Package 'ToPASeq'

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Title Topology-based pathway analysis of RNA-seq data

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Description Implementation of methods for topology-based pathway analysis of RNA-seq data. This includes Topological Analysis of Pathway Phenotype Association (TAPPA; Gao and Wang, 2007), PathWay Enrichment Analysis (PWEA; Hung et al., 2010), and the Pathway Regulation Score (PRS; Ibrahim et al., 2012).

Depends R(>= 3.5.0), graphite

Imports Rcpp, graph, methods

Suggests BiocStyle, EnrichmentBrowser, airway, knitr, rmarkdown

LinkingTo Rcpp

LazyData yes

License AGPL-3

biocViews GeneExpression, RNASeq, DifferentialExpression, GraphAndNetwork, Pathways, NetworkEnrichment, Visualization

VignetteBuilder knitr

RoxygenNote 6.0.1

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R topics documented:

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Description

prs

This function implements the PRS method to analyze pathway enrichment of gene expression data. For PRS, a gene weight correspond to the number of downstream differentially expressed genes.

Usage

```
prs(de, all, pwys, nperm = 1000)
```

prsWeights(pwy, de, all)

Arguments

de	A named numeric vector containing log2 fold-changes of the differentially expressed genes. Recommended names are Entrez gene IDs.
all	A character vector with the gene IDs in the reference set. If the data was obtained from a gene expression experiment, this set will contain all genes measured in the experiment. This vector should contain *all* names of the de argument.
pwys	A linkS4class{PathwayList} containing the pathways that should be analyzed for enrichment.
nperm	Integer. Number of permutations.
рwy	A linkS4class{Pathway} for which the weights should be computed.

Value

A data.frame with normalized score and p-value for each pathway analyzed.

Author(s)

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References

Ibrahim et al. (2012) A topology-based score for pathway enrichment. J Comput Biol, 19(5):563-73.

See Also

pathways

Examples

```
# pathways
library(graphite)
pwys <- pathways("hsapiens","kegg")[1:10]</pre>
```

expression data

all <- nodes(pwys[[1]]) nds <- sample(all, 30) de <- setNames(rnorm(30), nds)

executing PRS
prsWeights(pwys[[1]], de, all)
prs(de, all, pwys, nperm=100)

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