### Description

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

### Usage

```r
ngene(object)
```

### Arguments

- `object`: Object of class DataGeNET.Psy.

### Value

The number of unique genes

### Methods (by class)

- DataGeNET.Psy: Get number of genes

### Examples

```r
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 the panther-class these genes belongs to.

### Usage

```r
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

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

Description

This function allows to create a variety of plots for DataGeNET.Psy and JaccardIndexPsy objects.

Usage

```r
## 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, please visit psygenet.org. Argument ... can be filled with specific argument depending on the type of plot:

<table>
<thead>
<tr>
<th>Type</th>
<th>Argument</th>
<th>Color</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>GDA network</td>
<td>geneColor</td>
<td>Yellow</td>
<td>Determines the color of the gene nodes</td>
</tr>
<tr>
<td></td>
<td>diseaseColor</td>
<td>Blue</td>
<td>Determines the color of the disease nodes</td>
</tr>
<tr>
<td>GDA heatmap</td>
<td>highColor</td>
<td>Blue</td>
<td>Determines the color of the heatmap for the highest value</td>
</tr>
<tr>
<td></td>
<td>lowColor</td>
<td>White</td>
<td>Determines the color of the heatmap for the lowest value</td>
</tr>
<tr>
<td>GDCA network</td>
<td>ei0color</td>
<td>Yellow</td>
<td>Determines the color for those associations with evidence index 0</td>
</tr>
<tr>
<td></td>
<td>eiAmbiguitycolor</td>
<td>Grey</td>
<td>Determines the color for those associations with evidence index greater than 0</td>
</tr>
<tr>
<td></td>
<td>ei1color</td>
<td>Blue</td>
<td>Determines the color for those associations with evidence index 1</td>
</tr>
<tr>
<td>GDCA heatmap</td>
<td>AUDcolor</td>
<td>#FF3C32</td>
<td></td>
</tr>
<tr>
<td></td>
<td>BDcolor</td>
<td>#FFC698</td>
<td></td>
</tr>
<tr>
<td></td>
<td>DEPcolor</td>
<td>#9BE75E</td>
<td></td>
</tr>
<tr>
<td></td>
<td>SCHZcolor</td>
<td>#1F6024</td>
<td></td>
</tr>
<tr>
<td></td>
<td>CUDcolor</td>
<td>#5AB69C</td>
<td></td>
</tr>
<tr>
<td></td>
<td>SIDEPcolor</td>
<td>#50B8D6</td>
<td></td>
</tr>
<tr>
<td></td>
<td>CanUDcolor</td>
<td>#5467C3</td>
<td></td>
</tr>
<tr>
<td></td>
<td>SYPSYcolor</td>
<td>#A654C3</td>
<td></td>
</tr>
<tr>
<td>geneColor</td>
<td>Orange</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Examples

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

Description

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

Usage

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

- **x**: Object of class JaccardIndexPsy.
- **y**: NOT USED
- **cutOff**: Number to filter the shown results.
- **zero.remove**: By default 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.

Value

A plot for JaccardIndexPsy.

Examples

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

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.
psygenetDisease

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

Description
Given the name of one or multiple diseases and retrieves 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
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 '<'
meaning greater 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,
please visit psygenet.org.

Examples
d.sch <- psygenetDisease("schizophrenia", "ALL")
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, retrieves the pmids and sentences for each gene-disease association from PsyGeNET and creates an object of type `DataGeNET.Psy`.

**Usage**

```r
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**

```r
diseasesOfInterest <- c("Bipolar Disorder","Depressive Disorder, Major")
psyDisSen <- psygenetDiseaseSentences(diseaseList = diseasesOfInterest, database = "ALL")
```

psygenetGene

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

**Description**

Given the name of one or multiple gene and retrieves 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 '<' meaning greater 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, please visit psygenet.org.

Examples

d.alch <- psygenetGene( "ALDH2", "ALL" )

---------------------------------------------------------------------
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, retrieves 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)
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

genesOfInterest <- c("PECR", "ADH1C", "CAST", "ERAP1", "PPP2R2B", "ESR1", "GATA4", "CDH13")
psyGeneSen <- psygenetGeneSentences( geneList = genesOfInterest, database = "ALL")

---

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

Description

A dataset obtained from PsyGeNET after being queried with psygenetGene using 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
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.
cutOff 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

Examples

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

## End(Not run)
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

Examples

length(universe)
universe[1:10]
Index

* datasets
  qr, 17
  universe, 19
DataGeNET.Psy-class, 2
DataGeNET.Psy-plot
  (plot,DataGeNET.Psy,ANY-method), 11
enrichedPD, 3, 13
extract, 4, 13
extract,DataGeNET.Psy-method (extract), 4
eXtract,JaccardIndexPsy-method
  (extract), 4
extractSentences, 5, 13
extractSentences,DataGeNET.Psy-method
  (extractSentences), 5
geneAttrPlot, 6
getUMLs, 7
jaccardEstimation, 7, 13
JaccardIndexPsy-class, 8
JaccardIndexPsy-plot
  (plot,JaccardIndexPsy,ANY-method), 12
ndisease, 9
ndisease,DataGeNET.Psy-method
  (ndisease), 9
ngene, 10
ngene,DataGeNET.Psy-method (ngene), 10
pantherGraphic, 10, 13
plot,DataGeNET.Psy,ANY-method, 11
plot,JaccardIndexPsy,ANY-method, 12
psygenet2r, 13
psygenet2r-package (psygenet2r), 13
psygenetDisease, 13, 14
psygenetDiseaseSentences, 15
psygenetGene, 13, 15
psygenetGeneSentences, 13, 16
qr, 17
runTest, 18
topAnatEnrichment, 18
universe, 19