Package ‘diffUTR’

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Type Package

Title diffUTR: Streamlining differential exon and 3’ UTR usage

Version 1.10.0

Depends R (>= 4.0)

Description The diffUTR package provides a uniform interface and plotting functions for limma/edgeR/DEXSeq -powered differential bin/exon usage. It includes in addition an improved version of the limma::diffSplice method. Most importantly, diffUTR further extends the application of these frameworks to differential UTR usage analysis using poly-A site databases.

Imports S4Vectors, SummarizedExperiment, limma, edgeR, DEXSeq, GenomicRanges, Rsubread, ggplot2, rtracklayer, ComplexHeatmap, ggrepel, stringi, methods, stats, GenomeInfoDb, dplyr, matrixStats, IRanges, ensembldb, viridisLite

Suggests BiocStyle, knitr, rmarkdown

biocViews GeneExpression

BugReports https://github.com/ETHZ-INS/diffUTR

VignetteBuilder knitr

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Description

addNormalizedAssays

Usage

addNormalizedAssays(se, readLength = 50L)

Arguments

se
readLength

A bin-wise 'SummarizedExperiment' as produced by countFeatures
Used as a minimum width to estimate read density (default 50).

Value

The 'se' object with populated 'logcpm' and 'logNormDensity' assays.

Examples

data(example_bin_se)
example_bin_se <- addNormalizedAssays(example_bin_se)
countFeatures

countFeatures

Description

countFeatures

Usage

countFeatures(
  bamfiles,
  bins,
  strandSpecific = 0,
  readLength = 50L,
  allowMultiOverlap = TRUE,
  inclNormalized = TRUE,
  tmpDir = tempdir(),
  ...
)

Arguments

bamfiles       A vector of paths to bam files
bins          A GRanges of bins in which to count reads (or path to a rds file containing such an object
strandSpecific Pased to 'Rsubread::featureCounts'
readLength     Used as a minimum width to estimate read density.
allowMultiOverlap Passed to 'Rsubread::featureCounts'
inclNormalized Logical; whether to include normalized assays (needed for plotting)
tmpDir         Passed to 'Rsubread::featureCounts'
...            Passed to 'Rsubread::featureCounts'

Value

A RangedSummarizedExperiment-class

Examples

data("example_gene_annotation", package="diffUTR")
bins <- prepareBins(example_gene_annotation)
bam_files <- list.files(system.file("extdata", package="diffUTR"),
  pattern="bam$", full=TRUE)
# not run
# se <- countFeatures(bam_files, bins, verbose=FALSE)
deuBinPlot

Description

deuBinPlot

Usage

deuBinPlot(
  se,
  gene,
  type = c("summary", "condition", "sample"),
  intronSize = 2,
  exonSize = c("sqrt", "linear", "log"),
  y = NULL,
  condition = NULL,
  size = "type",
  lineSize = 1,
  colour = NULL,
  alpha = NULL,
  removeAmbiguous = TRUE,
  minDensityRatio = 0.1
)

Arguments

se A bin-wise SummarizedExperiment as produced by countFeatures and including bin-level tests (i.e. having been passed through one of the DEU wrappers such as diffSpliceWrapper or DEXSeqWrapper)
gene The gene of interest
type Either 'summary' (plot DEU summary), 'sample' (plot sample-wise data), or 'condition' (plot data aggregate by condition)
intronSize Intron plot size. If <=3, intron size will be this fraction of the mean exon size. If >3, each intron will have the given size.
exonSize Scaling for exon sizes, either 'sqrt', 'log', or 'linear'.
y Value to plot on the y-axis. If 'type="summary"', this should be a column of 'rowData(se)', otherwise should be an assay name of 'se'.
condition The colData column containing the samples’ condition.
size rowData variable to use to determine the thickness of the bins.
lineSize Size of the line connecting the bins. Use ‘lineSize=0’ to omit the line.
colour rowData variable to use to determine the colour of the bins. If 'type="condition"', can also be "condition"; if 'type="sample"' can be any colData column.
alpha Alpha level, passed to ggplot.
**diffSplice2**

removeAmbiguous

Logical; whether to remove bins that are gene-ambiguous (i.e. overlap multiple genes).

minDensityRatio

Minimum ratio of read density (with respect to the gene’s average) for a bin to be plotted.

**Value**

A ggplot object

**Examples**

data(example_bin_se)
se <- diffSpliceWrapper(example_bin_se, ~condition)
deuBinPlot(se, "Jund")

**Description**

This is a small improvement to the **diffSplice** function written by Gordon Smyth and Charity Law.

**Usage**

diffSplice2(fit, geneid, exonid = NULL, robust = FALSE, verbose = TRUE)

**Arguments**

- **fit**: an **MArrayLM-class** fitted model object produced by `lmFit` or ‘contrasts.fit’, with rows corresponding to exons.
- **geneid**: gene identifiers (as in **diffSplice**)  
- **exonid**: exon identifiers (as in **diffSplice**)  
- **robust**: logical, should the estimation of the empirical Bayes prior parameters be robustified against outlier sample variances?  
- **verbose**: logical, if TRUE will output some diagnostic information

**Value**

An **MArrayLM-class** object containing both exon level and gene level tests. Results are sorted by geneid and by exonid within gene.
Examples

```r
library(SummarizedExperiment)
library(edgeR)
data(example_bin_se)
se <- example_bin_se
design <- model.matrix(~condition, data=as.data.frame(colData(se)))
.dds <- calcNormFactors(DGEList(assays(se)$counts))
.dds <- voom(dds, design)
.dds <- lmFit(dds, design)
res <- diffSplice2(dds, geneid=rowData(se)$gene, exonid=row.names(se))
topSplice(res)
```

Description

Wrappers around commonly-used DEU methods (`diffSpliceDGE`, `DEXSeq` and an improved version of `diffSplice`)

Usage

```r
diffSpliceDGEWrapper(  
  se,  
  design,  
  coef = NULL,  
  QLF = TRUE,  
  robust = TRUE,  
  countFilter = TRUE,  
  excludeTypes = NULL  
)

diffSpliceWrapper(  
  se,  
  design,  
  coef = NULL,  
  robust = TRUE,  
  improved = TRUE,  
  countFilter = TRUE,  
  excludeTypes = NULL  
)

DEXSeqWrapper(  
  se,  
  design = ~sample + exon + condition:exon,  
  reducedModel = ~sample + exon,  
  excludeTypes = NULL,  
)```
Arguments

se
A bin-wise SummarizedExperiment as produced by countFeatures.

design
A formula (using columns of 'colData(se)') or (for 'diffSpliceWrapper' or 'diffSpliceDGEWrapper' only) a model.matrix.

design
The coefficient to be tested (ignored for 'DEXSeqWrapper').

QLF
Logical; whether to use edgeR’s quasi-likelihood negative binomial (applicable only to 'diffSpliceDGEWrapper').

robust
Logical; whether to use robust fitting for the dispersion trend (ignored for 'DEXSeqWrapper').

countFilter
Logical; whether to filter out low-count bins (ignored for 'DEXSeqWrapper').

excludeTypes
A vector of bin types to ignore for testing. To test for any kind of differential usage, leave empty. To test for differential UTR usage, use 'excludeTypes=c("CDS","non-coding")' (or see geneLevelStats for more options).

improved
Logical; whether to use diffSplice2 instead of the original diffSplice (default TRUE).

reducedModel
A reduced formula (applicable only to 'DEXSeqWrapper').

... Further arguments (passed to ‘testForDEU’ and ‘estimateExonFoldChanges’) of ‘DEXSeq’. Can for instance be used to enable multithreading, by passing ‘BPPARAM=BiocParallel::MulticoreParam(ncores)’.

Value

The ‘se’ object with additional rowData columns contain bin (i.e. exon) -level statistics, and a metadata slot containing gene level p-values.

Examples

library(SummarizedExperiment)
data(example_bin_se)
se <- diffSpliceWrapper(example_bin_se, ~condition)
head(rowData(se))

Description

An object produced by countFeatures containing small subset of genes from mouse hippocampal slices undergoing Forskolin-induced long-term potentiation (GSE84643).
Value

a ‘RangedSummarizedExperiment’

References

https://www.nature.com/articles/s41598-017-17407-w

example_gene_annotation

Example gene annotation

Description

An example gene annotation containing only a small subset of mouse genes.

Value

a ‘GRanges’ object

geneBinHeatmap
geneBinHeatmap

description

A wrapper around ‘ComplexHeatmap’.

Usage

geneBinHeatmap(
  se,
  gene,
  what = NULL,
  anno_rows = c("type", "logWidth", "meanLogDensity", "log10PValue", "geneAmbiguous"),
  anno_columns = c(),
  anno_colors = list(),
  removeAmbiguous = FALSE,
  merge_legends = TRUE,
  cluster_columns = FALSE,
  minDensityRatio = 0.1,
  left_annotation = NULL,
  top_annotation = NULL,
  ...)
)
geneLevelStats

Arguments

- se: A bin-wise SummarizedExperiment as produced by `countFeatures`
- gene: The gene of interest
- what: Type of values (i.e. assay) to plot
- anno_rows: Row annotation columns (i.e. columns of ‘rowData(se)’) to plot
- anno_columns: Column annotation columns (i.e. columns of ‘colData(se)’) to plot
- anno_colors: Annotation colors, as a list named with the row/column annotations, see `SingleAnnotation` for details. Ignored if ‘left_annotation’ and/or ‘top_annotation’ are given directly.
- removeAmbiguous: Logical; whether to remove bins that are gene-ambiguous (i.e. overlap multiple genes).
- merge_legends: Logical; whether to merge legends. This effectively calls `draw(..., merge_legends=TRUE)` around the heatmap.
- cluster_columns: Logical; whether to cluster columns (passed to `Heatmap`)
- minDensityRatio: Minimum ratio of read density (with respect to the gene’s average) for a bin to be plotted.
- left_annotation: Passed to `Heatmap`, overrides `anno_rows`.
- top_annotation: Passed to `Heatmap`, overrides `anno_columns`.
- ...: Passed to ‘ComplexHeatmap’ (see `Heatmap`)

Value

A `Heatmap`

Examples

```r
data(example_bin_se)
se <- diffSpliceWrapper(example_bin_se, ~condition)
geneBinHeatmap(se, "Jund")
```

Description

Aggregates bin-level statistics to the gene-level
Usage

geneLevelStats(
  se,
  coef = NULL,
  excludeTypes = NULL,
  includeTypes = NULL,
  returnSE = TRUE,
  minDensityRatio = 0.1,
  minWidth = 20,
  excludeGeneAmbiguous = TRUE
)

Arguments

se           A 'RangedSummarizedExperiment' containing the results of one of the DEU wrappers.
coef         The coefficients tested (if the model included more than one term).
excludeTypes Vector of bin types to exclude.
includeTypes Vector of bin types to include (overrides 'excludeTypes')
returnSE     Logical; whether to return the updated 'se' object (default), or the gene-level table.
minDensityRatio Minimum ratio of read density (with respect to the gene’s average) for a bin to be included.
minWidth     Minimum bin width to include
excludeGeneAmbiguous Logical; whether to exclude bins which are ambiguous (i.e. can be from different genes)

Value

If 'returnSE=TRUE' (default), returns the 'se' object with an updated ‘metadata(se)$geneLevel’ slot, otherwise returns the gene-level data.frame.

Examples

library(SummarizedExperiment)
data(example_bin_se)
se <- diffSpliceWrapper(example_bin_se, ~condition)
se <- geneLevelStats(se, includeTypes="3UTR")
head(metadata(se)$geneLevel)
Description

plotTopGenes

Usage

plotTopGenes(se, n = 10, FDR = 0.05, diffUTR = FALSE, alpha = 1, ...)

Arguments

se A bin-wise SummarizedExperiment as produced by countFeatures and including bin-level tests (i.e. having been passed through one of the DEU wrappers such as diffSpliceWrapper or DEXSeqWrapper)
n The maximum number of genes for which to plot labels
FDR The FDR threshold above which to plot labels
diffUTR Logical; if FALSE, uses absolute coefficients (appropriate for normal differential exon usage); if TRUE, uses non-absolute (i.e. changes should be in the same direction across significant bins) and width-weighted scores (i.e. larger bins have more weight) – this is relevant only when testing UTR usage.
alpha Points transparency
...
Passed to geom_label_repel; this can for instance be used to increase ‘max.overlaps’ when not all desired gene labels are displayed)

Value

A ggplot

Examples

data(example_bin_se)
se <- diffSpliceWrapper(example_bin_se, ~condition)
plotTopGenes(se)
Description

prepareBins

Usage

prepareBins(g,
    APA = NULL,
    onlyMainChr = TRUE,
    removeAntisense = TRUE,
    chrStyle = NULL,
    maxUTRbinSize = 15000,
    codingOnly = FALSE,
    genewise = FALSE,
    stranded = FALSE,
    verbose = TRUE)

Arguments

g A GRanges (or path to RDS file containing a GRanges) or path to a gtf file or EnsDb object containing the gene annotation.

APA A GRanges (or path to a GRanges in RDS format) or bed file containing the alternative poly-A site database

onlyMainChr Logical; whether to keep only main chromosomes

removeAntisense Logical; whether to remove antisense APA sites

chrStyle Chromosome notation to convert to (default no conversion)

maxUTRbinSize Max width of new alternative UTR bins

codingOnly Logical, whether to keep only coding transcripts

genewise Logical, whether annotation should be flattened genewise

stranded Logical, whether to perform disjoin in a stranded fashion.

verbose Logical, whether to print run information

Details

See the vignette for more details.

Value

A ‘GRanges’ object.
Author(s)
Stefan Greber

Examples
data(example_gene_annotation)
bins <- prepareBins(example_gene_annotation)

---

rn6_PAS  Poly-A sites compendium for Rattus Norvegicus (Rno6)

Description
These are the sites from polyA_DB release 3.2, downloaded from https://exon.apps.wistar.org/PolyA_DB/v3/download/3.2/rat_pas.zip, and lifted over to Rno6.

Value
a 'GRanges' object

---

simesAggregation  simesAggregation

Description
Simes p-value correction and aggregation, adapted from link[limma]{diffSplice}

Usage
simesAggregation(p.value, geneid)

Arguments
p.value  A vector of p-values
geneid  A vector of group labels such as gene identifiers

Value
A named vector of aggregated p-values

Examples
p <- runif(50)
genes <- sample(LETTERS, 50, replace=TRUE)
simesAggregation(p, genes)
addNormalizedAssays, 2
countFeatures, 2, 3, 4, 7, 9, 11
deuBinPlot, 4
dEUwrappers (diffSpliceDGEWrapper), 6
DEXSeq, 6
DEXSeqWrapper, 4, 11
DEXSeqWrapper (diffSpliceDGEWrapper), 6
diffSplice, 5–7
diffSplice2, 5, 7
diffSpliceDGE, 6
diffSpliceDGEWrapper, 6
diffSpliceWrapper, 4, 11
diffSpliceWrapper (diffSpliceDGEWrapper), 6
example_bin_se, 7
eample_gene_annotation, 8
geneBinHeatmap, 8
geneLevelStats, 7, 9
gem_label_repel, 11
Heatmap, 9
lmFit, 5
plotTopGenes, 11
prepareBins, 12
RangedSummarizedExperiment-class, 3
rn6_PAS, 13
simesAggregation, 13
SingleAnnotation, 9