

Package ‘DEHOGT’

September 13, 2024

Type Package

Title Differentially Expressed Heterogeneous Overdispersion Gene Test
for Count Data

Version 0.99.0

Description Implements a generalized linear model approach for detecting differentially expressed genes across treatment groups in count data. The package supports both quasi-Poisson and negative binomial models to handle over-dispersion, ensuring robust identification of differential expression. It allows for the inclusion of treatment effects and gene-wise covariates, as well as normalization factors for accurate scaling across samples. Additionally, it incorporates statistical significance testing with options for p-value adjustment and log₂ fold range thresholds, making it suitable for RNA-seq analysis as described in by Xu et al., (2024) <[doi:10.1371/journal.pone.0300565](https://doi.org/10.1371/journal.pone.0300565)>.

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Encoding UTF-8

Depends R (>= 3.5.0)

Imports doParallel, foreach, MASS,

Suggests knitr, rmarkdown, BiocStyle

biocViews GeneExpression, DifferentialExpression, StatisticalMethod,
Regression, Normalization

VignetteBuilder knitr

RoxygenNote 7.3.2

URL <https://github.com/ahshen26/DEHOGT>

BugReports <https://github.com/ahshen26/DEHOGT/issues>

NeedsCompilation no

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|-------------|--|

Description

Differentially Expressed Heterogeneous Overdispersion Genes Testing for Count Data This script implements the main function of the proposed method in the above paper

Usage

```
dehogt_func(
  data,
  treatment,
  norm_factors = NULL,
  covariates = NULL,
  dist = "qpois",
  padj = TRUE,
  pval_thre = 0.05,
  l2fc = FALSE,
  l2fc_thre = 1,
  num_cores = 1
)
```

Arguments

| | |
|--------------|---|
| data | A matrix of gene expression data where rows represent genes and columns represent samples. |
| treatment | A vector specifying the treatment conditions for each sample. |
| norm_factors | An optional vector of normalization factors for each sample. Default is NULL, which assumes equal normalization factors. |
| covariates | An optional matrix of gene-wise covariates. Default is NULL. |
| dist | The distribution family for the GLM. Can be "qpois" for quasi-Poisson or "neg-bin" for negative binomial. Default is "qpois". |

| | |
|------------------------|---|
| <code>padj</code> | Logical value indicating whether to adjust p-values using the Benjamini-Hochberg (BH) procedure. Default is TRUE. |
| <code>pval_thre</code> | The threshold for identifying differentially expressed genes based on adjusted p-values. Default is 0.05. |
| <code>l2fc</code> | Logical value indicating whether to consider log ₂ fold change for identifying differentially expressed genes. Default is FALSE. |
| <code>l2fc_thre</code> | The threshold for log ₂ fold change in identifying differentially expressed genes. Default is 1. |
| <code>num_cores</code> | The number of CPU cores to use for parallel computing. Default is 1. |

Value

A list containing:

| | |
|---------------------|--|
| <code>DE_idx</code> | A logical vector indicating differentially expressed genes. |
| <code>pvals</code> | A numeric vector of p-values for each gene. |
| <code>log2fc</code> | A numeric vector of log ₂ fold changes for each gene. |

Examples

```
# simulate gene expression data
data <- matrix(rpois(1000, 10), nrow = 100, ncol = 10)
# simulate random treatment assignments
treatment <- sample(0:1, 10, replace = TRUE)
# Run main function with parallel computing using 2 cores
result <- dehogt_func(data, treatment, num_cores = 2)
```

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