Package 'Path2PPI'

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Type Package

Title Prediction of pathway-related protein-protein interaction
networks

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Description Package to predict protein-protein interaction (PPI) networks in
target organisms for which only a view information about
PPIs is available. Path2PPI predicts PPI networks based on sets of
proteins which can belong to a certain pathway from well-established model
organisms. It helps to combine and transfer information of a certain
pathway or biological process from several reference organisms to one
target organism. Path2PPI only depends on the sequence similarity of the
involved proteins.

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Depends R (>= 3.2.1), igraph (>= 1.0.1), methods

Suggests knitr, rmarkdown, RUnit, BiocGenerics, BiocStyle

VignetteBuilder knitr

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R topics documented:

addReference   *Add reference species*

Description

Adds reference species to an object from the class Path2PPI.

Usage

```
addReference(path2ppi, taxName, taxId, proteins, irefindex, homologs)
```

Arguments

path2ppi   An object of the class Path2PPI.

taxName   A character string giving the taxonomy name.

taxId   A character string giving the taxonomy identifier.

proteins   Either a character vector with the identifiers of the proteins which are involved
             in the corresponding pathway or a character vector with the protein names or
             aliases, respectively, named by the protein identifiers.

irefindex   Either a data frame, representing the iRefIndex table of the current reference
             species, e.g. loaded previously via read.table, or the corresponding file name
             of the iRefIndex file.

homologs   Either a data frame representing the results of the BLAST search (e.g. loaded
             previously via read.table) or the corresponding file name of the BLAST result
             file.

Details

This method searches for all relevant interactions in the data frame or file defined in iRefIndex.
There are different and often ambiguous protein identifiers defined in an iRefIndex file, and the
putative "major" identifiers are not necessarily those defined in the corresponding "major" columns
"uidA" and "uidB". Furthermore, iRefIndex also contains protein complexes. Hence, Path2PPI
applies an advanced search algorithm to automatically find relevant interactions associated with the
pathway or the proteins of interest, respectively. The user does not have to predefine the identifiers
types (Uniprot, Swissprot, Ensemble etc.), since these types are often unambiguously assigned.
The algorithm searches for each identifier in 10 columns where any type of identifier or accession
number is defined ("uidA", "altA", "OriginalReferenceA", "FinalReferenceA", "aliasA", "uidB",
"altB", "OriginalReferenceB", "FinalReferenceB" and "aliasB"). Additionally, it searches for each
complex which contains one or more of the predefined proteins. Subsequently, each homologous
relationship which is not relevant for the previously found interactions is declined. The results of
these searches are centralized in the Path2PPI object and can be visualized using the appropriate
methods (e.g. showReferences)

Value

An object from the class `Path2PPI` with attached reference species.

Author(s)

Oliver Philipp `<MolBI-software@bioinformatik.uni-frankfurt.de>`

See Also

[showReferences](), [removeReference]()

Examples

```
data(ai) #Load test data set

ppi <- Path2PPI("Autophagy induction", "Podospora anserina", "5145")
ppi

ppi <- addReference(ppi, "Homo sapiens", "9606", human.ai.proteins,
                    human.ai.irefindex, pa2human.ai.homologs)
ppi <- addReference(ppi, "Saccharomyces cerevisiae (S288c)", "559292",
                    yeast.ai.proteins, yeast.ai.irefindex,
                    pa2yeast.ai.homologs)
ppi
```

---

ai   *Data set to predict autophagy induction in* Podospora anserina

---

Description

This data set consists of all data files necessary to predict the putative interactions of the induction step of autophagy in *Podospora anserina* by means of the corresponding PPIs in human and yeast.

Usage

```
data("ai")
```

Format

human.ai.irefindex: Data frames with 1694 observations of 54 variables. yeast.ai.irefindex: Data frames with 3840 observations of 54 variables. pa2human.ai.homologs: Data frames with 261 observations of 12 variables. pa2yeast.ai.homologs: Data frames with 98 observations of 12 variables. human.ai.proteins: Named character vector with 5 elements. yeast.ai.proteins: Named character vector with 7 elements.

Details

Data frames `human.ai.irefindex` and `yeast.ai.irefindex` consists of all relevant interactions of the corresponding iRefIndex files. The two data frames `pa2human.ai.homologs` and `pa2yeast.ai.homologs` are the necessary parts of the result files from the BLAST searches of the *P. anserina* proteom against the proteoms of human and yeast. The named character vectors `human.ai.proteins` and `yeast.ai.proteins` consists of the proteins involved in the induction process of autophagy in human and yeast.

Value

Four data frames and two named character vectors (see above).

References

Camacho, C. et al. (2009). BLAST+: architecture and applications. BMC Bioinformatics, 10(1), 421.

Razick, S. et al. (2008). iRefIndex: a consolidated protein interaction database with provenance. BMC Bioinformatics, 9(1), 405.

Examples

```
data(ai)
```

---

getHybridNetwork   *Get hybrid network of the predicted PPI*

Description

Get the hybrid network of the previously predicted PPI. The hybrid network consists of all relevant interactions from the reference species, the predicted interactions in the target species and all relevant homologous relationships.

Usage

```
getHybridNetwork(path2ppi, igraph = FALSE)
```

Arguments

path2ppi   An object of the class `Path2PPI`.

igraph   Logical; if `TRUE` then the hybrid network is given as igraph-object. Otherwise a data frame, consisting of each interaction and homologous relationship, will be returned.

Value

See `igraph` argument.

Author(s)

Oliver Philipp <MolBI-software@bioinformatik.uni-frankfurt.de>

See Also

[getPPI](#)

Examples

```
data(ai) #Load test data set

ppi <- Path2PPI("Autophagy induction", "Podospora anserina", "5145")

ppi <- addReference(ppi, "Homo sapiens", "9606", human.ai.proteins,
                    human.ai.irefindex, pa2human.ai.homologs)
ppi <- addReference(ppi, "Saccharomyces cerevisiae (S288c)", "559292",
                    yeast.ai.proteins, yeast.ai.irefindex,
                    pa2yeast.ai.homologs)

ppi <- predictPPI(ppi)

#Return the hybrid network as data frame
hybrid <- getHybridNetwork(ppi)

#Return the hybrid network as igraph object
hybrid <- getHybridNetwork(ppi,igraph=TRUE)
```

---

getPPI	*Get predicted PPI*

Description

Get the predicted PPI of an `Path2PPI` object consisting of each predicted interaction and protein in the target species.

Usage

```
getPPI(path2ppi, raw=FALSE, igraph=FALSE)
```

Arguments

path2ppi	An object of the class `Path2PPI`.
raw	Logical; if `TRUE` then the detailed view of the predicted PPI will be returned. That means that each predicted interaction deduced from each reference species is given. In contrast, `FALSE` leads to the actually predicted and combined PPI where no redundancies occur.
igraph	Logical; if `TRUE` then the returned PPI is given as igraph-object. Otherwise a data frame with each predicted interaction will be returned.

Value

See `igraph` argument.

Author(s)

Oliver Philipp <MolBI-software@bioinformatik.uni-frankfurt.de>

See Also

getHybridNetwork

Examples

```
data(ai) #Load test data set

ppi <- Path2PPI("Autophagy induction", "Podospora anserina", "5145")

ppi <- addReference(ppi, "Homo sapiens", "9606", human.ai.proteins,
                    human.ai.irefindex, pa2human.ai.homologs)
ppi <- addReference(ppi, "Saccharomyces cerevisiae (S288c)", "559292",
                    yeast.ai.proteins, yeast.ai.irefindex,
                    pa2yeast.ai.homologs)

ppi <- predictPPI(ppi)

#Get the predicted PPI as data frame.
network <- getPPI(ppi)

#Get the detailed predicted PPI as data frame.
network.raw <- getPPI(ppi,raw=TRUE)
```

---

homologyScore   *Computes homology scores*

Description

Computes the homology scores based on the BLAST E-value. This function is used by the `predictPPI` method to compute homology scores to decide whether an interaction in a reference species is adopted to the target species (see package vignette for a detailed description). It can be used to test which E-values lead to which scores given a predefined E-value range.

Usage

```
homologyScore(e.value, h.range)
```

Arguments

e.value   One BLAST E-value or a numeric vector with different BLAST E-values

h.range   Numeric vector consisting of two values. The first value indicates the lower bound (smallest E-value). Each E-value which is equal or less than this bound is scored with 1. The second value indicates the upper bound (biggest E-value). Each E-value which is equal or greater than this bound is scored with 0.

Details

Uses a linear function to map the E-value v to the range $[l, u]$ where l is the lower and u the upper bound:

$$s(v) = |m \log_{10}(v) + b|$$

$$m = \frac{1}{\log_{10}(l) - \log_{10}(u)}$$

$$b = -(m \log_{10}(u))$$

Value

Numeric vector containing the scores.

Author(s)

Oliver Philipp <MolBI-software@bioinformatik.uni-frankfurt.de>

See Also

[predictPPI](#)

Examples

```
l <- 1e-100                              #lower bound
u <- 1e-20                               #upper bound
h.range <- c(l,u)                        #define range
e.values <- c(1e-20,1e-40,1e-60,1e-80,1e-100) #some BLAST E-values

homologyScore(e.values,h.range)
```

Path2PPI-class   *Class* "Path2PPI"

Description

An instance of the class `Path2PPI` is the major object in the Path2PPI package. It manages all reference species and the target species. The prediction algorithm is implemented in this class as well.

Usage

```
Path2PPI(...)
```

Arguments

...   Argument list (see Note below).

Value

An instance of the class `Path2PPI`.

Slots

pathway: Object of class "character"
targetSpecies: Object of class ".TargetSpecies"
referenceContainer: Object of class ".ReferenceContainer"
h.thresh: Object of class "numeric"
h.range: Object of class "numeric"
i.thresh: Object of class "numeric"
consider.complexes: Object of class "logical"
max.complex.size: Object of class "numeric"
raw.ppi: Object of class "data.frame"
ppi: Object of class "data.frame"

Methods

  addReference signature(path2ppi = "Path2PPI")

  getHybridNetwork signature(path2ppi = "Path2PPI")

  getPPI signature(path2ppi = "Path2PPI")

  initialize signature(.Object = "Path2PPI")

  plot.Path2PPI signature(x = "Path2PPI")

  predictPPI signature(path2ppi = "Path2PPI")

  removeReference signature(path2ppi = "Path2PPI")

  show signature(object = "Path2PPI")

  showInteraction signature(path2ppi = "Path2PPI")

  showReferences signature(path2ppi = "Path2PPI")

Note

Arguments to Path2PPI() and the new method are obligatory and must be named if they differ from this order:

pathway	A character string with the name of the pathway which has to be predicted.
targetName	A character string giving the taxonomy name of the target species.
targetId	A character string giving the taxonomy identifier of the target species.

Author(s)

Oliver Philipp <MolBI-software@bioinformatik.uni-frankfurt.de>

Examples

```
ppi <- Path2PPI("Autophagy induction", "Podospora anserina", "5145")
ppi
```

---

  plot.Path2PPI   *Plots the predicted PPI*

Description

Plots the predicted PPI in three different ways. Depending on the type argument it manages the specific layout settings and finally uses the plot function of the igraph package.

Usage

```
## S3 method for class 'Path2PPI'
plot(x, type = "ppi", multiple.edges = FALSE,
                scores = FALSE, species.colors = c(),
                vertices.opacity=0.8, use.identifiers=FALSE,
                protein.labels = NA, show.legend = TRUE,
                vertices.coordinates = NA, return.coordinates = FALSE,
                tkplot=FALSE,...)
```

Arguments

x	An object from the class `Path2PPI` where the PPI network already has been predicted.
type	Character string. Which graph type to plot. "ppi": plots only the predicted PPI. "hybrid": plots the hybrid network which consists of all relevant interactions from the reference species, the predicted interactions in the target species and all relevant homologous relationships.
multiple.edges	Logical. Is only considered if `type="ppi"`. If TRUE then each reference interaction is depicted in the species-specific color (raw mode), in contrast, if set to `FALSE` only the finalized / combined interactions are depicted.
scores	Logical. If TRUE the edge scores will be shown.
species.colors	Named vector, to specify the species colors. If no value is given then default colors are used.
vertices.opacity	Numeric value between 0 and 1 defining the opacity of the vertices.
use.identifiers	Logical. If TRUE then only the proteins identifiers are used as the vertex labels.
protein.labels	Named vector to define the labels of the vertices. If no value is given then the protein identifiers are used. The vector does not have to be complete, i.e. not each protein has to be defined.
show.legend	Logical. If TRUE then a legend is depicted.
vertices.coordinates	Data frame containing the coordinates of the vertices. If no value is given then coordinates are computed using the [layout.auto](#) function.
return.coordinates	Logical. If TRUE the coordinates of the vertices are returned.
tkplot	Logical. If TRUE the graph is drawn in the interactive graph drawing facility [tkplot](#).
...	Additional plotting parameters.

Details

The argument `return.coordinates` only works correctly if `tkplot=FALSE`. If you want to get the coordinates of the tkplot device use [tkplot.getcoords](#).

Value

If `return.coordinates=TRUE` the coordinates of the vertices are returned.

Note

If you want to export the plotted graph to postscript you have to consider that the default font family is set to *sans* for vertex and edge labels. Please change the default font family of postscript to *sans* before you call the plot method: ps.options(family="sans"). Additionally, you have to consider that the default value for `vertices.opacity` is set to *0.8* in order to enhance the visibility of the graph, since some edges may be hidden by the vertices. Postscript does not support semi-transparencies. Hence, please change the `vertices.opacity` argument to *1* if you want to export the graph using postscript.

Author(s)

Oliver Philipp <MolBI-software@bioinformatik.uni-frankfurt.de>

See Also

predictPPI, igraph for other plotting parameters

Examples

```
data(ai) #Load test data set

ppi <- Path2PPI("Autophagy induction", "Podospora anserina", "5145")
ppi <- addReference(ppi, "Homo sapiens", "9606", human.ai.proteins,
                    human.ai.irefindex, pa2human.ai.homologs)
ppi <- addReference(ppi, "Saccharomyces cerevisiae (S288c)", "559292",
                    yeast.ai.proteins, yeast.ai.irefindex,
                    pa2yeast.ai.homologs)

ppi <- predictPPI(ppi,h.range=c(1e-60,1e-20))

#Plot the predicted PPI with the default settings and return
#the coordinates of the vertices
set.seed(12)
coordinates <- plot(ppi, return.coordinates=TRUE)

#Plot the predicted PPI and show each underlying reference interaction.
#Use different species specific colors. To compare both graphs,
#use the coordinates computed before
plot(ppi,multiple.edges=TRUE,vertices.coordinates=coordinates)

#Plot the corresponding hybrid network with predefined species colors.
#Also define some labels for the proteins of the target species.
#Keep in mind: You can not use the data in "coordinates" since
#the hybrid network consists of more vertices than the default PPI
set.seed(40)
target.labels<-c("B2AE79"="PaTOR","B2AXK6"="PaATG1",
                 "B2AUW3"="PaATG17","B2AM44"="PaATG11",
                 "B2AQV0"="PaATG13","B2B5M3"="PaVAC8")
species.colors <- c("5145"="red","9606"="blue","559292"="green")
plot(ppi,type="hybrid",species.colors=species.colors,
protein.labels=target.labels)
```

---

predictPPI   *Prediction of the PPI*

Description

Major method of the Path2PPI class to predict the final PPI in the target species using the information available from the stored reference species. Different values for the arguments of this method can lead to different PPI networks, differing in the degree of reliability and strictness.

Usage

```
predictPPI(path2ppi, mode="both", h.thresh=1e-05,
          h.range=c(1e-100, 1e-20), i.thresh=0.7,
          consider.complexes=FALSE, max.complex.size=5,
          decline.self.interaction.ref=FALSE,
          decline.self.interaction.tar=TRUE,
          verbose=TRUE)
```

Arguments

path2ppi   An object of the class `Path2PPI`.

mode   Which interaction from the reference species should be taken into account. "both": both interactors of an interaction has to be in the initial protein list previously inserted by the user (recommended if it is a large network or many proteins were initially defined, respectively). "one": only one of the interactors of each reference interaction has to be in the initial protein list (may lead to very large networks).

h.thresh   E-value cutoff at which each homologous relationship definitely will be declined (see also `h.range` argument).

h.range   Numeric vector consisting of two values. The first value indicates the lower border (smallest E-value). Each E-value which is equal or less than this border is scored with 1 (best). The second value indicates the upper border (biggest E-value). Each E-value which is equal or greater than this border is scored with 0 (worst).

i.thresh   Numeric. Threshold for accepted interactions. If the computed prediction score for an interaction is less than `i.thresh` it will be declined.

consider.complexes

      Logical. If `TRUE` then interactions are also considered which actually indicate an association of the current protein to one bigger protein complex. This may lead to very large networks if mode="one" since all other proteins of this complex are considered as well, i.e., each protein in such complexes are considered to interact with each other protein of this complex. If mode="both" then each protein of an complex has to be in the initial protein list to consider each interaction (see details).

max.complex.size

      Numeric. Is only considered if `consider.complexes=TRUE`. The maximum size of complexes to be considered.

decline.self.interaction.ref

      Logical. If `TRUE` then all self interactions from reference species are declined.

decline.self.interaction.tar

      Logical. If `TRUE` then all predicted self interactions in target species are declined.

verbose   Logical. `FALSE` hides messages in the output.

Details

Difference of `h.thresh` and `h.range`: If only one protein in the target species was found to be homologous to a current reference species protein and this homology was rated with an E-value which is equal or smaller than `h.thresh` it is scored with 1 (even if the E-value is larger than the upper border of `h.range`). See package vignette for more details.

Use the complex arguments with care, since each complex may lead to a vast amount of interactions, i.e., each protein is considered to interact with each other of this complex; e.g. if there are 10 proteins involved in one complex, this would lead to 10 over 2 = 45 interactions.

Value

An object of the class `Path2PPI` with predicted PPI.

Author(s)

Oliver Philipp <MolBI-software@bioinformatik.uni-frankfurt.de>

See Also

[plot.Path2PPI](), [homologyScore]()

Examples

```
data(ai) #Load test data set

ppi <- Path2PPI("Autophagy induction", "Podospora anserina", "5145")
ppi <- addReference(ppi, "Homo sapiens", "9606", human.ai.proteins,
                    human.ai.irefindex, pa2human.ai.homologs)
ppi <- addReference(ppi, "Saccharomyces cerevisiae (S288c)", "559292",
                    yeast.ai.proteins, yeast.ai.irefindex,
                    pa2yeast.ai.homologs)

#Using the default settings leads to 8 predicted interactions in the
#target species
ppi <- predictPPI(ppi)

#Consider complexes where each complex is allowed to be up to 10 proteins
#large. For this smaller pathway only one more interaction was predicted when
#considering larger complexes.
ppi <- predictPPI(ppi,consider.complexes=TRUE,max.complex.size=10)

#We can be less strict and decrease h.range what obviously increases the
#number of predicted interactions to 13
ppi <- predictPPI(ppi,h.range=c(1e-60,1e-20))
```

---

removeReference   *Remove reference species*

Description

Remove reference species previously attached to an object from the class `Path2PPI`.

Usage

```
removeReference(path2ppi, species)
```

Arguments

path2ppi	An object from the class `Path2PPI`.
species	Either a number between 1 and the number of stored reference species or a character string with the taxonomy id of the reference species to remove.

Value

An object of the class `Path2PPI` with removed reference species `species`.

Author(s)

Oliver Philipp <MolBI-software@bioinformatik.uni-frankfurt.de>

See Also

showReferences, addReference

Examples

```
data(ai) #Load test data set

ppi <- Path2PPI("Autophagy induction", "Podospora anserina", "5145")

ppi <- addReference(ppi, "Homo sapiens", "9606", human.ai.proteins,
                    human.ai.irefindex, pa2human.ai.homologs)
ppi <- addReference(ppi, "Saccharomyces cerevisiae (S288c)", "559292",
yeast.ai.proteins, yeast.ai.irefindex, pa2yeast.ai.homologs)

#Remove second reference species
ppi <- removeReference(ppi,2)

#Remove reference species with taxonomy id "9606"
ppi <- removeReference(ppi,"9606")
```

---

showInteraction   *Information about an interaction*

Description

Use `showInteraction` to get detailed information about one interaction of the predicted PPI.

Usage

```
showInteraction(path2ppi, interaction, mode="default", verbose=TRUE)
```

Arguments

path2ppi	An object from the class `Path2PPI`.
interaction	Character vector consisting of the identifiers of the two interactors.
mode	Character string. Which information of this interaction is requested. "default": only the predicted interaction and some major information are provided. "detailed": all interactions deduced from each reference species with this interaction is provided. "references": each reference interaction of the current interaction with some major information. "references.detailed": each reference interaction of the current interaction with all available information (extracted from the corresponding iRefIndex data set).
verbose	Logical. `FALSE` hides messages in the output.

Value

Data frame with the requested information defined in `mode`.

Author(s)

Oliver Philipp <MolBI-software@bioinformatik.uni-frankfurt.de>

See Also

[plot.Path2PPI,showReferences](#)

Examples

```
data(ai) #Load test data set

ppi <- Path2PPI("Autophagy induction", "Podospora anserina", "5145")
ppi <- addReference(ppi, "Homo sapiens", "9606", human.ai.proteins,
                    human.ai.irefindex, pa2human.ai.homologs)
ppi <- addReference(ppi, "Saccharomyces cerevisiae (S288c)", "559292",
                    yeast.ai.proteins, yeast.ai.irefindex,
                    pa2yeast.ai.homologs)

ppi <- predictPPI(ppi,h.range=c(1e-60,1e-20))

interaction <- showInteraction(ppi,interaction=c("B2AT71","B2AE79"),
mode="detailed")
interaction
```

---

showReferences   *Information about reference species*

Description

Get information about the currently stored reference species. If indicated by `returnValue` a data frame - containing information about each protein or interaction - is provided as well.

Usage

```
showReferences(path2ppi, species = NA, returnValue = NA)
```

Arguments

path2ppi	An object from the class `Path2PPI`.
species	Either a number between 1 and the number of stored reference species or a character string with the taxonomy id. If no value for `species` is given then information about each stored reference species is provided.
returnValue	Character value indicating whether to return a value. "proteins": a data frame containing the proteins associated with the pathway of interest in the corresponding reference species. "interactions": a data frame containing all processed, relevant and non-redundant interactions. "irefindex": a data frame containing all relevant interactions in the raw irefindex format. Is only reasonable if `species` is defined. If no value for `returnValue` is given then only general information is provided.

Value

See description for `returnValue`

Author(s)

Oliver Philipp <MolBI-software@bioinformatik.uni-frankfurt.de>

See Also

[addReference](#), [removeReference](#), [showInteraction](#)

Examples

```
data(ai) #Load test data set

ppi <- Path2PPI("Autophagy induction", "Podospora anserina", "5145")

ppi <- addReference(ppi, "Homo sapiens", "9606", human.ai.proteins,
                    human.ai.irefindex, pa2human.ai.homologs)
ppi <- addReference(ppi, "Saccharomyces cerevisiae (S288c)", "559292",
                    yeast.ai.proteins, yeast.ai.irefindex,
                    pa2yeast.ai.homologs)

#Get general information about each stored reference species
showReferences(ppi)

#Get general information about reference species with the taxonomy id "9606"
showReferences(ppi, species="9606")

#Get all proteins associated with the pathway of interest
#and previously given by the user
proteins <- showReferences(ppi, species="9606", returnValue="proteins")

#Get all processed and non-redundant interactions previously
#determined to be relevant for the pathway of interest
interactions <- showReferences(ppi, species="9606", returnValue="interactions")

#Get all relevant interactions in the detailed irefindex format
irefindex <- showReferences(ppi, species="9606", returnValue="irefindex")
```

Index