

Package ‘SBGNview.data’

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Title Demo gene expression datasets for SBGNview package

Description This package contains:

1. A microarray gene expression dataset from a human breast cancer study.
2. A RNA-Seq gene expression dataset from a mouse study on IFNG knockout.
3. ID mapping tables between gene IDs and SBGN-ML file glyph IDs.
4. Percent of orthologs detected in other species of the genes in a pathway. Cutoffs of this percentage for defining if a pathway exists in another species.
5. XML text of SBGN-ML files for all pre-collected pathways.

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Depends R (>= 3.6)

License AGPL-3

Collate data.R

LazyData FALSE

Imports knitr, rmarkdown

Suggests SummarizedExperiment

RoxygenNote 6.1.1

VignetteBuilder knitr

biocViews ExperimentData, CancerData, BreastCancerData,
MicroarrayData, GEO, RNASeqData

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cancer.ds

*A demo microarray dataset from a cancer study***Description**

A demo microarray dataset from a cancer study

Usage

cancer.ds

Format

A SummarizedExperiment object.

Details

This dataset is constructed using the first three columns of data `**gse16873.d**` in package `**pathview**` (i.e. columns "DCIS_1", "DCIS_2" and "DCIS_3"). The original values were used without additional processing. It is constructed for showing SGBNview's visualization ability, not for data analysis. Each column in the assay table is a pair of cancer-v.s.-control samples. The value of a gene in a column is the log fold change of this gene in the corresponding pair of cancer-v.s.-control samples.

IFNg

*RNA-Seq result from a mouse IFNG knockout experiment***Description**

RNA-Seq result from a mouse IFNG knockout experiment

Usage

IFNg

Format

A SummarizedExperiment object.

Details

This RNA-Seq dataset contains RNA abundance table of two groups: IFNG knockout mice and wild type mice. RNA abundance values were log2 transformed. For demo purpose, data of 4 IFNG knockout mice and 4 wild type mice were included. The experiment and data processing was described in this work: Greer, Renee L., Xiaoxi Dong, et al. "Akkermansia muciniphila mediates negative effects of IFNG on glucose metabolism." *Nature communications* 7 (2016): 13329.

mapping

Mapping table between two types of IDs

Description

Mapping table between two types of IDs

Usage

ENZYME_pathway.id

hsa_KO_ENTREZID

hsa_pathwayCommons_ENTREZID

mmu_KO_ENSEMBL

chebi_pathway.id

mmu_KO_ENTREZID

chebi_CompoundName

CompoundName_pathwayCommons

kegg.ligand_pathwayCommons

hsa_pathwayCommons_ENSEMBL

mmu_pathwayCommons_ENTREZID

KO_pathway.id

KO_pathwayCommons

SYMBOL_pathway.id

pathwayCommons_SYMBOL

chebi_pathwayCommons

mmu_pathwayCommons_ENSEMBL

hsa_ENTREZID_pathwayCommons

Format

A matrix with two columns: the ID mapping between two types of IDs.

Details

Each dataset contains a mapping table. There are several types of ID pairs, such as molecule ID \leftrightarrow pathway_glyph_ID, molecule ID \leftrightarrow pathway ID, and molecule ID \leftrightarrow KEGG ortholog ID. molecule ID \leftrightarrow pathway_glyph_ID tables are extracted from Biopax files. For example: <http://www.pathwaycommons.org/archives/PC3/v10/PathwayCommons10.reactome.BIOPAX.owl.gz>. Glyph IDs are extracted from the ID of each XML element "Protein". Its matching molecule ID is extracted from the corresponding XML child element "UnificationXref". See more details and examples in vignette 'SBGNview.data.vignette'

pathway.completeness.cutoff.info

Cutoffs of pathway completeness used for defining existence of pathway in a species

Description

Cutoffs of pathway completeness used for defining existence of pathway in a species

Usage

pathway.completeness.cutoff.info

Format

A matrix

Details

PathwayCommons only annotated human pathways, we mapped pathwayCommons' genes to other species using KEGG ortholog annotation. As a result, not all of the genes have corresponding genes in another species. We call the percentage of mapped genes the "coverage or completeness" in the species. To determine if a pathway exists in a species, we use a cutoff for this completeness. This cutoff is selected using the following approach: 1. A pathway has different completeness in different species thus form a completeness vector across all species (vector C). 2. Use a completeness cutoff we can define whether this pathway "exists" in a species, thus form a label vector E (a pathway "Exist" or "not Exist" across all species). 3. Use one way ANOVA to calculate F statistic of completeness between the two groups ("Exist" or "not Exist"), thus one cutoff will have one F statistic. 4. Try different cutoffs(unique completeness values in vector C) and select the one with the largest F statistic, i.e. the cutoff that can maximize the difference between "Exist" and "not Exist" groups. This is not a perfect way to define if a pathway exists in a species, but can serve as a filter criteria.

pathway.species.pct_Mapped

Pathway completeness in a species

Description

Pathway completeness in a species

Usage

pathway.species.pct_Mapped

Format

A matrix

Details

PathwayCommons only annotated human pathways, we mapped pathwayCommons' genes to other species using KEGG ortholog annotation. As a result, not all of the genes have corresponding genes in another species. We call the percentage of mapped genes the "coverage or completeness" in the species.

sbgn.xmls

XML code of a SBGN-ML file

Description

XML code of a SBGN-ML file

Usage

sbgn.xmls

Format

A list of character strings

Details

Each string is the full XML code of a SBGN-ML file. It includes glyphs and arcs of a SBGN map.

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