Package ‘paxtoolsr’

February 27, 2024

<table>
<thead>
<tr>
<th>Type</th>
<th>Package</th>
</tr>
</thead>
<tbody>
<tr>
<td>Title</td>
<td>Access Pathways from Multiple Databases Through BioPAX and Pathway Commons</td>
</tr>
<tr>
<td>Version</td>
<td>1.36.0</td>
</tr>
<tr>
<td>Date</td>
<td>2021-10-25</td>
</tr>
<tr>
<td>Imports</td>
<td>utils, httr, igraph, plyr, rjson, R.utils, jsonlite, readr, rappdirs</td>
</tr>
<tr>
<td>Depends</td>
<td>R (&gt;= 3.2), rJava (&gt;= 0.9-8), methods, XML</td>
</tr>
<tr>
<td>Suggests</td>
<td>testthat, knitr, BiocStyle, formatR, rmarkdown, RColorBrewer, foreach, doSNOW, parallel, org.Hs.eg.db, clusterProfiler</td>
</tr>
<tr>
<td>SystemRequirements</td>
<td>Java (&gt;= 1.6)</td>
</tr>
<tr>
<td>License</td>
<td>LGPL-3</td>
</tr>
<tr>
<td>Description</td>
<td>The package provides a set of R functions for interacting with BioPAX OWL files using Paxtools and the querying Pathway Commons (PC) molecular interaction database. Pathway Commons is a project by the Memorial Sloan-Kettering Cancer Center (MSKCC), Dana-Farber Cancer Institute (DFCI), and the University of Toronto. Pathway Commons databases include: BIND, BioGRID, CORUM, CTD, DIP, DrugBank, HPRD, HumanCyc, IntAct, KEGG, MirTarBase, Panther, PhosphoSitePlus, Reactome, RECON, TRANSFAC.</td>
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</tbody>
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| VignetteBuilder | knitr |
| LazyData        | false |
| biocViews       | GeneSetEnrichment, GraphAndNetwork, Pathways, Software, SystemsBiology, NetworkEnrichment, Network, Reactome, KEGG |
| URL             | https://github.com/BioPAX/paxtoolsr |
| Encoding        | UTF-8 |
| RoxygenNote     | 7.1.1 |
| git_url         | https://git.bioconductor.org/packages/paxtoolsr |
| git_branch      | RELEASE_3_18 |
| git_last_commit | efd04f2 |
| git_last_commit_date | 2023-10-24 |
Repository  Bioconductor 3.18
Date/Publication  2024-02-26
Author  Augustin Luna [aut, cre]
Maintainer  Augustin Luna <lunaa@cbio.mskcc.org>

R topics documented:

addAttributeList .................................................. 3
convertDataFrameListsToVectors ................................. 4
convertSifToGmt ................................................... 4
downloadFile ....................................................... 5
downloadPc2 ......................................................... 6
downloadSignedPc .................................................. 7
fetch ................................................................. 8
filterSif ............................................................. 8
getcacheFiles ....................................................... 10
getErrorMessage .................................................. 11
getNeighbors ....................................................... 11
getPc ................................................................. 12
getPcDatabaseNames ............................................. 13
getPcUrl ............................................................. 14
getShortestPathSif ............................................... 14
getSifInteractionCategories ...................................... 15
graphPc .............................................................. 16
integrateBiopax ..................................................... 17
loadSifInIgraph ................................................... 18
mapAttributes ....................................................... 18
mapValues ........................................................... 19
mergeBiopax .......................................................... 20
pcDirections ......................................................... 21
pcFormats ............................................................ 21
pcGraphQueries ..................................................... 22
processPcRequest .................................................. 23
readBiopax ........................................................... 23
readGmt ............................................................... 24
readPcPathwaysInfo ............................................... 25
readSbgn ............................................................. 25
readSif ............................................................... 26
readSifnx ............................................................. 26
searchListofVectors ............................................... 27
searchPc ............................................................. 28
splitSifnxByPathway ............................................... 30
summarize ........................................................... 30
summarizeSif ........................................................ 31
toCytoscape ........................................................ 32
toGSEA ............................................................... 32
toLevel3 ............................................................. 33
addAttributeList

Add attributes using a list of vectors to an igraph object

Description

Add attributes using a list of vectors to an igraph object

Usage

addAttributeList(g, attr, l)

Arguments

g an igraph object

tag the name of the attribute

l the list of vectors

Value

the modified igraph object

Examples

library(igraph)
g <- barabasi.game(20)
g <- set_vertex_attr(g, "name", value=LETTERS[1:20])
g <- addAttributeList(g, "isProt",
    list(A=TRUE, B=FALSE, C=TRUE, D=TRUE, E=FALSE))
**convertDataFrameListsToVectors**

*Convert columns with list in data.frame to vector*

**Description**

Convert columns with list in data.frame to vector

**Usage**

convertDataFrameListsToVectors(df, delimiter = ";")

**Arguments**

- **df**: a data.frame
- **delimiter**: a delimiter to concatenate (DEFAULT: ;)

**Value**

a data.frame without list columns

**Note**

Lists as columns are useful programmatically, but cause issue in writing output to text-based files

**Examples**

```
df <- data.frame(id = 1:2, name = c("Jon", "Mark"),
                 children = I(list(c("Mary", "James"), c("Greta", "Sally"))))
df <- convertDataFrameListsToVectors(df)
```

**convertSifToGmt**

*Convert SIF to GMT*

**Description**

Convert SIF to GMT

**Usage**

convertSifToGmt(sif, name = "gmt", returnSmallMolecules = FALSE)
Arguments

- **sif**: a data.frame representing a SIF (Simple Interaction Format)
- **name**: the name of the gene set
- **returnSmallMolecules**: a boolean whether to return genes or small molecules in the gene set

Value

- a list with one entry being a vector

Examples

```r
sif <- readSif(system.file("extdata", "test_sif.txt", package="paxtoolsr"))
gmt <- convertSifToGmt(sif)
```

---

### downloadFile

#### Description

Check Cache and Download File

#### Usage

```r
downloadFile(
  baseUrl,
  fileName,
  destDir = NULL,
  cacheEnv = "PAXTOOLSR_CACHE",
  verbose = FALSE
)
```

#### Arguments

- **baseUrl**: a string, entire download URL except filename
- **fileName**: a string, the filename of file to be downloaded
- **destDir**: a string, the path where a file should be saved
- **cacheEnv**: a string, environment variable pointing to specific cache
- **verbose**: show debugging information

#### Details

Description of file formats: [http://www.pathwaycommons.org/pc2/formats](http://www.pathwaycommons.org/pc2/formats)
downloadPc2

**Value**

a boolean TRUE if the file was downloaded or already exists, FALSE otherwise

**See Also**

readSif, readBiopax, readSbgn, readSifnx, readGmt

**Examples**

downloadFile("http://google.com/", fileName="index.html", destDir=tempdir())

downloadPc2

*Download Pathway Commons files (uses menu and cache)*

**Description**

Download Pathway Commons files (uses menu and cache)

**Usage**

downloadPc2(
  selectedFileName = NULL,
  destDir = NULL,
  returnNames = NULL,
  version,
  verbose = FALSE,
  ...
)

**Arguments**

- **selectedFileName**
  a string, a name of a file to skip the interactive selection
- **destDir**
  a string, the destination directory for the file to be downloaded (Default: NULL).
  If NULL, then file will be downloaded to cache directory at Sys.getenv("PAXTOOLS_CACHE")
- **returnNames**
  return a vector of names matching the given regular expression
- **version**
  a version number for a previous version of Pathway Commons data; versions 3 and above. Parameter set as version="8". Available versions "http://www.pathwaycommons.org/archives/PC2/"
- **verbose**
  a flag to display debugging information (Default: FALSE)
- **...**
  additional parameters to send to corresponding read* methods

**Value**

an R object using one of the read* methods provided in this package corresponding to the file downloaded
downloadSignedPC

## Not run:
```r
downloadPc2(version="8")
downloadPc2(version="8", returnNames="ext.*sif")
downloadPc2("PathwayCommons.8.inoh.GSEA.hgnc.gmt.gz", version="8", verbose=TRUE)
```

## End(Not run)

### Description

Download a SIF file containing only signed interactions

### Usage

```r
downloadSignedPC(destDir = NULL, forceCache = FALSE)
```

### Arguments

- `destDir`: a string, the destination directory for the file to be downloaded (Default: NULL). If NULL, then file will be downloaded to cache directory at `Sys.getenv("PAXTOOLSR_CACHE")`
- `forceCache`: a boolean to force the use of a cached version (DEFAULT: FALSE); the current host of the file (GitHub) does not support the LAST-MODIFIED header

### Value

a SIF containing interactions that are considered signed (i.e. interactions causing an increase on decrease in a molecular species)

### Examples

```r
# downloadSignedPC()
```
fetch

*Fetch a set of IDs from a BioPAX OWL file*

**Description**

This function will create a subsetted object with specified URIs.

**Usage**

```
fetch(inputFile, outputFile = NULL, idList)
```

**Arguments**

- `inputFile`: a string of the name of the input BioPAX OWL file
- `outputFile`: a string with the name of the output BioPAX OWL file
- `idList`: a vector of IDs from the BioPAX OWL file

**Details**

Only entities in the input BioPAX file will be used in the fetch. IDs used must be URIs for the entities of interest. Additional properties such as cross-references for fetched entities will be included in the output.

**Value**

an XMLInternalDocument representing a BioPAX OWL file

**Examples**

```
outFile <- tempfile()
ids <- c("http://identifiers.org/uniprot/P36894",
          "http://identifiers.org/uniprot/Q13873")
results <- fetch(system.file("extdata", "REACT_12034-3.owl", package="paxtoolsr"),
                 outFile, ids)
```

filterSif

*Keep interactions in SIF network based on certain criteria*

**Description**

Keep interactions in SIF network based on certain criteria
filterSif

Usage

filterSif(
  sif,
  ids = NULL,
  interactionTypes = NULL,
  dataSources = NULL,
  interactionPubmedIds = NULL,
  pathwayNames = NULL,
  mediatorIds = NULL,
  edgelist = NULL,
  idsBothParticipants = FALSE,
  edgelistCheckReverse = TRUE,
  verbose = FALSE
)

Arguments

sif a binary SIF as a data.frame with three columns: "PARTICIPANT_A", "INTERACTION_TYPE", "PARTICIPANT_B"
ids a vector of IDs to be kept
interactionTypes a vector of interaction types to be kept (List of interaction types: http://www.pathwaycommons.org/pc2/formats)
dataSources a vector of data sources to be kept. For Extended SIF.
interactionPubmedIds a vector of Pubmed IDs to be kept. For Extended SIF.
pathwayNames a vector of pathway names to be kept. For Extended SIF.
mediatorIds a vector of mediator IDs to be kept. For Extended SIF. Mediator IDs are the full BioPAX objects that were simplified to interaction given in the SIF. For Extended SIF.
edgelist a two-column data.frame where each row is an interaction to be kept. Directionality is ignored (e.g. Edge A B will return interactions A B and B A from SIF)
idsBothParticipants a boolean whether both interaction participants should be in a given interaction when using the ids parameter; TRUE if both (DEFAULT: TRUE)
edgelistCheckReverse a boolean whether to check for edges in the reverse order (DEFAULT: TRUE)
verbose Show debugging information (DEFAULT: FALSE)

Value

filtered interactions with three columns: "PARTICIPANT_A", "INTERACTION_TYPE", "PARTICIPANT_B". The intersection of multiple filters is returned.
Examples

```r
results <- readSif(system.file("extdata", "test_sif.txt", package="paxtoolsr"))
intTypes <- c("controls-state-change-of", "controls-expression-of", "catalysis-precedes")
filteredNetwork <- filterSif(results, intTypes)

tmp <- readSifnx(system.file("extdata", "test_sifnx_250.txt", package="paxtoolsr"))
results <- filterSif(tmp$edges, ids=c("CHEBI:17640", "MCM3"))
results <- filterSif(tmp$edges, dataSources=c("INOH", "KEGG"))
results <- filterSif(tmp$edges, dataSources=c("IntAct"), ids=c("CHEBI:17640", "MCM3"))
results <- filterSif(tmp$edges, pathwayNames=c("Metabolic pathways"))
results <- filterSif(tmp$edges,
mediatorIds=c("http://purl.org/pc2/8/MolecularInteraction_1452626895158"))
results <- filterSif(tmp$edges, interactionPubmedId="17654400")

tmp <- readSifnx(system.file("extdata", "test_sifnx_250.txt", package="paxtoolsr"))
edgelist <- read.table(system.file("extdata", "test_edgelist.txt", package="paxtoolsr"),
sep="\t", header=FALSE, stringsAsFactors=FALSE)
results <- filterSif(tmp$edges, edgelist=edgelist)
```

getCacheFiles

List files in cache directory

Description

List files in cache directory

Usage

getCacheFiles()

Value

a vector of the files in the cache directory

Examples

getCacheFiles()
getErrorMessage  
*Get Error Message for a Pathway Commons Error*

**Description**
Get Error Message for a Pathway Commons Error

**Usage**
getErrorMessage(code)

**Arguments**
code  
a three digit numerical error code

**Value**
an error message for the code

**Examples**
```
results <- getErrorMessage("452")
```

getNeighbors  
*Get the neighbors of a set of IDs in a BioPAX file*

**Description**
This function retrieves a set of neighbors for a set of IDs in a BioPAX file.

**Usage**
getNeighbors(inputFile, outputFile = NULL, idList)

**Arguments**
inputFile  
a string with the name of the input BioPAX OWL file
outputFile  
a string with the name of the output BioPAX OWL file
idList  
a vector of IDs from the BioPAX OWL file

**Details**
Only entities in the input BioPAX file will be searched for neighbors. IDs used must be URIs for the entities of interest.
Value

an XMLInternalDocument representing a BioPAX OWL file

Examples

```r
outFile <- tempfile()
results <- getNeighbors(system.file("extdata",
  "raf_map_kinase_cascade_reactome.owl", package="paxtoolsr"),
  outFile,
  c("HTTP://WWW.REACTOME.ORG/BIOPAX/48887#PROTEIN2360_1_9606",
    "HTTP://WWW.REACTOME.ORG/BIOPAX/48887#PROTEIN1631_1_9606"))
```

getPc  

*Get Pathway Commons BioPAX elements*

Description

This command retrieves full pathway information for a set of elements such as pathway, interaction or physical entity given the RDF IDs.

Usage

```r
getcP(uri, format = "BIOPAX", verbose = FALSE, ...)
```

Arguments

- **uri**: a vector that includes valid/existing BioPAX element’s URI (RDF ID; for utility classes that were "normalized", such as entity references and controlled vocabularies, it is usually a Identifiers.org URL. Multiple IDs are allowed per query, for example, c("http://identifiers.org/uniprot/Q06609", "http://identifiers.org/uniprot/Q549Z0") See also about MIRIAM and Identifiers.org in details.
- **format**: output format (Default: BIOPAX). Valid options can be found using pcFormats
- **verbose**: a boolean, display the command used to query Pathway Commons
- **...**: additional arguments to read* methods that handle data from Pathway Commons

Details

Get commands only retrieve the BioPAX elements that are directly mapped to the ID. Use the "traverse query to traverse BioPAX graph and obtain child/owner elements.

Information on MIRIAM and Identifiers.org [http://www.pathwaycommons.org/pc2/#miriam](http://www.pathwaycommons.org/pc2/#miriam)

Value

a XMLInternalDocument object
getPcDatabaseNames

See Also

pcFormats

Examples

uri <- "http://identifiers.org/uniprot/O14503"
#results <- getPc(uri)

uri <- c("http://identifiers.org/uniprot/O14503", "http://identifiers.org/uniprot/Q9P2X7")
#results <- getPc(uri, verbose=TRUE)

getPcDatabaseNames Get a Pathway Commons Databases

Description

Get a Pathway Commons Databases

Usage

getPcDatabaseNames(version)

Arguments

version PC2 version

Value

a names of databases that can be used as part of queries

Examples

getPcDatabaseNames(version=10)
getPcUrl  
*Get base Pathway Commons URL*

**Description**

Get base Pathway Commons URL

**Usage**

getPcUrl()

**Details**

paxtoolsr will support versions Pathway Commons 5 and later. Old versions of the webservice will not be operational. Users can parse older BioPAX outputs as an alternative.

**Value**

a string with base Pathway Commons URL

**Examples**

```r
url <- getPcUrl()
```

---

getShortestPathSif  
*Get the shortest between two IDs (HGNC or CHEBI)*

**Description**

Get the shortest between two IDs (HGNC or CHEBI)

**Usage**

```r
getShortestPathSif(
  sif,
  idA,
  idB,
  mode = c("all", "out", "in"),
  weights = NULL,
  verbose = FALSE,
  filterFun, ...
)
```
getSifInteractionCategories

Arguments

- `sif` a SIF network
- `idA` HGNC or CHEBI (CHEBI:XXXXX) ID
- `idB` HGNC or CHEBI (CHEBI:XXXXX) ID
- `mode` see shortest_paths() in igraph
- `weights` see shortest_paths() in igraph
- `verbose` a boolean whether to show debugging information
- `filterFun` a function to filter multiple paths of the same length
- `...` additional arguments passed on to filterFun

Value

a data.frame representing a SIF network

Examples

```r
idA <- "DAP3"
idB <- "RPS16"
sif <- readSif(system.file("extdata", "test_sif_shortestPath.txt", package="paxtoolsr"))
filterFun <- function(vpaths) { idx <- sample(1:length(vpaths), 1); return(vpaths[[idx]]) }
m1 <- getShortestPathSif(sif, idA, idB, mode="all", weights=NULL, filterFun=filterFun, verbose=TRUE)
```

getSifInteractionCategories

*Get a list of categories of SIF interactions*

Description

Get a list of categories of SIF interactions

Usage

getSifInteractionCategories()

Details

Description of interaction types: http://www.pathwaycommons.org/pc2/formats Categories provided: BetweenProteins, BetweenProteinsOther (often from high-throughput experiments), BetweenProteinSmallMolecule, BetweenSmallMolecules, SignedInteractions

Value

a list of interactions in categories
Examples

```r
sifCat <- getSifInteractionCategories()
sifCat[["BetweenProteins"]]
```

### Description

This function will retrieve a set of BioPAX elements given a graph query match.

### Usage

```r
graphPc(
  kind,
  source,
  target = NULL,
  direction = NULL,
  limit = NULL,
  format = NULL,
  datasource = NULL,
  organism = NULL,
  verbose = FALSE
)
```

### Arguments

- **kind**: graph query. Valid options can be found using `pcGraphQueries` See Details for information on graph queries.
- **source**: source object’s URI/ID. Multiple source URIs/IDs are allowed per query, for example c("http://identifiers.org/uniprot/Q06609", "http://identifiers.org/uniprot/Q549Z0") See a note about MIRIAM and Identifiers.org in details.
- **target**: [Required for PATHSFROMTO graph query] target URI/ID. Multiple target URIs are allowed per query; for example c("http://identifiers.org/uniprot/Q06609", "http://identifiers.org/uniprot/Q549Z0") See a note about MIRIAM and Identifiers.org in details.
- **direction**: [Optional, for NEIGHBORHOOD and COMMONSTREAM algorithms] - graph search direction. Valid options: `pcDirections`.
- **limit**: graph query search distance limit (default: 1).
- **format**: output format. Valid options: `pcFormats`.
- **datasource**: datasource filter (same as for 'search').
- **organism**: organism filter (same as for 'search').
- **verbose**: a boolean, display the command used to query Pathway Commons.
integrateBiopax

Value
depending on the the output format a different object may be returned. pcFormats

See Also
pcFormats, pcDirections

Examples
source <- "http://identifiers.org/uniprot/O14503"
#results <- graphPc(source=source, kind="neighborhood", format="TXT")

integrateBiopax Integrate two BioPAX OWL files (DEPRECATED)

Description
This function merges two BioPAX OWL files

Usage
integrateBiopax(inputFile1, inputFile2, outputFile = NULL)

Arguments
inputFile1 a string of the name of the input BioPAX OWL file
inputFile2 a string of the name of the input BioPAX OWL file
outputFile a string of the name of the output integrated BioPAX OWL file

Details
This method is deprecated. Use mergeBiopax instead.

Value
an XMLInternalDocument representing a BioPAX OWL file

See Also
mergeBiopax

Examples
outFile <- tempfile()
results <- integrateBiopax(system.file("extdata", "raf_map_kinase_cascade_reactome.owl",
 package="paxtoolsr"),
 system.file("extdata", "dna_replication.owl", package="paxtoolsr"),
 outFile)
loadSifInIgraph  \hspace{2cm} Load SIF as igraph Network

Description
Load SIF as igraph Network

Usage
loadSifInIgraph(sif, directed = TRUE)

Arguments
- sif: a binary SIF as a data.frame with three columns: "PARTICIPANT_A", "INTER-ACTION_TYPE", "PARTICIPANT_B"
- directed: a boolean weather the returned graph should be directed (DEFAULT: TRUE)

Details
Users are likely to run into issues if the input SIF has factor levels

Value
a directed igraph network with interaction types

Examples
results <- readSif(system.file("extdata", "test_sif.txt", package="paxtoolsr"))
g <- loadSifInIgraph(results)

mapAttributes  \hspace{2cm} Map Attributes from igraph to Cytoscape JSON

Description
Map Attributes from igraph to Cytoscape JSON

Usage
mapAttributes(attr.names, all.attr, i)

Arguments
- attr.names: names of attributes
- all.attr: all attributes
- i: index
mapValues

Value

attributes

Note

From https://github.com/idekerlab/cy-rest-R/blob/17f748426bb5e48ba4075b9d97318ad582b250da/utility/cytoscape_util.R

mapValues  Map values from One Vector to Another

Description

Map values from One Vector to Another

Usage

mapValues(data, oldValue, newValue)

Arguments

data a vector of strings where values will be replaced
oldValue a vector that matches values in the data vector
newValue a vector of new values that will replace the old values

Value

return the vector with the mapped values. If there was no corresponding entry then replace it with an NA.

Examples

data <- c("A", "B", "C", "X", "Y", "Z")
oldValue <- LETTERS[1:20]
newValue <- letters[1:20]
results <- mapValues(data, oldValue, newValue)
mergeBiopax  

Merges two BioPAX OWL files

Description

This function merges two BioPAX OWL files

Usage

mergeBiopax(inputFile1, inputFile2, outputFile = NULL)

Arguments

- inputFile1: a string of the name of the input BioPAX OWL file
- inputFile2: a string of the name of the input BioPAX OWL file
- outputFile: a string of the name of the output merged BioPAX OWL file (Optional)

Details

Only entities that share IDs will be merged. No additional merging occurs on cross-references. Merging may result in warning messages caused as a result of redundant actions being checked against by the Java library; these messages may be ignored.

Value

an XMLInternalDocument representing a BioPAX OWL file

Examples

```r
outFile <- tempfile()
results <- mergeBiopax(system.file("extdata", "raf_map_kinase_cascade_reactome.owl", package="paxtoolsr"),
system.file("extdata", "dna_replication.owl", package="paxtoolsr"),
outFile)
```
**pcDirections**

**Description**
A simple function to see valid options

**Usage**
```
pcDirections()
```

**Details**
- BOTHSTREAM where the current entity can either be the source or target of an interaction
- DOWNSTREAM where the current entity can only be the source
- UPSTREAM where the current entity can only be the target

**Value**
acceptable Pathway Commons directions

**Examples**
```
pcDirections()
```

**pcFormats**

**Description**
A simple function to see valid options

**Usage**
```
pcFormats()
```

**Details**
See references.

**Value**
acceptable Pathway Commons formats
References

Output Formats Description: http://www.pathwaycommons.org/pc2/help/formats.html

Examples

pcFormats()

<table>
<thead>
<tr>
<th>pcGraphQueries</th>
<th>Acceptable Pathway Commons Graph Queries</th>
</tr>
</thead>
</table>

Description

A simple function to see valid options

Usage

pcGraphQueries()

Details

- COMMONSTREAM searches common downstream or common upstream of a specified set of entities based on the given directions within the boundaries of a specified length limit
- NEIGHBORHOOD searches the neighborhood of given source set of nodes
- PATHSBEWEEN finds the paths between specific source set of states or entities within the boundaries of a specified length limit
- PATHSFROMTO finds the paths from a specific source set of states or entities to a specific target set of states or entities within the boundaries of a specified length limit

Value

acceptable Pathway Commons graph queries

Examples

pcGraphQueries()
**processPcRequest**  
*Process Pathway Commons request in various formats*

**Description**  
Process Pathway Commons request in various formats

**Usage**  
processPcRequest(content, format, ...)

**Arguments**
- content: a string, content to be processed
- format: a string, the type of format
- ...: other arguments passed to read* methods for reading different formats

**Value**  
an R object using one of the read* methods provided in this package corresponding to the format

**See Also**
- `pcFormats`

**Examples**
```
fileName <- system.file("extdata", "test_biopax.owl", package="paxtoolsr")
content <- readChar(fileName, file.info(fileName)$size)
results <- processPcRequest(content, "BIOPAX")
```

**readBiopax**  
*Read BioPAX files as XML documents*

**Description**  
Read BioPAX files as XML documents

**Usage**  
readBiopax(inputFile)

**Arguments**
- inputFile: an inputFile
readGmt

Description

This function will read in gene sets in the GMT format into a named list.

Usage

readGmt(inputFile, removePrefix = FALSE, returnInfo = FALSE)

Arguments

<table>
<thead>
<tr>
<th>Argument</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>inputFile</td>
<td>an inputFile</td>
</tr>
<tr>
<td>removePrefix</td>
<td>Pathway Commons genesets are prefixed with a NCBI organism taxonomy number (e.g. 9606 for humans); this is a boolean whether to remove the prefix (default: FALSE)</td>
</tr>
<tr>
<td>returnInfo</td>
<td>a boolean whether to return information on genesets; these results are returned a list of two items: 1) basic GMT results and 2) datasource, organism, and id type information for each gene set (default: FALSE)</td>
</tr>
</tbody>
</table>

Value

a named list where each entry corresponds to a gene set or a list described in the returnInfo parameter

Examples

```r
results <- readBiopax(system.file("extdata", "biopax3-short-metabolic-pathway.owl", package="paxtoolsr"))
```

```r
results <- readGmt("test_PathwayCommons12.kegg.hgnc.gmt", package="paxtoolsr")
results <- readGmt("test_PathwayCommons12.netpath.hgnc.gmt", package="paxtoolsr")
```

```r
f1 <- system.file("extdata", "test_PathwayCommons12.kegg.hgnc.gmt", package="paxtoolsr")
f2 <- system.file("extdata", "test_PathwayCommons12.netpath.hgnc.gmt", package="paxtoolsr")
```

```r
results <- readGmt(f1)
results <- readGmt(f2)
results <- readGmt(f1, removePrefix=TRUE)
results <- readGmt(f2, returnInfo=TRUE)
```
readPcPathwaysInfo  
Read in Pathway Commons Pathways Information

Description

Read in Pathway Commons Pathways Information

Usage

readPcPathwaysInfo(inputFile = NULL, version = NULL)

Arguments

inputFile an inputFile; if NULL then retrieve the current pathways.txt; see details (default: NULL)
version a version number for a previous version of Pathway Commons data; versions 3 and above. Parameter set as version="8". Available versions "http://www.pathwaycommons.org/archives/PC2/"

Details

This file is generally found as pathways.txt.gz (e.g. http://www.pathwaycommons.org/archives/PC2/current/pathways.txt.gz)

Value

a data.frame

Examples

inputFile <- system.file("extdata", "pathways.txt.gz", package="paxtoolsr")
results <- readPcPathwaysInfo(inputFile, version="8")

readSbgn  
Read SBGN files as XML documents

Description

Read SBGN files as XML documents

Usage

readSbgn(inputFile)

Arguments

inputFile an inputFile
**readSif**

**Description**
Read in a binary SIF file

**Usage**
readSif(inputFile)

**Arguments**
- inputFile: an inputFile

**Value**
a data.frame with the interactions in the binary SIF format

**Examples**
results <- readSif(system.file("extdata", "test_sif.txt", package="paxtoolsr"))

---

**readSifnx**

**Description**
Read in an Extended SIF file

**Usage**
readSifnx(inputFile)

**Arguments**
- inputFile: an inputFile

**Value**
an XMLInternalDocument

**Examples**
results <- readSifnx(system.file("extdata", "test_sbf.xml", package="paxtoolsr"))
Details

SIFNX files from Pathway Commons commonly come a single file that includes a tab-delimited sections for nodes and another for edges. The sections are separated by an empty lines. These sections must be split before they are read.

Value

a list with nodes and edges entries

Examples

```r
results <- readSifnx(system.file("extdata", "test_sifnx.txt", package="paxtoolsr"))
```

searchListOfVectors  

Search List of Vectors

Description

Search List of Vectors

Usage

```r
searchListOfVectors(q, lst)
```

Arguments

- **q**: query vector
- **lst**: list of vectors to search

Details

Taken from: http://stackoverflow.com/questions/11002391/fast-way-of-getting-index-of-match-in-list

Value

a list of vectors with the same length as the query vector, each list entry will have indices for lst where there was a match with the query vector. Return NA if there were no matches.
Examples

```r
lst <- list(1:3, 3:5, 3:7)
q <- c(3, 5)
results <- searchListOfVectors(q, lst)
names(results) <- q
```

```r
lst <- list(LETTERS[1:3], LETTERS[3:5], LETTERS[3:7])
q <- c("C", "E")
searchListOfVectors(q, lst)
```

```r
lst <- list(LETTERS[3], LETTERS[4:6])
q <- "C"
searchListOfVectors(q, lst)
```

```r
lst <- list(LETTERS[3], LETTERS[4:6])
q <- c("C")
searchListOfVectors(q, lst)
```

```r
lst <- list(LETTERS[3], LETTERS[4:6])
q <- c("C", "E")
searchListOfVectors(q, lst)
```

```r
lst <- list(LETTERS[3], LETTERS[4:6])
q <- "Z"
searchListOfVectors(q, lst)
```

---

**searchPc**

*Search Pathway Commons*

**Description**

This command provides a text search using the Lucene query syntax.

**Usage**

```r
searchPc(
  q,
  page = 0,
  datasource = NULL,
  organism = NULL,
  type = NULL,
  verbose = FALSE
)
```

**Arguments**

- `q`  
  a keyword, name, external identifier, or a Lucene query string.
searchPc

page        an integer giving the search result page number (N>=0, default: 0)
datasource  a vector that is a filter by data source (use names or URIs of pathway data sources
            or of any existing Provenance object). If multiple data source values are specified,
            a union of hits from specified sources is returned. For example, datasource
            as c("reactome", "pid") returns hits associated with Reactome or PID.
organism    a vector that is an organism filter. The organism can be specified either by offi-
            cial name, e.g. "homo sapiens" or by NCBI taxonomy id, e.g. "9606". Similar
            to data sources, if multiple organisms are declared a union of all hits from speci-
            fied organisms is returned. For example organism as c("9606", "10016") returns
            results for both human and mice. Only humans, "9606" is officially supported.
type        BioPAX class filter. See Details.
verbose     a boolean, display the command used to query Pathway Commons

Details

Indexed fields were selected based on most common searches. Some of these fields are direct
BioPAX properties, others are composite relationships. All index fields are (case-sensitive):comment,
ecnumber, keyword, name, pathway, term, xrefdb, xrefid, dataSource, and organism. The pathway
field maps to all participants of pathways that contain the keyword(s) in any of its text fields. This
field is transitive in the sense that participants of all sub-pathways are also returned. Finally, key-
word is a transitive aggregate field that includes all searchable keywords of that element and its
child elements - e.g. a complex would be returned by a keyword search if one of its members has
a match. Keyword is the default field type. All searches can also be filtered by data source and or-
ganism. It is also possible to restrict the domain class using the 'type' parameter. This query can be
used standalone or to retrieve starting points for graph searches. Search strings are case insensitive
unless put inside quotes.

BioPAX classes can be found at http://www.pathwaycommons.org/pc2/#biopax_types

Value

an XMLInternalDocument with results

Examples

query <- "Q06609"
#results <- searchPc(query)

query <- "glycolysis"
#results <- searchPc(query, type="Pathway")
**splitSifnxByPathway**  
*Splits SIFNX entries into individual pathways*

**Description**
Splits SIFNX entries into individual pathways

**Usage**

```r
splitSifnxByPathway(edges, parallel = FALSE)
```

**Arguments**

- `edges` a data.frame with SIF content with the additional column "PATHWAY_NAMES". "PATHWAY_NAMES" should include pathway names delimited with a semicolon `;`.
- `parallel` a boolean that will parallelize the process; requires foreach/doSNOW/parallel packages

**Details**
This method can be slow; ~1.5 minutes for 150K+ rows. Has a parallelized method to speed things up.

**Value**

a list of where each entry is a vector of row indicies for a given pathway

---

**summarize**  
*Summarize a BioPAX file*

**Description**
This function provides a summary of BioPAX classes.

**Usage**

```r
summarize(inputFile)
```

**Arguments**

- `inputFile` a string of the name of the input BioPAX OWL file

**Details**
BioPAX classes are defined by the BioPAX specification: [http://www.biopax.org/](http://www.biopax.org/)
summarizeSif

Value
list with BioPAX class counts

Examples
summary <- summarize(system.file("extdata", "raf_map_kinase_cascade.reactome.owl", package="paxtoolsr"))

summarizeSif Summarize a SIF Network

Description
Summarize a SIF Network

Usage
summarizeSif(sif)

Arguments
sif a binary SIF as a data.frame with three columns: "PARTICIPANT_A", "INTERACTION_TYPE", "PARTICIPANT_B"

Value
a list containing a count of the unique genes in the SIF and counts for the interaction types in the network

Examples
results <- readSif(system.file("extdata", "test.sif.txt", package="paxtoolsr"))
summarizeSif(results)
toCytoscape  

Convert igraph to Cytoscape JSON

Description

Convert igraph to Cytoscape JSON

Usage

```
toCytoscape(igraphobj)
```

Arguments

- `igraphobj`: an igraph object

Value

a JSON object

Note

From https://github.com/idekerlab/cy-rest-R/blob/17f748426bb5e48ba4075b9d97318ad582b250da/utility/cytoscape_util.R

Examples

```
library(igraph)
g <- barabasi.game(20)
json <- toCytoscape(g)
```

toGSEA  

Converts a BioPAX OWL file to a GSEA GMT gene set

Description

This function converts pathway information stored as BioPAX files into the the GSEA .gmt format.

Usage

```
toGSEA(
    inputFile,
    outputFile = NULL,
    database = "uniprot",
    crossSpeciesCheckFlag = TRUE
)
```
toLevel3

Convert a PSIMI or older BioPAX OWL file to BioPAX Level 3

description

This file will convert PSIMI or older BioPAX objects to BioPAX Level 3

Usage

toLevel3(inputFile, outputFile = NULL)

Arguments

InputFile       a string of the name of the input file
outputFile      a string of the name of the output BioPAX OWL file

toLevel3

Arguments

inputFile   a string of the name of the input OWL file
outputFile  a string of the name of the output file
database    a string of the name of the identifier type to be included (e.g. "HGNC Symbol")
crossSpeciesCheckFlag
            a boolean that ensures participant protein is from same species

Details

The GSEA GMT format is a tab-delimited format where each row represents a gene set. The first
column is the gene set name. The second column is a brief description. Other columns for each row
contain genes in the gene set; these rows may be of unequal lengths.

Value

see readGmt()

Examples

outFile <- tempfile()
results <- toGSEA(system.file("extdata", "biopax3-short-metabolic-pathway.owl",
package="paxtoolsr"),
outFile,
"uniprot",
crossSpeciesCheckFlag=TRUE)

see also

readGmt()
Examples

```r
inputFile <- system.file("extdata", "raf_map_kinase_cascade_reactome.owl", package="paxtoolsr")
outFile <- tempfile()
results <- toLevel3(inputFile, outFile)
```

---

**topPathways**

*Retrieve top pathways*

Description

This command returns all "top" pathways.

Usage

topPathways(q = NULL, datasource = NULL, organism = NULL, verbose = FALSE)

Arguments

- `q` [Optional] a keyword, name, external identifier, or a Lucene query string, like in 'search', but the default is '*' (match all).
- `datasource` filter by data source (same as for 'search').
- `organism` organism filter (same as for 'search').
- `verbose` a boolean, display the command used to query Pathway Commons

Details

Pathways that are neither 'controlled' nor 'pathwayComponent' of another process.

Value

A data frame with the following columns:

- `uri` URI ID for the pathway
- `biopaxClass` the type of BioPAX object
- `name` a human readable name
- `dataSource` the `dataSource` for the pathway
- `organism` an organism identifier
- `pathway URI ID` for the pathway

Examples

```r
#results <- topPathways(q="TP53", datasource="panther")
```
### toSBGN

**Convert a BioPAX OWL file to SBGNML**

#### Description

This function will convert a BioPAX OWL file into the Systems Biology Graphical Notation (SBGN) Markup Language (SBGNML) XML representation.

#### Usage

```r
toSBGN(inputFile, outputFile = NULL)
```

#### Arguments

- `inputFile` a string of the name of the input BioPAX OWL file
- `outputFile` a string of the name of the output SBGNML file

#### Details

Objects in the SBGNML format are laid out using a Compound Spring Embedder (CoSE) layout.

#### Value

see `readSbgn()`

#### References


#### Examples

```r
outFile <- tempfile()
results <- toSBGN(system.file("extdata", "biopax3-short-metabolic-pathway.owl", package="paxtoolsr"), outFile)
```
toSif

Convert a BioPAX OWL file to SIF

Description

Convert a BioPAX OWL file to a binary SIF file

Usage

toSif(inputFile, outputFile = NULL)

Arguments

inputFile a string of the name of the input BioPAX OWL file
outputFile a string of the name of the output SIF file (Optional)

Details

Information on SIF conversion is provided on the Pathway Commons site: http://www.pathwaycommons.org/pc2/

Value

see readSif()

Examples

outFile <- tempfile()
results <- toSif(system.file("extdata", "raf_map_kinase_cascade_reactome.owl", package="paxtoolsr"), outFile)

---

toSifnx

Converts BioPAX OWL file to extended binary SIF representation

Description

Converts BioPAX OWL file to extended binary SIF representation

Usage

toSifnx(inputFile, outputFile = tempfile(), idType = "uniprot")
Arguments

- **inputFile**: a string with the name of the input BioPAX OWL file
- **outputFile**: a string with the name of the output file for SIFNX information
- **idType**: a string either "hgnc" or "uniprot" (DEFAULT: uniprot, more common)

Details

Information on SIF conversion is provided on the Pathway Commons site: [http://www.pathwaycommons.org/pc2/](http://www.pathwaycommons.org/pc2/). Also, this is a Java-based methods, it is best to use full paths.

Value

see readSifnx()

Examples

```r
code:
inputFile <- system.file("extdata", "raf_map_kinase_cascade_reactome.owl", package="paxtoolsr")
results <- toSifnx(inputFile=inputFile)
```

---

**traverse**

*Access Pathway Commons using XPath-type expressions*

Description

This command provides XPath-like access to the Pathway Commons.

Usage

```r
code:
traverse(uri, path, verbose = FALSE)
```

Arguments

- **uri**: a BioPAX element URI - specified similarly to the 'GET' command above). Multiple IDs are allowed (uri=...&uri=...&uri=...).
- **path**: a BioPAX propery path in the form of property1[type1]/property2[type2]; see properties, inverse properties, Paxtools, org.biopax.paxtools.controller.PathAccessor.
- **verbose**: a boolean, display the command used to query Pathway Commons
Details

With traverse users can explicitly state the paths they would like to access. The format of the path query is in the form: [Initial Class]/[property1]:[classRestriction(optional)]/[property2]... A "*" sign after the property instructs path accessor to transitively traverse that property. For example, the following path accessor will traverse through all physical entity components within a complex: "Complex/component*/entityReference/xref:UnificationXref" The following will list display names of all participants of interactions, which are components (pathwayComponent) of a pathway (note: pathwayOrder property, where same or other interactions can be reached, is not considered here): "Pathway/pathwayComponent:Interaction/participant*/displayName" The optional parameter classRestriction allows to restrict/filter the returned property values to a certain subclass of the range of that property. In the first example above, this is used to get only the Unification Xrefs. Path accessors can use all the official BioPAX properties as well as additional derived classes and parameters in paxtools such as inverse parameters and interfaces that represent anonymous union classes in OWL. (See Paxtools documentation for more details).

Value

an XMLInternalDocument with results

References

Paxtools Documentation: http://www.biopax.org/m2site/

Examples

uri <- "http://identifiers.org/uniprot/P38398"
#results <- traverse(uri=uri, path="ProteinReference/organism/displayName")

validate

Validate BioPAX files

Description

This function validates BioPAX files for errors.

Usage

validate(
  inputFile,
  outputFile = NULL,
  type = c("xml", "html", "biopax"),
  autoFix = FALSE,
  onlyErrors = FALSE,
  maxErrors = NULL,
  notStrict = FALSE
)
validate

Arguments

- **inputFile**: a string of the name of the input BioPAX OWL file
- **outputFile**: a string of the name of the output file containing validation results
- **type**: a string denoting the type of output: xml (default), html, biopax
- **autoFix**: a boolean that determines if the input file should be fixed automatically. Errors that can be automatically fixed include generating displayName properties from names, inferring organism, and inferring dataSource
- **onlyErrors**: a boolean of whether to only display errors
- **maxErrors**: a integer denoting the number of errors to return
- **notStrict**: a boolean of whether to be strict in validation (default: FALSE)

Details

See the publication by Rodchenkov, et al. for information on the BioPAX validator. See http://biopax.baderlab.org/validator for additional information on validator. See http://biopax.baderlab.org/validator/errorTypes.html for information on error types.

Value

an XMLInternalDocument is returned if type is set to "xml" otherwise the location of the outputfile is returned.

References


Examples

```r
outFile <- tempfile()
rawDoc <- validate(system.file("extdata", "raf_map_kinase_cascade_reactome.owl", package="paxtoolsr"), onlyErrors=TRUE)
```
**Index**

* **paxtools**
  - addAttributeList, 3
  - convertDataFrameListsToVectors, 4
  - convertSifToGmt, 4
  - downloadFile, 5
  - downloadPc2, 6
  - fetch, 8
  - filterSif, 8
  - getCacheFiles, 10
  - getErrorMessage, 11
  - getNeighbors, 11
  - getPc, 12
  - getPcDatabaseNames, 13
  - getPcUrl, 14
  - getShortestPathSif, 14
  - getSifInteractionCategories, 15
  - graphPc, 16
  - integrateBiopax, 17
  - loadSifInIgraph, 18
  - mapAttributes, 18
  - mapValues, 19
  - mergeBiopax, 20
  - pcDirections, 21
  - pcFormats, 21
  - pcGraphQueries, 22
  - processPcRequest, 23
  - readBiopax, 23
  - readGmt, 24
  - readPcPathwaysInfo, 25
  - readSbggn, 25
  - readSif, 26
  - readSifnx, 26
  - searchListOfVectors, 27
  - searchPc, 28
  - splitSifnxByPathway, 30
  - summarize, 30
  - summarizeSif, 31
  - toCytoscape, 32
  - toGSEA, 32
  - toLevel3, 33
  - topPathways, 34
  - toSBGN, 35
  - toSif, 36
  - toSifnx, 36
  - traverse, 37
  - validate, 38

  - addAttributeList, 3
  - convertDataFrameListsToVectors, 4
  - convertSifToGmt, 4
  - downloadFile, 5
  - downloadPc (downloadPc2), 6
  - downloadPc2, 6
  - downloadSignedPC, 7
  - fetch, 8
  - filterSif, 8
  - getCacheFiles, 10
  - getErrorMessage, 11
  - getNeighbors, 11
  - getPc, 12
  - getPcDatabaseNames, 13
  - getPcUrl, 14
  - getShortestPathSif, 14
  - getSifInteractionCategories, 15
  - graphPc, 16
  - integrateBiopax, 17
  - loadSifInIgraph, 18
  - mapAttributes, 18
  - mapValues, 19
  - mergeBiopax, 20
  - pcDirections, 16, 17
  - pcFormats, 12, 13, 16, 17, 21, 23
INDEX

pcGraphQueries, 16, 22
processPcRequest, 23

readBiopax, 6, 23
readGmt, 6, 24
readPcPathwaysInfo, 25
readSbgn, 6, 25
readSif, 6, 26
readSifnx, 6, 26

searchListOfVectors, 27
searchPc, 28
splitSifnxByPathway, 30
summarize, 30
summarizeSif, 31

toCytoscape, 32
toGSEA, 32
toLevel3, 33
topPathways, 34
toSBGN, 35
toSif, 36
toSifnx, 36
traverse, 37

validate, 38