
countFeatures	<i>countFeatures</i>
---------------	----------------------

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	Passed 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)
```

diffSpliceDGEWrapper *DEUwrappers*

Description

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

Usage

```
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.
coef	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').

geneBinHeatmap	<i>geneBinHeatmap</i>
----------------	-----------------------

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,  
  ...  
)
```

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

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

geneLevelStats	<i>geneLevelStats</i>
----------------	-----------------------

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)
```

plotTopGenes

plotTopGenes

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 (ie 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)
```

```
prepareBins      prepareBins
```

Description

prepareBins

Usage

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

Arguments

<code>g</code>	A GRanges (or path to RDS file containing a GRanges) or path to a gtf file or EnsDb object containing the gene annotation.
<code>APA</code>	A GRanges (or path to a GRanges in RDS format) or bed file containing the alternative poly-A site database
<code>onlyMainChr</code>	Logical; whether to keep only main chromosomes
<code>removeAntisense</code>	Logical; whether to remove antisense APA sites
<code>chrStyle</code>	Chromosome notation to convert to (default no conversion)
<code>maxUTRbinSize</code>	Max width of new alternative UTR bins
<code>codingOnly</code>	Logical, whether to keep only coding transcripts
<code>genewise</code>	Logical, whether annotation should be flattened genewise
<code>stranded</code>	Logical, whether to perform disjoint in a stranded fashion.
<code>verbose</code>	Logical, whether to print run information

Details

See the vignette for more details.

Value

A ‘GRanges’ object.

Author(s)

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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)
```

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