Package ‘CHRONOS’

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Title CHRONOS: A time-varying method for microRNA-mediated sub-pathway enrichment analysis
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Description A package used for efficient unraveling of the inherent dynamic properties of pathways. MicroRNA-mediated subpathway topologies are extracted and evaluated by exploiting the temporal transition and the fold change activity of the linked genes/microRNAs.
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VignetteBuilder knitr
biocViews SystemsBiology, GraphAndNetwork, Pathways, KEGG

R topics documented:

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CHRONOSrun

Default run of CHRONOS

Description
Default run of CHRONOS

Usage
CHRONOSrun(mRNAexp, mRNAlabel, miRNAexp, pathType, subType, measures, thresholds, org, export, verbose, miRNAinteractions)

Arguments
mRNAexp  mRNA expressions filename located in CHRONOS/extdata/Input
mRNAlabel  mRNA nomenclature (for supported types see convertNomenclature)
miRNAexp  miRNA expressions filename located in CHRONOS/extdata/Input
pathType  Pathway type (‘Metabolic’, ‘Non-Metabolic’, ‘All’ or vector of pathway ids)
subType  Subpathway type (‘Linear’, ‘Non-Linear’, ‘All’)
measures  Include subpathway structural and functional aspects (‘TRUE’, ‘FALSE’)
thresholds  Subscore, mirscore and p-value thresholds
c(‘pvalue’=pvalue, ‘subscore’=subscore, ‘mirscore’=mirscore)
org  KEGG organism identifier
export  Export file type (’.xlsx’, .txt’)
verbose  Show informative messages (TRUE/FALSE).
miRNAinteractions  Edgelist of miRNA-mRNA interactions.

Details
• Imports gene and miRNA expressions from CHRONOS/extdata/Input/<mRNAexpFile>.txt and CHRONOS/extdata/Input/<miRNAexpFile>.txt
• Downloads all available pathways for the specified organism from KEGG.
• Creates pathway graphs from downloaded KGML files.
• Extracts linear subpathways from metabolic and non metabolic graphs.
• Extracts non linear subpathways from metabolic and non metabolic graphs.
convertMiRNAAnnotations

- Downloads miRecords miRNA-mRNA interactions.
- Scores and evaluates (linear and non-linear) subpathways to extract significant results.
- Organism identifier.
- Visualizes most the significant results (’.xlsx’ or ’.txt’).
- Display informative messages (TRUE/FALSE).
- User-defined miRNA-mRNA interactions can be supplied in the form of an edgelist with two columns. If no such information is available, a missing or a NULL argument forces the use of default interactions by using `downloadMiRecords`.

Value

Examples

# Default run

```r
load(system.file('extdata', 'Examples//data.RData', package='CHRONOS'))
res <- CHRONOSrun(mRNAexp=mRNAexpr,
                   mRNAlabel='entrezgene',
                   miRNAexp=miRNAexpr,
                   pathType=c('04915', '04917', '04930', '05031'),
                   org='hsa',
                   subType='Linear',
                   thresholds=c('subScore'=0.4, 'mirScore'=0.4),
                   miRNAinteractions=miRNAinteractions)
```

convertMiRNAAnnotations

*Conform miRNA annotations to the ones currently used by miRecords.*

Description

Conform miRNA annotations to the ones currently used by miRecords.

Usage

`convertMiRNAAnnotations(org, miRNAs, update)`

Arguments

<table>
<thead>
<tr>
<th>org</th>
<th>KEGG organism identifier.</th>
</tr>
</thead>
<tbody>
<tr>
<td>miRNAs</td>
<td>Vector of miRNAs identifiers.</td>
</tr>
<tr>
<td>update</td>
<td>Update annotation mapper with latest annotation changes.</td>
</tr>
</tbody>
</table>

Details

Determine which miRNAs are incompatible with miRecords annotations and retrieve the suitable ones from www.mirbase.org.
convertNomenclature

Value
.

Examples

data <- c('hsa-let-7g-5p', 'hsa-miR-154-5p', 'hsa-miR-376b-3p')

correctMiRNANomenclature(org='hsa', miRNAs=data)

convertNomenclature  Convert genes identifier nomenclature.

Description
Convert genes identifier nomenclature.

Usage
convertNomenclature(ids, org, from, to)

Arguments
ids  Vector of gene identifiers
org  KEGG organism identifier
from Initial identifier type
to A vector of final identifier types

Details
EntrezGene ID 'entrezgene'
Ensembl Gene ID 'ensembl_gene_id'
Ensemble Transcript ID 'ensembl_transcript_id'
Ensemble Protein ID 'ensembl_peptide_id'
HGNC ID 'hgnc_id'
HGNC Symbol 'hgnc_symbol'
HGNC Transcript name 'hgnc_transcript_name'
Refseq mRNA ID 'refseq_mrna'
Refseq Protein ID 'refseq_peptide'
UniProt/Swissprot Accession 'uniprot_swissprot_accession'
UniProt/Swissprot ID 'uniprot_swissprot'
UniGene ID 'unigene'
UniProt Genename ID 'uniprot_genename'

Value
Vector of converted gene identifiers
createPathwayGraphs

Examples

# Identifiers to be converted
ids <- c('5091', '5105')

# Convert to HGNC ID, Ensembl Gene ID and Uniprot Genename ID
from <- 'entrezgene'
to <- c('hgnc_symbol', 'ensembl_gene_id', 'uniprot_genename')

## Not run: res <- convertNomenclature(ids=ids, org='hsa', from=from, to=to)

createPathwayGraphs  Convert KEGG Pathways to Gene-Gene Network Graphs.

Description

Convert KEGG Pathways to Gene-Gene Network Graphs.

Usage

createPathwayGraphs(org, pathways, edgeTypes, doubleEdges, choice, groupMode)

Arguments

org  KEGG organism identifier.
pathways  Vector of KEGG pathway identifiers.
edgeTypes  Vector of edge types mappings.
doubleEdges  Specify which edgeTypes should be considered bidirectional.
choice  Create metabolic graph either by using relations or reactions from KGML file ('reactions', 'relations')
groupMode  'expand' to consider each group member a node, or 'collapse' to consider all components’ genes as a node

Details

KEGG pathways consist of nodes each one containing one or more genes. Thus, two kinds of adjacency matrices are created. The compact adjacency matrix retains the groupings and stores edge types between genes and genes, genes and groups of genes or between group of genes. The expanded adjacency matrix stores edge type information between individual genes.

Value

A list containing a list of compact adjacency matrices, a list of expanded adjacency matrices, and list detailing all nodes, edges and interaction types.

References

Examples

# Download Insulin Signaling Pathway
pathways <- c('04915', '04917', '04930', '05031')
paths <- downloadPathways(org='hsa', pathways=pathways)

# Create pathway graph
graphs <- createPathwayGraphs(org='hsa', pathways=paths)

downloadKEGGPathwayList

Retrieve all available pathways for an organism.

Description
Retrieve all available pathways for an organism.

Usage
downloadKEGGPathwayList(org)

Arguments
org KEGG organism identifier.

Details

Value
Data frame of pathway ids and names.

References
• http://www.genome.jp/kegg/pathway.html

Examples

# Load extracted linear subpathways from toy data
load(system.file('extdata','Examples//data.RData', package='CHRONOS'))

# Retrieve all available hsa pathways
## Not run: pathways <- downloadKEGGPathwayList(org='hsa')
downloadMiRecords  

**Download miRNA-mRNA interactions for an organism.**

**Description**

Download miRNA-mRNA interactions for an organism.

**Usage**

```r
downloadMiRecords(org, pn, update, databases)
```

**Arguments**

- `org`  KEGG organism identifier.
- `pn`  Number of databases that verify miRNA-mRNA interactions.
- `update`  Download preprocessed data (update=FALSE) or new data from miRecords (update=TRUE).
- `databases`  Specify which miRNA-mRNA interaction databases will be used.

**Details**

miRecords is a resource for animal miRNA-target interactions. The Predicted Targets component of miRecords is an integration of predicted miRNA targets produced by 11 established miRNA target prediction tools, namely DIANA-microT, MicroInspector, miRanda, MirTarget2, miTarget, NBmiRTar, PicTar, PITA, RNA22, RNAhybrid, and TargetScan/TargetScanS.

**Value**

Downloaded data is stored in CHRONOS/extdata/Downloads/miRecords/<org>/miRNATargets.RData

**References**

- [http://c1.accurascience.com/miRecords](http://c1.accurascience.com/miRecords)

**Examples**

```r
# Load extracted linear subpathways from toy data
load(system.file('extdata', 'Examples//data.RData', package='CHRONOS'))

## Not run: downloadMiRecords(org='hsa', pn=5, update=FALSE, databases='All')
```
downloadPathways  

Download KEGG pathways in KGML format.

Description

Download KEGG pathways in KGML format.

Usage

downloadPathways(org, pathways)

Arguments

<table>
<thead>
<tr>
<th>Argument</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>org</td>
<td>KEGG organism identifier</td>
</tr>
<tr>
<td>pathways</td>
<td>Download pathways for specified organism:</td>
</tr>
<tr>
<td></td>
<td>'All' All organism pathways</td>
</tr>
<tr>
<td></td>
<td>'Metabolic' Metabolic pathways</td>
</tr>
<tr>
<td></td>
<td>'Non-Metabolic' Non metabolic pathways</td>
</tr>
<tr>
<td></td>
<td>&lt;vector of indexes&gt; Using indexes from downloadKEGGPathwayList</td>
</tr>
<tr>
<td></td>
<td>&lt;vector of names&gt; Using pathway identifiers (i.e. c('00010', '00020'))</td>
</tr>
</tbody>
</table>

Details

KEGG (Kyoto Encyclopedia of Genes and Genomes) is a database resource for understanding high-level functions and utilities of the biological system such as the cell, the organism and the ecosystem, from molecular-level information, especially large-scale molecular datasets generated by genome sequencing and other high-throughput experimental technologies.

Files are downloaded in CHRONOS/extdata/Downloads/KEGG/<org> folder. Downloading is skipped for existing files.

Value

Downloaded data is stored in CHRONOS/extdata/Downloads/KEGG/<org>

References


Examples

# View all available hsa pathways
pathways <- downloadKEGGPathwayList(org='hsa')

# Download pathway KGML files
pathways <- c('04915', '04917', '04930', '05031')

pathways <- downloadPathways(org='hsa', pathways=pathways)
Description

Linear subpathway extraction from pathway graphs

Usage

\[
\text{extractLinearSubpathways}(\text{graphs}, \text{pathways}, a, b, \text{filter}, \text{export}, \text{groupMode}, \text{verbose})
\]

Arguments

graphs Pathway graphs as returned from \text{createPathwayGraphs}.
pathways The subset of pathways from whom subpathways are to be extracted. If missing, all pathway graphs are used.
a Minimum subpathway length.
b Maximum subpathway length.
filter Filter the subpaths with user genes (TRUE).
export Exports subpaths in CHRONOS/extdata/Output/Subpaths/Linear/<org> folder. Available formats are '.txt' and/or '.RData'.
groupMode Expand paralogues ('expand') or collapse them to a single entry ('collapse').
verbose Display informative messages (TRUE). Requires previous execution of \text{importExpressions}.

Details

Subpath filtering supports the removal of subpaths that have at least one member not belonging to the set of user supplied genes. These genes are extracted from the user’s mRNA expressions matrix. Thus, the execution of \text{importExpressions} is a prerequisite.

To extract linear subpathways from a pathway graph, all possible start and end nodes are considered. A start node has only outgoing edges while an end node only has incoming edges. For each such pair, all linear subpathways are found by traversing the corresponding graph. Since the initial pathway graph’s nodes contain one or more genes, resulting subpathways consist of bins of one or more genes. These subpaths are expanded to subpathways with one gene per bin in order to obtain usable subpathways.

Value

Returns a list consisting of

- A matrix of linear subpathways (subpaths)
- A list of processed pathway graphs adjacency matrices (adjMats)
- A list of processed pathway genes and interactions between them (lexicon)
Examples

# Load pathway graphs from toy data
load(system.file('extdata', 'Examples//data.RData', package='CHRONOS'))

# Extract linear subpathways
linSubs <- extractLinearSubpathways(graphs=graphs)

extractNonLinearSubpathways

Non linear subpathway extraction from pathway graphs

Description

Non linear subpathway extraction from pathway graphs

Usage

extractNonLinearSubpathways(graphs, pathways, a, b, k, filter, groupMode, export, verbose)

Arguments

graphs Pathway graphs as returned from createPathwayGraphs.
pathways The subset of pathways from whom subpathways are to be extracted. If missing, all pathway graphs are used.
a Minimum subpathway length.
b Maximum subpathway length.
k Clique size.
filter Filter the subpaths with user genes (TRUE).
groupMode Expand paralogues (‘expand’) or collapse them to a single entry (‘collapse’).
export Exports subpaths in CHRONOS/extdata/Output/Subpaths/Non-Linear/ <org> folder. Available formats are ‘.txt’ and/or ‘.RData’.
verbose Display informative messages (TRUE) Requires previous execution of importExpressions.

Value

Returns a list consisting of

- A matrix of linear subpathways (subpaths)
- A list of processed pathway graphs adjacency matrices(adjMats)
- A list of processed pathway genes and interactions between them (lexicon)

To extract non linear subpaths from a pathway graph, all interactions between nodes of belonging to k-cliques are found. The ones that correspond.
To extract non linear subpaths from a pathway graph, all interactions between nodes of belonging to k-cliques are found. The ones that correspond to actual interactions between genes make up the non linear subpath.
getEdgeTypes

Examples

# Load pathway graphs from toy data
load(system.file('extdata', 'Examples//data.RData', package='CHRONOS'))

# Extract linear subpathways
nliSubs <- extractNonLinearSubpathways(graphs=graphs)

getEdgeTypes

Map various types of gene-gene interactions in KGML files to edge types in corresponding pathway graphs.

Description

Map various types of gene-gene interactions in KGML files to edge types in corresponding pathway graphs.

Usage

getEdgeTypes(type)

Arguments

type  A vector of interaction types.

Details

Edge types

activation 1 inhibition 2 apathetic 3 no interaction 4

Default interaction - edge type mapping

<p>| | | | | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>01</td>
<td>unknown</td>
<td>3</td>
<td>02</td>
<td>activation</td>
</tr>
<tr>
<td>03</td>
<td>inhibition</td>
<td>2</td>
<td>04</td>
<td>binding/association</td>
</tr>
<tr>
<td>05</td>
<td>expression</td>
<td>1</td>
<td>06</td>
<td>repression</td>
</tr>
<tr>
<td>07</td>
<td>phosphorylation</td>
<td>3</td>
<td>08</td>
<td>dephosphorylation</td>
</tr>
<tr>
<td>09</td>
<td>ubiquitination</td>
<td>3</td>
<td>10</td>
<td>dissociation</td>
</tr>
<tr>
<td>11</td>
<td>indirect effect</td>
<td>3</td>
<td>12</td>
<td>state change</td>
</tr>
<tr>
<td>13</td>
<td>compound</td>
<td>3</td>
<td>14</td>
<td>hidden compound</td>
</tr>
<tr>
<td>16</td>
<td>missing interaction</td>
<td>3</td>
<td>16</td>
<td>activation_phosphorylation</td>
</tr>
<tr>
<td>17</td>
<td>activation_dephosphorylation</td>
<td>1</td>
<td>18</td>
<td>activation_ubiquitination</td>
</tr>
<tr>
<td>19</td>
<td>activation_indirect effect</td>
<td>1</td>
<td>20</td>
<td>activation_binding/association</td>
</tr>
<tr>
<td>21</td>
<td>activation_inhibition</td>
<td>3</td>
<td>22</td>
<td>activation_methylation</td>
</tr>
<tr>
<td>23</td>
<td>inhibition_phosphorylation</td>
<td>2</td>
<td>24</td>
<td>inhibition_dephosphorylation</td>
</tr>
<tr>
<td>25</td>
<td>inhibitionUbiquitination</td>
<td>2</td>
<td>26</td>
<td>inhibition_indirect effect</td>
</tr>
<tr>
<td>27</td>
<td>inhibition_binding/association</td>
<td>2</td>
<td>28</td>
<td>inhibition_expression</td>
</tr>
<tr>
<td>29</td>
<td>inhibition_methylation</td>
<td>2</td>
<td>30</td>
<td>compound_expression</td>
</tr>
<tr>
<td>31</td>
<td>compound_activation</td>
<td>1</td>
<td>32</td>
<td>compound_inhibition</td>
</tr>
</tbody>
</table>
importExpressions

33 compound_activation_indirect effect 1
34 compound_activation_phosphorylation 1
35 phosphorylation_indirect effect 3
36 phosphorylation_binding/association 3
37 phosphorylation_dissociation 3
38 dephosphorylation_indirect effect 3
39 binding/association_missing interaction 3
40 binding/association_indirect effect 3
41 expression_indirect effect 1
42 repression_indirect effect 2
43 ubiquitination_inhibition 2
44 dissociation_missing interaction 3
45 indirect_effect_phosphorylation 3
46 activation_phosphorylation_binding/association 1
47 activation_phosphorylation_indirect effect 1

Value

If an interaction type has been supplied, the corresponding edge types are returned. If not, the complete mapping is returned.

Examples

# Example 1

# Retreive edge types for phosphorylation and dephosphorylation.
getEdgeTypes(c(7,8))

# Example 2

# Returns a data frame containing the interaction - edge type mapper.
types <- getEdgeTypes()

# Set phosphorylation to inhibition.
types[8,2] <- 2

importExpressions \hspace{1cm} \textit{Import gene and miRNA expressions from}

Description

Import gene and miRNA expressions from

Usage

importExpressions(data, type, sep, org, mRNAomenclature)
pathwayMeasures

Arguments

data          Expressions data filename or matrix.
type          Expressions data type. (or mRNA expressions, type=<nomenType>). Available gene expression nomenclature can be found in convertNomenclature. For miRNA expressions, type='miRNA'.
sep            File delimiter.
org            KEGG organism identifier
mRNAnomenclature

Details

• Import gene expressions data from CHRONOS/extdata/Input/<userFile>.txt or a supplied matrix.
• Import miRNA expressions data from CHRONOS/extdata/Input/<userFile>.txt or a supplied matrix.

Value

Examples

# Example
load(system.file("extdata", "Examples//data.RData", package="CHRONOS"))
importExpressions(data=mRNAexpr, type='mRNA',
                  mRNAnomenclature='entrezgene', sep='\t', org='hsa')
importExpressions(data=miRNAexpr, type='miRNA', sep='\t', org='hsa')

pathwayMeasures               Pathway structural and functional aspects

Description

Pathway structural and functional aspects

Usage

pathwayMeasures(graphs)

Arguments

graphs          Pathway graphs as returned from createPathwayGraphs.

Details

Structural and functional aspects of a pathway are calculated in respect to all organism pathways.
Value

Matrix with pathness, betweenness centrality and degree values for each gene in the pathway graphs at its columns.

Examples

# Load pathway graphs from toy data
load(system.file('extdata', 'Examples/data.RData', package='CHRONOS'))

# Calculate pathway structural and functional aspects
measures <- pathwayMeasures(graphs)

scoreSubpathways

Evaluate subpathways using an interacting scoring scheme (IS) for each time point.

Description

Evaluate subpathways using an interacting scoring scheme (IS) for each time point.

Usage

scoreSubpathways(subpathways, filters, measures, parameters, miRNAinteractions)

Arguments

subpathways Subpaths as returned from extractLinearSubpathways and extractNonLinearSubpathways.
filters Named vector of filters used for subpathway evaluation. Values denote corresponding thresholds.
    pvalue Statistical evaluation
    measures Structural and functional evaluation
    subScore mRNA-mRNA interaction scoring
    mirScore miRNA-mRNA interaction scoring
measures Subpathway structural and functional aspects as returned from pathwayMeasures.
parameters C.K.T parameters of scoring scheme.
miRNAinteractions An edgelist of miRNA-mRNA interactions used to override downloaded interactions from miRecords.

Details

...

Value
subpathwayKEGGmap

Create links to KEGG pathway map with highlighted subpathways.

Description

Create links to KEGG pathway map with highlighted subpathways.

Usage

subpathwayKEGGmap(subpathways, type, openInBrowser)

Arguments

- subpathways: Subpathways as returned by extractLinearSubpathways or extractNonLinearSubpathways
- type: Subpathway type (Linear, Non-Linear)
- openInBrowser: Open link in default browser.

Value

Vector of links of KEGG pathway maps.

References


Examples

# Load extracted subpathways from toy data
load(system.file('extdata', 'Examples/data.RData', package='CHRONOS'))

# Import mRNA expressions
mRNAexpr <- importExpressions(data=mRNAexpr, type='mRNA', org='hsa')

# Score extracted linear subpathways
filters <- c('subScore'=0.4)
linSubsScored <- scoreSubpathways(subpathways=linSubs, filters=filters)
Examples

# Load extracted linear subpathways from toy data
load(system.file('extdata', 'Examples//data.RData', package='CHRONOS'))

# Opening selected subpathways in default browser
subs <- linSubs$subpaths[1:3,]
subpathwayKEGGmap(subpathways=subs, type='Linear', openInBrowser=FALSE)

subpathwayMiRNAs

Create a circulat plot of a subpathway and the miRNAs that target it.

Description

Create a circulat plot of a subpathway and the miRNAs that target it.

Usage

subpathwayMiRNAs(summary, subIdx, timePoints)

Arguments

summary Output from scoreSubpathways
subIdx Subpathway index
timePoints Time points to include in visualization, default to all.

Value

.

Examples

# Load scored subpathways from toy data
load(system.file('extdata', 'Examples//data.RData', package='CHRONOS'))
# Visualize one or more subpathways.
subpathwayMiRNAs(summary=linSubsScored, subIdx=2)
visualizeResults

**visualizeResults**  Visualize results in tabular form (txt, xlsx)

**Description**

Visualize results in tabular form (txt, xlsx)

**Usage**

```r
visualizeResults(summary, export, expand, colors, from, to)
```

**Arguments**

- `summary` Evaluation results as returned from `scoreSubpathways`
- `export` `.xlsx' exports a xlsx file and '.txt' a .txt file.
- `expand` TRUE if each subpathway member and miRNA belongs to a single cell, FALSE if all subpathway members belong to one cell and miRNAs to another cell.
- `colors` The color scheme used in subScores heatmap.
- `from` Primary annotation `convertNomenclature`. Defaults to EntrezGene ID.
- `to` Secondary annotation `convertNomenclature`

**Value**

A txt or a xlsx file in CHRONOS/extdata/Output/Scores/Linear/<org> or CHRONOS/extdata/Output/Scores/Non-Linear/<org>

**Examples**

```r
# Load scored subpathways from toy data
load(system.file("extdata", "Examples\data.RData", package="CHRONOS"))

visualizeResults(linSubsScored, export='txt')
```
<table>
<thead>
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<th>Index</th>
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</thead>
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<td>convertNomenclature, 2, 4, 13, 17</td>
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<td>createPathwayGraphs, 5, 9, 10, 13</td>
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<td>downloadKEGGPathwayList, 6, 8</td>
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