

# Package ‘extraChIPs’

September 25, 2023

**Version** 1.4.6

**Title** Additional functions for working with ChIP-Seq data

**Description** This package builds on existing tools and adds some simple but extremely useful capabilities for working with ChIP-Seq data. The focus is on detecting differential binding windows/regions. One set of functions focusses on set-operations retaining mcols for GRanges objects, whilst another group of functions are to aid visualisation of results. Coercion to tibble objects is also included.

**License** GPL-3

**Encoding** UTF-8

**URL** <https://github.com/smped/extraChIPs>

**BugReports** <https://github.com/smped/extraChIPs/issues>

**Depends** BiocParallel, R (>= 4.2.0), GenomicRanges, ggplot2 (>= 3.4.0), SummarizedExperiment, tibble

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extraChIPs-package      *extraChIPs: A package for enabling and extending ChIP-Seq analysis*

---

## Description

The package provides three categories of important functions: Range-based, Visualisation and Convenience functions, with most centred around GenomicRanges objects

## Range-based Functions

Many of the range-based functions included in this package have a focus on retaining the `mcols` information whilst manipulating the ranges, such as `reduceMC()` which not only reduces the Ranges, but collapses the `mcols` into vectors or `IRanges::CompressedList` objects. Key function from this group are:

- `reduceMC()`, `setdiffMC()`, `intersectMC()`, `unionMC()`, `distinctMC()` and `chopMC()`
- `bestOverlap()` and `propOverlap()` provide simple output easily able to be added as a column within the `mcols` element
- `as_tibble()` coerces a GRanges object to a `tibble::tibble`.
- `colToRanges()` enables parsing of a single column to a GRanges object, setting all other columns as the `mcols` element.
- `stitchRanges()` merges nearby ranges setting barrier ranges which cannot be crossed when merging
- `partitionRanges()` break apart one set of ranges by another
- `dualFilter()` filters ranges from sliding windows using a guide set of reference ranges where signal is confidently expected
- `mergeByCol()` merges overlapping ranges, as produced by sliding windows
- `mapByFeature()` is able to map a set of GRanges to the most appropriate gene, using any optional combination of promoters, enhancers and HiC interactions
- `grlToSE()` takes selected columns from a GRangesList and sets them as assays within a `SummarizedExperiment::RangedSummarizedExperiment` object. Used for combining peak intensities or results across multiple ChIP targets.

### Visualisation Functions

- `plotHFGC()` is a wrapper to Gviz plotting functions, able to take any combination of HiC, Features, Genes and Coverage (i.e. BigWig) and plot a specified range.
- `plotOverlaps()` visualises overlapping ranges as an UpSet plot or Venn Diagram
- `plotProfileHeatmap()` plots the average signal around a set of ranges, as prepared by `getProfileData()`
- `plotPie()` and `plotSplitDonut()` enable simple comparison across multiple annotation columns within a GRanges object.
- `plotAssayDensities()`, `plotAssayPCA()` and `plotAssayRle()` provide simple interfaces to plotting key values from a `SummarizedExperiment::RangedSummarizedExperiment`.

### Convenience Functions

- `fitAssayDiff()` enables differential signal analysis on a SummarizedExperiment object
- `collapseGenes()` prints a vector of genes for an rmarkdown document, using italics.
- `importPeaks()` imports large numbers of broadPeak or narrowPeak files
- `makeConsensus()` forms consensus peaks from overlapping ranges within a GRangesList()
- `voomWeightsFromCPM()` allows creation of an `limma::EList` object as would be created from counts by `limma::voom()`, but using `edgeR::cpm()` values as input.

### Author(s)

Stephen Pederson

---

.ec\_HMP

*This is a modified version of `harmonicmeanp::p.hmp` developed by Prof Daniel Wilson, and hardwired to simply return a combined asymptotically exact HMP. Hardwiring like this gives a 10-fold speed-up. Further modifications may be possible, but this seems enough for now*

---

### Description

This is a modified version of `harmonicmeanp::p.hmp` developed by Prof Daniel Wilson, and hardwired to simply return a combined asymptotically exact HMP. Hardwiring like this gives a 10-fold speed-up. Further modifications may be possible, but this seems enough for now

### Usage

```
.ec_HMP(p, w)
```

### Arguments

p	vector of p-values
w	vector of weights

---

.ec\_HMP\_adj                      *Similar to the above, this produces the FWER-controlled version in a streamlined way*

---

### Description

Similar to the above, this produces the FWER-controlled version in a streamlined way

### Usage

```
.ec_HMP_adj(p, w, L)
```

### Arguments

p	vector of p-values
w	vector of weights
L	Number of global tests

---

.makeFinalProfileHeatmap  
*Make a profile heatmap*

---

### Description

Make a profile heatmap with optional summary panel at the top

### Usage

```
.makeFinalProfileHeatmap(  
  data,  
  x = NULL,  
  y = NULL,  
  fill = NULL,  
  colour = NULL,  
  linetype = NULL,  
  facet_x = NULL,  
  facet_y = NULL,  
  summary_fun = c("mean", "median", "min", "max", "none"),  
  rel_height = 0.3,  
  x_lab = NULL,  
  y_lab = NULL,  
  fill_lab = NULL,  
  ...  
)
```

**Arguments**

data	A data.frame or tibble in long form
x, y	The values mapped to the x & y axes
fill	The column used for heatmap colours
colour, linetype	Columns used for the summary plot in the top panel
facet_x, facet_y	Columns used to facet the plot along these axes
summary_fun	Function used to create the summary value at each position
rel_height	The relative height of the top panel
x_lab, y_lab, fill_lab	_labels added to x/y-axes & the fill legend
...	Passed to facet_grid

**Details**

The workhorse function for generating the final heatmap Expects a single data.frame in long form with requisite columns

**Value**

A ggplot2 object

---

.mapFeatures

*Map ranges to genes using features as an anchor*

---

**Description**

Map ranges to genes using features as an anchor

**Usage**

```
.mapFeatures(.gr, .feat, .genes, .cols, .gr2feat, .feat2gene, ...)
```

**Arguments**

.gr	The ranges to map onto
.feat	Features to use for mapping
.genes	GRanges object containing gene-level information
.cols	The columns from .genes to map onto .gr
.gr2feat	The maximum distance between ranges and features
.feat2gene	The maximum distance between features & genes
...	Passed to findOverlaps and subsetByOverlaps

**Value**

A data.frame

---

.mapGi *Map ranges to genes via Interactions*

---

**Description**

Map ranges to genes via Interactions

**Usage**

```
.mapGi(.gr, .gi, .genes, .cols, .gr2gi, .gi2gene, ...)
```

**Arguments**

.gr	The ranges to map onto
.gi	GInteractions object
.genes	GRanges object containing gene-level information
.cols	The columns from .genes to map onto .gr
.gr2gi	The maximum distance between ranges and anchors
.gi2gene	The maximum distance between anchors & genes
...	Passed to findOverlaps

**Value**

data.frame of mapped ranges

---

.mapWithin *Map ranges to all genes within a set distance*

---

**Description**

Map ranges to all genes within a set distance

**Usage**

```
.mapWithin(.gr, .genes, .cols, .within, ...)
```

**Arguments**

.gr	The ranges to map onto
.genes	GRanges object containing gene-level information
.cols	The columns from .genes to map onto .gr
.within	The maximum distance between ranges & genes
...	Passed to findOverlaps

**Value**

A data.frame

---

as_tibble	<i>Convert to a tibble</i>
-----------	----------------------------

---

**Description**

Convert multiple Genomic objects to tibbles

**Usage**

```
## S3 method for class 'DataFrame'
as_tibble(x, rangeAsChar = TRUE, ...)

## S3 method for class 'GenomicRanges'
as_tibble(x, rangeAsChar = TRUE, name = "range", ...)

## S3 method for class 'Seqinfo'
as_tibble(x, ...)

## S3 method for class 'GInteractions'
as_tibble(x, rangeAsChar = TRUE, suffix = c(".x", ".y"), ...)

## S3 method for class 'SummarizedExperiment'
as_tibble(x, rangeAsChar = TRUE, ...)

## S3 method for class 'TopTags'
as_tibble(x, ...)
```

**Arguments**

x	A Genomic Ranges or DataFrame object
rangeAsChar	Convert any GRanges element to a character vector
...	Passed to <code>tibble::as_tibble()</code>
name	Name of column to use for ranges. Ignored if rangeAsChar = FALSE
suffix	Suffix appended to column names for anchor1 and anchor2 of a GInteractions object. Only used if specifying rangeAsChar = FALSE



**Details**

Quick and dirty conversion into a tibble.

By default, GenomicRanges will be returned with the range as a character column called range and all mcols parsed as the remaining columns. Seqinfo information will be lost during coercion.

Given that names for ranges are considered as rownames in the mcols element, these can be simply parsed by setting rownames = "id" in the call to as\_tibble()

When coercing a DataFrame, any Compressed/SimpleList columns will be coerced to S3 list columns. Any GRanges columns will be returned as a character column, losing any additional mcols from these secondary ranges

Coercion of SummarizedExperiment objects will be performed on the rowRanges() element, whilst for a GInteractions object, both anchors will returned with the default suffixes .x and .y

Defined as an S3 method for consistency with existing tidy methods

**Value**

A [tibble](#)

**Examples**

```
gr <- GRanges("chr1:1-10")
gr$p_value <- runif(1)
names(gr) <- "range1"
gr
as_tibble(gr)
as_tibble(gr, rownames = "id")
as_tibble(mcols(gr))
as_tibble(seqinfo(gr))

hic <- InteractionSet::GInteractions(gr, GRanges("chr1:201-210"))
hic$id <- "interaction1"
as_tibble(hic)
```

---

bestOverlap

*Find the best overlap between GRanges*

---

**Description**

Find the best overlap between ranges

**Usage**

```
bestOverlap(x, y, ...)

## S4 method for signature 'GRanges,GRanges'
bestOverlap(
```

```

    x,
    y,
    var = NULL,
    ignore.strand = FALSE,
    missing = NA_character_,
    min_prop = 0.01,
    ...
)

## S4 method for signature 'GRanges,GRangesList'
bestOverlap(
  x,
  y,
  ignore.strand = FALSE,
  missing = NA_character_,
  min_prop = 0.01,
  ...
)

```

### Arguments

<code>x</code>	a GRanges object
<code>y</code>	a named GRangesList or GRanges object with mcol as reference category
<code>...</code>	Not used
<code>var</code>	The variable to use as the category. Not required if <code>y</code> is a GRangesList
<code>ignore.strand</code>	logical(1) Passed to <a href="#">findOverlaps</a>
<code>missing</code>	Value to assign to ranges with no overlap
<code>min_prop</code>	Threshold below which overlaps are discarded

### Details

This finds the category in the subject GRanges (`y`) which has the best overlap with the query GRanges (`x`). The aim is to produce a character vector for best classifying the query GRanges using an external set of features (e.g. promoters, enhancers etc). If the subject (`y`) is a GRanges object, the values in the specified column will be used as the category. If the subject (`y`) is a GRangesList, the names of the list will be used to provide the best match

### Value

Character vector the same length as the supplied GRanges object

### Examples

```

gr <- GRanges("chr1:1-10")
gr_cat <- GRanges(c("chr1:2-10", "chr1:5-10"))
gr_cat$category <- c("a", "b")
propOverlap(gr, gr_cat)
bestOverlap(gr, gr_cat, var = "category")

```

```
gr1 <- splitAsList(gr_cat, gr_cat$category)
lapply(gr1, function(x) propOverlap(gr, x))
bestOverlap(gr, gr1)
```

---

chopMC

*Keep unique ranges and collapse mcols*

---

## Description

Keep unique ranges by 'chopping' mcols

## Usage

```
chopMC(x, simplify = TRUE)
```

## Arguments

x	A GenomicRanges object
simplify	logical(1)

## Details

This function finds unique ranges and chops **all** mcols in a manner similar to [chop](#). Chopped columns will be returned as CompressedList columns, unless simplify = TRUE (the default). In this case, columns will be returned as vectors where possible.

## Value

A GRanges object

## Examples

```
gr <- GRanges(rep(c("chr1:1-10"), 2))
gr$id <- paste0("range", seq_along(gr))
gr$gene <- "gene1"
gr
chopMC(gr)
```

---

collapseGenes	<i>Collapse a vector of gene names</i>
---------------	--

---

### Description

Collapse a vector of gene names

### Usage

```
collapseGenes(  
  x,  
  sort = TRUE,  
  dedup = TRUE,  
  format = "_",  
  sep = ", ",  
  last = " and ",  
  numeric = TRUE,  
  width = Inf,  
  ...  
)
```

### Arguments

x	character vector representing gene names
sort	logical(1) Should the names be sorted alphabetically
dedup	logical(1) Should duplicate names be removed
format	character string for markdown formatting of each element
sep	separator between vector elements
last	character string to place before the last element
numeric	logical(1) sort digits numerically, instead of as strings
width	The maximum width of the string before truncating to ...
...	passed to <a href="#">str_sort</a>

### Details

Convenience function to collapse a vector of gene names into a character/glue object of length 1. By default, symbols are deduplicated, sorted alpha-numerically and italicised with an underscore.

### Value

a glue object

### Examples

```
genes <- c("FOXP3", "BRCA1", "TP53")  
collapseGenes(genes)
```

---

colToRanges	<i>Coerce a column to a GRanges object</i>
-------------	--

---

**Description**

Coerce a column to a GRanges object from a rectangular object

**Usage**

```
colToRanges(x, ...)

## S4 method for signature 'DataFrame'
colToRanges(x, var, seqinfo = NULL, ...)

## S4 method for signature 'GRanges'
colToRanges(x, var, ...)

## S4 method for signature 'data.frame'
colToRanges(x, var, seqinfo = NULL, ...)
```

**Arguments**

x	A data-frame or GRanges object containing the column to coerce
...	Used to pass arguments to lower-level functions
var	The name of the column to coerce
seqinfo	A seqinfo object to be applied to the new GRanges object. Ignored if the column is already a GRanges object

**Details**

Take a data.frame-like object and coerce one column to a GRanges object, setting the remainder as the mcols. A particularly useful application of this is when you have a GRanges object with one mcol being a secondary GRanges object.

Alternatively, if you have a data.frame with GRanges represented as a character column, this provides a simple method of coercion. In this case, no Seqinfo element will be applied to the GRanges element.

**Value**

A GenomicRanges object

**Examples**

```
set.seed(73)
x <- GRanges(c("chr1:1-10", "chr1:6-15", "chr1:51-60"))
seqinfo(x) <- Seqinfo("chr1", 60, FALSE, "Example")
df <- data.frame(logFC = rnorm(3), logCPM = rnorm(3,8), p = 10^-rexp(3))
```

```
mcols(x) <- df
gr <- mergeByCol(x, col = "logCPM", pval = "p")
colToRanges(gr, "keyval_range")
```

---

cytobands

*Cytogenetic bands*

---

## Description

Cytogenetic bands for GRCh37/hg19 and GRCh38/hg38

## Usage

```
data(grch37.cytobands)
```

```
data(grch38.cytobands)
```

## Format

Cytogenetic bands for standard chromosomes from GRCh37, in the format required by [Ideogram-Track](#). A data.frame with 5 columns:

**chrom** Chromosome

**chromStart** Starting position for each cytogenetic band

**chromEnd** End position for each cytogenetic band

**name** Name for each band, e.g. p.36.33

**gieStain** Staining pattern

An object of class data.frame with 862 rows and 5 columns.

## Source

<https://hgdownload.soe.ucsc.edu/goldenPath/hg19/database/cytoBand.txt.gz>

<https://hgdownload.soe.ucsc.edu/goldenPath/hg38/database/cytoBand.txt.gz>

## Examples

```
data(grch37.cytobands)
head(grch37.cytobands)
```

```
data(grch38.cytobands)
head(grch38.cytobands)
```

---

defineRegions                      *Define Genomic Regions Based on Gene Annotations*

---

## Description

Use gene, transcript and exon annotations to define genomic regions

## Usage

```
defineRegions(
  genes,
  transcripts,
  exons,
  promoter = c(2500, 500),
  upstream = 5000,
  intron = TRUE,
  proximal = 10000,
  simplify = FALSE,
  cols = c("gene_id", "gene_name", "transcript_id", "transcript_name"),
  ...
)
```

## Arguments

genes, transcripts, exons	GRanges objects defining each level of annotation
promoter	Numeric vector defining upstream and/or downstream distances for a promoter. Passing a single value will define a symmetrical promoter The first value represents the upstream range
upstream	The distance from a TSS defining an upstream promoter
intron	logical(1) Separate gene bodies into introns and exons. If intron = FALSE gene bodies will simply be defined as gene bodies
proximal	Distance from a gene to be considered a proximal intergenic region. If set to 0, intergenic regions will simply be considered as uniformly intergenic
simplify	Passed internally to reduceMC and setdiffMC
cols	Column names to be retained from the supplied annotations
...	Not used

## Details

Using GRanges annotated as genes, transcripts and exons this function will define ranges uniquely assigned to a region type using a hierarchical process. By default, these region types will be (in order) 1) Promoters, 2) Upstream Promoters, 3) Exons, 4) Introns, 5) Proximal Intergenic and 6) Distal Intergenic.

Setting `intron = FALSE` will replace introns and exons with a generic "Gene Body" annotation. Setting `proximal = 0` will return all intergenic regions (not previously annotated as promoters or upstream promoters) to an "Intergenic" category

Notably, once a region has been defined, it is excluded from all subsequent candidate regions.

Any columns matching the names provided in `cols` will be returned, and it is assumed that the gene/transcript/exon ranges will contain informative columns in the `mcols()` element.

## Value

A `GRangesList`

## Examples

```
## Define two exons for two transcripts
sq <- Seqinfo(seqnames = "chr1", seqlengths = 50000)
e <- c("chr1:20001-21000", "chr1:29001-29950", "chr1:22001-23000", "chr1:29001-30000")
e <- GRanges(e, seqinfo = sq)
mcols(e) <- DataFrame(
  gene_id = "Gene1", transcript_id = c("Trans1", "Trans1", "Trans2", "Trans2")
)

## Define the transcript ranges
t <- unlist(endoapply(split(e, e$transcript_id), range))
t$gene_id <- "Gene1"
t$transcript_id <- names(t)
names(t) <- NULL

## Summarise to gene level
g <- reduceMC(t)
g$transcript_id <- NA_character_

## Now annotate the regions
regions <- defineRegions(genes = g, transcripts = t, exons = e)
sort(unlist(regions))

## Alternatively, collapse gene body and intergenic ranges
regions <- defineRegions(
  genes = g, transcripts = t, exons = e, intron = FALSE, proximal = 0
)
sort(unlist(regions))
```

---

distinctMC

*Keep distinct ranges and mcols*

---

## Description

Keep distinct ranges by including `mcols`



**Usage**

```
distinctMC(x, ..., .keep_all = FALSE)
```

**Arguments**

x	A GRanges object
...	<data-masking> Passed to <a href="#">distinct</a>
.keep_all	If TRUE, keep all columns in x

**Details**

Wrapper to [distinct](#) for GRanges objects. Finds unique ranges and mccols in combination and retains only the distinct combinations, in keeping with the `dplyr` function.

Will default to `unique(granges(x))` if no columns are provided

**Value**

A GRanges object

**Examples**

```
gr <- GRanges(rep(c("chr1:1-10"), 2))
gr$id <- paste0("range", seq_along(gr))
gr$gene <- "gene1"
gr
distinctMC(gr)
distinctMC(gr, gene)
distinctMC(gr, gene, .keep_all = TRUE)
```

---

dualFilter

*Apply two filters to sliding windows*


---

**Description**

Apply two filters to counts generated using sliding windows

**Usage**

```
dualFilter(
  x,
  bg,
  ref,
  q = 0.5,
  logCPM = TRUE,
  keep.totals = TRUE,
  BPPARAM = bpparam()
)
```

**Arguments**

x	RangedSummarizedExperiment containing sample counts
bg	RangedSummarizedExperiment containing background/input counts
ref	GRanges object containing ranges where signal is expected
q	The upper percentile of the reference ranges expected to be returned when tuning the filtering criteria
logCPM	logical(1) Add a logCPM assay to the returned data
keep.totals	logical(1) Keep the original library sizes or replace using only the retained windows
BPPARAM	Settings for running in parallel

**Details**

This function will take sliding (or tiling) windows for its input as a [RangedSummarizedExperiment](#) object. The dual strategy of applying [filterWindowsControl](#) and [filterWindowsProportion](#) will then be applied. A set of reference ranges for which signal is expected is used to refine the filtering criteria.

Cutoff values are found for both signal relative to input and overall signal, such that the  $100 \times q\%$  of the (sliding) windows which overlap a reference range will be returned, along with any others which match the dual filtering criteria. In general, higher values of  $q$  will return more windows as those with weak signal and a marginal overlap with a reference range will be returned. Lower values will ensure that fewer windows, generally with the strongest signal, are retained. Cutoff values for both criteria are added to the metadata element of the returned object.

If setting `bg = NULL` the [filterWindowsControl](#) step will be ignored and only the [filterWindowsProportion](#) will be used. This should only be performed if no Input sample is available.

**Please note** that the any `.bam` files referred to in the supplied objects **must** be accessible to this function. It will not run on a separate machine or file structure to that which the original sliding windows were prepared. Please see the example/vignette for runnable code.

**Value**

A [RangedSummarizedExperiment](#) which is a filtered subset of the original object. If requested the assay "logCPM" will be added (TRUE by default)

**Examples**

```
## Taken from the differential_binding vignette
library(tidyverse)
library(Rsamtools)
library(csaw)
library(BiocParallel)
library(rtracklayer)
## For this function we need a set of counts using sliding windows and the
## original BamFiles from which they were taken
## First we'll set up the bam file list
bfl <- system.file(
  "extdata", "bam", c("ex1.bam", "ex2.bam", "input.bam"), package = "extraChIPs"
```

```

) %>%
BamFileList() %>%
setNames(c("ex1", "ex2", "input"))

## Then define the readParam settings for csaw::readParam()
rp <- readParam(
  pe = "none",
  dedup = TRUE,
  restrict = "chr10"
)

## Now we can form our sliding window object with the counts.
wincounts <- windowCounts(
  bam.files = bfl,
  spacing = 60,
  width = 180,
  ext = 200,
  filter = 1,
  param = rp
)

## As this is a subset of reads, add the initial library sizes for accuracy
## Note that this step is not normally required
wincounts$totals <- c(964076L, 989543L, 1172179L)

## We should also update the metadata for our counts
wincounts$sample <- colnames(wincounts)
wincounts$treat <- as.factor(c("ctrl", "treat", NA))
colData(wincounts)

## The function dualFilter requires a set of peaks which will guide the
## filtering step. This indicate where genuine signal is likely to be found
## and will perform the filtering based on a) signal above the input, and
## b) The overall signal level, using the guide set of peaks to inform the
## cutoff values for inclusion
peaks <- import.bed(
  system.file("extdata", "peaks.bed.gz", package = "extraChIPs")
)
filtcounts <- dualFilter(
  x = wincounts[, !is.na(wincounts$treat)],
  bg = wincounts[, is.na(wincounts$treat)],
  ref = peaks,
  q = 0.8 # Better to use q = 0.5 on real data
)

```

**Description**

Various example datasets for demonstrating analysis and visualisation strategies. Generation of all datasets is documented in `system.file("script/ex_datasets.md", package = "extraChIPs")`

**ex\_genes** Simple GRanges object with complete ranges for each gene

**ex\_trans** Exon & transcript level information prepared for plotting with Gviz or plotHFGC()

**ex\_prom** Regions defined as promoters

**ex\_hic** Example HiC interactions

**Usage**

```
data(ex_trans)
```

```
data(ex_genes)
```

```
data(ex_prom)
```

```
data(ex_hic)
```

**Format**

GRanges and GInteractions objects

All annotations are from GRCh37

An object of class GRanges of length 4.

An object of class GRanges of length 9.

An object of class GInteractions of length 1.

**Examples**

```
data(ex_trans)
ex_trans
```

---

fitAssayDiff

*Detect Differential ChIP Signal*

---

**Description**

Detect differential ChIP signal using one of many approaches

**Usage**

```
fitAssayDiff(x, ...)

## S4 method for signature 'SummarizedExperiment'
fitAssayDiff(
  x,
  assay = "counts",
  design = NULL,
  coef = NULL,
  lib.size = "totals",
  method = c("qlf", "lt"),
  norm = c("none", "TMM", "RLE", "TMMwsp", "upperquartile"),
  groups = NULL,
  fc = 1,
  lfc = log2(fc),
  asRanges = FALSE,
  offset = NULL,
  null = c("interval", "worst.case"),
  ...
)
```

**Arguments**

x	a SummarizedExperiment object
...	Passed to <a href="#">calcNormFactors</a> , <a href="#">estimateDisp</a> and <a href="#">glmQLFit</a> when method = "qlf". If method = "lt", instead passed to <a href="#">lmFit</a> , <a href="#">treat</a> , <a href="#">eBayes</a>
assay	The assay to use for analysis
design	The design matrix to use for analysis
coef	The required column from the design matrix
lib.size	The column within the colData element which contains the library size information. If set to NULL, column summaries will be used.
method	the analytic method to be used. Can be 'qlf' which will fit counts using the <a href="#">glmQLFit</a> strategy , or 'lt' which fits the <a href="#">limma-trend</a> model on logCPM, or pre-processed logCPM values
norm	The normalisation strategy to use when running the glmQLF models. The value 'none' relies solely on library-size normalisation, and is the default. All methods available in <a href="#">calcNormFactors</a> are implemented. Ignored when using method = "lt"
groups	character(1) If a column name is supplied here, group-based normalisation will be applied to GLM models treating data in this column as a grouping factor. Ignored when using method = "lt"
fc, lfc	Thresholds passed to <a href="#">treat</a> or <a href="#">glmTreat</a>
asRanges	logical(1). By default, the returned object will be a SummarizedExperiment object with the results added to the rowData element. Setting asRanges = TRUE will only return the GRanges object from this element

offset	If provided will be used as the offset when the DGEList object is created during model fitting
null	Passed to <a href="#">glmTreat</a>

## Details

Starting with a SummarizedExperiment object this function fits either a [glmQLFit](#) model to count data, or the [limma-trend](#) model to logCPM data.

If fitting Generalised Linear Models via [glmQLFit](#), options for normalisation are "none", which normalises to library size. Existing library sizes are commonly found in the "totals" column of the colData element and this is attempted by default. All methods provided in [calcNormFactors](#) are also implemented, with the added possibility of normalising within groups instead of across the entire dataset. To enable this, the column with the grouping factor is expected to be in the colData element and is simply called by column name. No normalisation is applied when using the limma-trend model, as this allows for previous normalisation strategies to be performed on the data.

Normalising to ChIP Input samples, or using offsets is not yet implemented.

Either range-based hypothesis testing is implemented using [glmTreat](#) or [treat](#). Setting fc to 1 (or lfc to 0) will default to a point-based null hypothesis, equivalent to either [glmQLFTest](#) (method = "qlf") or [eBayes](#) (method = "l").

It should also be noted that this is primarily a convenience function and if requiring intermediate output from any setps, then these can be run individually as conventionally specified.

## Value

A SummarizedExperiment object with results set as the rowData element. Any existing columns not contained in the differential ChIP results will be retained. Results from testing will contain logCPM, logFC, PValue and the t/F statistic as appropriate, along with an FDR-adjusted p-value

## Examples

```
nrows <- 200; ncols <- 6
counts <- matrix(runif(nrows * ncols, 1, 1e4), nrows)
colnames(counts) <- paste0("Sample_", seq_len(ncols))
df <- DataFrame(treat = c("A", "A", "A", "B", "B", "B"))
df$treat <- as.factor(df$treat)
se <- SummarizedExperiment(
  assays = SimpleList(counts = counts), colData = df
)
X <- model.matrix(~treat, colData(se))
se <- fitAssayDiff(se, design = X, lib.size = NULL)
rowData(se)
```

---

fixed\_width\_datasets *Datasets for the Fixed-Width Vignette*

---

### Description

GRangesList of peaks and SummarizedExperiment of counts All were saved during initial vignette preparation at [https://github.com/smped/extraChIPs\\_vignette/blob/main/differential\\_signal\\_fixed.Rmd](https://github.com/smped/extraChIPs_vignette/blob/main/differential_signal_fixed.Rmd)

### Usage

```
data(se)
```

```
data(peaks)
```

### Format

An object of class RangedSummarizedExperiment with 188 rows and 6 columns.

An object of class CompressedGRangesList of length 6.

### Examples

```
data(se)
se
data(peaks)
peaks
```

---

getProfileData *Get Profile Data surrounding specified ranges*

---

### Description

Get coverage Profile Data surrounding specified ranges

### Usage

```
getProfileData(x, gr, ...)
```

```
## S4 method for signature 'BigWigFile,GenomicRanges'
```

```
getProfileData(
  x,
  gr,
  upstream = 2500,
  downstream = upstream,
  bins = 100,
  mean_mode = "w0",
  log = TRUE,
```

```

    offset = 1,
    ...
)

## S4 method for signature 'BigWigFileList,GenomicRanges'
getProfileData(
  x,
  gr,
  upstream = 2500,
  downstream = upstream,
  bins = 100,
  mean_mode = "w0",
  log = TRUE,
  offset = 1,
  BPPARAM = SerialParam(),
  ...
)

## S4 method for signature 'character,GenomicRanges'
getProfileData(
  x,
  gr,
  upstream = 2500,
  downstream = upstream,
  bins = 100,
  mean_mode = "w0",
  log = TRUE,
  offset = 1,
  ...
)

```

### Arguments

x	A BigWigFile or BigWiFileList
gr	A GRanges object
...	Passed to <a href="#">normalizeToMatrix</a>
upstream	The distance to extend upstream from the centre of each range within gr
downstream	The distance to extend downstream from the centre of each range within gr
bins	The total number of bins to break the extended ranges into
mean_mode	The method used for calculating the score for each bin. See <a href="#">normalizeToMatrix</a> for details
log	logical(1) Should the returned values be log2-transformed
offset	Value added to data if log-transforming. Ignored otherwise
BPPARAM	Passed internally to <a href="#">bplapply</a>



**Details**

This will take all provided ranges and set as identical width ranges, extending by the specified amount both up and downstream of the centre of the provided ranges. By default, the ranges extensions are symmetrical and only the upstream range needs to be specified, however this parameterisation allows for non-symmetrical ranges to be generated.

These uniform width ranges will then be used to extract the value contained in the score field from one or more BigWigFiles. Uniform width ranges are then broken into bins of equal width and the average score found within each bin.

The binned profiles are returned as a DataFrameList called `profile_data` as a column within the resized GRanges object. Column names in each DataFrame are `score`, `position` and `bp`.

If passing a BigWigFileList, profiles will be obtained in series by default. To run in parallel pass a [MulticoreParam](#) object to the BPPARAM argument.

**Value**

GRanges or GrangesList with column `profile_data`, as described above

**Examples**

```
bw <- system.file("tests", "test.bw", package = "rtracklayer")
gr <- GRanges("chr2:1000")
pd <- getProfileData(bw, gr, upstream = 500, bins = 10)
pd
pd$profile_data
```

---

grlToSE

*Set columns from a GRangesList as Assays in a SummarizedExperiment*

---

**Description**

Move one or more columns from a GRangesList elements into assays in a RangesSummarizedExperiment

**Usage**

```
grlToSE(x, ...)

## S4 method for signature 'GRangesList'
grlToSE(
  x,
  assayCols = c(),
  metaCols = c(),
  keyvals = c(),
  by = c("min", "max"),
```

```

    ...,
    ignore.strand = FALSE
  )

```

### Arguments

x	A GRangesList
...	Passed to <a href="#">reduce</a>
assayCols	Columns to move to separate assays
metaCols	Columns to move to mcols within the rowRanges element
keyvals	The value to use when choosing representative values
by	How to choose by keyvals
ignore.strand	logical(1). Whether the strand of the input ranges should be ignored or not.

### Details

Given a GRangesList which would commonly represent multiple samples, reduce any overlapping ranges into a consensus range, setting any metadata columns to be retained as separate assays. These columns may contain values such as coverage, p-values etc.

Additional columns can also be placed as rowData columns where the original values are better suited to information about the consensus range rather than the sample (or GRangesList element).

Only one value for each range will be retained, and these are chosen using the value provided as the keyvals, taking either the min or max value in this column as the representative range.

### Value

A RangedSummarizedExperiment

### Examples

```

a <- GRanges("chr1:1-10")
a$feature <- "Gene"
a$p <- 0.1
b <- GRanges(c("chr1:6-15", "chr1:15"))
b$feature <- c("Gene", "Promoter")
b$p <- c(0.5, 0.01)
grl <- GRangesList(a = a, b = b)
grl
se <- grlToSE(
  grl, assayCols = "p", metaCols = "feature", keyvals = "p", by = "min"
)
assay(se, "p")
rowRanges(se)

```

---

importPeaks	<i>Import peaks</i>
-------------	---------------------

---

## Description

Import peaks in narrowPeak or broadPeak format

## Usage

```
importPeaks(
  x,
  type = c("narrow", "broad"),
  blacklist,
  seqinfo,
  pruning.mode = c("coarse", "error"),
  sort = TRUE,
  setNames = TRUE,
  ...
)
```

## Arguments

x	One or more files to be imported. All files must be of the same type, i.e. narrow or broad
type	The type of peaks to be imported
blacklist	A set of ranges to be excluded
seqinfo	A seqinfo object to be applied to the GRanges objects
pruning.mode	How to handle conflicts if supplying a seqinfo object. Defaults to pruning.mode = "coarse". Only "coarse" and "error" are implemented. See <a href="#">seqinfo</a> .
sort	logical. Should the ranges be sorted during import
setNames	logical Set basename(x) as the name
...	passed to sort

## Details

Peaks are imported from either narrowPeak or broadPeak format as GenomicRanges objects.

## Value

A GRangesList

**Examples**

```
f1 <- system.file(
  c("extdata/ER_1.narrowPeak", "extdata/ER_2.narrowPeak"),
  package = "extraChIPs"
)
peaks <- importPeaks(f1)
peaks
```

---

makeConsensus	<i>Make a set of consensus peaks</i>
---------------	--------------------------------------

---

**Description**

Make a set of consensus peaks based on the number of replicates

**Usage**

```
makeConsensus(
  x,
  p = 0,
  var = NULL,
  method = c("union", "coverage"),
  ignore.strand = TRUE,
  simplify = FALSE,
  min_width = 0,
  ...
)
```

**Arguments**

x	A GRangesList
p	The minimum proportion of samples (i.e. elements of x) required for a peak to be retained in the output. By default all merged peaks will be returned
var	Additional columns in the mcols element to retain
method	Either return the union of all overlapping ranges, or the regions within the overlapping ranges which are covered by the specified proportion of replicates. When using p = 0, both methods will return identical results
ignore.strand, simplify, ...	Passed to <a href="#">reduceMC</a> or <a href="#">intersectMC</a> internally
min_width	Discard any regions below this width

**Details**

This takes a list of GRanges objects and forms a set of consensus peaks.

When using method = "union" the union ranges of all overlapping peaks will be returned, using the minimum proportion of replicates specified. When using method = "coverage", only the regions within each overlapping range which are 'covered' by the minimum proportion of replicates specified are returned. This will return narrower peaks in general, although some artefactual very small ranges may be included (e.g. 10bp). Careful setting of the min\_width parameter may be very helpful for these instances. It is also expected that setting method = "coverage" should return the region within each range which is more likely to contain the true binding site for the relevant ChIP targets

**Value**

GRanges object with mcols containing a logical vector for every element of x, along with the column n which adds all logical columns. These columns denote which replicates contain an overlapping peak for each range

If any additional columns have been requested using var, these will be returned as CompressedList objects as produced by reduceMC() or intersectMC().

**See Also**

[reduceMC](#) [intersectMC](#)

**Examples**

```
data("peaks")
## The first three replicates are from the same treatment group
gr1 <- peaks[1:3]
names(gr1) <- gsub("_peaks.", "", names(gr1))
makeConsensus(gr1)
makeConsensus(gr1, p = 2/3, var = "score")

## Using method = 'coverage' finds ranges based on the intersection
makeConsensus(gr1, p = 2/3, var = "score", method = "coverage")
```

**Description**

Map Genomic Ranges to genes using defined regulatory features

**Usage**

```
mapByFeature(
  gr,
  genes,
  prom,
  enh,
  gi,
  cols = c("gene_id", "gene_name", "symbol"),
  gr2prom = 0,
  gr2enh = 0,
  gr2gi = 0,
  gr2gene = 1e+05,
  prom2gene = 0,
  enh2gene = 1e+05,
  gi2gene = 0,
  ...
)
```

**Arguments**

<code>gr</code>	GRanges object with query ranges to be mapped to genes
<code>genes</code>	GRanges object containing genes (or any other nominal feature) to be assigned
<code>prom</code>	GRanges object defining promoters
<code>enh</code>	GRanges object defining Enhancers
<code>gi</code>	GInteractions object defining interactions. Mappings from interactions to genes should be performed as a separate prior step.
<code>cols</code>	Column names to be assigned as <code>mcols</code> in the output. Columns must be minimally present in <code>genes</code> . If all requested columns are found in any of <code>prom</code> , <code>enh</code> or <code>gi</code> , these pre-existing mappings will be preferentially used. Any columns not found in utilised reference objects will be ignored.
<code>gr2prom</code>	The maximum permissible distance between a query range and any ranges defined as promoters
<code>gr2enh</code>	The maximum permissible distance between a query range and any ranges defined as enhancers
<code>gr2gi</code>	The maximum permissible distance between a query range and any ranges defined as GInteraction anchors
<code>gr2gene</code>	The maximum permissible distance between a query range and genes (for ranges not otherwise mapped)
<code>prom2gene</code>	The maximum permissible distance between a range provided in <code>prom</code> and a gene
<code>enh2gene</code>	The maximum permissible distance between a range provided in <code>enh</code> and a gene
<code>gi2gene</code>	The maximum permissible distance between a GInteractions anchor (provided in <code>gi</code> ) and a gene
<code>...</code>	Passed to <code>findOverlaps</code> and <code>overlapsAny</code> internally

## Details

This function is able to utilise feature-level information and long-range interactions to enable better mapping of regions to genes. If provided, this essentially maps from ranges to genes using the regulatory features as a framework. The following sequential strategy is used:

1. Ranges overlapping a promoter are assigned to that gene
2. Ranges overlapping an enhancer are assigned to **all genes** within a specified distance
3. Ranges overlapping a long-range interaction are assigned to all genes connected by the interaction
4. Ranges with no gene assignment from the previous steps are assigned to *all overlapping genes* or the nearest gene within a specified distance

If information is missing for one of these steps, the algorithm will simply proceed to the next step. If no promoter, enhancer or interaction data is provided, all ranges will be simply mapped by step 4. Ranges can be mapped by any or all of the first three steps, but step 4 is mutually exclusive with the first 3 steps.

Distances between each set of features and the query range can be individually specified by modifying the `gr2prom`, `gr2enh`, `gr2gi` or `gr2gene` parameters. Distances between features and genes can also be set using the parameters `prom2gene`, `enh2gene` and `gi2gene`.

Additionally, if previously defined mappings are included with any of the `prom`, `enh` or `gi` objects, this will be used in preference to any obtained from the `genes` object.

## Value

A GRanges object with added `mcols` as specified

## Examples

```
## Define some genes
genes <- GRanges(c("chr1:2-10:*", "chr1:25-30:-", "chr1:31-40:+"))
genes$gene_id <- paste0("gene", seq_along(genes))
genes
## Add a promoter for each gene
prom <- promoters(genes, upstream = 1, downstream = 1)
prom
## Some ranges to map
gr <- GRanges(paste0("chr1:", seq(0, 60, by = 15)))
gr

## Map so that any gene within 25bp of the range is assigned
mapByFeature(gr, genes, gr2gene = 25)

## Now use promoters to be more accurate in the gene assignment
## Given that the first range overlaps the promoter of gene1, this is a
## more targetted approach. Similarly for the third range
mapByFeature(gr, genes, prom, gr2gene = 25)
```

---

mergeByCol	<i>Merge sliding windows using a specified column</i>
------------	---

---

**Description**

Merge sliding windows using a specified column

**Usage**

```
mergeByCol(x, ...)

## S4 method for signature 'GenomicRanges'
mergeByCol(
  x,
  df = NULL,
  col,
  by = c("max", "median", "mean", "min"),
  logfc = "logFC",
  pval = "P",
  inc_cols,
  p_adj_method = "fdr",
  merge_within = 1L,
  ignore_strand = TRUE,
  ...
)

## S4 method for signature 'RangedSummarizedExperiment'
mergeByCol(
  x,
  df = NULL,
  col,
  by = c("max", "median", "mean", "min"),
  logfc = "logFC",
  pval = "P",
  inc_cols,
  p_adj_method = "fdr",
  merge_within = 1L,
  ignore_strand = FALSE,
  ...
)
```

**Arguments**

x	A GenomicRanges or SummarizedExperiment object
...	Not used
df	A data.frame-like object containing the columns of interest. If not provided, any columns in the mcols() slot will be used.



col	The column to select as representative of the merged ranges
by	The method for selecting representative values
logfc	Column containing logFC values
pval	Column containing p-values
inc_cols	Any additional columns to return. Output will always include columns specified in the arguments col, logfc and pval. Note that values from any additional columns will correspond to the selected range returned in keyval_range
p_adj_method	Any of <a href="#">p.adjust.methods</a>
merge_within	Merge any ranges within this distance
ignore_strand	Passed internally to <a href="#">reduce</a> and <a href="#">findOverlaps</a>

### Details

This merges sliding windows using the values in a given column to select representative values for the subsequent merged windows. Values can be chosen from the specified column using any of `min()`, `max()`, `mean()` or `median()`, although `max()` is strongly recommended when specifying values like logCPM. Once a representative range is selected using the specified column, values from columns specified using `inc_cols` are also returned. In addition to these columns, the range from the representative window is returned in the `mcols` element as a `GRanges` object in the column `keyval_range`.

Merging windows using either the logFC or p-value columns is not implemented.

If adjusted p-values are requested an additional column names the same as the initial p-value, but tagged with the adjustment method, will be added. In addition, using the p-value from the selected window, the number of windows with lower p-values are counted by direction and returned in the final object. The selected window will always be counted as up/down regardless of significance as the p-value for this column is taken as the threshold. This is a not dissimilar approach to [cluster-direction](#).

If called on a `SummarizedExperiment` object, the function will be applied to the `rowRanges` element.

### Value

A Genomic Ranges object

### Examples

```
x <- GRanges(c("chr1:1-10", "chr1:6-15", "chr1:51-60"))
set.seed(1001)
df <- DataFrame(logFC = rnorm(3), logCPM = rnorm(3,8), p = rexp(3, 10))
mergeByCol(x, df, col = "logCPM", pval = "p")
mcols(x) <- df
x
mergeByCol(x, col = "logCPM", pval = "p")
```

---

`mergeByHMP`*Merge Sliding Windows using the Harmonic Mean P*

---

**Description**

Merge overlapping windows using harmonic mean p-values from significance testing

**Usage**

```
mergeByHMP(x, ...)  
  
## S4 method for signature 'GenomicRanges'  
mergeByHMP(  
  x,  
  df = NULL,  
  w = NULL,  
  logfc = "logFC",  
  pval = "P",  
  cpm = "logCPM",  
  inc_cols = NULL,  
  p_adj_method = "fdr",  
  merge_within = 1L,  
  ignore_strand = TRUE,  
  ...  
)  
  
## S4 method for signature 'RangedSummarizedExperiment'  
mergeByHMP(  
  x,  
  df = NULL,  
  w = NULL,  
  logfc = "logFC",  
  pval = "P",  
  cpm = "logCPM",  
  inc_cols = NULL,  
  p_adj_method = "fdr",  
  merge_within = 1L,  
  ignore_strand = FALSE,  
  ...  
)
```

**Arguments**

<code>x</code>	GenomicRanges object
<code>...</code>	Not used

<code>df</code>	data.frame with results of differential binding analysis performed using a sliding window strategy. If not provided, the columns in the <code>mcols()</code> element of <code>x</code> will be used
<code>w</code>	vector of weights to applied when calculating harmonic mean p-values
<code>logfc, pval, cpm</code>	Column names for the values holding window specific estimates of change in binding ( <code>logfc</code> ), overall signal intensity ( <code>cpm</code> ) and the significance from statistical testing ( <code>pval</code> )
<code>inc_cols</code>	(Optional) Character vector of any additional columns in <code>df</code> to return. Values will correspond to the range in the <code>keyval_range</code> column
<code>p_adj_method</code>	One of <code>p.adjust.methods</code> or "fwer". If "fwer" is specified the adjusted harmonic-mean p-value will be returned in a form which strictly controls the experiment-wide FWER. Please see <code>vignette("harmonicmeanp")</code> for more details
<code>merge_within</code>	Merge any non-overlapping windows within this distance
<code>ignore_strand</code>	Passed internally to <code>reduce</code> and <code>findOverlaps</code>

### Details

When using sliding windows to test for differential signal, overlapping windows can be merged based on the significance of results. `mergeByHMP()` merges overlapping windows using the asymptotically exact harmonic mean p-value `p.hmp` from the individual, window-level tests. This tests the Null Hypothesis that there is no significance amongst the initial set of p-values, and returns a summarised value which controls the FDR within a set of tests (Wilson, PNAS, 2019). Multilevel testing across the set of results is currently implemented using `p_adj_method = "fwer"`

Given that the harmonic mean p-value is calculated from the inverse p-values, these are used to provide a *weighted average* of expression and logFC values in the returned object. Any weights provided in `w` are ignored for these values as they are simple representative estimates. The representative range returned in `keyval_range` corresponds to the window with the lowest p-value.

The total number of windows is also returned in the final object, with the summarised values `n_up` and `n_down` indicating the number of windows with raw p-values below the calculated harmonic mean p-value, and with the corresponding direction of change.

The column containing the harmonic mean p-values is returned as 'hmp'. An additional column with adjusted hmp-values is returned with the suffix '\_\*' added where the p-value adjustment method is added after the underscore.

### Value

A `GenomicRanges` object with merged ranges from the original object along with summarised or representative values from the relevant columns. The range corresponding to a representative values is also returned as described above

### Examples

```
x <- GRanges(c("chr1:1-10", "chr1:6-15", "chr1:51-60"))
set.seed(1001)
df <- DataFrame(logFC = rnorm(3), logCPM = rnorm(3,8), p = rexp(3, 10))
mergeByHMP(x, df, pval = "p")
```

```
mcols(x) <- df
x
mergeByHMP(x, pval = "p", p_adj_method = "fwer")
```

---

mergeBySig

*Merge overlapping ranges based on p-values*


---

### Description

Merge overlapping windows using p-values from significance testing

### Usage

```
mergeBySig(x, ...)

## S4 method for signature 'GenomicRanges'
mergeBySig(
  x,
  df = NULL,
  logfc = "logFC",
  pval = "P",
  cpm = "logCPM",
  inc_cols,
  p_adj_method = "fdr",
  alpha = 0.05,
  method = c("combine", "best", "minimal"),
  merge_within = 1L,
  ignore_strand = TRUE,
  ...
)

## S4 method for signature 'RangedSummarizedExperiment'
mergeBySig(
  x,
  df = NULL,
  logfc = "logFC",
  pval = "P",
  cpm = "logCPM",
  inc_cols,
  p_adj_method = "fdr",
  alpha = 0.05,
  method = c("combine", "best", "minimal"),
  merge_within = 1L,
  ignore_strand = TRUE,
  ...
)
```

**Arguments**

<code>x</code>	GenomicRanges object
<code>...</code>	Passed to all <code>csaw</code> functions being wrapped
<code>df</code>	<code>data.frame</code> with results of differential binding analysis performed using a sliding window strategy. If not provided, the columns in the <code>mcols()</code> element of <code>x</code> will be used
<code>logfc, pval, cpm</code>	Column names for the values holding window specific estimates of change in binding ( <code>logfc</code> ), overall signal intensity ( <code>cpm</code> ) and the significance from statistical testing ( <code>pval</code> )
<code>inc_cols</code>	(Optional) Character vector of any additional columns in <code>df</code> to return
<code>p_adj_method</code>	One of <code>p.adjust.methods</code>
<code>alpha</code>	Significance threshold to apply during internal calculations
<code>method</code>	Shorthand versions for which <code>csaw</code> strategy to use for merging windows. Choose from 'combine' ( <a href="#">combineTests</a> ), 'best' ( <a href="#">getBestTest</a> ) or 'minimal' ( <a href="#">minimalTests</a> ).
<code>merge_within</code>	Merge any non-overlapping windows within this distance
<code>ignore_strand</code>	Passed internally to <a href="#">reduce</a> and <a href="#">findOverlaps</a>

**Details**

When using sliding windows to test for differential signal, overlapping windows can be merged based on the significance of results. `mergeBySig()` is a wrapper to the functions [combineTests](#), [getBestTest](#) and [minimalTests](#), using each function's approach to finding a representative window. The returned object differs from those returned by the original functions in that the description of windows as 'up', 'down' or mixed is omitted and the genomic range corresponding to the representative window is also returned. Column names also correspond to those in the original object.

An additional column with adjusted p-values is returned. This column retains the same name as the original but with the suffix `'_*'` added where the p-value adjustment method is added after the underscore.

**Value**

A `GenomicRanges` object with overlapping ranges from the original object merged and representative values returned. The range corresponding to the representative values is also returned

**Examples**

```
x <- GRanges(c("chr1:1-10", "chr1:6-15", "chr1:51-60"))
set.seed(1001)
df <- Dataframe(logFC = rnorm(3), logCPM = rnorm(3,8), p = rexp(3, 10))
mcols(x) <- df
mergeBySig(x, pval = "p", method = "combine")
mergeBySig(x, pval = "p", method = "best")
mergeBySig(x, pval = "p", method = "min")
```

---

partitionRanges      *Partition a set of Genomic Ranges*

---

### Description

Partition a set of Genomic Ranges by another

### Usage

```
partitionRanges(x, y, ...)

## S4 method for signature 'GRanges,GRanges'
partitionRanges(
  x,
  y,
  y_as_both = TRUE,
  ignore.strand = FALSE,
  simplify = TRUE,
  suffix = c(".x", ".y"),
  ...
)
```

### Arguments

x, y	GenomicRanges objects
...	Not used
y_as_both	logical(1) If there are any unstranded regions in y, should these be assigned to both strands. If TRUE unstranded regions can be used to partition stranded regions
ignore.strand	If set to TRUE, then the strand of x and y is set to "*" prior to any computation.
simplify	Pass to chopMC and simplify mcols in the output
suffix	Added to any shared column names in the provided objects

### Details

The query set of ranges can be broken in regions which strictly overlap a second set of ranges. The complete set of mcols from both initial objects will included in the set of partitioned ranges

### Value

A GRanges object

**Examples**

```
x <- GRanges(c("chr1:1-10", "chr1:6-15"))
x$id <- paste0("range", seq_along(x))
x
y <- GRanges(c("chr1:2-5", "chr1:6-12"))
y$id <- paste0("range", seq_along(y))
y
partitionRanges(x, y)
```

---

plotAssayDensities      *Plot Densities for any assay within a SummarizedExperiment*

---

**Description**

Plot Densities for any assay within a SummarizedExperiment

**Usage**

```
plotAssayDensities(x, ...)

## S4 method for signature 'SummarizedExperiment'
plotAssayDensities(
  x,
  assay = "counts",
  colour = NULL,
  linetype = NULL,
  group,
  trans = NULL,
  n_max = Inf,
  ...
)
```

**Arguments**

x	A SummarizedExperiment object
...	Passed to <a href="#">density</a>
assay	An assay within x
colour	The column in colData to colour lines by. To remove any colours, set this argument to NULL
linetype	Any optional column in colData used to determine linetype
group	Used by <a href="#">geom_line</a> . Defaults to the sample names but setting to NULL will over-write this and only groups specified by colour or linetype will be drawn
trans	character(1). Any transformative function to be applied to the data before calculating the density, e.g. trans = "log2"
n_max	Maximum number of points to use when calculating densities

**Details**

Uses ggplot2 to create a density plot for all samples within the selected assay

**Value**

A ggplot2 object. Scales and labels can be added using conventional ggplot2 syntax.

**Examples**

```
data("se")
se$treatment <- c("E2", "E2", "E2", "E2DHT", "E2DHT", "E2DHT")
## Plot individual samples
plotAssayDensities(se, colour = "treatment")
## Plot combined within treatment groups
plotAssayDensities(se, colour = "treatment", group = NULL)
## Use a data transformation
plotAssayDensities(se, trans = "log1p", colour = "treatment")
```

---

plotAssayHeatmap      *Draw a heatmap from a single SummarizedExperiment assay*

---

**Description**

Use ggplot2 to create a heatmap from a SummarizedExperiment object

**Usage**

```
plotAssayHeatmap(x, ...)
```

```
## S4 method for signature 'SummarizedExperiment'
plotAssayHeatmap(
  x,
  assay = "counts",
  by_x = "colnames",
  facet_x = NULL,
  ysideline = FALSE,
  yside_col = NULL,
  trans = NULL,
  n_max = 100,
  ...
)
```



**Arguments**

<code>x</code>	a SummarizedExperiment object
<code>...</code>	Not used
<code>assay</code>	the assay to take values from
<code>by_x</code>	the parameter to use for the x-axis. Will default to column names but should be one value per sample, such as an additional column containing shortened sample labels.
<code>facet_x</code>	column from <code>colData(x)</code> which will be used to group samples along the x-axis
<code>yside_line</code>	logical(1) Draw a line across the side of the y-axis summarising values for each range
<code>yside_col</code>	column from <code>colData(x)</code> to group and colour the lines drawn on the side of the y-axis. If grouping by treatment or replicate, the mean values will be shown
<code>trans</code>	character(1). Any transformative function to be applied to the data before calculating the density, e.g. <code>trans = "log2"</code>
<code>n_max</code>	Maximum number of ranges to draw

**Details**

Draw a heatmap containing selected values from an assay within a SummarizedExperiment object. Columns within the `colData` element of the object can be used to facet along the x-axis (e.g. treatment groups). The maximum number of points is set to be 100, although this can be changed easily should the plot require more ranges to be drawn.

The averages across any grouping of samples can be drawn as a line plot on the side of the y-axis by setting `yside_line = TRUE`, with groups as specified in `yside_col`. This feature is added for the specific context of neighbouring or overlapping ranges, and as such may be less informative in any other scenario

The returned object is a `ggplot2` object so scales can easily be added after heatmap creation using `scale_fill_*` for the main heatmap, and `scale_colour_*` for any groupings along the y-axis

**Value**

A `ggplot2` object. Scales and labels can be added using conventional `ggplot2` syntax.

**Examples**

```
nrows <- 10; ncols <- 4
counts <- matrix(runif(nrows * ncols, 1, 1e4), nrows)
colnames(counts) <- paste0("Sample_", seq_len(ncols))
df <- DataFrame(treat = c("A", "A", "B", "B"))
se <- SummarizedExperiment(
  assays = SimpleList(counts = counts),
  colData = df
)
rowRanges(se) <- GRanges(paste0("chr1:", seq_len(nrows)))
plotAssayHeatmap(se, facet_x = "treat")
```

---

plotAssayPCA

*Plot PCA For any assay within a SummarizedExperiment*


---

### Description

Plot PCA for any assay within a SummarizedExperiment object

### Usage

```
plotAssayPCA(x, ...)

## S4 method for signature 'SummarizedExperiment'
plotAssayPCA(
  x,
  assay = "counts",
  colour = NULL,
  shape = NULL,
  size = NULL,
  label = NULL,
  show_points = TRUE,
  pc_x = 1,
  pc_y = 2,
  trans = NULL,
  n_max = Inf,
  tol = sqrt(.Machine$double.eps),
  rank = NULL,
  ...
)
```

### Arguments

x	An object containing an assay slot
...	Passed to <a href="#">geom_text</a>
assay	The assay to perform PCA on
colour	The column name to be used for colours
shape, size	The column name(s) to be used for determining the shape or size of points
label	The column name to be used for labels
show_points	logical(1). Display the points. If TRUE any labels will repel. If FALSE, labels will appear at the exact points
pc_x	numeric(1) The PC to plot on the x-axis
pc_y	numeric(1) The PC to plot on the y-axis
trans	character(1). Any transformative function to be applied to the data before performing the PCA, e.g. trans = "log2"
n_max	Subsample the data to this many points before performing PCA

tol	Any rows with variance below this value will be excluded prior to passing to <a href="#">prcomp</a> . All rows are scaled and centred by default
rank	Passed to <a href="#">prcomp</a>

**Details**

Uses `ggplot2` to create a PCA plot for the selected assay. Any numerical transformation prior to performing the PCA can be specified using the `trans` argument

**Value**

A `ggplot2` object

**Examples**

```
data("se")
se$treatment <- c("E2", "E2", "E2", "E2DHT", "E2DHT", "E2DHT")
se$sample <- colnames(se)
plotAssayPCA(se, trans = "log1p", colour = "treatment", label = "sample")
plotAssayPCA(
  se, trans = "log1p", colour = "treatment", label = "sample",
  show_points = FALSE
)
```

---

plotAssayRle

*Plot RLE for a given assay within a SummarizedExperiment*


---

**Description**

Plot RLE for a given assay within a `SummarizedExperiment`

**Usage**

```
plotAssayRle(x, ...)

## S4 method for signature 'SummarizedExperiment'
plotAssayRle(
  x,
  assay = "counts",
  colour = NULL,
  fill = NULL,
  rle_group = NULL,
  by_x = NULL,
  n_max = Inf,
  trans = NULL,
  ...
)
```

**Arguments**

x	A SummarizedExperiment object
...	Passed to <a href="#">geom_boxplot</a>
assay	The assay to plot
colour	Column from colData(x) to outline the boxplots
fill	Column from colData(x) to fill the boxplots
rle_group	Column from colData(x) to calculate RLE within groups Commonly an alternative sample label.
by_x	Boxplots will be drawn by this grouping variable from colData(x). If not specified, the default values will be colnames(x)
n_max	Maximum number of points to plot
trans	character(1). NUerical transformation to apply to the data prior to RLE calculation

**Details**

Uses ggplot2 to create an RLE plot for the selected assay. Any numerical transformation prior to performing the RLE can be specified using the trans argument

**Value**

A ggplot2 object

**Examples**

```
data("se")
se$treatment <- c("E2", "E2", "E2", "E2DHT", "E2DHT", "E2DHT")
se$sample <- colnames(se)
## A conventional RLE Plot using all samples
plotAssayRle(se, trans = "log1p", fill = "treatment")
## Calculate RLE within groups
plotAssayRle(se, trans = "log1p", fill = "treatment", rle_group = "treatment")
# Or show groups combined
plotAssayRle(se, trans = "log1p", fill = "treatment", by_x = "treatment")
```

---

plotHFGC

---

*Plot a Genomic Region showing HiC, Features, Genes and Coverage*


---

**Description**

Plot a region with showing HiC, Features, Genes and Coverage

**Usage**

```

plotHFGC(
  gr,
  hic,
  features,
  genes,
  coverage,
  annotation,
  zoom = 1,
  shift = 0,
  max = 1e+07,
  axistrack = TRUE,
  cytobands,
  covtype = c("1", "heatmap"),
  linecol = c(),
  gradient = hcl.colors(101, "viridis"),
  hiccol = list(anchors = "lightblue", interactions = "red"),
  featcol,
  genecol,
  annotcol,
  highlight = "blue",
  hicsize = 1,
  featsize = 1,
  genesize = 1,
  covsize = 4,
  annotsize = 0.5,
  hicname = "HiC",
  featname = "Features",
  featstack = c("full", "hide", "dense", "squish", "pack"),
  collapseTranscripts = "auto",
  maxTrans = 12,
  ylim = NULL,
  ...,
  fontsize = 12,
  cex.title = 0.8,
  rotation.title = 0,
  col.title = "white",
  background.title = "lightgray",
  title.width = 1.5
)

```

**Arguments**

<code>gr</code>	The range(s) of interest. Must be on a single chromosome
<code>hic</code>	Any HiC interactions to be included as a GenomicInteractions object. If not supplied, no HiC track will be drawn.
<code>features</code>	A named GRangesList or list of GRangesList objects. Each GRangesList should contain features in each element which will drawn on the same track. If provid-

ing a list, each GRangesList within the list will drawn on a separate track. If this argument is not specified, no feature track will be drawn. Features will be drawn with colours provided in featcol.

genes	A GRanges object with exon structure for each transcript/gene. If not included, no track will be drawn for gene/transcript structure
coverage	A named list of BigWigFileList objects containing the primary tracks to show coverage for. Each list element will be drawn on a separate track, with elements within each BigWigFileList shown on the same track. List names will become track names. Alternatively, a single BigWigFileList will plot all individual files on separate tracks. If not included, no coverage tracks will be drawn.
annotation	Annotations for the coverage track(s). A single GRangesList if coverage is a BigWigListList. If coverage is supplied as a list of BigWigFileLists, a named list of GRangesList objects for each coverage track being annotated. Names must match those given for coverage.
zoom	Multiplicative factor for zooming in and out
shift	Shift the plot. Applied after zooming
max	The maximum width of the plotting region. Given that the width of the final plotting window will be determined by any HiC interactions, this argument excludes any interactions beyond this distance. Plotting can be somewhat slow if any long range interactions are included. Ignored if no HiC interactions are supplied.
axistrack	logical. Add an AxisTrack()
cytobands	Cytogenetic bands to be displayed on each chromosome. See data('grch37.cytobands') for the correct format. Only drawn if a cytobands data.frame is provided.
covtype	The plot type for coverage. Currently only lines ("l") and heatmaps ("heatmap") are supported
linecol	If passing a BigWigFileList to coverage, a vector of colours. If passing a list of BigWigFileList objects to coverage, a list of colours with structure that matches the object being passed to coverage, i.e. a named list of the same length, with elements whose length matches each BigWigFileList. Only used if covtype = "l".
gradient	Colour gradient for heatmaps
hiccol	list with names "anchors" and "interactions". Colours are passed to these elements
featcol	Named vector (or list) of colours for each feature. Must be provided if drawing features
genecol	Named vector (or list) of colours for each gene category
annotcol	Colours matching the coverage annotations
highlight	Outline colour for the highlight track. Setting this to NULL will remove the highlight
hicsize, featsize, genesize, covsize, annotsize	Relative sizes for each track (hic, features, genes, coverage & annotation)
hicname, featname	Names displayed in the LHS panel

featstack	Stacking for the future track
collapseTranscripts	Passed to <a href="#">GeneRegionTrack</a> for the genes track. Defaults to "auto" for automatic setting. If the number of transcripts to be plotted is > maxtrans, the argument will be automatically set to "meta", otherwise this will be passed as FALSE which will show all transcripts.
maxTrans	Only used if collapseTranscripts is set to "auto".
ylim	If a numeric vector, this will be passed to all coverage tracks. Alternatively, a named list of y-limits for each coverage track with names that match those in each element of the coverage list.
...	Passed to <a href="#">DataTrack</a> for the <b>coverage tracks</b> only. Useful arguments may be things like legend
fontsize	Applied across all tracks
cex.title	Passed to all tracks
rotation.title	Passed to all tracks
col.title	Passed to all tracks
background.title	Passed to all tracks
title.width	Expansion factor passed to <a href="#">plotTracks</a> , and used to widen the panels on the LHS of all tracks. Can have unpredictable effects on the font size of y-axis limits due to the algorithm applied by plotTracks

## Details

Convenience function for plotting a common set of tracks. All tracks are optional. For more fine control, users are advised to simply use Gviz directly.

The primary tracks defined in this function are H (HiC), F (features), G (genes), and C (coverage). Axis and Ideogram tracks are an additional part of this visualisation, with the Ideogram also being optional

Use all tracks specific to this dataset to generate a simple visualisation. In descending order the tracks displayed will be:

1. HiC Interactions (if supplied)
2. Regulatory features
3. Genes/genes
4. Coverage tracks as supplied

All tracks are optional and will simply be omitted if no data is supplied. See individual sections below for a more detailed explanation of each track

If wanting a single track of genes, simply pass a GRanges object in the format specified for a [GeneRegionTrack](#). Passing a GRangesList with the same format will yield an individual track for each list element, with each track shown by default as a separate colour. This can be used for showing Up/Down-regulated genes, or Detected/Undetected genes.

If passing a BigWigFileList for the coverage track, each file within the object will be drawn on a separate track. If specified, the same y-limits will be applied to each track. If passing a list of

BigWigFileList objects, each list element will be drawn as a single track with the individual files within each BigWigFileList overlaid within each track.

Cytogenetic band information must be in the structure required by [IdeogramTrack](#), with data for both GRCh37 and GRCh38 provided in this package ([grch37.cytobands](#), [grch38.cytobands](#)).

A highlight overlay over the GRanges provided as the `gr` argument will be added if a colour is provided. If set to NULL, no highlight will be added.

### Value

A Gviz object

### Displaying HiC Interactions

The available arguments for displaying HiC Interactions are defined below. If `hic` is supplied, a single [InteractionTrack](#) will be added displaying all interactions with an anchor within the range specified by `gr`. Only interactions with an anchor explicitly overlapping `gr` will be shown. If no interactions are found within `gr`, the track will not be displayed. The **plotting range will expand to incorporate these interactions**, with the parameter `max` providing an upper limit on the displayed range.

**hic** This is the `GInteractions` object required for inclusion of a HiC track in the final output. Will be ignored if not supplied

**hiccol** Determines the colours used for display of anchors and interactions

**hicsize** Relative size of the track compared to others

**hicname** The name to display on the LHS panel

**max** The maximum width of the plotted region. If multiple long-range interactions are identified, this provides an upper limit for the display. This defaults to 10Mb.

### Displaying Features

If wanting to add an [AnnotationTrack](#) with regions defined as 'features', the following arguments are highly relevant. All are ignored if `features` is not provided.

**features** A named `GRangesList`. Each element will be considered as a separate feature and drawn as a block in a distinct colour. Any `mcols` data will be ignored.

**featcol** A **named** vector (or list) providing a colour for each element of features

**featname** The name to display on the LHS panel

**featstack** Stacking to be applied to all supplied features

**featsize** Relative size of the track compared to others

### Displaying Genes And Transcripts

To display genes or transcripts, simply provide a single `GRanges` object if you wish to display all genes on a single track. The `mcols` element of this object should contain the columns `feature`, `gene`, `exon`, `transcript` and `symbol` as seen on the [GeneRegionTrack](#) help page.

Alternatively, a `GRangesList` can be provided to display genes on separate tracks based on their category. This can be useful for separating and colouring Up/Down regulated genes in a precise



way. All elements should be as described above. Again, all parameters associated with this track-set will be ignored if no object is supplied to this argument.

**genes** A GRanges or GRangesList object as described above

**genecol** A single colour if supplying a GRanges object, or a **named** vector/list of colours matching the GRangesList

**genesize** Relative size of the track compared to others

**collapseTranscripts** Passed to all tracks. See the GeneRegionTrack section in [settings](#) for detail regarding possible arguments. If genes is a GRangesList, can be a **named** vector/list with names matching the names of the genes object.

### Displaying Coverage Tracks

This section contains the most flexibility and can take two types of input. The first option is a BigWigFileList, which will lead to each BigWig file being plotted on its own track. An alternative is a list of BigWigFileList objects. In this case, each list element will be plotted as a separate track, with all individual BigWig files within each list element overlaid within the relevant track.

In addition to the coverage tracks, annotations can be added to each BigWigFileList in the form of coloured ranges, indicating anything of the user's choice. Common usage may be to indicate regions with binding of a ChIP target is found to be detected, unchanged, gained or lost.

**coverage** A BigWigFileList or list of BigWigFileList objects. A single BigWigFileList will be displayed with each individual file on a separate track with independent y-axes. Each element of the BigWigFileList **must be named** and these names will be displayed on the LHS panels. A list of BigWigFileList objects will be displayed with each list element as a separate track, with any BigWig files overlaid using the same y-axis. The list **must be named** with these names displayed on the LHS panel. Each internal BigWig within a BigWigFileList must also be named.

**covtype** Currently only lines (covtype = "l") and heatmaps (covtype = "heatmap") are supported. Colours can be specified using the arguments below

**linecol** Can be a single colour applied to all tracks, or a *named* vector (or list) of colours. If coverage is a single BigWigFileList, these names should match the names of this object exactly. If coverage is a list of BigWigFileList objects, linecol should be a list with matching names. Each element of this list should also be a **named** vector with names that exactly match those of each corresponding BigWigFileList.

**gradient** A colour gradient applied to all heatmap tracks. No specific structure is required beyond a vector of colours.

**covsize** Relative size of the tracks compared to others

**ylim** Can be a vector of length 2 applied to all coverage tracks. Alternatively, if passing a list of BigWigFileList objects to the coverage argument, this can be a **named** list of numeric vectors with names matching coverage

**annotation** Each BigWigFileList needs annotations to be passed to this argument as a **named** GRangesList, with names being used to associate unique colours with that set of ranges. If coverage is a BigWigFileList a simple GRangesList would be supplied and a single 'annotation' track will appear at the top of the set of coverage tracks. If coverage is a list, then a **named** list of GRangesList objects should be supplied, with each being displayed above the corresponding track from the coverage object.

**annotcol** A vector of colours corresponding to all names within all GRangesList elements supplied as annotation. It is assumed that the same colour scheme will be applied to all annotation tracks and, as such, the colours should **not** be provided as a list which matches the coverage tracks. Instead, every named element anywhere in the annotation GRanges, across all of the tracks must be included as a colour

**annotsize** Relative size of the tracks compared to others

## Examples

```
library(rtracklayer)
## Make sure we have the cytobands active
data(grch37.cytobands)

## Prepare the HiC, promoter & transcript information
data(ex_hic, ex_trans, ex_prom)
ex_features <- GRangesList(Promoter = ex_prom)
featcol <- c(Promoter = "red")

## Prepare the coverage
f1 <- system.file(
  "extdata", "bigwig", c("ex1.bw", "ex2.bw"), package = "extraChIPs"
)
bwfl <- BigWigFileList(f1)
names(bwfl) <- c("ex1", "ex2")
bw_col <- c(ex1 = "#4B0055", ex2 = "#007094")

## Define the plotting range
gr <- GRanges("chr10:103862000-103900000")

## Now create the basic plot
plotHFGC(
  gr,
  hic = ex_hic, features = ex_features, genes = ex_trans, coverage = bwfl,
  featcol = featcol, linecol = bw_col, cytobands = grch37.cytobands
)

plotHFGC(
  gr,
  hic = ex_hic, features = ex_features, genes = ex_trans, coverage = bwfl,
  featcol = featcol, linecol = bw_col, cytobands = grch37.cytobands,
  maxTrans = 1
)
```

---

plotOverlaps

*Plot Overlaps Between List Elements*

---

## Description

Plot Overlaps between list elements as an upset or Venn diagram

**Usage**

```

plotOverlaps(x, ...)

## S4 method for signature 'GRangesList'
plotOverlaps(
  x,
  type = c("auto", "venn", "upset"),
  var = NULL,
  f = c("mean", "median", "max", "min", "sd"),
  set_col = NULL,
  ...,
  .sort_sets = "ascending",
  min.gapwidth = 1L,
  ignore.strand = TRUE
)

## S4 method for signature 'list'
plotOverlaps(
  x,
  type = c("auto", "venn", "upset"),
  set_col = NULL,
  ...,
  .sort_sets = "ascending"
)

```

**Arguments**

x	GRangesList of S3 list able to be coerced to character vectors
...	Passed to <a href="#">draw.pairwise.venn</a> (or <a href="#">draw.single/triple.venn</a> ) for Venn Diagrams, and to <a href="#">upset</a> for UpSet plots
type	The type of plot to be produced
var	Column to summarised as a boxplot in an upper panel (UpSet plot only)
f	Summarisation function. Must return a single value from any numeric vector
set_col	Colours to be assigned to each set
.sort_sets	passed to <code>sort_sets</code> in <a href="#">upset</a>
min.gapwidth, ignore.strand	Passed to <a href="#">reduce</a>

**Details**

This function should give the capability to show overlaps for any number of replicates or groups, or a list of items such as gene names. For  $n = 2$ , a scaled Venn Diagram will be produced, however no scaling is implemented for  $n = 3$

UpSet plots are possible for any lists with length  $> 1$ , and are the only implemented possibility for lists  $> 3$ .

If the input is a GRangesList an additional boxplot can be requested using any numeric column within the existing mcols() element. Values will be summarised across all elements using the requested function and the boxplot will be included as an upper panel above the intersections

### Value

Either a VennDiagram (i.e. grid) object, or a ComplexUpset plot

### Examples

```
## Examples using a list of character vectors
ex <- list(
  x = letters[1:5], y = letters[c(6:15, 26)], z = letters[c(2, 10:25)]
)
plotOverlaps(ex, type = "upset")
plotOverlaps(ex, type = "venn", set_col = 1:3, alpha = 0.3)
plotOverlaps(ex, type = "upset", set_col = 1:3, labeller = stringr::str_to_title)
plotOverlaps(ex[1:2])

## GRangesList object will produce a boxplot of summarised values in the
## upper panel
data("peaks")
gr1 <- peaks[1:3]
names(gr1) <- gsub("_peaks.", "", names(gr1))
plotOverlaps(gr1, type = 'upset', var = 'score', f = 'max')

## If only two samples are present, a VennDiagram will be produced
plotOverlaps(gr1[1:2], set_col = c("green", "blue"))
```

---

plotPie

*Draw Pie Graphs based on one or more columns*

---

### Description

Draw Pie Graphs based one or more data.frame columns

### Usage

```
plotPie(object, ...)

## S4 method for signature 'GRanges'
plotPie(object, scale_by = c("n", "width"), ...)

## S4 method for signature 'DataFrame'
plotPie(object, ...)

## S4 method for signature 'data.frame'
plotPie(
```

```

object,
fill,
x,
y,
scale_by,
scale_factor = 1000,
width = 0.8,
total_geom = c("label", "text", "none"),
total_glue = "{comma(N})",
total_colour = "black",
total_fill = "white",
total_alpha = 1,
total_size = 3,
min_p = 0.01,
max_p = 1,
cat_geom = c("label", "text", "none"),
cat_glue = "{.data[[fill]]}\n{comma(n, 1)}\n({percent(p, 0.1})",
cat_colour = "black",
cat_fill = "white",
cat_size = 3,
cat_alpha = 1,
cat_adj = 0,
...
)

```

### Arguments

object	An object (data.frame)
...	Not used
scale_by	Scale the counts by this column. In this case of a GRanges object this defaults to the count (scale_by = "n") but can also be specified as being width of each range (scale_by = "width"). If choosing width, width will be displayed in Kb
fill	The category/column used to fill the slices of the pie charts
x	The second (optional) category/column to place along the x-axis
y	The final (optional) category/column to place along the y-axis
scale_factor	When scaling by another column, such as width, totals will be divided by this value, with 1000 being the default to provide output in kb.
width	Scale the width of all pies
total_geom	The geom_* to use for the totals at the centre of each pie. Setting this to 'none' will disable totals
total_glue	<a href="#">glue</a> syntax to use for the totals in the centre of each pie. The column 'N' will produce the totals and any other values or formatting may be added here.
total_colour, total_fill, total_alpha, total_size	Colour, fill, alpha and size for the main totals in the centre of each pie chart

min_p	The minimum proportion of the total required for adding labels. Effectively removes labels from pie charts with few members. Alternatively when only one column is specified, categories below this will not be shown around the edge of the plot
max_p	only display labels for segments representing less than this proportion of the total.
cat_geom	The geom_* to use for category labels corresponding to each slice of the pie. Setting this to 'none' will disable category labels
cat_glue	<a href="#">glue</a> syntax to use for the category labels corresponding to each slice of the pie charts. The columns 'n' and 'p' can be used to print totals and proportions for each slice.
cat_colour, cat_fill, cat_size, cat_alpha	Colour, fill, size and alpha for category labels
cat_adj	Adjust category labels

### Details

Using a `data.frame` as input, this function will draw pie graphs based on one or more columns, by simply counting the values in combination across these columns. One column must be selected for the fill as a bare minimum, with up to three being possible. Additional columns can be set for the x-axis to draw a series of pie-graphs in a row, with a further column able to be added to layout a series of pie graphs in a grid

If only one column/category is chosen, category labels will be added around the edge of the plot

If `show_total = TRUE` the overall counts for each pie graph will be added in the centre using [geom\\_label](#). Parameters for these labels are customisable

### Value

A `ggplot2` object able to be customised with colour scales and themes.

Also note that the `$data` element of the returned object will contain the `data.frame` used for plotting. The additional column `label_radians` represents the mid-point of each pie slice and can be used for manually adding labels to each pie. Only applies when plotting across the x or y axes

### Examples

```
set.seed(200)
df <- data.frame(
  feature = sample(
    c("Promoter", "Enhancer", "Intergenic"), 200, replace = TRUE
  ),
  TF1 = sample(c("Up", "Down", "Unchanged"), 200, replace = TRUE),
  TF2 = sample(c("Up", "Down", "Unchanged"), 200, replace = TRUE),
  w = rexp(200)
)
plotPie(df, fill = "feature", total_glue = "N = {comma(N)}")
plotPie(
  df, fill = "feature", scale_by = "w", total_geom = "none",
  cat_glue = "{percent(p)}", cat_size = 5
)
```

```

)
plotPie(df, fill = "feature", x = "TF1")
plotPie(
  df, fill = "feature", x = "TF1", y = "TF2", min_p = 0.02,
  total_geom = "none", cat_glue = "{n} / {N}"
) +
scale_fill_viridis_d() +
theme_bw()

## And using a GRanges object
data("ex_prom")
gr <- ex_prom
mcols(gr) <- df[seq_along(gr),]
## Show values by counts
plotPie(gr, fill = "feature", total_size = 5)
## Show values scaled by width of each range
plotPie(
  gr, fill = "feature", scale_by = "width", total_glue = "{round(N, 1)}kb",
  cat_glue = "{percent(p, 0.1)}", cat_size = 4, total_size = 5
)

```

---

plotProfileHeatmap     *Draw a coverage Profile Heatmap*

---

## Description

Plot a coverage Profile Heatmap across multiple ranges

## Usage

```

plotProfileHeatmap(object, ...)

## S4 method for signature 'GenomicRangesList'
plotProfileHeatmap(
  object,
  profileCol,
  xValue = "bp",
  fillValue = "score",
  facetX = NULL,
  facetY = NULL,
  colour = facetY,
  linetype = NULL,
  summariseBy = c("mean", "median", "min", "max", "none"),
  xLab = xValue,
  yLab = NULL,
  fillLab = fillValue,
  relHeight = 0.3,

```

```

    ...
  )

## S4 method for signature 'GenomicRanges'
plotProfileHeatmap(
  object,
  profileCol,
  xValue = "bp",
  fillValue = "score",
  facetX = NULL,
  facetY = NULL,
  colour = facetY,
  linetype = NULL,
  summariseBy = c("mean", "median", "min", "max", "none"),
  xLab = xValue,
  yLab = NULL,
  fillLab = fillValue,
  relHeight = 0.3,
  ...
)

```

### Arguments

object	A GRanges or GRangesList object
...	Passed to <code>facet_grid</code> internally. Can be utilised for switching panel strips or passing a labeller function
profileCol	Column name specifying where to find the profile DataFrames
xValue, fillValue	Columns within the profile DataFrames for heatmaps
facetX, facetY	Column used for facetting across the x- or y-axis respectively
colour	Column used for colouring lines in the summary panel. Defaults to any column used for facetY
linetype	Column used for linetypes in the summary panel
summariseBy	Function for creating the summary plot in the top panel. If set to 'none', no summary plot will be drawn. Otherwise the top panel will contain a line-plot representing this summary value for each x-axis bin
xLab, yLab, fillLab	Labels for plotting aesthetics. Can be overwritten using <code>labs()</code> on any returned object
relHeight	The relative height of the top summary panel. Represents the fraction of the plotting area taken up by the summary panel.

### Details

Convenience function for plotting coverage heatmaps across a common set of ranges, shared between one or more samples. These are most commonly the coverage values from merged samples



within a treatment group. The input data structure is based on that obtained from `getProfileData`, and can be provided either as a `GRanges` object (generally for one sample) or as a `GRangesList`.

A 'profile `DataFrame`' here refers to a `data.frame` (or `tibble`, or `DataFrame`) with a coverage value in one column that corresponds to a genomic bin of a fixed size denoted in another, as generated by `getProfileData`. Given that multiple ranges are most likely to be drawn, each profile data frame must be the same size in terms of the number of bins, each of which represent a fixed number of nucleotides. At a minimum this is a two column data frame although `getProfileData` will provide three columns for each specified genomic region.

If using a `GRangesList`, each list element will be drawn as a separate panel by default. These panels will appear in the same order as the list elements of the `GRangesList`, although this can easily be overwritten by passing a column name to the `facetX` argument. The default approach will add the original element names as the column "name" which can be seen in the `$data` element of any resultant `ggplot` object produced by this function.

### Value

A `ggplot2` object, able to be customised using standard `ggplot2` syntax

### Examples

```
library(rtracklayer)
fl <- system.file(
  "extdata", "bigwig", c("ex1.bw", "ex2.bw"), package = "extraChIPs"
)
bwfl <- BigWigFileList(fl)
names(bwfl) <- c("ex1", "ex2")

gr <- GRanges(
  c(
    "chr10:103880281-103880460", "chr10:103892581-103892760",
    "chr10:103877281-103877460"
  )
)
pd <- getProfileData(bwfl, gr)
plotProfileHeatmap(pd, "profile_data") +
  scale_fill_viridis_c(option = "inferno", direction = -1) +
  labs(fill = "Coverage")
```

### Description

Create Donut charts based on one or two columns in a data frame

**Usage**

```
plotSplitDonut(object, ...)

## S4 method for signature 'GRanges'
plotSplitDonut(object, scale_by = c("n", "width"), ...)

## S4 method for signature 'DataFrame'
plotSplitDonut(object, ...)

## S4 method for signature 'data.frame'
plotSplitDonut(
  object,
  inner,
  outer,
  scale_by = NULL,
  scale_factor = 1000,
  r_centre = 0.5,
  r_inner = 1,
  r_outer = 1,
  total_glue = "{comma(N)}",
  total_size = 5,
  total_colour = "black",
  inner_glue = "{inner} {.data[[inner]]}\n{percent(p,0.1)}",
  outer_glue = "{outer} {.data[[outer]]}\n{percent(p,0.1)}",
  total_label = c("label", "text", "none"),
  inner_label = c("label", "text", "none"),
  outer_label = c("label", "text", "none"),
  label_alpha = 1,
  inner_label_alpha = NULL,
  outer_label_alpha = NULL,
  label_size = 3,
  inner_label_size = NULL,
  outer_label_size = NULL,
  label_colour = "black",
  inner_label_colour = NULL,
  outer_label_colour = NULL,
  min_p = 0.05,
  inner_min_p = NULL,
  outer_min_p = NULL,
  max_p = 1,
  inner_max_p = NULL,
  outer_max_p = NULL,
  explode_inner = NULL,
  explode_outer = NULL,
  explode_query = c("AND", "OR"),
  explode_x = 0,
  explode_y = 0,
  explode_r = 0,
```

```

    nudge_r = 0.5,
    expand = 0.1,
    inner_palette = NULL,
    outer_palette = NULL,
    inner_legend = TRUE,
    outer_legend = TRUE,
    layout = c(main = area(1, 1, 6, 6), lg1 = area(2, 7), lg2 = area(4, 7)),
    ...
)

```

## Arguments

object	A GRanges or data.frame-like object
...	Not used
scale_by	Column to scale values by. If provided, values in this column will be summed, instead of simply counting entries. Any label in the centre of the plot will also reflect this difference
inner	Column name to create the inner ring
outer	Column name to create the outer ring, subset by the inner ring
scale_factor	When scaling by another column, such as width, totals will be divided by this value, with 1000 being the default to provide output in kb.
r_centre	The radius of the hole in the centre. Setting to zero will create a Pie chart
r_inner, r_outer	The radii of the inner/outer rings
total_glue	<a href="#">glue</a> -syntax for formatting the total which appears in the centre of the plot. Internally, the value N will be calculated and as such, this value should appear within this argument.
total_size	Label size total number of entries in the centre of the plot.
total_colour	Label colour for the summary total in the centre
inner_glue, outer_glue	<a href="#">glue</a> -syntax for formatting labels which appear on each inner/outer segment Internally, the values n and p will be calculated as totals and proportions of the total. As such, these values can appear within this argument, as well as the fields described in the details
total_label, inner_label, outer_label	Can take values 'text', 'label' or 'none'. If setting one the first two values, the labelling function <code>geom_*</code> will be called, otherwise no label will be drawn
label_alpha, inner_label_alpha, outer_label_alpha	transparency for labels
label_size, inner_label_size, outer_label_size	Size of all text labels
label_colour, inner_label_colour, outer_label_colour	Takes any colour specification, with the additional option of 'palette'. In this special case, the same palette as is used for each segment will be applied.

min_p, inner_min_p, outer_min_p	only display labels for segments representing greater than this proportion of the total. If inner/outer values are specified, the values in min_p will be ignored for that layer
max_p, inner_max_p, outer_max_p	only display labels for segments representing less than this proportion of the total. If inner/outer values are specified, the values in max_p will be ignored for that layer
explode_inner, explode_outer	Regular expressions from either the inner or outer ring for which segments will be 'exploded'
explode_query	Setting to AND and specifying values for both the inner and outer ring will require matches in both categories
explode_x, explode_y	Numeric values for shifting exploded values
explode_r	Radius expansion for exploded values
nudge_r	Radius expansion for labels in the outer ring
expand	Passed to <a href="#">expansion</a> for both x and y axes
inner_palette	Colour palette for the inner ring
outer_palette	Optional colour palette for the outer ring
inner_legend, outer_legend	logical(1). Show legends for either layer
layout	Passed to <a href="#">plot_layout</a>

## Details

Using a data.frame or GRanges object, this function enables creation of a Pie/Donut chart with an inner and outer ring. The function itself is extremely flexible allowing for separate colour palettes in the inner and outer rings, as well as highly customisable labels.

Sections can be exploded using a value from the inner ring or outer ring separately, or in combination by setting `explode_query = "AND"`. Exploded sections can be shifted by expanding the radius (`explode_r`), or along the x/y co-ordinates using `explode_x/y`, allowing for detailed placement of sections.

If only the inner palette is specified, segments in the outer ring will be assigned the same colours as the inner segments, but with increased transparency. Only a single legend will be drawn in this scenario. If an outer palette is specified, both colour palettes are completely distinct and two distinct legends will be drawn. The placement of these legends, along with the larger donut plot, can be manually specified by providing a layout as defined in [plot\\_layout](#). Names are not required on this layout, but may be beneficial for code reproducibility.

The inner label denoting the total can also be heavily customised using the [glue](#) syntax to present the calculated value N along with any additional text, such as 'kb' if scaling GenomicRanges by width. The same approach can be taken for the inner and outer labels, where totals are held in the value n, proportions are held in the value p and the values corresponding to each segment can be accessed using `.data[[inner]]` or `.data[[outer]]`. Column titles can be added using `{inner}/{outer}`. Values from the inner segments can be added to the outer labels using this strategy enabling a wide variety of labelling approaches to be utilised.

**Value**

A patchwork object consisting of both ggplot2 objects and legend grobs

**Examples**

```
set.seed(200)
df <- data.frame(
  feature = sample(
    c("Promoter", "Enhancer", "Intergenic"), 200, replace = TRUE
  ),
  TF1 = sample(c("Up", "Down", "Unchanged"), 200, replace = TRUE),
  TF2 = sample(c("Up", "Down", "Unchanged"), 200, replace = TRUE)
)
## The standard plot
plotSplitDonut(df, inner = "TF1", outer = "TF2", inner_legend = FALSE)

## Adding an exploded section along with an outer palette & customisation
plotSplitDonut(
  df, inner = "TF1", outer = "feature", total_label = "none",
  inner_label_alpha = 0.5, r_centre = 0,
  outer_glue = "{.data[[outer]]}\n(n = {n})", outer_label = "text",
  explode_inner = "Up", explode_outer = "Prom|Enh",
  explode_query = "AND", explode_r = 0.4,
  inner_palette = hcl.colors(3, "Spectral", rev = TRUE),
  outer_palette = hcl.colors(3, "Cividis")
)
```

---

propOverlap

*Find the proportions of an overlapping range*


---

**Description**

Find the proportion of a query reange which overlaps the subject

**Usage**

```
propOverlap(x, y, ...)
```

```
## S4 method for signature 'GRanges,GRanges'
propOverlap(x, y, ignore.strand = FALSE, ...)
```

**Arguments**

x, y            A GenomicRanges object

...            Not used

ignore.strand   If set to TRUE, then the strand of x and y is set to "\*" prior to any computation.

**Details**

This behaves similarly to [overlapsAny](#) except the proportion of the query range which overlaps one or more subject ranges is returned instead of a logical vector

**Value**

Numeric vector the same length as x

**Examples**

```
x <- GRanges("chr1:1-10")
y <- GRanges("chr1:1-5")
propOverlap(x, y)
propOverlap(y, x)
```

---

 reduceMC

*Reduce ranges retaining mcols*


---

**Description**

Reduce ranges retaining mcols

**Usage**

```
reduceMC(x, ignore.strand = FALSE, simplify = TRUE, ...)
```

**Arguments**

x	A GenomicRanges object
ignore.strand	If set to TRUE, then the strand of x and y is set to "*" prior to any computation.
simplify	logical(1). Attempt to simplify returned columns where possible
...	Passed to <a href="#">reduce</a>

**Details**

This function extends [reduce](#) so that **all** mcols are returned in the output. Where the reduced ranges map to multiple ranges in the original range, mcols will be returned as CompressedList columns.

If simplify = TRUE columns will be returned as vectors where possible.

**Value**

A GRanges object

**Examples**

```
x <- GRanges(c("chr1:1-10:+", "chr1:6-12:-"))
x$id <- c("range1", "range2")
reduceMC(x)
reduceMC(x, ignore.strand = TRUE)
```

---

setoptsMC

*Perform set operations retaining mcols*


---

**Description**

Perform set operations retaining all mcols from the query range

**Usage**

```
setdiffMC(x, y, ...)

intersectMC(x, y, ...)

unionMC(x, y, ...)

## S4 method for signature 'GRanges,GRanges'
setdiffMC(x, y, ignore.strand = FALSE, simplify = TRUE, ...)

## S4 method for signature 'GRanges,GRanges'
intersectMC(x, y, ignore.strand = FALSE, simplify = TRUE, ...)

## S4 method for signature 'GRanges,GRanges'
unionMC(x, y, ignore.strand = FALSE, simplify = TRUE, ...)
```

**Arguments**

<code>x, y</code>	GenomicRanges objects
<code>...</code>	Not used
<code>ignore.strand</code>	If set to TRUE, then the strand of x and y is set to "*" prior to any computation.
<code>simplify</code>	logical(1) If TRUE, any List columns will be returned as vectors where possible. This can only occur if single, unique entries are present in all initial elements.

**Details**

This extends the methods provided by [setdiff](#), [intersect](#) and [union](#) so that mcols from x will be returned as part of the output.

Where output ranges map back to multiple ranges in x, CompressedList columns will be returned. By default, these will be simplified if possible, however this behaviour can be disabled by setting `simplify = FALSE`.

All columns will be returned which can also be time-consuming. A wise approach is to only provide columns you require as part of the query ranges `x`.

If more nuanced approaches are required, the returned columns can be further modified by many functions included in the `plyranges` package, such as `mutate()`.

### Value

A `GRanges` object with all `mcols` returned from the original object. If a range obtained by `setdiff` maps back to two or more ranges in the original set of `Ranges`, `mcols` will be returned as [CompressedList](#) columns

### Examples

```
x <- GRanges("chr1:1-100:+")
x$id <- "range1"
y <- GRanges(c("chr1:51-60:+", "chr1:21-30:-"))
setdiffMC(x, y)
setdiffMC(x, y, ignore.strand = TRUE)

# The intersection works similarly
intersectMC(x, y)

# Union may contain ranges not initially in x
unionMC(x, y)
unionMC(x, y, ignore.strand = TRUE)
```

---

stitchRanges

*Stitch Ranges within a given distance*

---

### Description

Stitch together ranges within a given distance, using excluded ranges as barriers that cannot be crossed

### Usage

```
stitchRanges(x, exclude, maxgap = 12500L, ignore.strand = TRUE)
```

### Arguments

<code>x</code>	Ranges to be stitched together
<code>exclude</code>	Ranges to exclude
<code>maxgap</code>	The maximum distance between ranges to be stitched
<code>ignore.strand</code>	logical



**Details**

Stitches together ranges within a given distance, using any ranges provided for exclusion as barriers between stitched ranges. This may be particularly useful if wanting to stitch enhancers whilst excluding promoters.

All inputs and outputs are Genomic Ranges objects

**Value**

A GRanges object

**Examples**

```
x <- GRanges(c("chr1:1-10", "chr1:101-110", "chr1:201-210", "chr2:1-10"))
y <- GRanges("chr1:200:+")
stitchRanges(x, exclude = y, maxgap = 100)
```

---

voomWeightsFromCPM      *Estimate voom precision weights directly From CPM values*

---

**Description**

Estimate voom precision weights directly From CPM values

**Usage**

```
voomWeightsFromCPM(
  cpm,
  design = NULL,
  w0 = NULL,
  lib.size = NULL,
  isLogCPM = TRUE,
  span = 0.5,
  ...
)
```

**Arguments**

cpm	Matrix of CPM or logCPM values
design	The design matrix for the experiment
w0	Initial vector of sample weights. Should be calculated using <a href="#">arrayWeights</a>
lib.size	Initial library sizes. Must be provided as these are not estimable from CPM values
isLogCPM	logical(1). Indicates whether the data is log2 transformed already. Most commonly (e.g. if using the output of <code>cpm</code> ) it will be,
span	Width of the smoothing window used for the lowess mean-variance trend. Expressed as a proportion between 0 and 1.
...	Passed to <code>lmFit</code> internally

## Details

This function takes CPM or logCPM values and estimates the precision weights as would be done by providing counts directly to the `voom` function. Using this function enables the use of logCPM values which have been normalised using other methods such as Conditional-Quantile or Smooth-Quantile Normalisation.

The precision weights are returned as part of the `EList` output, and these are automatically passed to the function `lmFit` during model fitting. This will ensure that the mean-variance relationship is appropriate for the linear modelling steps as performed by `limma`.

Initial sample weights can be passed to the function, and should be calculated using `arrayWeights` called on the normalised logCPM values. The returned sample weights will be different to these, given that the function `voomWithQualityWeights` performs two rounds of estimation. The first is on the initial data, with the inappropriate mean-variance relationship, whilst the second round is after incorporation of the precision weights.

## Value

An object of class `EList` as would be output by `voom`. Importantly, there will be no `genes` element, although this can be added later. Similarly, the returned `targets` element will only contain sample names and library sizes. This can be incorporated with any other metadata as required.

Plotting data is always returned, noting the the value `sx` has been offset by the library sizes and will be simple logCPM values. As such, the fitted `Amean` is also returned in this list element.

If initial sample weights were provided, modified weights will also be returned, as the initial function `voomWithQualityWeights` performs two rounds of estimation of sample weights. Here we would simply provide the initial weights a priori, with the second round performed within the function. Importantly, this second round of sample weight estimation uses the precision weights ensuring the correct mean-variance relationship is used for the final estimation of sample weights

## Examples

```
bamFiles <- system.file("exdata", c("rep1.bam", "rep2.bam"), package="csaw")
wc <- csaw::windowCounts(bamFiles, filter=1)
cpm <- edgeR::cpm(wc, log = TRUE)
e1 <- voomWeightsFromCPM(cpm, lib.size = wc$totals)
```

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