

Package ‘seqsetvis’

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Type Package

Title Set Based Visualizations for Next-Gen Sequencing Data

Version 1.12.0

Description seqsetvis enables the visualization and analysis of sets of genomic sites in next gen sequencing data.
Although seqsetvis was designed for the comparison of multiple ChIP-seq samples, this package is domain-agnostic and allows the processing of multiple genomic coordinate files (bed-like files) and signal files (bigwig files pileups from bam file).

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Encoding UTF-8

LazyData true

Suggests BiocFileCache, BiocManager, BiocStyle, ChIPpeakAnno, covr, cowplot, knitr, rmarkdown, testthat

Depends R (>= 3.6), ggplot2

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biocViews Software, ChIPSeq, MultipleComparison, Sequencing, Visualization

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seqsetvis-package

*easy awesome peak set vis TESTING seqsetvis allows you to...***Description**

2 steps [ssvOverlapIntervalSets](#). [ssvFetchBigwig](#). Otherwise refer to the vignettes to see

Author(s)

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<code>.expand_cigar_dt</code>	<i>Expand intermediate bam fetch by cigar codes</i>
-------------------------------	---

Description

see [sam specs](#) for cigar details

Usage

```
.expand_cigar_dt(cigar_dt, op_2count = c("M", "D", "=", "X"))
```

Arguments

<code>cigar_dt</code>	data.table with 5 required named columns in any order. <code>c("which_label", "seq-names", "strand", "start", "cigar")</code>
<code>op_2count</code>	Cigar codes to count. Default is alignment (M), deletion (D), match (=), and mismatch (X). Other useful codes may be skipped regions for RNA splicing (N). The locations of any insterions (I) or clipping/padding (S, H, or P) will be a single bp immediately before the interval.

Value

data.table with cigar entries expanded

<code>.expand_cigar_dt_recursive</code>	<i>Expand intermediate bam fetch by cigar codes</i>
---	---

Description

see [sam specs](#) for cigar details

Usage

```
.expand_cigar_dt_recursive(cigar_dt)
```

Arguments

<code>cigar_dt</code>	data.table with 5 required named columns in any order. <code>c("which_label", "seq-names", "strand", "start", "cigar")</code>
-----------------------	---

Value

data.table with cigar entries expanded

<code>.rm_dupes</code>	<i>Remove duplicate reads based on stranded start position. This is an over-simplification. For better duplicate handling, duplicates must be marked in bam and flag passed to fetchBam() ... for ScanBamParam</i>
------------------------	--

Description

`flag = scanBamFlag(isDuplicate = FALSE)`

Usage

`.rm_dupes(reads_dt, max_dupes)`

Arguments

<code>reads_dt</code>	data.table of reads as loaded by fetchBam
<code>max_dupes</code>	maximum allowed positional duplicates

Value

`reads_dt` with duplicated reads over `max_dupes` removed

<code>.rm_dupesPE</code>	<i>Remove duplicate paired-end reads based on start and end position. This is an over-simplification. For better duplicate handling, duplicates must be marked in bam and flag passed to fetchBamPE() ... for ScanBamParam</i>
--------------------------	--

Description

`flag = scanBamFlag(isDuplicate = FALSE)`

Usage

`.rm_dupesPE(reads_dt, max_dupes)`

Arguments

<code>reads_dt</code>	data.table of reads as loaded by fetchBamPE
<code>max_dupes</code>	maximum allowed positional duplicates

Value

`reads_dt` with duplicated reads over `max_dupes` removed

```
add_cluster_annotation
      add_cluster_annotation
```

Description

adds rectangle boxes proportional to cluster sizes of heatmap with optional labels.

Usage

```
add_cluster_annotation(
  cluster_ids,
  p = NULL,
  xleft = 0,
  xright = 1,
  rect_colors = c("black", "gray"),
  text_colors = rev(rect_colors),
  show_labels = TRUE,
  label_angle = 0,
  row_ = "id",
  cluster_ = "cluster_id"
)
```

Arguments

<code>cluster_ids</code>	Vector of cluster ids for each item in heatmap. Should be sorted by plot order for heatmap.
<code>p</code>	Optionally an existing ggplot to add annotation to.
<code>xleft</code>	left side of cluster annotation rectangles. Default is 0.
<code>xright</code>	right side of cluster annotation rectangles. Default is 1.
<code>rect_colors</code>	colors of rectangle fill, repeat to match number of clusters. Default is c("black", "gray").
<code>text_colors</code>	colors of text, repeat to match number of clusters. Default is reverse of rect_colors.
<code>show_labels</code>	logical, should rectangles be labelled with cluster identity. Default is TRUE.
<code>label_angle</code>	angle to add clusters labels at. Default is 0, which is horizontal.
<code>row_</code>	variable name mapped to row, likely id or gene name for ngs data. Default is "id" and works with ssvFetch* outputs.
<code>cluster_</code>	variable name to use for cluster info. Default is "cluster_id".

Value

A ggplot with cluster annotations added.

Examples

```
#simplest uses
add_cluster_annotation(factor(c(rep("A", 3), "B")))
p = ggplot() + coord_cartesian(xlim = c(0,10))
add_cluster_annotation(factor(c(rep("A", 3), "B")), p)

#intended use with ssvSignalHeatmap
clust_dt = ssvSignalClustering(CTCF_in_10a_profiles_dt, nclust = 3)
assign_dt = unique(clust_dt[, .(id, cluster_id)])[order(id)]
p_heat = ssvSignalHeatmap(clust_dt, show_clusterBars = FALSE)
add_cluster_annotation(assign_dt$cluster_id, p_heat,
  xleft = -400, xright = -360, rect_colors = rainbow(3), text_colors = "gray")
p_clusters = add_cluster_annotation(assign_dt$cluster_id,
  rect_colors = rainbow(3), text_colors = "gray")
#specialized use as plot outside of heatmap
assemble_heatmap_clusterBars(plots = list(p_clusters, p_heat), rel_widths = c(1, 3))
```

append_ynorm

*append_ynorm***Description**

see [calc_norm_factors](#) for normalization details.

Usage

```
append_ynorm(
  full_dt,
  value_ = "y",
  cap_value_ = "y_cap_value",
  norm_value_ = "y_norm",
  by1 = "id",
  by2 = "sample",
  aggFUN1 = max,
  aggFUN2 = function(x) quantile(x, 0.95),
  cap_dt = NULL,
  do_not_cap = FALSE,
  force_append = FALSE
)
```

Arguments

full_dt	a data.table, as returned by <code>ssvFetch*(..., return_data.table = TRUE)</code> .
value_	character, attribute in full_dt to normalzie.
cap_value_	character, new attribute name specifying values to cap to.
norm_value_	character, new attribute name specifying normalized values.
by1	character vector, specifies attributes relevant to step 1.

by2	character vector, specifies attributes relevant to step 1 and 2.
aggFUN1	function called on value_ with by = c(by1, by2) in step 1.
aggFUN2	function called on result of aggFUN1 with by = by2 in step 2.
cap_dt	optionally, provide user generated by2 to cap_value_ mapping
do_not_cap	if TRUE, normalized values are not capped to 1. Default is FALSE.
force_append	if TRUE, any previous cap_value or norm_value is overridden. Default is FALSE.

Value

data.table, full_dt with cap_value_ and norm_value_ values appended.

Examples

```
append_ynorm(CTCF_in_10a_profiles_dt)
append_ynorm(CTCF_in_10a_profiles_dt,
  aggFUN1 = mean, aggFUN2 = function(x)quantile(x, .5))
```

applySpline	<i>applies a spline smoothing to a tidy data.table containing x and y values.</i>
-------------	---

Description

applySpline Is intended for two-dimensional tidy data.tables, as returned by ssvFetchBigwig

Usage

```
applySpline(dt, n, x_ = "x", y_ = "y", by_ = "", splineFun = stats::spline)
```

Arguments

dt	a tidy data.table containing two-dimensional data
n	the number of interpolation points to use per input point, see ?spline. n must be > 1.
x_	the variable name of the x-values
y_	the variable name of the y-values
by_	optionally, any variables that provide grouping to the data. default is none. see details.
splineFun	a function that accepts x, y, and n as arguments and returns a list of length 2 with named elements x and y. stats::spline by default. see stats::spline for details.

Details

by_ is quite powerful. If by_ = c('gene_id', 'sample_id'), splines will be calculated individually for each gene in each sample. alternatively if by_ = c('gene_id')

Value

a newly derived data.table that is n times longer than original.

See Also

[ssvFetchBigwig](#)

Examples

```
#data may be blockier than we'd like
ggplot(CTCF_in_10a_profiles_dt[, list(y = mean(y)), by = list(sample, x)] +
  geom_line(aes(x = x, y = y, color = sample)))

#can be smoothed by applying a spline (think twice about doing so,
#it may look prettier but may also be deceptive or misleading)

splined_smooth = applySpline(CTCF_in_10a_profiles_dt, n = 10,
  y_ = 'y', by_ = c('id', 'sample'))
ggplot(splined_smooth[, list(y = mean(y)), by = list(sample, x)] +
  geom_line(aes(x = x, y = y, color = sample)))
```

assemble_heatmap_cluster_bars

assemble_heatmap_cluster_bars

Description

assemble_heatmap_cluster_bars

Usage

```
assemble_heatmap_cluster_bars(plots, ...)
```

Arguments

plots	list of plots as returned from ssvSignalHeatmap.ClusterBars when return_unassembled_plots = TRUE
...	arguments passed to cowplot::plot_grid

Value

A grob produced by cowplot::plot_grid

Examples

```
plots = ssvSignalHeatmap.ClusterBars(CTCF_in_10a_profiles_gr, return_unassembled_plots = TRUE)
assemble_heatmap_cluster_bars(plots)
```

Bcell_peaks	<i>4 random peaks for paired-end data</i>
-------------	---

Description

matches `system.file("extdata/Bcell_PE.mm10.bam", package = "seqsetvis")`

Format

GRanges length 4

Details

this is included only for testing `ssvFetchBamPE` functions.

calc_norm_factors	<i>calc_norm_factors</i>
-------------------	--------------------------

Description

Calculate normalization factors in a two step process:

Usage

```
calc_norm_factors(
  full_dt,
  value_ = "y",
  cap_value_ = "y_cap_value",
  by1 = "id",
  by2 = "sample",
  aggFUN1 = max,
  aggFUN2 = function(x) quantile(x, 0.95)
)
```

Arguments

full_dt	a data.table, as returned by <code>ssvFetch*(..., return_data.table. = TRUE)</code>
value_	character, attribute in full_dt to normalzie.
cap_value_	character, new attribute name specifying values to cap to.
by1	character vector, specifies attributes relevant to step 1.
by2	character vector, specifies attributes relevant to step 1 and 2.
aggFUN1	function called on value_ with by = c(by1, by2) in step 1.
aggFUN2	function called on result of aggFUN1 with by = by2 in step 2.

Details

- 1) summarize every region for each sample (default summary function is max)
- 2) calculate a value to cap each sample to based on regions (default is 95th quantile).

The underlying assumption here is that meaningful enrichment is present at the majority of regions provided. If prevalence varies by a specific factor, say ChIP-seq targets with different characteristics - ie. when analyzing TSSes for H3K4me3 and an infrequent transcription factor it is more appropriate to specify appropriate quantile cutoffs per factor.

Value

data.table mapping by2 to cap_value_.

Examples

```
calc_norm_factors(CTCF_in_10a_profiles_dt)
calc_norm_factors(CTCF_in_10a_profiles_dt,
  aggFUN1 = mean, aggFUN2 = function(x)quantile(x, .5))
```

centerAtMax	<i>centers profile of x and y. default is to center by region but across all samples.</i>
-------------	---

Description

centerAtMax locates the coordinate x of the maximum in y and shifts x such that it is zero at max y.

Usage

```
centerAtMax(
  dt,
  x_ = "x",
  y_ = "y",
  by_ = "id",
  view_size = NULL,
  trim_to_valid = TRUE,
  check_by_dupes = TRUE,
  x_precision = 3,
  replace_x = TRUE
)
```

Arguments

dt	data.table
x_	the variable name of the x-values. default is 'x'
y_	the variable name of the y-values default is 'y'

by_	optionally, any variables that provide grouping to the data. default is none. see details.
view_size	the size in x_ to consider for finding the max of y_. if length(view_size) == 1, range will be c(-view_size, view_size). if length(view_size) > 1, range will be range(view_size). default value of NULL uses complete range of x.
trim_to_valid	valid x_ values are those with a set y_ value in all by_ combinations
check_by_dupes	default assumption is that there should be on set of x_ for a by_ instance. if this is not the case and you want to disable warnings about set this to FALSE.
x_precision	numerical precision of x, default is 3.
replace_x	logical, default TRUE. if TRUE x_ will be replaced with position relative to summit. if FALSE x_ will be preserved and x_summitPosition added.

Details

character. by_ controls at the level of the data centering is applied. If by_ is "" or NULL, a single max position will be determined for the entire dataset. If by_ is "id" (the default) then each region will be centered individually across all samples.

Value

data.table with x (or xnew if replace_x is FALSE) shifted such that x = 0 matches the maximum y-value define by by_ grouping

Examples

```
centerAtMax(CTCF_in_10a_profiles_gr, y_ = 'y', by_ = 'id',
  check_by_dupes = FALSE)
#it's a bit clearer what's happening with trimming disabled
#but results are less useful for heatmaps etc.
centerAtMax(CTCF_in_10a_profiles_gr, y_ = 'y', by_ = 'id',
  check_by_dupes = FALSE, trim_to_valid = FALSE)
#specify view_size to limit range of x values considered, prevents
#excessive data trimming.
centerAtMax(CTCF_in_10a_profiles_gr, y_ = 'y', view_size = 100, by_ = 'id',
  check_by_dupes = FALSE)
```

centerFixedSizeGRanges

Transforms set of GRanges to all have the same size.

Description

centerFixedSizeGRanges First calculates the central coordinate of each GRange in grs and extends in both direction by half of fixed_size

Usage

```
centerFixedSizeGRanges(grs, fixed_size = 2000)
```

Arguments

grs	Set of GRanges with inconsistent and/or incorrect size
fixed_size	The final width of each GRange returned.

Value

Set of GRanges after resizing all input GRanges, either shortened or lengthened as required to match fixed_size

Examples

```
library(GenomicRanges)
grs = GRanges("chr1", IRanges(1:10+100, 1:10*3+100))
centered_grs = centerFixedSizeGRanges(grs, 10)
width(centered_grs)
```

centerGRangesAtMax	<i>Centers query GRanges at maximum signal in prof_dt.</i>
--------------------	--

Description

Centers query GRanges at maximum signal in prof_dt.

Usage

```
centerGRangesAtMax(prof_dt, qgr, x_ = "x", y_ = "y", by_ = "id", width = 1)
```

Arguments

prof_dt	a GRanges or data.table as returned by ssvFetch*.
qgr	the GRanges used to query ssvFetch* as the qgr argument.
x_	positional variable. Should almost always be the default, "x".
y_	the signal value variable. Likely the default value of "y" but could be "y_norm" if append_ynorm was applied to data.
by_	region identifier variable. Should almost always be the default, "id".
width	Desired width of final regions. Default is 1.

Value

a GRanges with same mcols as qgr that has been centered based on signal in prof_dt and with regions of specified width.

Examples

```
centerGRangesAtMax(CTCF_in_10a_profiles_dt, CTCF_in_10a_overlaps_gr)
centerGRangesAtMax(CTCF_in_10a_profiles_gr, CTCF_in_10a_overlaps_gr)
```

chromHMM_demo_bw_states_gr

MCF10A CTCF profiles at 20 windows per chromHMM state, hg38.

Description

MCF10A CTCF profiles at 20 windows per chromHMM state, hg38.

Format

a GRanges object of length 4000 with 5 metadata columns sufficient for use with ggplot2

Details

part of [chromHMM_demo_data](#)

the result of `ssvFetchBigwig()` on the MCF10A_CTCF_FE.bw near 20 randomly selected windows per chromHMM state.

chromHMM_demo_chain_url

URL to download hg19ToHg38 liftover chain from UCSC

Description

URL to download hg19ToHg38 liftover chain from UCSC

Format

a character containing a URL

Details

file is gzipped .txt

part of [chromHMM_demo_data](#)

chromHMM_demo_data	<i>chromHMM state segmentation in the MCF7 cell line</i>
--------------------	--

Description

Vignette data for seqsetvis was downloaded directly from GEO series [GSE57498](#). This data is the state segmentation by chromHMM in the MCF7 cell line. chromHMM creates a hidden markov model by integrating several ChIP-seq samples, in this case:

- MCF7_H3K27ac_ChIP-Seq
- MCF7_H3K27me3_ChIP-Seq
- MCF7_H3K4me1_ChIP-Seq
- MCF7_H3K4me3_ChIP-Seq
- MCF7_RNApolIIp_ChIP-Seq

Data from GEO series [GSE57498](#) is from the publication [Taberlay PC et al. 2014](#)

Details

Contains:

- [chromHMM_demo_overlaps_gr](#)
- [chromHMM_demo_bw_states_gr](#)
- [chromHMM_demo_state_total_widths](#)
- [chromHMM_demo_state_colors](#)
- [chromHMM_demo_segmentation_url](#)
- [chromHMM_demo_chain_url](#)

chromHMM_demo_overlaps_gr	<i>overlap of MCF10A CTCF with MCF7 chromHMM states, hg38.</i>
---------------------------	--

Description

overlap of MCF10A CTCF with MCF7 chromHMM states, hg38.

Format

a GRanges object of length 98 with 10 logical metadata columns, 1 per state.

Details

part of [chromHMM_demo_data](#)

the result of `ssvOverlapIntervalSets()` on MCF10A CTCF peaks and MCF7 chromHMM states with `use_first = TRUE`

first (the MCF10A peaks) and `no_hit` columns have been removed each remaining column represents MCF10A peaks overlapping with a state.

`chromHMM_demo_segmentation_url`

URL to download hg19 MCF7 chromHMM segmentation

Description

URL to download hg19 MCF7 chromHMM segmentation

Format

a character containing a URL

Details

file is gzipped bed with name, score, itemRgb and thick meta columns

part of [chromHMM_demo_data](#)

`chromHMM_demo_state_colors`

original state name to color mappings stored in segmentation bed

Description

original state name to color mappings stored in segmentation bed

Format

a named character vector mapping states to hex colors

Details

part of [chromHMM_demo_data](#)

chromHMM_demo_state_total_widths
<i>state name to total width mappings, hg38</i>

Description

state name to total width mappings, hg38

Format

named numeric of total widths per state

Details

part of [chromHMM_demo_data](#)

clusteringKmeans	<i>perform kmeans clustering on matrix rows and return reordered matrix along with order matched cluster assignments. clusters are sorted using hclust on centers</i>
------------------	---

Description

perform kmeans clustering on matrix rows and return reordered matrix along with order matched cluster assignments. clusters are sorted using hclust on centers

Usage

clusteringKmeans(mat, nclust, centroids = NULL)

Arguments

- | | |
|-----------|--|
| mat | numeric matrix to cluster. |
| nclust | the number of clusters. |
| centroids | optional matrix with same columns as mat and one centroid per row to base clusters off of. Overrides any setting to nclust. Default of NULL results in randomly initialized k-means. |

Value

data.table with group variable indicating cluster membership and id variable that is a factor indicating order based on within cluster similarity

Examples

```
dt = data.table::copy(CTCF_in_10a_profiles_dt)
mat = data.table::dcast(dt, id ~ sample + x, value.var = "y" )
rn = mat$id
mat = as.matrix(mat[,-1])
rownames(mat) = rn
clust_dt = clusteringKmeans(mat, nclust = 3)
dt = merge(dt, clust_dt)
dt$id = factor(dt$id, levels = clust_dt$id)
dt[order(id)]
```

clusteringKmeansNestedHclust

perform kmeans clustering on matrix rows and return reordered matrix along with order matched cluster assignments clusters are sorted using hclust on centers the contents of each cluster are sorted using hclust

Description

perform kmeans clustering on matrix rows and return reordered matrix along with order matched cluster assignments clusters are sorted using hclust on centers the contents of each cluster are sorted using hclust

Usage

```
clusteringKmeansNestedHclust(
  mat,
  nclust,
  within_order_strategy = c("hclust", "sort")[2],
  centroids = NULL,
  manual_mapping = NULL
)
```

Arguments

mat	A wide format matrix
nclust	the number of clusters
within_order_strategy	one of "hclust" or "sort". if hclust, hierarchical clustering will be used. if sort, a simple decreasing sort of rowSums.
centroids	optional matrix with same columns as mat and one centroid per row to base clusters off of. Overrides any setting to nclust. Default of NULL results in randomly initialized k-means.
manual_mapping	optional named vector manually specifying cluster assignments. names should be item ids and values should be cluster names the items are assigned to. Default of NULL allows clustering to proceed.

Value

data.table with 2 columns of cluster info. id column corresponds with input matrix rownames and is sorted within each cluster using hierarchical clustering group column indicates cluster assignment

Examples

```
dt = data.table::copy(CTCF_in_10a_profiles_dt)
mat = data.table::dcast(dt, id ~ sample + x, value.var = "y" )
rn = mat$id
mat = as.matrix(mat[,-1])
rownames(mat) = rn
clust_dt = clusteringKmeansNestedHclust(mat, nclust = 3)
dt = merge(dt, clust_dt)
dt$id = factor(dt$id, levels = clust_dt$id)
dt[order(id)]
```

col2hex	<i>converts a valid r color name ("black", "red", "white", etc.) to a hex value</i>
---------	---

Description

converts a valid r color name ("black", "red", "white", etc.) to a hex value

Usage

```
col2hex(color_name)
```

Arguments

color_name character. one or more r color names.

Value

hex value of colors coded by colors()

Examples

```
col2hex(c("red", "green", "blue"))
col2hex(c("lightgray", "gray", "darkgray"))
```

<code>collapse_gr</code>	<i>collapse_gr</i>
--------------------------	--------------------

Description

collapse non-contiguous regions (i.e. exons) into a contiguous coordinate starting at 1. this is strand sensitive and intended for use with all exons of a single gene.

Usage

```
collapse_gr(genome_gr)
```

Arguments

`genome_gr` a GRanges of regions on a single chromosome. Regions are intended to be non-contiguous and may even overlap.

Value

a new GRanges object with same mcols as input with all intervals starting at 1 and no empty space between syntenic regions.

Examples

```
library(data.table)
library(GenomicRanges)
dev_dat = data.table(seqnames = "chrTest",
                     transcript_id = c(1, 1, 2, 2, 3, 3, 3),
                     start = c(5, 30, 8, 30, 2, 30, 40),
                     end = c(10, 35, 15, 38, 7, 35, 45),
                     strand = "+")

genome_gr = GRanges(dev_dat)
collapse_gr(genome_gr)

neg_gr = genome_gr
strand(neg_gr) = "-"
collapse_gr(neg_gr)
```

<code>convert_collapsed_coord</code>	<i>convert_collapsed_coord</i>
--------------------------------------	--------------------------------

Description

(preliminary implementation, sub-optimal)

Usage

```
convert_collapsed_coord(genome_gr, x)
```

Arguments

genome_gr	non-contiguous regions to collapse a la collapse_gr
x	numeric, positions within genome_gr to convert to collapsed coordinates.

Details

see [collapse_gr](#) for explanation of intended uses. this function translates all values of x from original genomic coordinates to new coordinate space created by [collapse_gr](#).

Value

numeric, positions of every value of x within collapse coordinates. values outside of collapsed regions (an intron or outside range) will be NA.

Examples

```
library(data.table)
library(GenomicRanges)
dev_dat = data.table(seqnames = "chrTest",
                     transcript_id = c(1, 1, 2, 2, 3, 3, 3),
                     start = c(5, 30, 8, 30, 2, 30, 40),
                     end = c(10, 35, 15, 38, 7, 35, 45),
                     strand = "+")

genome_gr = GRanges(dev_dat)
convert_collapsed_coord(genome_gr, start(genome_gr))
convert_collapsed_coord(genome_gr, end(genome_gr))
```

crossCorrByRle	<i>Calculate cross correlation by using shiftApply on read coverage Rle</i>
----------------	---

Description

Calculate cross correlation by using shiftApply on read coverage Rle

Usage

```
crossCorrByRle(
  bam_file,
  query_gr,
  max_dupes = 1,
  fragment_sizes = 50:300,
  read_length = NULL,
  flip_strand = FALSE,
  ...
)
```

Arguments

<code>bam_file</code>	character. Path to .bam file, must have index at .bam.bai.
<code>query_gr</code>	GRanges. Regions to calculate cross correlation for.
<code>max_dupes</code>	integer. Duplicate reads above this value will be removed.
<code>fragment_sizes</code>	integer. fragment size range to search for maximum correlation.
<code>read_length</code>	integer. Any values outside <code>fragment_range</code> that must be searched. If not supplied will be determined from <code>bam_file</code> . Set as NA to disable this behavior.
<code>flip_strand</code>	boolean. if TRUE strands that reads align to are swapped. This is typically only necessary if there was a mismatch between library chemistry and aligner settings. Default is FALSE.
<code>...</code>	arguments passed to <code>ScanBamParam</code>

Value

named list of results

Examples

```
bam_f = system.file("extdata/test.bam",
  package = "seqsetvis", mustWork = TRUE)
query_gr = CTCF_in_10a_overlaps_gr[1:2]
crossCorrByRle(bam_f, query_gr[1:2], fragment_sizes = seq(50, 300, 50))
```

CTCF_in_10a_bigWig_urls

FTP URL path for vignette data.

Description

FE bigWig tracks for CTCF ChIP-seq in a MCF10A progression model. See GEO series GSE98551 for details.

Format

named character vector of length 3

Details

part of [CTCF_in_10a_data](#)

CTCF_in_10a_data	<i>CTCF ChIP-seq in breast cancer cell lines</i>
------------------	--

Description

Vignette data for seqsetvis was downloaded directly from GEO series [GSE98551](#). This data is CTCF ChIP-seq from a model of breast cancer progression derived from the MCF10A cell line.

Data from GEO series [GSE98551](#) is from the publication [Fritz AJ et al. 2018](#)

Details

- Contains:
- [CTCF_in_10a_overlaps_gr](#)
 - [CTCF_in_10a_profiles_dt](#)
 - [CTCF_in_10a_bigWig_urls](#)
 - [CTCF_in_10a_narrowPeak_urls](#)

CTCF_in_10a_narrowPeak_grs	<i>list of GRanges that results in 100 random subset when overlapped</i>
----------------------------	--

Description

list of GRanges that results in 100 random subset when overlapped

Format

named character vector of length 3

Details

part of [CTCF_in_10a_data](#)

CTCF_in_10a_narrowPeak_urls
<i>FTP URL path for vignette data. from</i>

Description

macs2 peak calls for CTCF ChIP-seq in a MCF10A progression model. See GEO series GSE98551 for details.

Format

named character vector of length 3

Details

part of [CTCF_in_10a_data](#)

CTCF_in_10a_overlaps_gr
<i>100 randomly selected regions from overlapping CTCF peaks in 10a cell ChIP-seq</i>

Description

MACS2 narrowPeak calls on pooled biological replicates at pval 1e-5 and then 0.05 IDR filtered. IDR cutoffs determined by comparing top 150,000 pvalue sorted peak in replicates.

Format

GenomicRanges with 3 metadata columns of membership table

Details

See GEO series GSE98551 for details.

part of [CTCF_in_10a_data](#)

CTCF_in_10a_profiles_dt

Profiles for 100 randomly selected regions from overlapping CTCF peaks in 10a cell ChIP-seq Results from fetching bigwigs with CTCF_in_10a_overlaps_gr.

Description

A tidy data.table at window size 50 bp within 350 bp of peak center The variables are as follows:

Format

A tidy data.table of 2100 rows and 9 columns

Details

part of [CTCF_in_10a_data](#)

1. seqnames. chromosome for GRanges compatibility
2. start. start of interval
3. end. end of interval
4. width. width of interval
5. strand. leftover from GRanges.
6. id. unique identifier
7. y. fold-enrichment over input.
8. x. bp relative to center
9. sample. name of originating sample

CTCF_in_10a_profiles_gr

Profiles for 100 randomly selected regions from overlapping CTCF peaks in 10a cell ChIP-seq Results from CTCF_in_10a_overlaps_gr

Description

A tidy GRanges at window size 50 bp within 350 bp of peak center The variables are as follows:

Format

A tidy GRanges of 2100 rows and 4 metadata columns

Details

part of [CTCF_in_10a_data](#)

1. id. unique identifier
2. y. fold-enrichment over input.
3. x. bp relative to center
4. sample. name of originating sample

easyLoad_bed	<i>easyLoad_bed takes a character vector of file paths to bed plus files and returning named list of GRanges. Mainly a utility function for loading MACS2 narrowPeak and broadPeak.</i>
--------------	---

Description

easyLoad_bed takes a character vector of file paths to bed plus files and returning named list of GRanges. Mainly a utility function for loading MACS2 narrowPeak and broadPeak.

Usage

```
easyLoad_bed(
  file_paths,
  file_names = NULL,
  extraCols = character(),
  n_cores = getOption("mc.cores", 1)
)
```

Arguments

file_paths	character vector of paths to narrowPeak files. If named, those names will be used in output unless overridden by providing file_names.
file_names	character vector of names for output list. If not NULL will override any existing names for file_paths. Default is NULL.
extraCols	named character vector of classes. passed to rtracklayer::import for format = "BED". default is character().
n_cores	number of cores to use, uses mc.cores option if set or 1.

Value

a named list of GRanges loaded from file_paths

Examples

```
bed_f = system.file("extdata/test_loading.bed",
  package = "seqsetvis", mustWork = TRUE)
easyLoad_bed(bed_f, "my_bed")
```

easyLoad_broadPeak	<i>easyLoad_broadPeak takes a character vector of file paths to narrowPeak files from MACS2 and returns a named list of GRanges.</i>
--------------------	--

Description

easyLoad_broadPeak takes a character vector of file paths to narrowPeak files from MACS2 and returns a named list of GRanges.

Usage

```
easyLoad_broadPeak(
  file_paths,
  file_names = NULL,
  n_cores = getOption("mc.cores", 1)
)
```

Arguments

file_paths	character vector of paths to narrowPeak files. If named, those names will be used in output unless overridden by providing file_names.
file_names	character vector of names for output list. If not NULL will override any existing names for file_paths. Default is NULL.
n_cores	number of cores to use, uses mc.cores option if set or 1.

Value

a named list of GRanges loaded from file_paths

Examples

```
bp_f = system.file("extdata/test_loading.broadPeak",
  package = "seqsetvis", mustWork = TRUE)
easyLoad_broadPeak(bp_f, "my_broadPeak")
```

easyLoad_narrowPeak	<i>easyLoad_narrowPeak takes a character vector of file paths to narrowPeak files from MACS2 and returns a named list of GRanges.</i>
---------------------	---

Description

easyLoad_narrowPeak takes a character vector of file paths to narrowPeak files from MACS2 and returns a named list of GRanges.

Usage

```
easyLoad_narrowPeak(
  file_paths,
  file_names = NULL,
  n_cores = getOption("mc.cores", 1)
)
```

Arguments

<code>file_paths</code>	character vector of paths to narrowPeak files. If named, those names will be used in output unless overridden by providing <code>file_names</code> .
<code>file_names</code>	character vector of names for output list. If not NULL will override any existing names for <code>file_paths</code> . Default is NULL.
<code>n_cores</code>	number of cores to use, uses <code>mc.cores</code> option if set or 1.

Value

a named list of GRanges loaded from `file_paths`

Examples

```
np_f = system.file("extdata/test_loading.narrowPeak",
  package = "seqsetvis", mustWork = TRUE)
easyLoad_narrowPeak(np_f, "my_narrowPeak")
```

<code>easyLoad_seacr</code>	<i>easyLoad_seacr takes a character vector of file paths to seacr output bed files and returns a named list of GRanges.</i>
-----------------------------	---

Description

`easyLoad_seacr` takes a character vector of file paths to seacr output bed files and returns a named list of GRanges.

Usage

```
easyLoad_seacr(
  file_paths,
  file_names = NULL,
  n_cores = getOption("mc.cores", 1)
)
```

Arguments

file_paths	character vector of paths to seacr bed files. If named, those names will be used in output unless overridden by providing file_names.
file_names	character vector of names for output list. If not NULL will override any existing names for file_paths. Default is NULL.
n_cores	number of cores to use, uses mc.cores option if set or 1.

Value

a named list of GRanges loaded from file_paths

Examples

```
bed_f = system.file("extdata/test_loading.seacr.bed",
  package = "seqsetvis", mustWork = TRUE)
easyLoad_seacr(bed_f, "my_seacr")
```

expandCigar	<i>Expand cigar codes to GRanges</i>
-------------	--------------------------------------

Description

see [sam specs](#) for cigar details

Usage

```
expandCigar(
  cigar_dt,
  op_2count = c("M", "D", "=", "X"),
  return_data.table = FALSE
)
```

Arguments

cigar_dt	data.table with 5 required named columns in any order. c("which_label", "seq-names", "strand", "start", "cigar")
op_2count	Cigar codes to count. Default is alignment (M), deletion (D), match (=), and mismatch (X). Other useful codes may be skipped regions for RNA splicing (N). The locations of any insterions (I) or clipping/padding (S, H, or P) will be a single bp immediately before the interval.
return_data.table	if TRUE, a data.table is returned, else a GRanges. Default is FALSE.

Value

data.table with cigar entries expanded

Examples

```
qgr = CTCF_in_10a_overlaps_gr[1:5]
bam_file = system.file("extdata/test.bam", package = "seqsetvis", mustWork = TRUE)
raw_dt = ssvFetchBam(bam_file, qgr, return_unprocessed = TRUE)
expandCigar(raw_dt)
```

fetchBam	<i>fetch a bam file pileup with the ability to consider read extension to fragment size (fragLen)</i>
----------	---

Description

fetch a bam file pileup with the ability to consider read extension to fragment size (fragLen)

Usage

```
fetchBam(
  bam_f,
  qgr,
  fragLen = NULL,
  target_strand = c("*", "+", "-")[1],
  max_dupes = Inf,
  splice_strategy = c("none", "ignore", "add", "only", "splice_count")[1],
  flip_strand = FALSE,
  return_unprocessed = FALSE,
  ...
)
```

Arguments

bam_f	character or BamFile to load
qgr	GRanges regions to fetchs
fragLen	numeric, NULL, or NA. if numeric, supplied value is used. if NULL, value is calculated with fragLen_calcStranded (default) if NA, raw bam pileup with no cross strand shift is returned.
target_strand	character. if one of "+" or "-", reads are filtered to match. ignored if any other value.
max_dupes	numeric >= 1. duplicate reads by stranddd start position over this number are removed, Default is Inf.
splice_strategy	character, one of c("none", "ignore", "add", "only"). Default is "none" and split read alignments are assumed not present. fragLen must be NA for any other value to be valid. "ignore" will not count spliced regions. "add" counts spliced regions along with others, "only" will only count spliced regions and ignore others.

flip_strand if TRUE, strand alignment is flipped prior to fragLen extension. Default is FALSE.
 return_unprocessed boolean. if TRUE returns read alignment in data.table. Default is FALSE.
 ... passed to ScanBamParam(), can't be which or what.

Value

GRanges containing tag pileup values in score meta column. tags are optionally extended to fragment length (fragLen) prior to pile up.

fragLen_calcStranded	<i>calculate fragLen from a bam file for specified regions</i>
----------------------	--

Description

calculate fragLen from a bam file for specified regions

Usage

```
fragLen_calcStranded(
  bam_f,
  qgr,
  n_regions = 100,
  include_plot_in_output = FALSE,
  test_fragLen = seq(100, 400, 5),
  flip_strand = FALSE,
  ...
)
```

Arguments

bam_f character or BamFile. bam file to read from. .bai index file must be in same directory
 qgr GRanges. used as which for ScanBamParam. Can be NULL if it's REALLY important to load the entire bam, force_no_which = TRUE also required.
 n_regions numeric (integer) it's generally overkill to pull all regions at this stage and will slow calculation down. Default is 100.
 include_plot_in_output if TRUE ouptut is a list of fragLen and a ggplot showing values considered by calculation. Default is FALSE.
 test_fragLen numeric. The set of fragment lenghts to gather strand cross correlation for.
 flip_strand boolean. if TRUE strands that reads align to are swapped. This is typically only necessary if there was a mismatch between library chemistry and aligner settings. Default is FALSE.
 ... passed to Rsamtools::ScanBamParam, can't be which or what.

Value

numeric fragment length

Examples

```
bam_file = system.file("extdata/test.bam",
  package = "seqsetvis")
qgr = CTCF_in_10a_overlaps_gr[1:5]
fragLen_calcStranded(bam_file, qgr)
#if plot is included, a list is returned, item 2 is the plot
fragLen_calcStranded(bam_file, qgr,
  include_plot_in_output = TRUE)[[2]]
```

fragLen_fromMacs2Xls *parse fragLen from MACS2 output*

Description

parse fragLen from MACS2 output

Usage

```
fragLen_fromMacs2Xls(macs2xls_file)
```

Arguments

macs2xls_file character. an xls file output by MACS2 to parse frag length from

Value

numeric fragment length

Examples

```
xls_file = system.file("extdata/test_peaks.xls",
  package = "seqsetvis")
fragLen_fromMacs2Xls(xls_file)
```

getReadLength	<i>determine the most common read length for input bam_file. uses 50 randomly selected regions from query_gr. If fewer than 20 reads are present, loads all of query_gr.</i>
---------------	--

Description

determine the most common read length for input bam_file. uses 50 randomly selected regions from query_gr. If fewer than 20 reads are present, loads all of query_gr.

Usage

```
getReadLength(bam_file, query_gr)
```

Arguments

bam_file	indexed bam file
query_gr	GRanges to read from bam file

Value

numeric of most common read length.

Examples

```
qgr = CTCF_in_10a_overlaps_gr[1:5]
bam_file = system.file("extdata/test.bam", package = "seqsetvis", mustWork = TRUE)
getReadLength(bam_file, qgr)
```

ggellipse	<i>returns a ggplot with ellipses drawn using specified parameters used by ssvFeatureVenn and ssvFeatureEuler</i>
-----------	---

Description

uses eulerr's non-exported ellipse drawing coordinate function

Usage

```
ggellipse(
  xcentres,
  ycentres,
  r,
  r2 = r,
  phi = rep(0, length(xcentres)),
  circle_colors = NULL,
```

```

    group_names = LETTERS[seq_along(xcentres)],
    line_alpha = 1,
    fill_alpha = 0.3,
    line_width = 2,
    n_points = 200
  )

```

Arguments

<code>xcentres</code>	numeric x-coord of centers of ellipses
<code>ycentres</code>	numeric y-coord of centers of ellipses, must have same length as <code>xcentres</code>
<code>r</code>	numeric radius1 of ellipse, must have length of 1 or match length of <code>xcentres</code>
<code>r2</code>	numeric radius2 of ellipse, must have length of 1 or match length of <code>xcentres</code> . same as <code>r</code> by default.
<code>phi</code>	numeric phi of ellipse, must have length of 1 or match length of <code>xcentres</code> . 0 by default.
<code>circle_colors</code>	character of <code>rcolors</code> or hex colors or NULL. if null <code>safeBrew</code> of <code>Dark2</code> is used
<code>group_names</code>	character/factor names of color/fill groups. capital letters by default.
<code>line_alpha</code>	numeric [0,1] alpha of lines, 1 by default
<code>fill_alpha</code>	numeric [0,1] alpha of fill, .3 by default.
<code>line_width</code>	numeric > 0. passed to <code>size</code> . 2 by default
<code>n_points</code>	integer > 1. number of points to approximate circle with. 200 by default

Value

a ggplot containing ellipses

Examples

```

ggellipse(xcentres = c(1, 1, 2),
  ycentres = c(2, 1, 1),
  r = c(1, 2, 1))
ggellipse(xcentres = c(1, 1, 2),
  ycentres = c(2, 1, 1),
  r = c(1, 2, 1),
  fill_alpha = 0,
  group_names = paste("set", 1:3))
ggellipse(xcentres = c(1, 1, 2),
  ycentres = c(2, 1, 1),
  r = c(1, 2, 1),
  circle_colors = c("red", "orange", "yellow"),
  line_alpha = 0,
  group_names = paste("set", 1:3))

```

harmonize_seqlengths	<i>harmonize_seqlengths</i>
----------------------	-----------------------------

Description

ensures compatibility between seqlength of gr and bam_file based on header

Usage

```
harmonize_seqlengths(gr, bam_file)
```

Arguments

gr	GRanges, object to harmonize seqlengths for
bam_file	character, a path to a valid bam file

Value

gr with seqlengths matching bam_file

Examples

```
library(GenomicRanges)
gr = GRanges("chr1", IRanges(1, 100))
#seqlengths has not been set
seqlengths(gr)
bam = system.file("extdata/test.bam", package = "seqsetvis")
gr2 = harmonize_seqlengths(gr, bam)
#seqlengths now set
seqlengths(gr2)
```

make_clustering_matrix	<i>make_clustering_matrix</i>
------------------------	-------------------------------

Description

Create a wide matrix from a tidy data.table more suitable for clustering methods

Usage

```
make_clustering_matrix(
  tidy_dt,
  row_ = "id",
  column_ = "x",
  fill_ = "y",
  facet_ = "sample",
  max_rows = 500,
  max_cols = 100,
  clustering_col_min = -Inf,
  clustering_col_max = Inf,
  dcast_fill = NA
)
```

Arguments

<code>tidy_dt</code>	the tidy data.table to covert to a wide matrix. Must have entries for variables specified by <code>row_</code> , <code>column_</code> , <code>fill_</code> , and <code>facet_</code> .
<code>row_</code>	variable name mapped to row, likely peak id or gene name for ngs data
<code>column_</code>	variable mapped to column, likely bp position for ngs data
<code>fill_</code>	numeric variable to map to fill
<code>facet_</code>	variable name to facet horizontally by
<code>max_rows</code>	for speed rows are sampled to 500 by default, use <code>Inf</code> to plot full data
<code>max_cols</code>	for speed columns are sampled to 100 by default, use <code>Inf</code> to plot full data
<code>clustering_col_min</code>	numeric minimum for col range considered when clustering, default in <code>-Inf</code>
<code>clustering_col_max</code>	numeric maximum for col range considered when clustering, default in <code>Inf</code>
<code>dcast_fill</code>	value to supply to <code>dcast</code> fill argument. default is <code>NA</code> .

Value

A wide matrix version of input tidy data.table

Examples

```
mat = make_clustering_matrix(CTCF_in_10a_profiles_dt)
mat[1:5, 1:5]
```

prepare_fetch_GRanges *prepares GRanges for windowed fetching.*

Description

Deprecated and renamed as prepare_fetch_GRanges_width

Usage

```
prepare_fetch_GRanges(
  qgr,
  win_size,
  min_quantile = 0.75,
  target_size = NULL,
  skip_centerFix = FALSE
)
```

Arguments

qgr	GRanges to prepare
win_size	numeric window size for fetch
min_quantile	numeric [0,1], lowest possible quantile value. Only relevant if target_size is not specified.
target_size	numeric final width of qgr if known. Default of NULL leads to quantile based determination of target_size.
skip_centerFix	boolean, if FALSE (default) all regions will be resized GenomicRanges::resize(x, w, fix = "center") to a uniform size based on min_quantile to a width divisible by win_size.

Details

output GRanges parallels input with consistent width evenly divisible by win_size. Has warning if GRanges needed resizing, otherwise no warning and input GRanges is returned unchanged.

Value

GRanges, either identical to qgr or with suitable consistent width applied.

Examples

```
#use prepare_fetch_GRanges_width instead:
qgr = prepare_fetch_GRanges_width(CTCF_in_10a_overlaps_gr, win_size = 50)
#no warning if qgr is already valid for windowed fetching
prepare_fetch_GRanges_width(qgr, win_size = 50)
```

```
prepare_fetch_GRanges_names
```

Creates a named version of input GRanges using the same method seqsetvis uses internally to ensure consistency.

Description

If \$id is set, that value is used as name and duplicates are checked for.

Usage

```
prepare_fetch_GRanges_names(qgr, include_id = FALSE)
```

Arguments

qgr	input GRanges object the set/check names on
include_id	if TRUE, \$id is retained. Default is FALSE.

Value

and named GRanges based on input qgr.

Examples

```
qgr = seqsetvis::CTCF_in_10a_overlaps_gr
names(qgr) = NULL
#default is to paste "region_" and iteration along length of qgr
prepare_fetch_GRanges_names(qgr)
#id gets used is already set
qgr$id = paste0("peak_", rev(seq_along(qgr)), "_of_", length(qgr))
prepare_fetch_GRanges_names(qgr)
```

```
prepare_fetch_GRanges_width
```

prepares GRanges for windowed fetching.

Description

output GRanges parallels input with consistent width evenly divisible by win_size. Has warning if GRanges needed resizing, otherwise no warning and input GRanges is returned unchanged.

Usage

```
prepare_fetch_GRanges_width(
  qgr,
  win_size,
  min_quantile = 0.75,
  target_size = NULL,
  skip_centerFix = FALSE
)
```

Arguments

qgr	GRanges to prepare
win_size	numeric window size for fetch
min_quantile	numeric [0,1], lowest possible quantile value. Only relevant if target_size is not specified.
target_size	numeric final width of qgr if known. Default of NULL leads to quantile based determination of target_size.
skip_centerFix	boolean, if FALSE (default) all regions will be resized GenomicRanges::resize(x, w, fix = "center") to a uniform size based on min_quantile to a width divisible by win_size.

Value

GRanges, either identical to qgr or with suitable consistent width applied.

Examples

```
qgr = prepare_fetch_GRanges_width(CTCF_in_10a_overlaps_gr, win_size = 50)
#no warning if qgr is already valid for windowed fetching
prepare_fetch_GRanges_width(qgr, win_size = 50)
```

quantileGRangesWidth *Quantile width determination strategy*

Description

Returns the lowest multiple of win_size greater than min_quantile quantile of width(qgr)

Usage

```
quantileGRangesWidth(qgr, min_quantile = 0.75, win_size = 1)
```

Arguments

qgr	GRanges to calculate quantile width for
min_quantile	numeric [0,1] the minimum quantile of width in qgr
win_size	numeric/integer >=1, returned value will be a multiple of this

Value

numeric that is \geq min_quantile and evenly divisible by win_size

Examples

```
gr = CTCF_in_10a_overlaps_gr
quantileGRangesWidth(gr)
quantileGRangesWidth(gr, min_quantile = .5, win_size = 100)
```

safeBrew

allows RColorBrew to handle n values less than 3 and greater than 8 without warnings and return expected number of colors.

Description

allows RColorBrew to handle n values less than 3 and greater than 8 without warnings and return expected number of colors.

Usage

```
safeBrew(n, pal = "Dark2")
```

Arguments

n	integer value of number of colors to make palette for
pal	palette recognized by RColorBrewer

Value

a character vector of hex coded colors of length n from the color brewer palette pal

Examples

```
plot(1:2, rep(0, 2), col = safeBrew(2, "dark2"), pch = 16, cex = 6)
plot(1:12, rep(0, 12), col = safeBrew(12, "set1"), pch = 16, cex = 6)
plot(1:12, rep(0, 12), col = safeBrew(12, "set2"), pch = 16, cex = 6)
plot(1:12, rep(0, 12), col = safeBrew(12, "set3"), pch = 16, cex = 6)
```

set_list2memb	<i>convert a list of sets, each list item should be a character vector denoting items in sets</i>
---------------	---

Description

convert a list of sets, each list item should be a character vector denoting items in sets

Usage

```
set_list2memb(set_list)
```

Arguments

set_list a list of character vectors. default names will be added if missing

Value

converts list of characters/numeric to membership table matrix

shift_anchor	<i>orients the relative position of x's zero value and extends ranges to be contiguous</i>
--------------	--

Description

orients the relative position of x's zero value and extends ranges to be contiguous

Usage

```
shift_anchor(score_dt, window_size, anchor)
```

Arguments

score_dt data.table, GRanges() sufficient
 window_size numeric, window size used to generate score_dt
 anchor character, one of c("center", "center_unstranded", "left", "left_unstranded")

Value

score_dt with x values shifted appropriately and start and end extended to make ranges contiguous

ssvConsensusIntervalSets

Intersect a list of GRanges to create a single GRanges object of merged ranges including metadata describing overlaps per input GRanges.

Description

In contrast to ssvOverlapIntervalSets, only regions where a consensus of input grs are present are preserved and annotated.

Usage

```
ssvConsensusIntervalSets(grs, ext = 0, min_number = 2, min_fraction = 0.5, ...)
```

Arguments

grs	A list of GRanges
ext	An integer specifying how far to extend ranges before merging. in effect, ranges withing 2*ext of one another will be joined during the merge
min_number	An integer number specifying the absolute minimum of input grs that must overlap for a site to be considered consensus.
min_fraction	A numeric between 0 and 1 specifying the fraction of grs that must overlap to be considered consensus.
...	arguments passed to IRanges::findOverlaps, i.e. maxgap, minoverlap, type, select, invert.

Details

Only the most stringent of min_number or min_fraction will be applied.

Value

GRanges with metadata columns describing consensus overlap of input grs.

Examples

```
library(GenomicRanges)
a = GRanges("chr1", IRanges(1:7*10, 1:7*10))
b = GRanges("chr1", IRanges(5:10*10, 5:10*10))
ssvConsensusIntervalSets(list(a, b))
```

`ssvFactorizeMembTable` *Convert any object accepted by `ssvMakeMembTable` to a factor To avoid ambiguity,*

Description

see [ssvMakeMembTable](#)

Usage

```
ssvFactorizeMembTable(object)
```

Arguments

`object` a valid object for conversion to a membership table and then factor

Value

a 2 column ("id" and "group") data.frame. "id" is factor of item names if any or simply order of items. "group" is a factor of set combinations

Examples

```
ssvFactorizeMembTable(CTCF_in_10a_overlaps_gr)
ssvFactorizeMembTable(list(1:4, 2:3, 4:6))
```

`ssvFeatureBars` *bar plots of set sizes*

Description

bar plots of set sizes

Usage

```
ssvFeatureBars(
  object,
  show_counts = TRUE,
  bar_colors = NULL,
  counts_text_colors = NULL,
  return_data = FALSE
)
```

Arguments

<code>object</code>	passed to <code>ssvMakeMembTable</code> for conversion to membership table
<code>show_counts</code>	logical. should counts be displayed at the center of each bar. default is TRUE
<code>bar_colors</code>	character. rcolor or hex colors. default of NULL uses RColorBrewer Dark2. Will repeat to match number of samples.
<code>counts_text_colors</code>	character. rcolor or hex colors. default of NULL uses RColorBrewer Dark2. Will repeat to match number of samples.
<code>return_data</code>	logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.

Value

ggplot of bar plot of set sizes

Examples

```
ssvFeatureBars(list(1:3, 2:6))
ssvFeatureBars(CTCF_in_10a_overlaps_gr)
ssvFeatureBars(S4Vectors::mcols(CTCF_in_10a_overlaps_gr)[,2:3])
```

`ssvFeatureBinaryHeatmap`

binary heatmap indicating membership. heatmap is sorted by column left to right. change column order to reveal patterns

Description

binary heatmap indicating membership. heatmap is sorted by column left to right. change column order to reveal patterns

Usage

```
ssvFeatureBinaryHeatmap(
  object,
  raster_approximation = FALSE,
  true_color = "black",
  false_color = "#EFEFEF",
  raster_width_min = 1000,
  raster_height_min = 1000,
  return_data = FALSE
)
```

Arguments

object	passed to ssvMakeMembTable
raster_approximation	If TRUE, instead of standard ggplot, write temporary raster png image and re-draw that as plot background. default is FALSE
true_color	character. rcolor or hex color used for TRUE values. default is "black".
false_color	character. rcolor or hex color used for TRUE values. default is "#EFEFEF", a gray.
raster_width_min	raster width will be minimum multiple of number of columns over this number. ignored if raster_approximation is FALSE.
raster_height_min	raster height will be minimum multiple of number of rows over this number ignored if raster_approximation is FALSE
return_data	logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.

Value

ggplot using geom_tile of membership table sorted from left to right.

Examples

```
ssvFeatureBinaryHeatmap(list(1:3, 2:6))
# horizontal version
ssvFeatureBinaryHeatmap(list(1:3, 2:6)) + coord_flip() +
  theme(axis.text.x = element_blank(), axis.text.y = element_text())
ssvFeatureBinaryHeatmap(CTCF_in_10a_overlaps_gr)
ssvFeatureBinaryHeatmap(S4Vectors::mcols(CTCF_in_10a_overlaps_gr)[,2:3])
ssvFeatureBinaryHeatmap(S4Vectors::mcols(CTCF_in_10a_overlaps_gr)[,3:2])
```

ssvFeatureEuler

Try to load a bed-like file and convert it to a GRanges object

Description

Try to load a bed-like file and convert it to a GRanges object

Usage

```
ssvFeatureEuler(
  object,
  line_width = 2,
  shape = c("circle", "ellipse")[1],
  n_points = 200,
  fill_alpha = 0.3,
```

```
    line_alpha = 1,
    circle_colors = NULL,
    return_data = FALSE
  )
```

Arguments

object	A membership table
line_width	numeric, passed to size aesthetic to control line width
shape	shape argument passed to eulerr::euler
n_points	number of points to use for drawing ellipses, passed to eulerr::ellipse
fill_alpha	numeric [0,1], alpha value for circle fill
line_alpha	numeric [0,1], alpha value for circle line
circle_colors	colors to choose from for circles. passed to ggplot2 color scales.
return_data	logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.

Value

ggplot of venneuler results

Examples

```
ssvFeatureEuler(list(1:3, 2:6))
ssvFeatureEuler(CTCF_in_10a_overlaps_gr)
ssvFeatureEuler(S4Vectors::mcols(CTCF_in_10a_overlaps_gr)[,2:3])
```

ssvFeaturePie	<i>pie plot of set sizes</i>
---------------	------------------------------

Description

pie plot of set sizes

Usage

```
ssvFeaturePie(object, slice_colors = NULL, return_data = FALSE)
```

Arguments

object	object that ssvMakeMembTable can convert to logical matrix membership
slice_colors	colors to use for pie slices
return_data	logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.

Value

ggplot pie graph of set sizes

Examples

```
ssvFeaturePie(list(1:3, 2:6))
ssvFeaturePie(CTCF_in_10a_overlaps_gr)
ssvFeaturePie(S4Vectors::mcols(CTCF_in_10a_overlaps_gr)[,2:3])
```

ssvFeatureUpset

ssvFeatureUpset

Description

Uses the UpSetR package to create an upset plot of overlaps.

Usage

```
ssvFeatureUpset(
  object,
  return_UpSetR = FALSE,
  nsets = NULL,
  nintersects = 15,
  order.by = "freq",
  ...
)
```

Arguments

<code>object</code>	will be passed to ssvMakeMembTable for conversion to membership matrix
<code>return_UpSetR</code>	If TRUE, return the UpSetR object, The default is FALSE and results in a gg-plotted version compatible with cowplot etc.
<code>nsets</code>	Number of sets to look at
<code>nintersects</code>	Number of intersections to plot. If set to NA, all intersections will be plotted.
<code>order.by</code>	How the intersections in the matrix should be ordered by. Options include frequency (entered as "freq"), degree, or both in any order.
<code>...</code>	Additional parameters passed to upset in the UpSetR package.

Value

ggplot version of UpSetR plot

Examples

```
ssvFeatureUpset(list(1:3, 2:6))
ssvFeatureUpset(CTCF_in_10a_overlaps_gr)
ssvFeatureUpset(S4Vectors::mcols(CTCF_in_10a_overlaps_gr)[,2:3])
```

ssvFeatureVenn	<i>ggplot implementation of vennDiagram from limma package. currently limited at 3 sets. ssvFeatureUpset and ssvFeatureBinaryHeatmap are good options for more than 3 sets. ssvFeatureEuler can work too but can take a very long time to run for more than 5 or so.</i>
----------------	--

Description

ggplot implementation of vennDiagram from limma package. currently limited at 3 sets. ssvFeatureUpset and ssvFeatureBinaryHeatmap are good options for more than 3 sets. ssvFeatureEuler can work too but can take a very long time to run for more than 5 or so.

Usage

```
ssvFeatureVenn(
  object,
  group_names = NULL,
  counts_txt_size = 5,
  counts_as_labels = FALSE,
  show_outside_count = FALSE,
  line_width = 3,
  circle_colors = NULL,
  fill_alpha = 0.3,
  line_alpha = 1,
  counts_color = NULL,
  n_points = 200,
  return_data = FALSE
)
```

Arguments

object	will be passed to ssvMakeMembTable for conversion to membership matrix
group_names	useful if names weren't provided or were lost in creating membership matrix
counts_txt_size	font size for count numbers
counts_as_labels	if TRUE, geom_label is used instead of geom_text. can be easier to read.
show_outside_count	if TRUE, items outside of all sets are counted outside. can be confusing.
line_width	uses size aesthetic to control line width of circles.
circle_colors	colors to use for circle line colors. Uses Dark2 set from RColorBrewer by default.
fill_alpha	alpha value to use for fill, defaults to .3.
line_alpha	numeric [0,1], alpha value for circle line
counts_color	character. single color to use for displaying counts

`n_points` integer. number of points to approximate circle with. default is 200.

`return_data` logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.

Value

ggplot venn diagram

Examples

```
ssvFeatureVenn(list(1:3, 2:6))
ssvFeatureVenn(CTCF_in_10a_overlaps_gr)
ssvFeatureVenn(S4Vectors::mcols(CTCF_in_10a_overlaps_gr)[,2:3])
```

<code>ssvFetchBam</code>	<i>Iterates a character vector (ideally named) and calls <code>ssvFetchBam.single</code> on each. Appends grouping variable to each resulting <code>data.table</code> and uses <code>rbindlist</code> to efficiently combine results</i>
--------------------------	--

Description

`ssvFetchBam` iteratively calls `fetchWindowedBam.single`. See [ssvFetchBam.single](#) for more info.

Usage

```
ssvFetchBam(
  file_paths,
  qgr,
  unique_names = NULL,
  names_variable = "sample",
  file_attribs = NULL,
  win_size = 50,
  win_method = c("sample", "summary")[1],
  summary_FUN = stats::weighted.mean,
  fragLens = "auto",
  target_strand = c("*", "+", "-", "both")[1],
  flip_strand = FALSE,
  anchor = c("left", "left_unstranded", "center", "center_unstranded")[3],
  return_data.table = FALSE,
  max_dupes = Inf,
  splice_strategy = c("none", "ignore", "add", "only", "splice_count")[1],
  n_cores = getOption("mc.cores", 1),
  n_region_splits = 1,
  return_unprocessed = FALSE,
  force_skip_centerFix = FALSE,
  ...
)
```

Arguments

<code>file_paths</code>	character vector of <code>file_paths</code> to load from. Alternatively, <code>file_paths</code> can be a <code>data.frame</code> or <code>data.table</code> whose first column is a character vector of paths and additional columns will be used as metadata.
<code>qgr</code>	Set of <code>GRanges</code> to query. For valid results the width of each interval should be identical and evenly divisible by <code>win_size</code> .
<code>unique_names</code>	names to use in final <code>data.table</code> to designate source bigwig. Default is 'sample'
<code>names_variable</code>	The column name where <code>unique_names</code> are stored.
<code>file_attribs</code>	optional <code>data.frame</code> / <code>data.table</code> with one row per item in file paths. Each column will be a variable added to final tidy output.
<code>win_size</code>	The window size that evenly divides widths in <code>qgr</code> .
<code>win_method</code>	character. one of <code>c("sample", "summary")</code> . Determines if <code>viewGRangesWinSample_dt</code> or <code>viewGRangesWinSummary_dt</code> is used to represent each region in <code>qgr</code> .
<code>summary_FUN</code>	function. only relevant if <code>win_method</code> is "summary". passed to <code>viewGRangesWinSummary_dt</code> .
<code>fragLens</code>	numeric. The fragment length to use to extend reads. The default value "auto" causes an automatic calculation from 100 regions in <code>qgr</code> . NA causes no extension of reads to fragment size.
<code>target_strand</code>	character. One of <code>c("*", "+", "-")</code> . Controls filtering of reads by strand. Default of "*" combines both strands.
<code>flip_strand</code>	boolean. if TRUE strands are flipped.
<code>anchor</code>	character, one of <code>c("center", "center_unstranded", "left", "left_unstranded")</code>
<code>return_data.table</code>	logical. If TRUE the internal <code>data.table</code> is returned instead of <code>GRanges</code> . Default is FALSE.
<code>max_dupes</code>	numeric ≥ 1 . duplicate reads by strand and start position over this number are removed, Default is Inf.
<code>splice_strategy</code>	character, one of <code>c("none", "ignore", "add", "only", "splice_count")</code> . Default is "none" and spliced alignment are assumed not present. <code>fragLen</code> will be forced to be NA for any other value. "ignore" will not count spliced regions. "add" counts spliced regions along with others, "only" will only count spliced regions and ignore others.
<code>n_cores</code>	integer number of cores to use. Uses <code>mc.cores</code> option if not supplied.
<code>n_region_splits</code>	integer number of splits to apply to <code>qgr</code> . The query <code>GRanges</code> will be split into this many roughly equal parts for increased parallelization. Default is 1, no split.
<code>return_unprocessed</code>	boolean. if TRUE returns read alignment in <code>data.table</code> . Default is FALSE.
<code>force_skip_centerFix</code>	boolean, if TRUE all query ranges will be used "as is". This is already the case by default if <code>win_method == "summary"</code> but may have applications where <code>win_method == "sample"</code> .
<code>...</code>	passed to <code>Rsamtools::ScanBamParam()</code>

Details

if qgr contains the range chr1:1-100 and win_size is 10, values from positions chr1 5,15,25...85, and 95 will be retrieved from bw_file

Value

A tidy formatted GRanges (or data.table if specified) containing fetched values.

Examples

```
if(Sys.info()['sysname'] != "Windows"){
  library(GenomicRanges)
  bam_f = system.file("extdata/test.bam",
    package = "seqsetvis", mustWork = TRUE)
  bam_files = c("a" = bam_f, "b" = bam_f)
  qgr = CTCF_in_10a_overlaps_gr[1:5]
  bw_gr = ssvFetchBam(bam_files, qgr, win_size = 10)
  bw_gr2 = ssvFetchBam(as.list(bam_files), qgr, win_size = 10)

  bw_dt = ssvFetchBam(bam_files, qgr, win_size = 10,
    return_data.table = TRUE)
}
```

ssvFetchBam.single	<i>fetch a windowed version of a bam file, returns GRanges</i>
--------------------	--

Description

fetch a windowed version of a bam file, returns GRanges

Usage

```
ssvFetchBam.single(
  bam_f,
  qgr,
  win_size = 50,
  win_method = c("sample", "summary")[1],
  summary_FUN = stats::weighted.mean,
  fragLen = NULL,
  target_strand = c("*", "+", "-", "both")[1],
  anchor = c("left", "left_unstranded", "center", "center_unstranded")[3],
  return_data.table = FALSE,
  max_dupes = Inf,
  splice_strategy = c("none", "ignore", "add", "only", "splice_count")[1],
  flip_strand = FALSE,
  return_unprocessed = FALSE,
  force_skip_centerFix = FALSE,
  ...
)
```

Arguments

<code>bam_f</code>	character or BamFile to load
<code>qgr</code>	GRanges regions to fetchs
<code>win_size</code>	numeric ≥ 1 . pileup grabbed every <code>win_size</code> bp for <code>win_method</code> sample. If <code>win_method</code> is <code>summary</code> , this is the number of windows used (confusing, sorry).
<code>win_method</code>	character. one of <code>c("sample", "summary")</code> . Determines if <code>viewGRangesWinSample_dt</code> or <code>viewGRangesWinSummary_dt</code> is used to represent each region in <code>qgr</code> .
<code>summary_FUN</code>	function. only relevant if <code>win_method</code> is <code>"summary"</code> . passed to <code>viewGRangesWinSummary_dt</code> .
<code>fragLen</code>	numeric, NULL, or NA. if numeric, supplied value is used. if NULL, value is calculated with <code>fragLen_calcStranded</code> if NA, raw bam pileup with no cross strand shift is returned.
<code>target_strand</code>	character. if one of <code>"+"</code> or <code>"-"</code> , reads are filtered accordingly. ignored if any other value.
<code>anchor</code>	character, one of <code>c("center", "center_unstranded", "left", "left_unstranded")</code>
<code>return_data.table</code>	logical. If TRUE the internal <code>data.table</code> is returned instead of GRanges. Default is FALSE.
<code>max_dupes</code>	numeric ≥ 1 . duplicate reads by stranddd start position over this number are removed, Default is Inf.
<code>splice_strategy</code>	character, one of <code>c("none", "ignore", "add", "only", "splice_count")</code> . Default is <code>"none"</code> and spliced alignment are assumed not present. <code>fragLen</code> must be NA for any other value to be valid. <code>"ignore"</code> will not count spliced regions. <code>"add"</code> counts spliced regions along with others, <code>"only"</code> will only count spliced regions and ignore others.
<code>flip_strand</code>	if TRUE, strand alignment is flipped prior to <code>fragLen</code> extension. Default is FALSE.
<code>return_unprocessed</code>	boolean. if TRUE returns read alignment in <code>data.table</code> . Default is FALSE.
<code>force_skip_centerFix</code>	boolean, if TRUE all query ranges will be used <code>"as is"</code> . This is already the case by default if <code>win_method == "summary"</code> but may have applications where <code>win_method == "sample"</code> .
<code>...</code>	passed to <code>Rsamtools::ScanBamParam()</code>

Value

tidy GRanges (or `data.table` if specified) with pileups from bam file. pileup is calculated only every `win_size` bp.

ssvFetchBamPE	<i>ssvFetchBam for paired-end ChIP-seq files. Only concordant reads are considered, but this has been minimally tested, please verify.</i>
---------------	--

Description

Iterates a character vector (ideally named) and calls `ssvFetchBamPE.single` on each. Appends grouping variable to each resulting `data.table` and uses `rbindlist` to efficiently combine results

Usage

```
ssvFetchBamPE(
  file_paths,
  qgr,
  unique_names = NULL,
  win_size = 50,
  win_method = c("sample", "summary")[1],
  summary_FUN = stats::weighted.mean,
  anchor = c("left", "left_unstranded", "center", "center_unstranded")[3],
  names_variable = "sample",
  return_data.table = FALSE,
  max_dupes = Inf,
  n_cores = getOption("mc.cores", 1),
  n_region_splits = 1,
  min_isize = 1,
  max_isize = Inf,
  return_unprocessed = FALSE,
  force_skip_centerFix = FALSE,
  ...
)
```

Arguments

<code>file_paths</code>	character vector of <code>file_paths</code> to load from. Alternatively, <code>file_paths</code> can be a <code>data.frame</code> or <code>data.table</code> whose first column is a character vector of paths and additional columns will be used as metadata.
<code>qgr</code>	Set of <code>GRanges</code> to query. For valid results the width of each interval should be identical and evenly divisible by <code>win_size</code> .
<code>unique_names</code>	names to use in final <code>data.table</code> to designate source bigwig. Default is 'sample'
<code>win_size</code>	The window size that evenly divides widths in <code>qgr</code> .
<code>win_method</code>	character. one of <code>c("sample", "summary")</code> . Determines if viewGRangesWinSample_dt or viewGRangesWinSummary_dt is used to represent each region in <code>qgr</code> .
<code>summary_FUN</code>	function. only relevant if <code>win_method</code> is "summary". passed to viewGRangesWinSummary_dt .
<code>anchor</code>	character, one of <code>c("center", "center_unstranded", "left", "left_unstranded")</code>
<code>names_variable</code>	The column name where <code>unique_names</code> are stored.

```

return_data.table
    logical. If TRUE the internal data.table is returned instead of GRanges. Default
    is FALSE.
max_dupes
    numeric >= 1. duplicate reads by strand and start position over this number are
    removed, Default is Inf.
n_cores
    integer number of cores to use.
n_region_splits
    integer number of splits to apply to qgr. The query GRanges will be split into
    this many roughly equal parts for increased parallelization. Default is 1, no split.
min_isize
    integer. Read pairs must have an isize greater than or equal to this value. Default
    is 1.
max_isize
    integer. Read pairs must have an isize less than or equal to this value. Default is
    Inf.
return_unprocessed
    boolean. if TRUE returns read alignment in data.table. Default is FALSE.
force_skip_centerFix
    boolean, if TRUE all query ranges will be used "as is". This is already the
    case by default if win_method == "summary" but may have applications where
    win_method == "sample".
...
    passed to Rsamtools::ScanBamParam() Uses mc.cores option if not supplied.

```

Details

#' In contrast to ssvFetchBam, extension of reads to estimated fragment size is not an issue as each read pair represents a fragment of exact size.

ssvFetchBamPE iteratively calls `fetchWindowedBam.single`. See [ssvFetchBamPE.single](#) for more info.

if qgr contains the range chr1:1-100 and win_size is 10, values from positions chr1 5,15,25...85, and 95 will be retrieved from bw_file

Value

A tidy formatted GRanges (or data.table if specified) containing fetched values.

Examples

```

if(Sys.info()['sysname'] != "Windows"){
  library(GenomicRanges)
  bam_f = system.file("extdata/Bcell_PE.mm10.bam",
    package = "seqsetvis", mustWork = TRUE)
  bam_files = c("a" = bam_f, "b" = bam_f)
  data("Bcell_peaks")
  qgr = Bcell_peaks
  bw_gr = ssvFetchBamPE(bam_files, qgr, win_size = 10)
  bw_gr2 = ssvFetchBamPE(as.list(bam_files), qgr, win_size = 10)

  bw_dt = ssvFetchBamPE(bam_files, qgr, win_size = 10,
    return_data.table = TRUE)
}

```

`ssvFetchBamPE.single` *fetch a windowed version of a paired-end bam file, returns GRanges In contrast to `ssvFetchBam`, extension of reads to estimated fragment size is not an issue as each read pair represents a fragment of exact size.*

Description

fetch a windowed version of a paired-end bam file, returns GRanges In contrast to `ssvFetchBam`, extension of reads to estimated fragment size is not an issue as each read pair represents a fragment of exact size.

Usage

```
ssvFetchBamPE.single(
  bam_f,
  qgr,
  win_size = 50,
  win_method = c("sample", "summary")[1],
  summary_FUN = stats::weighted.mean,
  anchor = c("left", "left_unstranded", "center", "center_unstranded")[3],
  return_data.table = FALSE,
  max_dupes = Inf,
  min_isize = 1,
  max_isize = Inf,
  return_unprocessed = FALSE,
  force_skip_centerFix = FALSE,
  ...
)
```

Arguments

<code>bam_f</code>	character or BamFile to load
<code>qgr</code>	GRanges regions to fetchs
<code>win_size</code>	numeric >=1. pileup grabbed every <code>win_size</code> bp for <code>win_method</code> sample. If <code>win_method</code> is <code>summary</code> , this is the number of windows used (confusing, sorry).
<code>win_method</code>	character. one of <code>c("sample", "summary")</code> . Determines if viewGRangesWinSample_dt or viewGRangesWinSummary_dt is used to represent each region in <code>qgr</code> .
<code>summary_FUN</code>	function. only relevant if <code>win_method</code> is <code>"summary"</code> . passed to viewGRangesWinSummary_dt .
<code>anchor</code>	character, one of <code>c("center", "center_unstranded", "left", "left_unstranded")</code>
<code>return_data.table</code>	logical. If TRUE the internal <code>data.table</code> is returned instead of GRanges. Default is FALSE.
<code>max_dupes</code>	numeric >= 1. duplicate reads by strandd start position over this number are removed, Default is Inf.

`min_isize` integer. Read pairs must have an isize greater than or equal to this value. Default is 1.
`max_isize` integer. Read pairs must have an isize less than or equal to this value. Default is Inf.
`return_unprocessed` boolean. if TRUE returns read alignment in data.table. Default is FALSE.
`force_skip_centerFix` boolean, if TRUE all query ranges will be used "as is". This is already the case by default if `win_method == "summary"` but may have applications where `win_method == "sample"`.
`...` passed to `Rsamtools::ScanBamParam()`

Value

tidy GRanges (or data.table if specified) with pileups from bam file. pileup is calculated only every `win_size` bp.

<code>ssvFetchBigwig</code>	<i>Iterates a character vector (ideally named) and calls <code>ssvFetchBigwig.single</code> on each. Appends grouping variable to each resulting data.table and uses <code>rbindlist</code> to efficiently combine results.</i>
-----------------------------	---

Description

`ssvFetchBigwig` iteratively calls `fetchWindowedBigwig.single`. See [ssvFetchBigwig.single](#) for more info.

Usage

```
ssvFetchBigwig(
  file_paths,
  qgr,
  unique_names = NULL,
  names_variable = "sample",
  win_size = 50,
  win_method = c("sample", "summary")[1],
  summary_FUN = stats::weighted.mean,
  anchor = c("left", "left_unstranded", "center", "center_unstranded")[3],
  return_data.table = FALSE,
  n_cores = getOption("mc.cores", 1),
  n_region_splits = 1,
  force_skip_centerFix = FALSE
)
```

Arguments

<code>file_paths</code>	character vector of <code>file_paths</code> to load from. Alternatively, <code>file_paths</code> can be a <code>data.frame</code> or <code>data.table</code> whose first column is a character vector of paths and additional columns will be used as metadata.
<code>qgr</code>	Set of <code>GRanges</code> to query. For valid results the width of each interval should be identical and evenly divisible by <code>win_size</code> .
<code>unique_names</code>	names to use in final <code>data.table</code> to designate source bigwig.
<code>names_variable</code>	The column name where <code>unique_names</code> are stored. Default is 'sample'
<code>win_size</code>	The window size that evenly divides widths in <code>qgr</code> .
<code>win_method</code>	character. one of <code>c("sample", "summary")</code> . Determines if <code>viewGRangesWinSample_dt</code> or <code>viewGRangesWinSummary_dt</code> is used to represent each region in <code>qgr</code> .
<code>summary_FUN</code>	function. only relevant if <code>win_method</code> is "summary". passed to <code>viewGRangesWinSummary_dt</code> .
<code>anchor</code>	character, one of <code>c("center", "center_unstranded", "left", "left_unstranded")</code>
<code>return_data.table</code>	logical. If TRUE the internal <code>data.table</code> is returned instead of <code>GRanges</code> . Default is FALSE.
<code>n_cores</code>	integer number of cores to use. Uses <code>mc.cores</code> option if not supplied.
<code>n_region_splits</code>	integer number of splits to apply to <code>qgr</code> . The query <code>GRanges</code> will be split into this many roughly equal parts for increased parallelization. Default is 1, no split.
<code>force_skip_centerFix</code>	boolean, if TRUE all query ranges will be used "as is". This is already the case by default if <code>win_method == "summary"</code> but may have applications where <code>win_method == "sample"</code> .

Details

if `qgr` contains the range `chr1:1-100` and `win_size` is 10, values from positions `chr1 5,15,25...85`, and 95 will be retrieved from `bw_file`

Value

A tidy formatted `GRanges` (or `data.table` if specified) containing fetched values.

Examples

```
if(Sys.info()['sysname'] != "Windows"){
  library(GenomicRanges)
  bw_f = system.file("extdata/test_loading.bw",
    package = "seqsetvis", mustWork = TRUE)
  bw_files = c("a" = bw_f, "b" = bw_f)
  qgr = GRanges("chrTest", IRanges(1, 30))
  bw_gr = ssvFetchBigwig(bw_files, qgr, win_size = 10)
  bw_gr2 = ssvFetchBigwig(as.list(bw_files), qgr, win_size = 10)

  bw_dt = ssvFetchBigwig(bw_files, qgr, win_size = 10,
    return_data.table = TRUE)
}
```

ssvFetchBigwig.single *Fetch values from a bigwig appropriate for heatmaps etc.*

Description

ssvFetchBigwig.single Gets values for each region of the query GRanges (qgr). Values correspond to the center of each window of size win_size. A tidy formatted data.table object is returned suitable for plotting using ggplots.

Usage

```
ssvFetchBigwig.single(
  bw_file,
  qgr,
  win_size = 50,
  win_method = c("sample", "summary")[1],
  summary_FUN = stats::weighted.mean,
  anchor = c("left", "left_unstranded", "center", "center_unstranded")[3],
  return_data.table = FALSE,
  force_skip_centerFix = FALSE
)
```

Arguments

bw_file	The character vector path to bigwig files to read from.
qgr	Set of GRanges to query. For valid results the width of each interval should be identical and evenly divisible by win_size.
win_size	The window size that evenly divides widths in qgr.
win_method	character. one of c("sample", "summary"). Determines if viewGRangesWinSample_dt or viewGRangesWinSummary_dt is used to represent each region in qgr.
summary_FUN	function. only relevant if win_method is "summary". passed to viewGRangesWinSummary_dt .
anchor	character, one of c("center", "center_unstranded", "left", "left_unstranded")
return_data.table	logical. If TRUE the internal data.table is returned instead of GRanges. Default is FALSE.
force_skip_centerFix	boolean, if TRUE all query ranges will be used "as is". This is already the case by default if win_method == "summary" but may have applications where win_method == "sample".

Details

if qgr contains the range chr1:1-100 and win_size is 10, values from positions chr1 5,15,25...85, and 95 will be retrieved from bw_file

Value

A GRanges (or data.table if specified) containing fetched values.

ssvFetchGRanges	<i>Fetch coverage values for a list of GRanges.</i>
-----------------	---

Description

ssvFetchGRanges Gets coverage values for each region of the query GRanges (qgr). Values correspond to the center of each window of size win_size. A tidy formatted data.table object is returned suitable for plotting using ggplots.

Usage

```
ssvFetchGRanges(
  grs,
  qgr,
  file_attribs = data.frame(matrix(0, nrow = length(grs), ncol = 0)),
  unique_names = names(grs),
  names_variable = "sample",
  win_size = 50,
  win_method = c("sample", "summary")[1],
  summary_FUN = function(x, w) max(x),
  target_strand = c("*", "+", "-", "both")[1],
  use_coverage = NULL,
  attrib_var = "score",
  fill_value = 0,
  anchor = c("left", "left_unstranded", "center", "center_unstranded")[3],
  return_data.table = FALSE,
  n_cores = getOption("mc.cores", 1),
  force_skip_centerFix = FALSE
)
```

Arguments

grs	a list of GRanges for which to calculate coverage.
qgr	Set of GRanges to query. For valid results the width of each interval should be identical and evenly divisible by win_size.
file_attribs	data.frame (1 row per item in grs) containing attributes to append to results.
unique_names	The column name where unique_names are stored. Default is 'sample'
names_variable	The column name where unique_names are stored. Default is 'sample'
win_size	The window size that evenly divides widths in qgr.
win_method	character. one of c("sample", "summary"). Determines if viewGRangesWinSample_dt or viewGRangesWinSummary_dt is used to represent each region in qgr.

summary_FUN	function. only relevant if win_method is "summary". passed to viewGRangesWinSummary_dt .
target_strand	character. if one of "+" or "-", reads are filtered to match. ignored if any other value.
use_coverage	boolean or NULL, if TRUE, query regions are scored by the number of intervals overlapping. Default of NULL checks if attrib_var is "score" and uses coverage if so.
attrib_var	character, column in mcols of GRanges to pull values from. Default of "score" is compatible with internal coverage calculation or bedgraph-like files.
fill_value	numeric or character value to use where queried regions are empty. Default is 0 and appropriate for both calculated coverage and bedgraph/bigwig like files. Will automatically switch to "MISSING" if data is guessed to be qualitative.
anchor	character, one of c("center", "center_unstranded", "left", "left_unstranded")
return_data.table	logical. If TRUE the internal data.table is returned instead of GRanges. Default is FALSE.
n_cores	integer number of cores to use. Uses mc.cores option if not supplied.
force_skip_centerFix	boolean, if TRUE all query ranges will be used "as is". This is already the case by default if win_method == "summary" but may have applications where win_method == "sample".

Value

A tidy formatted GRanges (or data.table if specified) containing fetched values.

Examples

```
ssvFetchGRanges(CTCF_in_10a_narrowPeak_grs, CTCF_in_10a_overlaps_gr, win_size = 200)
```

ssvFetchSignal	<i>signal loading framework</i>
----------------	---------------------------------

Description

Does nothing unless load_signal is overridden to carry out reading data from file_paths (likely via the appropriate ssvFetch* function, ie. [ssvFetchBigwig](#) or [ssvFetchBam](#)

Usage

```
ssvFetchSignal(
  file_paths,
  qgr,
  unique_names = NULL,
  names_variable = "sample",
  file_attribs = NULL,
```

```

win_size = 50,
win_method = c("sample", "summary")[1],
return_data.table = FALSE,
load_signal = function(f, nam, qgr) {      warning("nothing happened, ",
      "supply a function to", "load_signal parameter.") },
n_cores = getOption("mc.cores", 1),
n_region_splits = 1,
force_skip_centerFix = FALSE
)

```

Arguments

<code>file_paths</code>	character vector of <code>file_paths</code> to load from. Alternatively, <code>file_paths</code> can be a <code>data.frame</code> or <code>data.table</code> whose first column is a character vector of paths and additional columns will be used as metadata.
<code>qgr</code>	<code>GRanges</code> of intervals to return from each file
<code>unique_names</code>	unique file ids for each file in <code>file_paths</code> . Default is names of <code>file_paths</code> vector
<code>names_variable</code>	character, variable name for column containing <code>unique_names</code> entries. Default is "sample"
<code>file_attribs</code>	optional <code>data.frame</code> / <code>data.table</code> with one row per item in file paths. Each column will be a variable added to final tidy output.
<code>win_size</code>	numeric/integer window size resolution to load signal at. Default is 50.
<code>win_method</code>	character. one of <code>c("sample", "summary")</code> . Determines if viewGRangesWinSample_dt or viewGRangesWinSummary_dt is used to represent each region in <code>qgr</code> .
<code>return_data.table</code>	logical. If TRUE <code>data.table</code> is returned instead of <code>GRanges</code> , the default.
<code>load_signal</code>	function taking <code>f</code> , <code>nam</code> , and <code>qgr</code> arguments. <code>f</code> is from <code>file_paths</code> , <code>nam</code> is from <code>unique_names</code> , and <code>qgr</code> is <code>qgr</code> . See details.
<code>n_cores</code>	integer number of cores to use. Uses <code>mc.cores</code> option if not supplied.
<code>n_region_splits</code>	integer number of splits to apply to <code>qgr</code> . The query <code>GRanges</code> will be split into this many roughly equal parts for increased parallelization. Default is 1, no split.
<code>force_skip_centerFix</code>	boolean, if TRUE all query ranges will be used "as is". This is already the case by default if <code>win_method == "summary"</code> but may have applications where <code>win_method == "sample"</code> .

Details

`load_signal` is passed `f`, `nam`, and `qgr` and is executed in the environment where `load_signal` is defined. See [ssvFetchBigwig](#) and [ssvFetchBam](#) for examples.

Value

A `GRanges` with values read from `file_paths` at intervals of `win_size`. Originating file is coded by `unique_names` and assigned to column of name `names_variable`. Output is `data.table` is `return_data.table` is TRUE.

Examples

```

library(GenomicRanges)
bam_f = system.file("extdata/test.bam",
  package = "seqsetvis", mustWork = TRUE)
bam_files = c("a" = bam_f, "b" = bam_f)
qgr = CTCF_in_10a_overlaps_gr[1:2]
qgr = resize(qgr, 500, "center")
load_bam = function(f, nam, qgr) {
  message("loading ", f, " ...")
  dt = seqsetvis::ssvFetchBam.single(bam_f = f,
    qgr = qgr,
    win_size = 50,
    fragLen = NULL,
    target_strand = "*",
    return_data.table = TRUE)
  dt[["sample"]] = nam
  message("finished loading ", nam, ".")
  dt
}
ssvFetchSignal(bam_files, qgr, load_signal = load_bam)

```

ssvMakeMembTable	<i>generic for methods to convert various objects to a logical matrix indicating membership of items (rows) in sets (columns)</i>
------------------	---

Description

generic for methods to convert various objects to a logical matrix indicating membership of items (rows) in sets (columns)

list of character vectors input

GRangesList input

GRanges with mcols input

DataFrame input

matrix of logicals, membership table

data.frame input, final output The final method for all inputs, checks column names and returns logical matrix

Usage

```
ssvMakeMembTable(object)
```

```
## S4 method for signature 'list'
ssvMakeMembTable(object)
```

```
## S4 method for signature 'GRangesList'
ssvMakeMembTable(object)
```

```
## S4 method for signature 'GRanges'
ssvMakeMembTable(object)

## S4 method for signature 'DataFrame'
ssvMakeMembTable(object)

## S4 method for signature 'matrix'
ssvMakeMembTable(object)

## S4 method for signature 'data.frame'
ssvMakeMembTable(object)
```

Arguments

object the object to convert. Supported types: list (of character or GRanges), GRanges with membership table metadata, GRangesList, data.frame/matrix/DataFrame of membership table

Value

a logical matrix indicating membership of items (rows) in sets (columns)

Examples

```
char_list = list(letters[1:3], letters[2:4])
ssvMakeMembTable(char_list)
library(GenomicRanges)
gr_list = list(GRanges("chr1", IRanges(1:3*2, 1:3*2)),
               GRanges("chr1", IRanges(2:4*2, 2:4*2)))
ssvMakeMembTable(gr_list)
library(GenomicRanges)
gr_list = list(GRanges("chr1", IRanges(1:3*2, 1:3*2)),
               GRanges("chr1", IRanges(2:4*2, 2:4*2)))
ssvMakeMembTable(GRangesList(gr_list))
gr = GRanges("chr1", IRanges(1:3*2, 1:3*2))
gr$set_a = c(TRUE, TRUE, FALSE)
gr$set_b = c(FALSE, TRUE, TRUE)
ssvMakeMembTable(gr)
gr = GRanges("chr1", IRanges(1:3*2, 1:3*2))
gr$set_a = c(TRUE, TRUE, FALSE)
gr$set_b = c(FALSE, TRUE, TRUE)
ssvMakeMembTable(mcols(gr))
memb_mat = matrix(c(TRUE, TRUE, FALSE, FALSE, TRUE, FALSE, TRUE, FALSE),
                  ncol = 2, byrow = FALSE)
ssvMakeMembTable(memb_mat)
memb_df = data.frame(a = c(TRUE, TRUE, FALSE, FALSE),
                     b = c(TRUE, FALSE, TRUE, FALSE))
ssvMakeMembTable(memb_df)
```

ssvOverlapIntervalSets

Intersect a list of GRanges to create a single GRanges object of merged ranges including metadata describing overlaps per input GRanges

Description

Intersect a list of GRanges to create a single GRanges object of merged ranges including metadata describing overlaps per input GRanges

Usage

```
ssvOverlapIntervalSets(grs, ext = 0, use_first = FALSE, ...)
```

Arguments

grs	A list of GRanges
ext	An integer specifying how far to extend ranges before merging. in effect, ranges withing 2*ext of one another will be joined during the merge
use_first	A logical. If True, instead of merging all grs, only use first and add metadata logicals for others.
...	arguments passed to IRanges::findOverlaps, i.e. maxgap, minoverlap, type, select, invert.

Value

GRanges with metadata columns describing overlap of input grs.

Examples

```
library(GenomicRanges)
a = GRanges("chr1", IRanges(1:7*10, 1:7*10))
b = GRanges("chr1", IRanges(5:10*10, 5:10*10))
ssvOverlapIntervalSets(list(a, b))
```

ssvSignalBandedQuantiles

plot profiles from bigwigs

Description

plot profiles from bigwigs

Usage

```
ssvSignalBandedQuantiles(
  bw_data,
  y_ = "y",
  x_ = "x",
  by_ = "fake",
  hsv_reverse = FALSE,
  hsv_saturation = 1,
  hsv_value = 1,
  hsv_grayscale = FALSE,
  hsv_hue_min = 0,
  hsv_hue_max = 0.7,
  hsv_symmetric = FALSE,
  n_quantile = 18,
  quantile_min = 0.05,
  quantile_max = 0.95,
  return_data = FALSE
)
```

Arguments

<code>bw_data</code>	a GRanges or data.table of bigwig signal. As returned from ssvFetchBam and ssvFetchBigwig
<code>y_</code>	the variable name in <code>bw_data</code> for y axis in plot
<code>x_</code>	the variable name in <code>bw_data</code> for x axis in plot
<code>by_</code>	the variable name in <code>bw_data</code> to facet on
<code>hsv_reverse</code>	logical, should color scale be reversed? default FALSE
<code>hsv_saturation</code>	numeric [0, 1] saturation for color scale. default 1
<code>hsv_value</code>	numeric [0, 1] value for color scale. default 1
<code>hsv_grayscale</code>	logical, if TRUE <code>gray()</code> is used instead of <code>rainbow()</code> . default FALSE
<code>hsv_hue_min</code>	numeric [0, <code>hsv_hue_max</code>) hue min of color scale
<code>hsv_hue_max</code>	numeric (<code>hsv_hue_min</code> , 1] hue max of color scale
<code>hsv_symmetric</code>	if TRUE, colorscale is symmetrical, default FALSE.
<code>n_quantile</code>	number of evenly size quantile bins
<code>quantile_min</code>	the lowest quantile start
<code>quantile_max</code>	the highest quantile end
<code>return_data</code>	logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.

Value

ggplot object using ribbon plots to show quantile distributions

Examples

```
#rainbow colors
qgr = CTCF_in_10a_profiles_gr
ssvSignalBandedQuantiles(qgr)
#grayscale
ssvSignalBandedQuantiles(qgr, hsv_grayscale = TRUE,
  hsv_symmetric = TRUE, hsv_reverse = TRUE)
#using "by_" per sample
ssvSignalBandedQuantiles(qgr, hsv_grayscale = TRUE,
  hsv_symmetric = TRUE, hsv_reverse = TRUE, by_ = "sample")
#adding spline smoothing
splined = applySpline(qgr, n = 10,
  by_ = c("id", "sample"))
ssvSignalBandedQuantiles(splined, n_quantile = 50,
  quantile_min = .25, quantile_max = .75,
  hsv_symmetric = TRUE, hsv_reverse = TRUE, by_ = "sample")
```

ssvSignalClustering	<i>Clustering as for a heatmap. This is used internally by ssvSignalHeatmap but can also be run before calling ssvSignalHeatmap for greater control and access to clustering results directly.</i>
---------------------	--

Description

Clustering is via k-means by default. The number of clusters is determined by `nclust`. Optionally, k-means can be initialized with a data.frame provided to `k_centroids`. As an alternative to k-means, a membership table from [ssvMakeMembTable](#) can be provided to determine logical clusters.

Usage

```
ssvSignalClustering(
  bw_data,
  nclust = NULL,
  k_centroids = NULL,
  memb_table = NULL,
  row_ = "id",
  column_ = "x",
  fill_ = "y",
  facet_ = "sample",
  cluster_ = "cluster_id",
  max_rows = 500,
  max_cols = 100,
  clustering_col_min = -Inf,
  clustering_col_max = Inf,
  within_order_strategy = c("hclust", "sort")[2],
  dcast_fill = NA
)
```

Arguments

<code>bw_data</code>	a GRanges or data.table of bigwig signal. As returned from ssvFetchBam and ssvFetchBigwig
<code>nclust</code>	Number of clusters. Defaults to 6 if <code>nclust</code> , <code>k_centroids</code> , and <code>memb_table</code> are not set.
<code>k_centroids</code>	data.frame of centroids for k-means clusters. Incompatible with <code>nclust</code> or <code>memb_table</code> .
<code>memb_table</code>	Membership table as from ssvMakeMembTable . Logical groups from membership table will be clusters. Incompatible with <code>nclust</code> or <code>k_centroids</code> .
<code>row_</code>	variable name mapped to row, likely id or gene name for ngs data. Default is "id" and works with <code>ssvFetch*</code> output.
<code>column_</code>	variable mapped to column, likely bp position for ngs data. Default is "x" and works with <code>ssvFetch*</code> output.
<code>fill_</code>	numeric variable to map to fill. Default is "y" and works with <code>ssvFetch*</code> output.
<code>facet_</code>	variable name to facet horizontally by. Default is "sample" and works with <code>ssvFetch*</code> output. Set to "" if data is not faceted.
<code>cluster_</code>	variable name to use for cluster info. Default is "cluster_id".
<code>max_rows</code>	for speed rows are sampled to 500 by default, use Inf to plot full data
<code>max_cols</code>	for speed columns are sampled to 100 by default, use Inf to plot full data
<code>clustering_col_min</code>	numeric minimum for col range considered when clustering, default in -Inf
<code>clustering_col_max</code>	numeric maximum for col range considered when clustering, default in Inf
<code>within_order_strategy</code>	one of "hclust" or "sort". if hclust, hierarchical clustering will be used. if sort, a simple decreasing sort of <code>rosSums</code> .
<code>dcast_fill</code>	value to supply to dcast fill argument. default is NA.

Details

Within each cluster, items will either be sorted by decreasing average signal or hierachically clustered; this is controlled via `within_order_strategy`.

Value

data.table of signal profiles, ready for `ssvSignalHeatmap`

Examples

```
clust_dt = ssvSignalClustering(CTCF_in_10a_profiles_gr)
ssvSignalHeatmap(clust_dt)

clust_dt2 = ssvSignalClustering(CTCF_in_10a_profiles_gr, nclust = 2)
ssvSignalHeatmap(clust_dt2)

#clustering can be targetted to a specific part of the region
```

```

clust_dt3 = ssvSignalClustering(CTCF_in_10a_profiles_gr, nclust = 2,
  clustering_col_min = -250, clustering_col_max = -150)
ssvSignalHeatmap(clust_dt3)
clust_dt4 = ssvSignalClustering(CTCF_in_10a_profiles_gr, nclust = 2,
  clustering_col_min = 150, clustering_col_max = 250)
ssvSignalHeatmap(clust_dt4)

```

ssvSignalHeatmap	<i>heatmap style representation of membership table. instead of clustering, each column is sorted starting from the left.</i>
------------------	---

Description

See [ssvSignalHeatmap.ClusterBars](#) for an alternative with more control over where the cluster bars appear.

Usage

```

ssvSignalHeatmap(
  bw_data,
  nclust = 6,
  perform_clustering = c("auto", "yes", "no")[1],
  row_ = "id",
  column_ = "x",
  fill_ = "y",
  facet_ = "sample",
  cluster_ = "cluster_id",
  max_rows = 500,
  max_cols = 100,
  fill_limits = NULL,
  clustering_col_min = -Inf,
  clustering_col_max = Inf,
  within_order_strategy = c("hclust", "sort")[2],
  dcast_fill = NA,
  return_data = FALSE,
  show_clusterBars = TRUE
)

```

Arguments

bw_data	a GRanges or data.table of bigwig signal. As returned from ssvFetchBam and ssvFetchBigwig
nclust	number of clusters
perform_clustering	should clustering be done? default is auto. auto considers if row_ has been ordered by being a factor and if cluster_ is a numeric.

row_	variable name mapped to row, likely id or gene name for ngs data. Default is "id" and works with ssvFetch* output.
column_	variable mapped to column, likely bp position for ngs data. Default is "x" and works with ssvFetch* output.
fill_	numeric variable to map to fill. Default is "y" and works with ssvFetch* output.
facet_	variable name to facet horizontally by. Default is "sample" and works with ssvFetch* output. Set to "" if data is not faceted.
cluster_	variable name to use for cluster info. Default is "cluster_id".
max_rows	for speed rows are sampled to 500 by default, use Inf to plot full data
max_cols	for speed columns are sampled to 100 by default, use Inf to plot full data
fill_limits	limits for fill legend. values will be cropped to this range if set. Default of NULL uses natural range of fill_.
clustering_col_min	numeric minimum for col range considered when clustering, default in -Inf
clustering_col_max	numeric maximum for col range considered when clustering, default in Inf
within_order_strategy	one of "hclust" or "sort". if hclust, hierarchical clustering will be used. if sort, a simple decreasing sort of rosSums.
dcast_fill	value to supply to dcast fill argument. default is NA.
return_data	logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.
show_clusterBars	if TRUE, show bars indicating cluster membership.

Value

ggplot heatmap of signal profiles, faceted by sample

Examples

```
#the simplest use
ssvSignalHeatmap(CTCF_in_10a_profiles_gr)
ssvSignalHeatmap(CTCF_in_10a_profiles_gr, show_clusterBars = FALSE)

#clustering can be done manually beforehand
clust_dt = ssvSignalClustering(CTCF_in_10a_profiles_gr, nclust = 3)
ssvSignalHeatmap(clust_dt)

ssvSignalHeatmap(clust_dt, max_rows = 20, max_cols = 7)
```

ssvSignalHeatmap.ClusterBars

heatmap style representation of membership table. instead of clustering, each column is sorted starting from the left.

Description

Compared to ssvSignalHeatmap, cluster_bars are displayed on the left once instead of for each facet

Usage

```
ssvSignalHeatmap.ClusterBars(
  bw_data,
  nclust = 6,
  perform_clustering = c("auto", "yes", "no")[1],
  row_ = "id",
  column_ = "x",
  fill_ = "y",
  facet_ = "sample",
  cluster_ = "cluster_id",
  max_rows = 500,
  max_cols = 100,
  fill_limits = NULL,
  clustering_col_min = -Inf,
  clustering_col_max = Inf,
  within_order_strategy = c("hclust", "sort")[2],
  dcast_fill = NA,
  return_data = FALSE,
  return_unassembled_plots = FALSE,
  rel_widths = c(1, 9),
  ...
)
```

Arguments

bw_data	a GRanges or data.table of bigwig signal. As returned from ssvFetchBam and ssvFetchBigwig
nclust	number of clusters
perform_clustering	should clustering be done? default is auto. auto considers if row_ has been ordered by being a factor and if cluster_ is a numeric.
row_	variable name mapped to row, likely id or gene name for ngs data. Default is "id" and works with ssvFetch* output.
column_	variable mapped to column, likely bp position for ngs data. Default is "x" and works with ssvFetch* output.
fill_	numeric variable to map to fill. Default is "y" and works with ssvFetch* output.

facet_	variable name to facet horizontally by. Default is "sample" and works with ssvFetch* output. Set to "" if data is not faceted.
cluster_	variable name to use for cluster info. Default is "cluster_id".
max_rows	for speed rows are sampled to 500 by default, use Inf to plot full data
max_cols	for speed columns are sampled to 100 by default, use Inf to plot full data
fill_limits	limits for fill legend. values will be cropped to this range if set. Default of NULL uses natural range of fill_.
clustering_col_min	numeric minimum for col range considered when clustering, default in -Inf
clustering_col_max	numeric maximum for col range considered when clustering, default in Inf
within_order_strategy	one of "hclust" or "sort". if hclust, hierarchical clustering will be used. if sort, a simple decreasing sort of rosSums.
dcast_fill	value to supply to dcast fill argument. default is NA.
return_data	logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.
return_unassembled_plots	logical. If TRUE, return list of heatmap and cluster-bar ggplots. Can be customized and passed to assemble_heatmap_cluster_bars
rel_widths	numeric of length 2. Passed to cowplot::plot_grid. Default is c(1, 9).
...	additional arguments passed to cowplot::plot_grid

Value

ggplot heatmap of signal profiles, faceted by sample

Examples

```
#the simplest use
ssvSignalHeatmap.ClusterBars(CTCF_in_10a_profiles_gr)
ssvSignalHeatmap.ClusterBars(CTCF_in_10a_profiles_gr, rel_widths = c(1, 5))

#clustering can be done manually beforehand
clust_dt = ssvSignalClustering(CTCF_in_10a_profiles_gr, nclust = 3)
ssvSignalHeatmap.ClusterBars(clust_dt)
```

ssvSignalLineplot	<i>construct line type plots where each region in each sample is represented</i>
-------------------	--

Description

construct line type plots where each region in each sample is represented

Usage

```
ssvSignalLineplot(
  bw_data,
  x_ = "x",
  y_ = "y",
  color_ = "sample",
  sample_ = "sample",
  region_ = "id",
  group_ = "auto_grp",
  line_alpha = 1,
  facet_ = "auto_facet",
  facet_method = facet_wrap,
  spline_n = NULL,
  return_data = FALSE
)
```

Arguments

<code>bw_data</code>	a GRanges or data.table of bigwig signal. As returned from ssvFetchBam and ssvFetchBigwig
<code>x_</code>	variable name mapped to x aesthetic, x by default.
<code>y_</code>	variable name mapped to y aesthetic, y by default.
<code>color_</code>	variable name mapped to color aesthetic, sample by default.
<code>sample_</code>	variable name, along with <code>region_</code> used to group and facet by default, change <code>group_</code> or <code>facet_</code> to override.
<code>region_</code>	variable name, along with <code>sample_</code> used to group and facet by default, change <code>group_</code> or <code>facet_</code> to override.
<code>group_</code>	group aesthetic keeps lines of <code>geom_path</code> from mis-connecting. <code>auto_grp</code> by default which combines <code>sample_</code> and <code>region_</code> . probably shouldn't change.
<code>line_alpha</code>	alpha value for lines. default is 1.
<code>facet_</code>	facetting divides up plots. <code>auto_facet</code> by default which combines <code>sample_</code> and <code>region_</code> . if overriding <code>facet_method</code> with <code>facet_grid</code> , make sure to include ~ between two variables, ie. "a~b", ".~b", "a~."
<code>facet_method</code>	ggplot2 facetting method or wrapper for same, <code>facet_wrap</code> by default.
<code>spline_n</code>	if not NULL, <code>applySpline</code> will be called with <code>n = spline_n</code> . default is NULL.
<code>return_data</code>	logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.

Value

ggplot of signal potentially facetted by region and sample

Examples

```
bw_gr = CTCF_in_10a_profiles_gr
ssvSignalLineplot(subset(bw_gr, bw_gr$id %in% seq_len(3)), facet_ = "sample")
ssvSignalLineplot(subset(bw_gr, bw_gr$id %in% seq_len(3)),
  facet_ = "sample~.",
  facet_method = facet_grid)
ssvSignalLineplot(subset(bw_gr, bw_gr$id %in% seq_len(3)),
  facet_ = paste("sample", "~", "id"), facet_method = facet_grid)
ssvSignalLineplot(subset(bw_gr, bw_gr$id %in% seq_len(3)))
ssvSignalLineplot(subset(bw_gr, bw_gr$id %in% seq_len(3)), facet_ = "id")
ssvSignalLineplot(subset(bw_gr, bw_gr$id %in% seq_len(3)),
  facet_ = "id", spline_n = 10)
```

ssvSignalLineplotAgg *aggregate line signals in a single line plot*

Description

aggregate line signals in a single line plot

Usage

```
ssvSignalLineplotAgg(
  bw_data,
  x_ = "x",
  y_ = "y",
  sample_ = "sample",
  color_ = sample_,
  group_ = sample_,
  agg_fun = mean,
  spline_n = NULL,
  return_data = FALSE
)
```

Arguments

bw_data	a GRanges or data.table of bigwig signal. As returned from ssvFetchBam and ssvFetchBigwig
x_	variable name mapped to x aesthetic, x by default.
y_	variable name mapped to y aesthetic, y by default.
sample_	variable name, along with region_ used to group by default,
color_	variable name mapped to color aesthetic, sample_ by default. change group_ to override.
group_	group aesthetic keeps lines of geom_path from mis-connecting. Most useful if you need to supply a variable to later facet upon. Defaults to value of sample_.
agg_fun	the aggregation function to apply by sample_ and x_, default is mean

spline_n if not NULL, applySpline will be called with n = spline_n. default is NULL.
 return_data logical. If TRUE, return value is no longer ggplot and is instead the data used to
 generate that plot. Default is FALSE.

Value

ggplot of signal aggregated with agg_fun() by sample.

Examples

```
bw_gr = CTCF_in_10a_profiles_gr
ssvSignalLineplotAgg(bw_gr) +
  labs(title = "agg regions by sample.")
ssvSignalLineplotAgg(CTCF_in_10a_profiles_gr, spline_n = 10) +
  labs(title = "agg regions by sample, with spline smoothing.")
ssvSignalLineplotAgg(subset(bw_gr, bw_gr$id %in% seq_len(10)),
  sample_ = "id", color_ = "id") +
  labs(title = "agg samples by region id (weird)")
ssvSignalLineplotAgg(subset(bw_gr, bw_gr$id %in% seq_len(10)), sample_ = "id",
  color_ = "id", spline_n = 10) +
  labs(title = "agg samples by region id (weird), with spline smoothing")
```

ssvSignalScatterplot *maps signal from 2 sample profiles to the x and y axis. axes are stan-*
 standard or "volcano" min XY vs fold-change Y/X

Description

maps signal from 2 sample profiles to the x and y axis. axes are standard or "volcano" min XY vs fold-change Y/X

Usage

```
ssvSignalScatterplot(
  bw_data,
  x_name,
  y_name,
  color_table = NULL,
  value_variable = "y",
  xy_variable = "sample",
  value_function = max,
  by_ = "id",
  plot_type = c("standard", "volcano")[1],
  show_help = FALSE,
  fixed_coords = TRUE,
  return_data = FALSE
)
```

Arguments

<code>bw_data</code>	a GRanges or data.table of bigwig signal. As returned from ssvFetchBam and ssvFetchBigwig
<code>x_name</code>	sample name to map to x-axis, must be stored in variable specified in <code>xy_variable</code>
<code>y_name</code>	sample name to map to y-axis, must be stored in variable specified in <code>xy_variable</code>
<code>color_table</code>	data.frame with 2 columns, one of which must be named "group" and gets mapped to color. The other column must be the same as <code>by_</code> parameter and is used for merging.
<code>value_variable</code>	variable name that stores numeric values for plotting, default is "y"
<code>xy_variable</code>	variable name that stores sample, must contain entires for <code>x_name</code> and <code>y_name</code>
<code>value_function</code>	a function to apply to <code>value_variable</code> in all combinations of <code>by_</code> per <code>x_name</code> and <code>y_name</code>
<code>by_</code>	variables that store individual measurement ids
<code>plot_type</code>	standard or volcano, default is "standard"
<code>show_help</code>	if TRUE overlay labels to aid plot interpretation, default is FALSE
<code>fixed_coords</code>	if TRUE coordinate system is 1:1 ratio, default is TRUE
<code>return_data</code>	logical. If TRUE, return value is no longer ggplot and is instead the data used to generate that plot. Default is FALSE.

Value

ggplot of points comparing signal from 2 samples

Examples

```
ssvSignalScatterplot(CTCF_in_10a_profiles_gr,
  x_name = "MCF10A_CTCF", y_name = "MCF10AT1_CTCF")
ssvSignalScatterplot(CTCF_in_10a_profiles_gr,
  x_name = "MCF10A_CTCF", y_name = "MCF10CA1_CTCF")

ssvSignalScatterplot(CTCF_in_10a_profiles_gr,
  x_name = "MCF10A_CTCF", y_name = "MCF10AT1_CTCF",
  value_function = median) + labs(title = "median FE in regions")

ssvSignalScatterplot(CTCF_in_10a_profiles_gr,
  x_name = "MCF10A_CTCF", y_name = "MCF10AT1_CTCF",
  plot_type = "volcano")

ssvSignalScatterplot(CTCF_in_10a_profiles_gr,
  x_name = "MCF10A_CTCF", y_name = "MCF10AT1_CTCF",
  plot_type = "volcano", show_help = TRUE)
```

test_peaks	<i>4 random peaks for single-end data and 4 control regions 30kb downstream from each peak.</i>
------------	---

Description

matches system.file("extdata/test_peaks.bam",package = "seqsetvis")

Format

GRanges length 8

Details

this is included only for testing ssvFetchBam functions.

viewGRangesWinSample_dt	<i>get a windowed sampling of score_gr</i>
-------------------------	--

Description

This method is appropriate when all GRanges in qgr are identical width and when it is practical to use a window_size smaller than features in genomic signal. For instance, when retrieving signal around peaks or promoters this method maintains a fixed genomic scale across regions. This allows meaningful comparison of peak widths can be made.

Usage

```
viewGRangesWinSample_dt(  
  score_gr,  
  qgr,  
  window_size,  
  attrib_var = "score",  
  fill_value = 0,  
  anchor = c("center", "center_unstranded", "left", "left_unstranded")[1]  
)
```

Arguments

score_gr	GRanges with a "score" metadata column.
qgr	regions to view by window.
window_size	qgr will be represented by value from score_gr every window_size bp.
attrib_var	character name of attribute to pull data from. Default is "score", compatible with with bigWigs or bam coverage.

fill_value	numeric or character value to use where queried regions are empty. Default is 0 and appropriate for both calculated coverage and bedgraph/bigwig like files. Will automatically switch to "MISSING" if data is guessed to be qualitative.
anchor	character. controls how x value is derived from position for each region in qgr. 0 may be the left side or center. If not unstranded, x coordinates are flipped for (-) strand. One of c("center", "center_unstranded", "left", "left_unstranded"). Default is "center".

Details

Summarizes score_gr by grabbing value of "score" every window_size bp. Columns in output data.table are: standard GRanges columns: seqnames, start, end, width, strand id - matched to names(score_gr). if names(score_gr) is missing, added as 1:length(score_gr). y - value of score from score_gr. x - relative bp position.

Value

data.table that is GRanges compatible

Examples

```
bam_file = system.file("extdata/test.bam",
  package = "seqsetvis")
qgr = CTCF_in_10a_overlaps_gr[seq_len(5)]
qgr = GenomicRanges::resize(qgr, width = 500, fix = "center")
bam_gr = seqsetvis::fetchBam(bam_file, qgr)
bam_dt = viewGRangesWinSample_dt(bam_gr, qgr, 50)

if(Sys.info()['sysname'] != "Windows"){
  bw_file = system.file("extdata/MCF10A-CTCF_FE_random100.bw",
    package = "seqsetvis")
  bw_gr = rtracklayer::import.bw(bw_file, which = qgr)
  bw_dt = viewGRangesWinSample_dt(bw_gr, qgr, 50)
}
```

viewGRangesWinSummary_dt

Summarizes signal in bins. The same number of bins per region in qgr is used and widths can vary in qgr, in contrast to [viewGRangesWinSample_dt](#) where width must be constant across regions.

Description

This function is most appropriate where features are expected to vary greatly in size and feature boundaries are important, ie. gene bodies, enhancers or TADs.

Usage

```
viewGRangesWinSummary_dt(
  score_gr,
  qgr,
  n_tiles = 100,
  attrib_var = "score",
  attrib_type = NULL,
  fill_value = 0,
  anchor = c("center", "center_unstranded", "left", "left_unstranded")[1],
  summary_FUN = stats::weighted.mean
)
```

Arguments

score_gr	GRanges with a "score" metadata column.
qgr	regions to view by window.
n_tiles	numeric >= 1, the number of tiles to use for every region in qgr.
attrib_var	character name of attribute to pull data from. Default is "score", compatible with bigWigs or bam coverage.
attrib_type	one of NULL, qualitative or quantitative. If NULL will attempt to guess by casting attrib_var attribute to character or factor. Default is NULL.
fill_value	numeric or character value to use where queried regions are empty. Default is 0 and appropriate for both calculated coverage and bedgraph/bigwig like files. Will automatically switch to "MISSING" if data is guessed to be qualitative.
anchor	character. controls how x value is derived from position for each region in qgr. 0 may be the left side or center. If not unstranded, x coordinates are flipped for (-) strand. One of c("center", "center_unstranded", "left", "left_unstranded"). Default is "center".
summary_FUN	function. used to aggregate score by tile. must accept x=score and w=width numeric vectors as only arguments. default is weighted.mean. limma::weighted.median is a good alternative.

Details

Columns in output data.table are: standard GRanges columns: seqnames, start, end, width, strand
 id - matched to names(score_gr). if names(score_gr) is missing, added as seq_along(score_gr). y -
 value of score from score_gr x - relative bp position

Value

data.table that is GRanges compatible

Examples

```
bam_file = system.file("extdata/test.bam",
  package = "seqsetvis")
qgr = CTCF_in_10a_overlaps_gr[1:5]
```

```
# unlike viewGRangesWinSample_dt, width is not fixed
# qgr = GenomicRanges::resize(qgr, width = 500, fix = "center")
bam_gr = seqsetvis::fetchBam(bam_file, qgr)
bam_dt = viewGRangesWinSummary_dt(bam_gr, qgr, 50)

if(Sys.info()['sysname'] != "Windows"){
  bw_file = system.file("extdata/MCF10A_CTCF_FE_random100.bw",
    package = "seqsetvis")
  bw_gr = rtracklayer::import.bw(bw_file, which = qgr)
  bw_dt = viewGRangesWinSummary_dt(bw_gr, qgr, 50)
}
```

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