Package ‘ggbio’

March 13, 2024

Version  1.50.0

Title  Visualization tools for genomic data

Description  The ggbio package extends and specializes the grammar of graphics for biological data. The graphics are designed to answer common scientific questions, in particular those often asked of high throughput genomics data. All core Bioconductor data structures are supported, where appropriate. The package supports detailed views of particular genomic regions, as well as genome-wide overviews. Supported overviews include ideograms and grand linear views. High-level plots include sequence fragment length, edge-linked interval to data view, mismatch pileup, and several splicing summaries.

Depends  methods, BiocGenerics, ggplot2 (>= 1.0.0)

Imports  grid, grDevices, graphics, stats, utils, gridExtra, scales, reshape2, gtable, Hmisc, biovizBase (>= 1.29.2), Biobase, S4Vectors (>= 0.13.13), IRanges (>= 2.11.16), GenomeInfoDb (>= 1.1.3), GenomicRanges (>= 1.29.14), SummarizedExperiment, Biostrings, Rsamtools (>= 1.17.28), GenomicAlignments (>= 1.1.16), BSgenome, VariantAnnotation (>= 1.11.4), rtracklayer (>= 1.25.16), GenomicFeatures (>= 1.29.11), OrganismDbi, GGally, ensemblpdb (>= 1.99.13), AnnotationDbi, AnnotationFilter, rlang

VignetteBuilder knitr


URL  https://lawremi.github.io/ggbio/

BugReports  https://github.com/lawremi/ggbio/issues

License  Artistic-2.0

LazyLoad  Yes
**R topics documented:**

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**arrangeGrobByParsingLegend**

*Arrange grobs by parse their legend.*

**Description**

Arrange grobs and parse their legend, then put it together on the right.

**Usage**

```r
arrangeGrobByParsingLegend(..., nrow = NULL, ncol = NULL,
                             widths = c(4, 1), legend.idx = NULL)
```
Arguments

... ggplot graphics.

nrow number of row for layout.

ncol number of columns for layout

widths width ratio for plot group and legend group.

legend.idx legend index you want to keep.

Value

a

Author(s)

Tengfei Yin

Examples

library(ggplot2)
p1 <- qplot(x = mpg, y = cyl, data = mtcars, color = carb)
p2 <- qplot(x = mpg, y = cyl, data = mtcars, color = wt)
p3 <- qplot(x = mpg, y = cyl, data = mtcars, color = qsec)
p4 <- qplot(x = mpg, y = cyl, data = mtcars, color = gear)
arrangeGrobByParsingLegend(p1, p2, p3, p4)
arrangeGrobByParsingLegend(p1, p2, p3, p4, ncol = 1)
arrangeGrobByParsingLegend(p1, p2, p3, p4, legend.idx = 2)

autoplot

Generic autoplot function

Description

autoplot is a generic function to visualize various data object, it tries to give better default graphics and customized choices for each data type, quick and convenient to explore your genomic data compare to low level ggplot method, it is much simpler and easy to produce fairly complicate graphics, though you may lose some flexibility for each layer.

Usage

## S4 method for signature 'GRanges'
autoplot(object, ..., chr, xlab, ylab, main, truncate.gaps = FALSE, truncate.fun = NULL, ratio = 0.0025, space.skip = 0.1, legend = TRUE, geom = NULL, stat = NULL, chr.weight = NULL, coord = c("default", "genome", "truncate_gaps"), layout = c("linear", "karyogram", "circle"))
## S4 method for signature 'GRangesList'
autoplot(object, ..., xlab, ylab, main, indName = "grl_name",
         geom = NULL, stat = NULL, coverage.col = "gray50",
         coverage.fill = coverage.col, group.selfish = FALSE)

## S4 method for signature 'IRanges'
autoplot(object, ..., xlab, ylab, main)

## S4 method for signature 'Seqinfo'
autoplot(object, ideogram = FALSE, ...)

## S4 method for signature 'GAlignments'
autoplot(object, ..., xlab, ylab, main, which,
         geom = NULL, stat = NULL)

## S4 method for signature 'BamFile'
autoplot(object, ..., which, xlab, ylab, main,
         bsgenome, geom = "line", stat = "coverage", method = c("raw",
         "estimate"), coord = c("linear", "genome"),
         resize.extra = 10, space.skip = 0.1, show.coverage = TRUE)

## S4 method for signature 'character'
autoplot(object, ..., xlab, ylab, main, which)

## S4 method for signature 'TxDbOREnsDb'
autoplot(object, which, ..., xlab, ylab, main, truncate.gaps = FALSE,
         truncate.fun = NULL, ratio = 0.0025,
         mode = c("full", "reduce"), geom = c("alignment"),
         stat = c("identity", "reduce"),
         names.expr = "tx_name", label = TRUE)

## S4 method for signature 'BSgenome'
autoplot(object, which, ..., xlab, ylab, main, geom = NULL)

## S4 method for signature 'Rle'
autoplot(object, ..., xlab, ylab, main, binwidth, nbin = 30,
         geom = NULL, stat = c("bin", "identity", "slice"),
         type = c("viewSums", "viewMins", "viewMaxs", "viewMeans"))

## S4 method for signature 'RleList'
autoplot(object, ..., xlab, ylab, main, nbin = 30, binwidth,
facetByRow = TRUE, stat = c("bin", "identity", "slice"),
geom = NULL, type = c("viewSums", "viewMins", "viewMaxs", "viewMeans")

## S4 method for signature 'matrix'
autoplot(object, ..., xlab, ylab, main,
        geom = c("tile", "raster"), axis.text.angle = NULL,
        hjust = 0.5, na.value = NULL,
        rownames.label = TRUE, colnames.label = TRUE,
        axis.text.x = TRUE, axis.text.y = TRUE)

## S4 method for signature 'ExpressionSet'
autoplot(object, ..., type = c("heatmap", "none",
        "scatterplot.matrix", "pcp", "MA", "boxplot",
        "mean-sd"), test.method =
        "t", rotate = FALSE, pheno.plot = FALSE, main_to_pheno =
        4.5, padding = 0.2)

## S4 method for signature 'RangedSummarizedExperiment'
autoplot(object, ..., type = c("heatmap", "link", "pcp", "boxplot", "scatterplot.matrix"), pheno.plot =
        FALSE, main_to_pheno = 4.5, padding = 0.2, assay.id = 1)

## S4 method for signature 'VCF'
autoplot(object, ...,
        xlab, ylab, main,
        assay.id,
        type = c("default", "geno", "info", "fixed"),
        full.string = FALSE,
        ref.show = TRUE,
        genome.axis = TRUE,
        transpose = TRUE)

## S4 method for signature 'OrganismDb'
autoplot(object, which, ...,
        xlab, ylab, main,
        truncate.gaps = FALSE,
        truncate.fun = NULL,
        ratio = 0.0025,
        geom = c("alignment"),
        stat = c("identity", "reduce"),
        columns = c("TXNAME", "SYMBOL", "TXID", "GENEID"),
        names.expr = "SYMBOL",
        label = TRUE,
        label.color = "gray40")

## S4 method for signature 'VRanges'

autoplot
autoplot

autoplot(object, ..., which = NULL,
    arrow = TRUE, indel.col = "gray30",
    geom = NULL,
    xlab, ylab, main)

## S4 method for signature 'TabixFile'
autoplot(object, which, ...)

Arguments

object          object to be plot.
columns         columns passed to method works for TxDb, EnsDb and OrganismDb.
label.color     when label turned on for gene model, this parameter controls label color.
arrow           arrow passed to geome_alignment function to control intron arrow attributes.
indel.col       indel colors.
ideogram        Weather to call plotIdeogram or not, default is FALSE, if TRUE, layout_karyogram will be called.
transpose       logical value, default TRUE, always make features from VCF as x, so we can use it to map to genomic position.
axis.text.angle axis text angle.
axis.text.x     logical value indicates whether to show x axis and labels or not.
axis.text.y     logical value indicates whether to show y axis and labels or not.
hjust            horizontal just for axis text.
rownames.label  logical value indicates whether to show rownames of matrix as y label or not.
colnames.label  logical value indicates whether to show colnames of matrix as y label or not.
na.value        color for NA value.
rotate          show pheno plot or not.
main_to_pheno   main matrix plot width to pheno plot width ratio.
padding         padding between plots.
assay.id        index for assay you are going to use.
geom            Geom to use (Single character for now). Please see section Geometry for details.
truncate.gaps   logical value indicate to truncate gaps or not.
truncate.fun    shrinkage function. Please see shrinkagefun in package biovizBase.
ratio           used in maxGap.
mode            Display mode for genomic features.
space.skip      space ratio between chromosome spaces in coordinate genome.
coord           Coodinate system.
chr.weight numeric vectors which sum to <1, the names of vectors has to be matched with seqnames in seqinfo, and you can only specify part of the seqnames, other lengths of chromosomes will be assined proportionally to their seqlengths, for example, you could specify chr1 to be 0.5, so the chr1 will take half of the space and other chromosomes squeezed to take left of the space.

legend A logical value indicates whether to show legend or not. Default is TRUE

which A GRanges object to subset the result, usually passed to the ScanBamParam function. For autoplot,EnsDb, which can in addition also be an object extending AnnotationFilter, an AnnotationFilterList combining such objects or a formula representing a filter expression. See examples below or documentation of AnnotationFilter for more details.

show.coverage A logical value indicates whether to show coverage or not. This is used for geom "mismatch.summary".

resize.extra A numeric value used to add buffer to intervals to compute stepping levels on.

bsgenome A BSgenome object. Only need for geom "mismatch.summary".

xlab x label.

ylab y label.

label logic value, default TRUE. To show label by the side of features.

facetByRow A logical value, default is TRUE, facet RleList by row. If FALSE, facet by column.

type For Rle/RleList, "raw" plot everything, so be careful, that would be pretty slow if you have too much data. For "viewMins", "viewMaxs", "viewMeans", "viewSums", require extra arguments to slice the object. so users need to at least provide lower, more details and control please refer the the manual of slice function in IRanges. For "viewMins", "viewMaxs", we use viewWhichMin and viewWhichMax to get x scale, for "viewMeans", "viewSums", we use window midpoint as x.

For ExpressionSet, plotting types.

layout Layout including linear, circular and karyogram. for GenomicRangesList, it only supports circular layout.

method method used for parsing coverage from bam files. 'estimate' use fast esitmated method and 'raw' use relatively slow parsing method.

test.method test method

... Extra parameters. Usually are those parameters used in autoplot to control aesthetics or geometries.

main title.

stat statistical transformation.

indName When coerce GRangesList to GRanges, names created for each group.

coverage.col coverage stroke color.

coverage.fill coverage fill color.

group.selfish Passed to addStepping, control whether to show each group as unique level or not. If set to FALSE, if two groups are not overlapped with each other, they will probably be layout in the same level to save space.
names.expr: names expression used for creating labels. For EnsDb objects either "tx_id", "gene_name" or "gene_id".

binwidth: width of the bins.

nbin: number of bins.

genome.axis: logical value, if TRUE, whenever possible, try to parse genomic position for each column(e.g. RangedSummarizedExperiment), show column as exactly the genomic position instead of showing them side by side and indexed from 1.

full.string: logical value. If TRUE, show full string of indels in plot for VCF.

ref.show: logical value. If TRUE, show REF in VCF at bottom track.

chr: characters indicates the seqnames to be subseted.

Value

A ggplot object, so you can use common features from ggplot2 package to manipulate the plot.

Introduction

autoplot is redefined as generic s4 method inside this package, user could use autoplot in the way they are familiar with, and we are also setting limitation inside this package, like

- scales X scales is always genomic coordinates in most cases, x could be specified as start/end/midpoint when it’s special geoms for interval data like point/line
- colors Try to use default color scheme defined in biovizBase package as possible as it can

Geometry

We have developed new geom for different objects, some of them may require extra parameters you need to provide. Some of the geom are more like geom + stat in ggplot2 package, e.g. "coverage.line" and "coverage.polygon". We simply combine them together, but in the future, we plan to make it more general.

This package is designed for only biological data, especially genomic data if users want to explore the data in a more flexible way, you could simply coerce the GRanges to a data.frame, then just use formal autoplot function in ggplot2, or autoplot generic for data.frame.

Some objects share the same geom so we introduce all the geom together in this section

Showing all the intervals as stepped rectangle, colored by strand automatically.
For TxDb or EnsDb objects, showing full model.

segment: Showing all the intervals as stepped segments, colored by strand automatically.

For object BSgenome, show nucleotides as colored segment.
For Rle/RleList, show histogram-like segments.

line: Showing interval as line, the interval data could also be just single position when start = end,
x is one of start/end/midpoint, y value is unquoted name in elementMetadata column names.
y value is required.

point: Showing interval as point, the interval data could also be just single position when start = end,
x is one of start/end/midpoint, y value is unquoted name in elementMetadata column names.
y value is required.
For object BSgenome, show nucleotides as colored point.
**coverage.line** Coverage showing as lines for interval data.

**coverage.polygon** Coverage showing as polygon for interval data.

**splice** Splicing summary. The size and width of the line and rectangle should represent the counts in each model. Need to provide model.

**single** For TxDb or EnsDb objects, showing single(reduced) model only.

**tx** For TxDb or EnsDb objects, showing transcripts isoforms.

**mismatch.summary** Showing color coded mismatched stacked bar to indicate the proportion of mismatching at each position, the reference is set to gray.

**text** For object BSgenome, show nucleotides as colored text.

**rectangle** For object BSgenome, show nucleotides as colored rectangle.

### Faceting

Faceting in ggbio package is a little different from ggplot2 in several ways

- The faceted column could only be seqnames or regions on the genome. So we limited the formula passing to facet argument, e.g something \~ seqnames, is accepted formula, you can change "something" to variable name in the elementMetadata. But you can not change the second part.

- Sometime, we need to view different regions, so we also have a facet_gr argument which accept a GRanges. If this is provided, it will override the default seqnames and use provided region to facet the graphics, this might be useful for different gene centric views.

### Author(s)

Tengfei Yin

### Examples

```r
set.seed(1)
N <- 1000
library(GenomicRanges)
gr <- GRanges(seqnames =
sample(c("chr1", "chr2", "chr3"),
      size = N, replace = TRUE),
  IRanges(
    start = sample(1:300, size = N, replace = TRUE),
    width = sample(70:75, size = N,replace = TRUE)),
  strand = sample(c("+", "-", "*"), size = N,
    replace = TRUE),
  value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
  sample = sample(c("Normal", "Tumor"),
    size = N, replace = TRUE),
  pair = sample(letters, size = N,
    replace = TRUE))

idx <- sample(1:length(gr), size = 50)
```
### code chunk number 3: default

```r
autoplot(gr[idx])
```

### code chunk number 4: bar-default-pre

```r
set.seed(123)
gr.b <- GRanges(seqnames = "chr1", IRanges(start = seq(1, 100, by = 10),
                width = sample(4:9, size = 10, replace = TRUE),
                score = rnorm(10, 10, 3), value = runif(10, 1, 100))
gr.b2 <- GRanges(seqnames = "chr2", IRanges(start = seq(1, 100, by = 10),
                width = sample(4:9, size = 10, replace = TRUE),
                score = rnorm(10, 10, 3), value = runif(10, 1, 100))
```

```r
gr.b <- c(gr.b, gr.b2)
```

```r
head(gr.b)
```

### code chunk number 5: bar-default

```r
p1 <- autoplot(gr.b, geom = "bar")
## use value to fill the bar
p2 <- autoplot(gr.b, geom = "bar", aes(fill = value))
tracks(default = p1, fill = p2)
```

### code chunk number 6: autoplot.Rnw:236-237

```r
autoplot(gr[idx], geom = "arch", aes(color = value), facets = sample ~ seqnames)
```

### code chunk number 7: gr-group

```r
gra <- GRanges("chr1", IRanges(c(1,7,20), end = c(4,9,30), group = c("a", "a", "b"))
## if you desn't specify group, then group based on stepping levels, and gaps are computed without
## considering extra group method
p1 <- autoplot(gra, aes(fill = group), geom = "alignment")
## when use group method, gaps only computed for grouped intervals.
## default is group.selfish = TRUE, each group keep one row.
## in this way, group labels could be shown as y axis.
p2 <- autoplot(gra, aes(fill = group, group = group), geom = "alignment")
## group.selfish = FALSE, save space
p3 <- autoplot(gra, aes(fill = group, group = group), geom = "alignment", group.selfish = FALSE)
tracks('non-group' = p1,'group.selfish = TRUE' = p2, 'group.selfish = FALSE' = p3)
```

### code chunk number 8: gr-facet-strand
autoplot(gr, stat = "coverage", geom = "area", facets = strand ~ seqnames, aes(fill = strand))

### code chunk number 9: gr-autoplot-circle
autoplot(gr[idx], layout = 'circle')

### code chunk number 10: gr-circle
seqlengths(gr) <- c(400, 500, 700)
values(gr)$to.gr <- gr[sample(1:length(gr), size = length(gr))]
idx <- sample(1:length(gr), size = 50)
gr <- gr[idx]
ggplot() + layout_circle(gr, geom = "ideo", fill = "gray70", radius = 7, trackWidth = 3) +
layout_circle(gr, geom = "bar", radius = 10, trackWidth = 4,
aes(fill = score, y = score)) +
layout_circle(gr, geom = "point", color = "red", radius = 14,
trackWidth = 3, grid = TRUE, aes(y = score)) +
layout_circle(gr, geom = "link", linked.to = "to.gr", radius = 6, trackWidth = 1)

### code chunk number 11: seqinfo-src
data(hg19Ideogram, package = "biovizBase")
seq <- seqinfo(hg19Ideogram)
seq

### code chunk number 12: seqinfo
autoplot(sq[paste0("chr", c(1:22, "X"))])

### code chunk number 13: ir-load
set.seed(1)
N <- 100
ir <- IRanges(start = sample(1:300, size = N, replace = TRUE),
width = sample(70:75, size = N, replace = TRUE))
## add meta data
df <- DataFrame(value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
sample = sample(c("Normal", "Tumor"),
size = N, replace = TRUE),
pair = sample(letters, size = N,
replace = TRUE))
values(ir) <- df
ir

# code chunk number 14: ir-exp
p1 <- autoplot(ir)
p2 <- autoplot(ir, aes(fill = pair)) + theme(legend.position = "none")
p3 <- autoplot(ir, stat = "coverage", geom = "line", facets = sample ~ .)
p4 <- autoplot(ir, stat = "reduce")
tracks(p1, p2, p3, p4)

# code chunk number 15: grl-simul
set.seed(1)
N <- 100
## simulated GRanges
gr <- GRanges(seqnames = sample(c("chr1", "chr2", "chr3"), size = N, replace = TRUE),
  IRanges(  
    start = sample(1:300, size = N, replace = TRUE),
    width = sample(30:40, size = N, replace = TRUE),
    strand = sample(c("+", "-", "*"), size = N, replace = TRUE),
    value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
    sample = sample(c("Normal", "Tumor"), size = N, replace = TRUE),
    pair = sample(letters, size = N, replace = TRUE))

grl <- split(gr, values(gr)$pair)

# code chunk number 16: grl-exp
p1 <- autoplot(grl, group.selfish = TRUE)
p2 <- autoplot(grl, group.selfish = TRUE, main.geom = "arrowrect", gap.geom = "segment")
tracks(p1, p2)

# code chunk number 17: grl-name
autoplot(grl, aes(fill = ..grl_name..))
## equal to
## autoplot(grl, aes(fill = grl_name))

```r
# code chunk number 18: rle-simul
library(IRanges)
set.seed(1)
lambda <- c(rep(0.001, 4500), seq(0.001, 10, length = 500),
            seq(10, 0.001, length = 500))

xVector <- rpois(1e4, lambda)
xRle <- Rle(xVector)
xRle
```

```r
# code chunk number 19: rle-bin
p1 <- autoplot(xRle)
p2 <- autoplot(xRle, nbin = 80)
p3 <- autoplot(xRle, geom = "heatmap", nbin = 200)
tracks('nbin = 30' = p1, "nbin = 80" = p2, "nbin = 200(heatmap)" = p3)
```

```r
# code chunk number 20: rle-id
p1 <- autoplot(xRle, stat = "identity")
p2 <- autoplot(xRle, stat = "identity", geom = "point", color = "red")
tracks('line' = p1, "point" = p2)
```

```r
# code chunk number 21: rle-slice
p1 <- autoplot(xRle, type = "viewMaxs", stat = "slice", lower = 5)
p2 <- autoplot(xRle, type = "viewMaxs", stat = "slice", lower = 5, geom = "heatmap")
tracks('bar' = p1, "heatmap" = p2)
```

```r
# code chunk number 22: rlel-simul
xRleList <- RleList(xRle, 2L * xRle)
xRleList
```

```r
# code chunk number 23: rlel-bin
```
p1 <- autoplot(xRleList)
p2 <- autoplot(xRleList, nbin = 80)
p3 <- autoplot(xRleList, geom = "heatmap", nbin = 200)
tracks('nbin = 30' = p1, "nbin = 80" = p2, "nbin = 200(heatmap)" = p3)

### code chunk number 24: rlel-id
### code chunk number 25: rlel-slice
### code chunk number 26: txdb
### code chunk number 27: txdb-visual
### EnsDb
## Fetching gene models from an EnsDb object.
library(EnsDb.Hsapiens.v75)
ensdb <- EnsDb.Hsapiens.v75
## We use a GenenameFilter to specifically retrieve all transcripts for that gene.
p1 <- autoplot(ensdb, which = GeneNameFilter("ALDOA"), names.expr = "tx_name:::gene_id")
## Instead of providing the GenenameFilter, we can also use filter expressions
p2 <- autoplot(ensdb, which = ~ genename == "ALDOA", stat = "reduce", color = "brown", fill = "brown")
tracks(full = p1, reduce = p2, heights = c(5, 1)) + ylab("")

## Alternatively, we can specify a GRangesFilter and display all genes
## that are (partially) overlapping with that genomic region:

```r
gr <- GRanges(seqnames=16, IRanges(30768000, 30770000), strand="+")
autoplot(ensdb, GRangesFilter(gr, "any"), names.expr="gene_name")
```

## Just submitting the GRanges object also works.

```r
autoplot(ensdb, gr, names.expr="gene_name")
```

## Or genes encoded on both strands.

```r
gr <- GRanges(seqnames = 16, IRanges(30768000, 30770000), strand = "+")
autoplot(ensdb, GRangesFilter(gr), names.expr="gene_name")
```

## Also, we can specify directly the gene ids and plot all transcripts of these
## genes (not only those overlapping with the region)

```r
autoplot(ensdb, GeneIdFilter(c("ENSG00000196118", "ENSG00000156873")))
```

### code chunk number 28: ga-load

```r
library(GenomicAlignments)
data("genesymbol", package = "biovizBase")
bamfile <- system.file("extdata", "SRR027894subRBM17.bam", package="biovizBase")
```

```r
which <- keepStandardChromosomes(genesymbol["RBM17"])
```

## need to set use.names = TRUE

```r
ga <- readGAlignments(bamfile,
   param = ScanBamParam(which = which),
   use.names = TRUE)
```

```r
```

### code chunk number 29: ga-exp

```r
p1 <- autoplot(ga)
p2 <- autoplot(ga, geom = "rect")
p3 <- autoplot(ga, geom = "line", stat = "coverage")
```

```r
tracks(default = p1, rect = p2, coverage = p3)
```

### code chunk number 30: bf-load (eval = FALSE)

```r
## library(Rsamtools)
## bamfile <- "./wgEncodeCaltechRnaSeqK562R1x75dAlignsRep1V2.bam"
## bf <- BamFile(bamfile)
```

### code chunk number 31: bf-est-cov (eval = FALSE)

```r
## autoplot(bamfile)
## autoplot(bamfile, which = c("chr1", "chr2"))
## autoplot(bf)
## autoplot(bf, which = c("chr1", "chr2"))
```
```
# data(genesymbol, package = "biovizBase")
# autoplot(bamfile, method = "raw", which = genesymbol["ALDOA"])
#
# library(BSgenome.Hsapiens.UCSC.hg19)
# autoplot(bf, stat = "mismatch", which = genesymbol["ALDOA"], bsgenome = Hsapiens)

###################################################
### code chunk number 32: char-bam (eval = FALSE)
###################################################
# bamfile <- "/wgEncodeCaltechRnaSeqK562R1x75dAlignsRep1V2.bam"
# autoplot(bamfile)

###################################################
### code chunk number 33: char-gr
###################################################
library(rtracklayer)

test_path <- system.file("tests", package = "rtracklayer")
test_bed <- file.path(test_path, "test.bed")

autoplot(test_bed, aes(fill = name))

###################################################
### matrix
###################################################
volcano <- volcano[20:70, 20:60] - 150

autoplot(volcano)

# special scale theme for 0-centered values

autoplot(volcano, geom = "raster") + scale_fill_fold_change()

# when a matrix has colnames and rownames label them by default

colnames(volcano) <- sort(sample(1:300, size = ncol(volcano), replace = FALSE))

autoplot(volcano) + scale_fill_fold_change()

rownames(volcano) <- letters[sample(1:24, size = nrow(volcano), replace = TRUE)]

autoplot(volcano)

# even with row/col names, you could also disable it and just use numeric index

autoplot(volcano, colnames.label = FALSE)

autoplot(volcano, rownames.label = FALSE, colnames.label = FALSE)

# don't want the axis has label??

autoplot(volcano, axis.text.x = FALSE)

autoplot(volcano, axis.text.y = FALSE)

# or totally remove axis

colnames(volcano) <- lapply(letters[sample(1:24, size = ncol(volcano), replace = TRUE)],

function(x){
    paste(rep(x, 7), collapse = "")
}
```
autoplot

## Oops, overlapped
autoplot(volcano)
## tweak with it.
autoplot(volcano, axis.text.angle = -45, hjust = 0)

## when character is the value
x <- sample(c(letters[1:3], NA), size = 100, replace = TRUE)
mx <- matrix(x, nrow = 5)
autoplot(mx)
## tile gives you a white margin
rownames(mx) <- LETTERS[1:5]
autoplot(mx, color = "white")
colnames(mx) <- LETTERS[1:20]
autoplot(mx, color = "white")
autoplot(mx, color = "white", size = 2)
## weird in aes(), though works
## default tile is flexible
autoplot(mx, aes(width = 0.6, height = 0.6))
autoplot(mx, aes(width = 0.6, height = 0.6), na.value = "white")
autoplot(mx, aes(width = 0.6, height = 0.6)) + theme_clear()

### Views

lambda <- c(rep(0.001, 4500), seq(0.001, 10, length = 500),
seq(10, 0.001, length = 500))
xVector <- dnorm(1:5e3, mean = 1e3, sd = 200)
xRle <- Rle(xVector)
v1 <- Views(xRle, start = sample(.4e3:.6e3, size = 50, replace = FALSE), width =1000)
autoplot(v1)
names(v1) <- letters[sample(1:24, size = length(v1), replace = TRUE)]
autoplot(v1)
autoplot(v1, geom = "tile", aes(width = 0.5, height = 0.5))
autoplot(v1, geom = "line")
autoplot(v1, geom = "line", aes(color = row)) + theme(legend.position = "none")
autoplot(v1, geom = "line", facets = NULL)
autoplot(v1, geom = "line", facets = NULL, alpha = 0.1)

### ExpressionSet

library(Biobase)
data(sample.ExpressionSet)
sample.ExpressionSet
set.seed(1)
## select 50 features
idx <- sample(seq_len(dim(sample.ExpressionSet)[1]), size = 50)
eset <- sample.ExpressionSet[idx,]
eset
autoplot(as.matrix(pData(eset)))
## default heatmap
p1 <- autoplot(eset)
p2 <- p1 + scale_fill_fold_change()
p2
autoplot(eset)
autoplot(eset, geom = "tile", color = "white", size = 2)
autoplot(eset, geom = "tile", aes(width = 0.6, height = 0.6))

autoplot(eset, pheno.plot = TRUE)
idx <- order(pData(eset)[,1])
eset2 <- eset[,idx]
autoplot(eset2, pheno.plot = TRUE)

## parallel coordainte plot
autoplot(eset, type = "pcp")

## boxplot
autoplot(eset, type = "boxplot")

## scatterplot.matrix
## slow, be carefull
##autoplot(eset[, 1:7], type = "scatterplot.matrix")

## mean-sd
autoplot(eset, type = "mean-sd")

########################################################################
### RangedSummarizedExperiment
########################################################################
library(SummarizedExperiment)

nrows <- 200; ncols <- 6
counts <- matrix(runif(nrows * ncols, 1, 1e4), nrows)
counts2 <- matrix(runif(nrows * ncols, 1, 1e4), nrows)
rowRanges <- GRanges(rep(c("chr1", "chr2"), c(50, 150)),
                      IRanges(floor(runif(200, 1e5, 1e6)), width=100),
                      strand=sample(c("+", "-"), 200, TRUE))
colData <- DataFrame(Treatment=rep(c("ChIP", "Input"), 3),
                      row.names=LETTERS[1:6])
sset <- SummarizedExperiment(assays=SimpleList(counts=counts, counts2 = counts2),
                             rowRanges=rowRanges, colData=colData)
autoplot(sset) + scale_fill_fold_change()
autoplot(sset, pheno.plot = TRUE)

########################################################################
### pcp
########################################################################
autoplot(sset, type = "pcp")

###################################################
### boxplot
###################################################
autoplot(sset, type = "boxplot")

###################################################
### scatterplot matrix
###################################################
##autoplot(sset, type = "scatterplot.matrix")

###################################################
### vcf
###################################################
## Not run:
library(VariantAnnotation)
vcffile <- system.file("extdata", "chr22.vcf.gz", package="VariantAnnotation")
vcf <- readVcf(vcffile, "hg19")
## default use type 'geno'
## default use genome position
autoplot(vcf)
## or disable it
autoplot(vcf, genome.axis = FALSE)
## not transpose
autoplot(vcf, genome.axis = FALSE, transpose = FALSE, rownames.label = FALSE)
autoplot(vcf)
## use
autoplot(vcf, assay.id = "DS")
## equivalent to
autoplot(vcf, assay.id = 2)
## doesn't work when assay.id cannot find
autoplot(vcf, assay.id = "NO")
## use AF or first
autoplot(vcf, type = "info")
## geom bar
autoplot(vcf, type = "info", aes(y = THETA))
autoplot(vcf, type = "info", aes(y = THETA, fill = VT, color = VT))
autoplot(vcf, type = "fixed")
autoplot(vcf, type = "fixed", size = 10) + xlim(c(50310860, 50310890)) + ylim(0.75, 1.25)
p1 <- autoplot(vcf, type = "fixed") + xlim(50310860, 50310890)
p2 <- autoplot(vcf, type = "fixed", full.string = TRUE) + xlim(50310860, 50310890)
tracks("full.string = FALSE" = p1, "full.string = TRUE" = p2)+
  scale_y_continuous(breaks = NULL, limits = c(0, 3))
p3 <- autoplot(vcf, type = "fixed", ref.show = FALSE) + xlim(50310860, 50310890) +
  scale_y_continuous(breaks = NULL, limits = c(0, 2))
p3
library(BSgenome.Hsapiens.UCSC.hg19)
data(genesymbol, package = "biovizBase")
p1 <- autoplot(Hsapiens, which = resize(genesymbol["ALDOA"], width = 50))
p2 <- autoplot(Hsapiens, which = resize(genesymbol["ALDOA"], width = 50), geom = "rect")
tracks(text = p1, rect = p2)

sessionInfo()

---

**geom_alignment**

Alignment geoms for GRanges object

**Description**

Show interval data as alignment.

**Usage**

```
## S4 method for signature 'GRanges'
geom_alignment(data, ..., xlab, ylab, main, facets = NULL, stat =
c("stepping", "identity"), range.geom = c("rect",
"arrowrect"), gap.geom = c("chevron", "arrow",
"segment"), rect.height = NULL, group.selfish = TRUE,
label = TRUE)

## S4 method for signature 'TxDbOREnsDb'
geom_alignment(data, ..., which, columns = c("tx_id", "tx_name",
"gene_id"), names.expr = "tx_name", facets = NULL,
truncate.gaps = FALSE, truncate.fun = NULL, ratio =
0.0025)

## S4 method for signature 'GRangesList'
geom_alignment(data, ..., which = NULL,
cds.rect.h = 0.25,
exon.rect.h = cds.rect.h,
utr.rect.h = cds.rect.h/2,
xlab, ylab, main,
facets = NULL, geom = "alignment",
```

```---

```
stat = c("identity", "reduce"),
range.geom = "rect",
gap.geom = "arrow",
utr.geom = "rect",
names.expr = NULL,
label = TRUE,
label.color = "gray40",
label.size = 3,
arrow.rate = 0.015,
length = unit(0.1, "cm"))

## S4 method for signature 'OrganismDb'
geom_alignment(data, ..., which,
columns = c("TXNAME", "SYMBOL", "TXID", "GENEID"),
names.expr = "SYMBOL",
facets = NULL,
truncate.gaps = FALSE,
truncate.fun = NULL, ratio = 0.0025
)

Arguments

data A GRanges, data.frame, TxDb or EnsDb object.
...
Extra parameters such as aes() passed.
which GRanges object to subset the TxDb or EnsDb object. For EnsDb: can also be a
single object extending AnnotationFilter, an AnnotationFilterList combining such objects or a filter expression in form of a formula.
cds.rect.h cds heights.
exon.rect.h exon heights.
utr.rect.h utr heights.
label.color label color.
label.size label size.
arrow.rate arrow rate.
length arrow length.
columns columns to get from object.
xlab Label for x
ylab Label for y
main Title for plot.
facets Faceting formula to use.
stat For GRanges: Character vector specifying statistics to use. "stepping" with randomly assigned stepping levels as y variable. "identity" allow users to specify y value in aes.
For TxDb: default "identity" give full gene model and "reduce" for reduced model.
Geom for 'gap' computed from the data you passed based on the group information.

gap.geom

rect.height

Half height of the arrow body.

group.selfish

Passed to addStepping, control whether to show each group as unique level or not. If set to FALSE, if two groups are not overlapped with each other, they will probably be layout in the same level to save space.

truncate.gaps

logical value indicate to truncate gaps or not.

truncate.fun

shrinkage function. Please see shrinkagefun in package biovizBase.

ratio

used in maxGap.

group

geometric object. only support "gene" now.

range.geom

geom for main intervals or exons.

utr.geom

geom for utr region.

names.expr

expression for showing y label.

label

logical value. Whether to label the intervals with names specified by argument names.expr.

Value

A 'Layer'.

Author(s)

Tengfei Yin

Examples

set.seed(1)
N <- 100
require(GenomicRanges)

## simmulated GRanges

gr <- GRanges(seqnames =
sample(c("chr1", "chr2", "chr3"),
    size = N, replace = TRUE),
IRanges(
    start = sample(1:300, size = N, replace = TRUE),
    width = sample(70:75, size = N,replace = TRUE)),
strand = sample(c("+", "-", "x"), size = N,
    replace = TRUE),
value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
sample = sample(c("Normal", "Tumor"),
    size = N, replace = TRUE),
pair = sample(letters, size = N,
    replace = TRUE))

## simmulated GRanges
Arch geoms for GRanges object

Description

Show interval data as arches.
Usage

## S4 method for signature 'data.frame'
geom_arch(data, ..., n = 25, max.height = 10)

## S4 method for signature 'GRanges'
geom_arch(data, ..., xlab, ylab, main, facets = NULL,
            rect.height = 0, n = 25, max.height = 10)

Arguments

data A GRanges or data.frame object.

... Extra parameters passed to autoplot function, aes mapping support height, x, xend.

  • xstart of the arches
  • xend of the arches
  • height of arches

xlab Label for x

ylab Label for y

main Title for plot.

n Integer values at which interpolation takes place to create 'n' equally spaced points spanning the interval ['min(x)', 'max(x)'].

facets Faceting formula to use.

rect.height When data is GRanges, this padding the arches from original y value to allow users putting arches 'around' the interval rectangles.

max.height Max height of all arches.

Details

To draw a interval data as arches, we need to provide a special geom for this purpose. Arches is popular in gene viewer or genomoe browser, when they try to show isoforms or gene model. `geom_arch`, just like any other `geom_*` function in ggplot2, you can pass aes() to it to map variable to height of arches.

Value

A 'Layer'.

Author(s)

Tengfei Yin
Examples

```r
set.seed(1)
N <- 100
library(GenomicRanges)

## simmulated GRanges
gr <- GRanges(seqnames = sample(c("chr1", "chr2", "chr3"),
                      size = N, replace = TRUE),
                      IRanges(
                        start = sample(1:300, size = N, replace = TRUE),
                        width = sample(70:75, size = N, replace = TRUE),
                        strand = sample(c("+", "-", "\*"), size = N,
                            replace = TRUE),
                        value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
                        sample = sample(c("Normal", "Tumor"),
                            size = N, replace = TRUE),
                        pair = sample(letters, size = N,
                            replace = TRUE))

## default
ggplot(gr) + geom_arch()
# or
ggplot() + geom_arch(gr)

## facetting and aesthetics
ggplot(gr) + geom_arch(aes(color = value, height = value, size = value),
                        alpha = 0.2, facets = sample ~ seqnames)
```

**geom_arrow**  
Arrow geoms for GRanges object

**Description**

Show interval data as arrows.

**Usage**

```r
## S4 method for signature 'GRanges'
geom_arrow(data, ..., xlab, ylab, main, 
            angle = 30, length = unit(0.12, "cm"), type = "open",
```
geom_arrow

stat = c("stepping", "identity"), facets = NULL,
arrow.rate = 0.03, group.selfish = TRUE)

Arguments

data A GRanges object.

... Extra parameters such as aes() passed.
xlab Label for x

ylab Label for y

main Title for plot.

angle The angle of the arrow head in degrees (smaller numbers produce narrower, pointier arrows). Essentially describes the width of the arrow head.

length A unit specifying the length of the arrow head (from tip to base).

type One of "open" or "closed" indicating whether the arrow head should be a closed triangle.

stat Character vector specifying statistics to use. "stepping" with randomly assigned stepping levels as y variable. "identity" allow users to specify y value in aes.

facets Faceting formula to use.

arrow.rate Arrow density of the arrow body.

group.selfish Passed to addStepping, control whether to show each group as unique level or not. If set to FALSE, if two groups are not overlapped with each other, they will probably be layout in the same level to save space.

Value

A 'Layer'.

Author(s)

Tengfei Yin

Examples

set.seed(1)
N <- 100
require(GenomicRanges)
## ======================================================================
## simmulated GRanges
## ======================================================================
gr <- GRanges(seqnames =
sample(c("chr1", "chr2", "chr3"),
        size = N, replace = TRUE),
IRanges(
    start = sample(1:300, size = N, replace = TRUE),
    width = sample(70:75, size = N,replace = TRUE)),
strand = sample(c("+", "-", "*"), size = N,
```r
replace = TRUE),
value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
sample = sample(c("Normal", "Tumor"),
size = N, replace = TRUE),
pair = sample(letters, size = N,
replace = TRUE))
```
**Description**

Show interval data as rectangle with a arrow head.

**Usage**

```r
## S4 method for signature 'GRanges'
geom_arrowrect(data, ..., xlab, ylab, main,
               facets = NULL, stat = c("stepping", "identity"),
               rect.height = NULL, arrow.head = 0.06,
               arrow.head.rate = arrow.head, arrow.head.fix = NULL,
               group.selfish = TRUE)
```

**Arguments**

- `data`: A GRanges object.
- `...`: Extra parameters such as aes() passed.
- `xlab`: Label for x
- `ylab`: Label for y
- `main`: Title for plot.
- `facets`: Faceting formula to use.
- `stat`: Character vector specifying statistics to use. "stepping" with randomly assigned stepping levels as y variable. "identity" allow users to specify y value in aes.
- `rect.height`: Half height of the arrow body.
- `arrow.head`: Arrow head to body ratio.
- `arrow.head.rate`: Arrow head to body ratio. same with arrow.head.
- `arrow.head.fix`: fixed length of arrow head.
- `group.selfish`: Passed to addStepping, control whether to show each group as unique level or not. If set to FALSE, if two groups are not overlapped with each other, they will probably be layout in the same level to save space.

**Value**

A 'Layer'.

**Author(s)**

Tengfei Yin

**Examples**

```r
set.seed(1)
N <- 100
require(GenomicRanges)
```
## simmulated GRanges
```
gr <- GRanges(seqnames =
sample(c("chr1", "chr2", "chr3"),
    size = N, replace = TRUE),
IRanges(
    start = sample(1:300, size = N, replace = TRUE),
    width = sample(70:75, size = N, replace = TRUE)),
strand = sample(c("+", "-", "x"), size = N,
    replace = TRUE),
value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
sample = sample(c("Normal", "Tumor"),
    size = N, replace = TRUE),
pair = sample(letters, size = N,
    replace = TRUE))
```

## default
```
ggplot(gr) + geom_arrowrect()
```

## or
```
ggplot() + geom_arrowrect(gr)
```

## facetting and aesthetics
```
ggplot(gr) + geom_arrowrect(facets = sample ~ seqnames, aes(color = strand, fill = strand))
```

## stat:identity
```
ggplot(gr) + geom_arrowrect(stat = "identity", aes(y = value))
```

## stat:stepping
```
ggplot(gr) + geom_arrowrect(stat = "stepping", aes(y = value, group = pair))
```

## group.selfish controls when
```
ggplot(gr) + geom_arrowrect(gr, stat = "stepping", aes(y = value, group = pair), group.selfish = FALSE)
```
**Description**

Show interval data as vertical bar, width equals to interval width and use 'score' or specified 'y' as y scale.

**Usage**

```r
## S4 method for signature 'ANY'
geom_bar(data, ...)
## S4 method for signature 'GRanges'
geom_bar(data, ..., xlab, ylab, main)
```

**Arguments**

- `data` Typically a GRanges or data.frame object.
- `...` Extra parameters such as aes() or color, size passed.
- `xlab` Label for x
- `ylab` Label for y
- `main` Title for plot.

**Details**

Useful for showing bed like files, when imported as GRanges, have a extra 'score' column, use it as default y, you could also specify y by using aes(y = ).

**Value**

A 'Layer'.

**Examples**

```r
## load
library(GenomicRanges)

## simul
set.seed(123)
gr.b <- GRanges(seqnames = "chr1", IRanges(start = seq(1, 100, by = 10),
        width = sample(4:9, size = 10, replace = TRUE),
        score = rnorm(10, 10, 3), value = runif(10, 1, 100))
gr.b2 <- GRanges(seqnames = "chr2", IRanges(start = seq(1, 100, by = 10),
        width = sample(4:9, size = 10, replace = TRUE),
        score = rnorm(10, 10, 3), value = runif(10, 1, 100))
gr.b <- c(gr.b, gr.b2)
## default use score as y

## bar
ggplot(gr.b) + geom_bar(aes(fill = value))
## or
ggplot() + geom_bar(gr.b, aes(fill = value))
ggplot(gr.b) + geom_bar(aes(y = value))
```
## geom_chevron

- **Description**
  
  Break normal intervals stored in GRanges object and show them as chevron, useful for showing model or splice summary.

- **Usage**
  
  ```r
  # S4 method for signature 'GRanges'
  geom_chevron(data, ..., xlab, ylab, main,
  offset = 0.1,
  facets = NULL,
  stat = c("stepping", "identity"),
  chevron.height.rescale = c(0.1, 0.8),
  group.selfish = TRUE)
  ```

- **Arguments**
  
  - **data**: A GRanges object.
  - **...**: Extra parameters passed to autoplot function.
  - **xlab**: Label for x
  - **ylab**: Label for y
  - **main**: Title for plot.
  - **offset**: A numeric value or characters. If it's numeric value, indicate how much you want the chevron to wiggle, usually the rectangle for drawing GRanges is of height unit 1, so it's better between -0.5 and 0.5 to make it nice looking. Unless you specify offset as one of those columns, this will use height of the chevron to indicate the columns. Of course you could use size of the chevron to indicate the column variable easily, please see the examples.
  - **facets**: Faceting formula to use.
  - **stat**: Character vector specifying statistics to use. "stepping" with randomly assigned stepping levels as y variable. "identity" allow users to specify y value in `aes`.
  - **chevron.height.rescale**: A numeric vector of length 2. When the offset parameters is a character which is one of the data columns, this parameter rescale the offset.
  - **group.selfish**: Passed to addStepping, control whether to show each group as unique level or not. If set to FALSE, if two groups are not overlapped with each other, they will probably be layout in the same level to save space.
Details

To draw a normal GRanges as Chevron, we need to provide a special geom for this purpose. Chevron is popular in gene viewer or genomoe browser, when they try to show isoforms or gene model. geom_chevron, just like any other geom_* function in ggplot2, you can pass aes() to it to use height of chevron or width of chevron to show statistics summary.

Value

A `Layer`.

Author(s)

Tengfei Yin

Examples

```r
set.seed(1)
N <- 100
require(GenomicRanges)

## simmulated GRanges
gr <- GRanges(seqnames =
  sample(c("chr1", "chr2", "chr3"),
         size = N, replace = TRUE),
  IRanges(
    start = sample(1:300, size = N, replace = TRUE),
    width = sample(70:75, size = N, replace = TRUE),
    strand = sample(c("+", "-", "*"), size = N,
                    replace = TRUE),
    value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
    sample = sample(c("Normal", "Tumor"), size = N,
                    replace = TRUE),
    pair = sample(letters, size = N,
                  replace = TRUE))

## default

ggplot(gr) + geom_chevron()
## or
ggplot() + geom_chevron(gr)

## facetting and aesthetics

ggplot(gr) + geom_chevron(facets = sample ~ seqnames, aes(color = strand))
```
## stat:identity
```
ggplot(gr) + geom_chevron(stat = "identity", aes(y = value))
```

## stat:stepping
```
ggplot(gr) + geom_chevron(stat = "stepping", aes(group = pair))
```

## group.selfish controls when
```
ggplot(gr) + geom_chevron(stat = "stepping", aes(group = pair), group.selfish = FALSE, 
                           xlab = "xlab", ylab = "ylab", main = "main")
```

```
p <- qplot(x = mpg, y = cyl, data = mtcars)
```

## offset
```
gr2 <- GRanges("chr1", IRanges(c(1, 10, 20), width = 5))
gr2.p <- gaps(gr2)
# resize to connect them
gr2.p <- resize(gr2.p, fix = "center", width = width(gr2.p)+2)

```
ggplot(gr2) + geom_rect() + geom_chevron(gr2.p)
```

## notice the rectangle height is 0.8
```
# offset = 0 just like a line
ggplot(gr2) + geom_rect() + geom_chevron(gr2.p, offset = 0)
```

## equal height
```
# ggplot(gr2) + geom_rect() + geom_chevron(gr2.p, offset = 0.4)
```

## chevron.height
```
values(gr2.p)$score <- c(100, 200)
ggplot(gr2) + geom_rect() + geom_chevron(gr2.p, offset = "score")
```

```
# chevron.height
# ggplot(gr2) + geom_rect() + geom_chevron(gr2.p, offset = "score",
# chevron.height.rescale = c(0.4, 10))
```
Description

Show interval data as rectangle.

Usage

```r
## S4 method for signature 'ANY'
geom_rect(data, ...)
## S4 method for signature 'GRanges'
geom_rect(data, ..., xlab, ylab, main,
facets = NULL, stat = c("stepping", "identity"),
rect.height = NULL,
group.selfish = TRUE)
```

Arguments

- `data` Typically a GRanges or data.frame object. When it’s data.frame, it’s simply calling ggplot2::geom_rect.
- `...` Extra parameters such as aes() or color, size passed.
- `xlab` Label for x
- `ylab` Label for y
- `main` Title for plot.
- `facets` Faceting formula to use.
- `stat` Character vector specifying statistics to use. "stepping" with randomly assigned stepping levels as y variable. "identity" allow users to specify y value in aes.
- `rect.height` Half height of the arrow body.
- `group.selfish` Passed to addStepping, control whether to show each group as unique level or not. If set to FALSE, if two groups are not overlapped with each other, they will probably be layout in the same level to save space.

Value

A 'Layer'.

Author(s)

Tengfei Yin
Examples

```r
set.seed(1)
N <- 100
require(GenomicRanges)

## simmulated GRanges
gr <- GRanges(seqnames = 
  sample(c("chr1", "chr2", "chr3"),
    size = N, replace = TRUE),
  IRanges(
    start = sample(1:300, size = N, replace = TRUE),
    width = sample(70:75, size = N,replace = TRUE)),
  strand = sample(c("+", ",-", "*") , size = N,
    replace = TRUE),
  value = rnorm(N, 10, 3),
  score = rnorm(N, 100, 30),
  sample = sample(c("Normal", "Tumor"),
    size = N, replace = TRUE),
  pair = sample(letters, size = N,
    replace = TRUE))

## data.frame call ggplot2::geom_rect
ggplot() + geom_rect(data = mtcars, aes(xmin = mpg, ymin = wt, xmax = mpg + 10, ymax = wt + 0.2,
    fill = cyl))

## default
ggplot(gr) + geom_rect()
# or
ggplot() + geom_rect(gr)

## facetting and aesthetics
ggplot(gr) + geom_rect(facets = sample ~ seqnames, aes(color = strand, fill = strand))

## stat:identity
ggplot(gr) + geom_rect(stat = "identity", aes(y = value))
```
## geom_segment

Segment geoms for GRanges object

### Description

Show interval data as segments.

### Usage

```r
## S4 method for signature 'ANY'
geom_segment(data, ...)

## S4 method for signature 'GRanges'
geom_segment(data, ..., xlab, ylab, main,
             facets = NULL, stat = c("stepping", "identity"),
             group.selfish = TRUE)
```

### Arguments

- **data**: A GRanges or data.frame object.
- **...**: Extra parameters such as aes() or color, size passed.
- **xlab**: Label for x
- **ylab**: Label for y
- **main**: Title for plot.
- **facets**: Faceting formula to use.
- **stat**: Character vector specifying statistics to use. "stepping" with randomly assigned stepping levels as y variable. "identity" allow users to specify y value in aes.
- **group.selfish**: Passed to addStepping, control whether to show each group as unique level or not. If set to FALSE, if two groups are not overlapped with each other, they will probably be layout in the same level to save space.

### Value

A 'Layer'.

---

```r
ggplot(gr) + geom_rect(stat = "stepping", aes(y = value, group = pair))

ggplot(gr) + geom_rect(stat = "stepping", aes(y = value, group = pair), group.selfish = FALSE)
```
Author(s)

Tengfei Yin

Examples

```r
set.seed(1)
N <- 100
require(GenomicRanges)

## simmulated GRanges
gr <- GRanges(seqnames =
sample(c("chr1", "chr2", "chr3"),
size = N, replace = TRUE),
IRanges(
    start = sample(1:300, size = N, replace = TRUE),
    width = sample(70:75, size = N, replace = TRUE),
    strand = sample(c("+", "-", "*"), size = N,
        replace = TRUE),
    value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
    sample = sample(c("Normal", "Tumor"),
        size = N, replace = TRUE),
    pair = sample(letters, size = N,
        replace = TRUE))

## data.frame call ggplot2::geom_segment
ggplot() + geom_segment(data = mtcars, aes(x = mpg, y = wt, xend = mpg + 10, yend = wt + 0.2,
        color = cyl))

## default
##
## ggplot(gr) + geom_segment()
## or
## ggplot() + geom_segment(gr)

## facetting and aesthetics
##
## ggplot(gr) + geom_segment(facets = sample ~ seqnames, aes(color = strand))

## stat:identity
```
GGbio

class ggbio

Description

A sub class of ggplot and gg class defined in ggplot2 package, used for ggbio specific methods.

Usage

GGbio(ggplot = NULL, data = NULL, fetchable = FALSE, blank = FALSE, ...)

Arguments

ggplot  
a ggplot or gg object.
data  
raw data
fetchable  
logical value, default FALSE, is there any fetch method available.
blank  
logical value, default FALSE, is this plot a blank plot.
...  
More properties passed to class like Cache.

Details

This class is defined to facilitate the ggbio-specific visualization method, especially when using ggplot to construct ggbio supported object, that will return a ggbio class. And internals tricks will help a lazy evaluation for following + method.

Value

A ggbio object.

Author(s)

Tengfei Yin
See Also

ggplot

Examples

p1 <- qplot()
g1 <- ggbio(p1)
class(g1)

Description

These methods extend ggplot to support several types of Bioconductor objects, as well as some base types like matrix. They return a ggbio object, which stores the original data object. Please check the corresponding method for mold to see how an object is coerced into a data.frame.

Usage

```r
## S3 method for class 'Vector'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())
## S3 method for class 'Seqinfo'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())
## S3 method for class 'ExpressionSet'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())
## S3 method for class 'RsamtoolsFile'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())
## S3 method for class 'TxDbOREnsDb'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())
## S3 method for class 'BSgenome'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())
## S3 method for class 'matrix'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())
## S3 method for class 'character'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())
## S3 method for class 'SummarizedExperiment'
ggplot(data, mapping = aes(), 
       assay.id = 1L, ..., environment = parent.frame())
```
## S3 method for class 'GAlignments'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())

## S3 method for class 'VCF'
ggplot(data, mapping = aes(), ..., 
       environment = parent.frame())

### Arguments
- **data**: original data object.
- **mapping**: the aesthetic mapping.
- **...**: other arguments passed to specific methods.
- **environment**: fall-back environment for evaluation of aesthetic symbols.
- **assay.id**: index of assay you are using when multiple assays exist.

### Details
The biggest difference for objects returned by `ggplot` in ggbio from ggplot2, is we always keep the original data copy, this is useful because in ggbio, our starting point is not always data.frame, many special statistical transformation is computed upon original data objects instead of coerced data.frame. This is a hack to follow ggplot2’s API while allow our own defined components to trace back to original data copy and do the transformation. For objects supported by `mold` we transform them to data.frame stored along the original data set, for objects which not supported by `mold` method, we only store the original copy for ggbio specific graphics.

`ggplot()` is typically used to construct a plot incrementally, using the `+` operator to add layers to the existing `ggplot` object. This is advantageous in that the code is explicit about which layers are added and the order in which they are added. For complex graphics with multiple layers, initialization with `ggplot` is recommended. You can always call `qplot` in package ggplot2 or `autoplot` in ggbio for convenient usage.

There are three common ways to invoke `ggplot`:

- `ggplot(df, aes(x, y, <other aesthetics>))`
- `ggplot(df)`
- `ggplot()`

The first method is recommended if all layers use the same data and the same set of aesthetics, although this method can also be used to add a layer using data from another data frame. The second method specifies the default data frame to use for the plot, but no aesthetics are defined up front. This is useful when one data frame is used predominantly as layers are added, but the aesthetics may vary from one layer to another. The third method initializes a skeleton `ggplot` object which is fleshed out as layers are added. This method is useful when multiple data frames are used to produce different layers, as is often the case in complex graphics.

The examples below illustrate how these methods of invoking `ggplot` can be used in constructing a graphic.
Value

a return ggbio object, which is a subclass of ggplot defined in ggplot2 package, but that's more, a `.data` list entry is stored with the returned object.

Author(s)

Tengfei Yin

See Also

mold, ggbio

Examples

```r
set.seed(1)
N <- 100
library(GenomicRanges)
## GRanges
gr <- GRanges(seqnames =
    sample(c("chr1", "chr2", "chr3"),
        size = N, replace = TRUE),
    IRanges(
        start = sample(1:300, size = N, replace = TRUE),
        width = sample(70:75, size = N, replace = TRUE)),
    strand = sample(c("+", "+", "+"), size = N,
        replace = TRUE),
    value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
    sample = sample(c("Normal", "Tumor"),
        size = N, replace = TRUE),
    pair = sample(letters, size = N,
        replace = TRUE))

## automatically facetting and assign y
## this must mean geom_rect support GRanges object
ggplot(gr) + geom_rect()
## use pure ggplot2's geom_rect, no auto facet
ggplot(gr) + ggplot2::geom_rect(aes(xmin = start, ymin = score,
    xmax = end, ymax = score + 1))

## GRangesList
grl <- split(gr, values(gr)$pair)
## IRanges
```
ir <- ranges(gr)
ggplot(ir) + geom_rect()
ggplot(ir) + layout_circle(geom = "rect")

## Seqinfo
seqlengths(gr) <- c(400, 500, 420)
ggplot(seqinfo(gr)) + geom_point(aes(x = midpoint, y = seqlengths))

## matrix
mx <- matrix(1:12, nrow = 3)
ggplot(mx, aes(x = x, y = y)) + geom_raster(aes(fill = value))
## row is the factor
ggplot(mx, aes(x = x, y = row)) + geom_raster(aes(fill = value))
colnames(mx)
colnames(mx) <- letters[1:ncol(mx)]
mx
## has extra 'colnames'
ggplot(mx, aes(x = x, y = row)) + geom_raster(aes(fill = colnames))
rownames(mx)
rownames(mx) <- LETTERS[1:nrow(mx)]
ggplot(mx, aes(x = x, y = row)) + geom_raster(aes(fill = rownames))
## please check autoplot, matrix for more control

## Views

## ExpressionSet
library(Biobase)
data(sample.ExpressionSet)
sample.ExpressionSet
set.seed(1)
## select 50 features
idx <- sample(seq_len(dim(sample.ExpressionSet)[1]), size = 50)
eset <- sample.ExpressionSet[idx,]
ggplot(eset) + geom_tile(aes(x = x, y = y, fill = value))
## please check autoplot.matrix method which gives you more control
ggplot(eset) + geom_tile(aes(x = x, y = y, fill = sex))
ggplot(eset) + geom_tile(aes(x = x, y = y, fill = type))

## Rle
library(IRanges)
lambda <- c(rep(0.001, 4500), seq(0.001, 10, length = 500),
            seq(10, 0.001, length = 500))
xVector <- rpois(1e4, lambda)
xRle <- Rle(xVector)
ggplot(xRle) + geom_tile(aes(x = x, y = y, fill = value))

## RleList
xRleList <- RleList(xRle, 2L * xRle)
xRleList
ggplot(xRleList) + geom_tile(aes(x = x, y = y, fill = value)) + facet_grid(group~.)
names(xRleList) <- c("a", "b")
ggplot(xRleList) + geom_tile(aes(x = x, y = y, fill = value)) + facet_grid(group~.)

## RangedSummarizedExperiment
library(SummarizedExperiment)
nrows <- 200; ncols <- 6
counts <- matrix(runif(nrows * ncols, 1, 1e4), nrows)
counts2 <- matrix(runif(nrows * ncols, 1, 1e4), nrows)
rowRanges <- GRanges(rep(c("chr1", "chr2"), c(50, 150)),
                      IRanges(floor(runif(200, 1e5, 1e6)), width=100),
                      strand=sample(c("+", "-"), 200, TRUE))
colData <- DataFrame(Treatment=rep(c("ChIP", "Input"), 3),
                      row.names=LETTERS[1:6])
sset <- SummarizedExperiment(assays=SimpleList(counts=counts,
                      counts2 = counts2),
                      rowRanges=rowRanges, colData=colData)
ggplot(sset) + geom_raster(aes(x = x, y = y, fill = value))

ggsave

Save a ggplot object or tracks with sensible defaults

Description

`ggsave` is a convenient function for saving a plot. It defaults to saving the last plot that you displayed, and for a default size uses the size of the current graphics device. It also guesses the type of graphics device from the extension. This means the only argument you need to supply is the filename.

Usage

    ggsave(filename, plot = last_plot(),
           device = default_device(filename), path = NULL,
           scale = 1, width = par("din")[1],
           height = par("din")[2], units = c("in", "cm", "mm"),
           dpi = 300, limitsize = TRUE, ...)

Arguments

filename file name/filename of plot
plot  plot to save, defaults to last plot displayed
device device to use, automatically extract from file name extension
path  path to save plot to (if you just want to set path and not filename)
scale  scaling factor
width width (defaults to the width of current plotting window)
height height (defaults to the height of current plotting window)
units units for width and height when either one is explicitly specified (in, cm, or mm)
dpi  dpi to use for raster graphics
limitsize when TRUE (the default), ggsave will not save images larger than 50x50 inches, to prevent the common error of specifying dimensions in pixels.
... other arguments passed to graphics device

Details

ggsave currently recognises the extensions eps/ps, tex (pictex), pdf, jpeg, tiff, png, bmp, svg and wmf (windows only).

---

**Grob-class**  
*Grob getter*

**Description**

’Grob’ class is a container for ’grob’ based object defined with grid system. Generic function Grob gets grob object supported by grid system, and make an instance of subclass of class ’Grob’. ’GrobList’ is a container of list of ’Grob’ object.

**Usage**

```r
## S4 method for signature 'gg'
Grob(x)
## S4 method for signature 'gtable'
Grob(x)
## S4 method for signature 'trellis'
Grob(x)
## S4 method for signature 'lattice'
Grob(x)
## S4 method for signature 'GGbio'
Grob(x)
```

**Arguments**

x  object of class: gg, gtable, trellis, lattice, GGbio.
### Ideogram

**Value**

A Grob object.

**Author(s)**

Tengfei Yin

---

**Description**

Plot single chromosome with cytobands.

**Usage**

```r
plotIdeogram(obj, subchr = NULL, zoom.region = NULL, which = NULL, xlab, ylab, main, xlabel = FALSE, color = "red", fill = "red", alpha = 0.7, zoom.offset = 0.2, size = 1, cytobands = TRUE, aspect.ratio = 1/20, genome)
```

```r
## constructor
Ideogram(obj, subchr = NULL, which = NULL, xlabel = FALSE, cytobands = TRUE, color = "red", fill = "red", alpha = 0.7, zoom.region = NULL, zoom.offset = 0.2, size = 1, aspect.ratio = 1/20, ..., genome)
```

**Arguments**

- `obj` A GenomicRanges object, which include extra information about cytobands, check biovizBase::isIdeogram.
- `subchr` A single character of chromosome names to show.
- `which` GRanges object to subset and highlight the ideogram.
- `zoom.region` A numeric vector of length 2 indicating zoomed region.
- `xlab` Label for x
- `ylab` Label for y
- `main` Title for plot.
- `xlabel` A logical value. Show the x label or not.
- `color` color for highlight region.
- `fill` fill color for highlight region.
- `alpha` alpha for highlight regio.
- `zoom.offset` zoomed highlights region offset around chromosome plotting region.
- `size` size for zoomed region rectangle boundary.
layout_circle  

Create a circle layout

Description

Create a circle layout.

Usage

```r
## S4 method for signature 'GRanges'
layout_circle(data, ...,
  geom = c("point", "line", "link", "ribbon", 
  "rect", "bar", "segment", "hist", "scale", "heatmap", "ideogram", 
  "text"),
  linked.to, radius = 10, trackWidth = 5,
  space.skip = 0.015, direction = c("clockwise", 
  "anticlockwise"), link.fun = function(x, y, n = 30)
  bezier(x, y, evaluation = n), rect.inter.n = 60, rank,
  ylim = NULL,
  scale.n = 60, scale.unit = NULL, scale.type = c("M", 
  "B", "sci"), grid.n = 5, grid.background = "gray70",
```
layout_circle

```
grid.line = "white", grid = FALSE, chr.weight = NULL)
```

## S4 method for signature 'missing'
layout_circle(data, ...)
circle(...)

**Arguments**

- **data**
  - A GRanges object.

- **...**
  - Extra parameters such as aesthetics mapping in aes(), or color, size, etc. For circle function, it passed to layout_circle.

- **geom**
  - The geometric object to use display the data.

- **linked.to**
  - Character indicates column that specifying end of the linking lines, that column should be a GRanges object.

- **radius**
  - Numeric value indicates radius. Default is 10.

- **trackWidth**
  - Numeric value indicates the track width.

- **space.skip**
  - Numeric value indicates the ratio of skipped region between chunks(chromosomes in GRanges) to the whole track space.

- **direction**
  - Space layout orders.

- **link.fun**
  - Function used for interpolate the linking lines. Default is Hmisc::bezier.

- **rect.inter.n**
  - n passed to interpolate function in rectangle transformation(from a rectangle) to a section in circular view.

- **rank**
  - For default equal trackWidth, use rank to specify the circle orders.

- **ylim**
  - Numeric range to control y limits.

- **scale.n**
  - Approximate number of ticks you want to show on the whole space. used when scale.unit is NULL.

- **scale.unit**
  - Unit used for computing scale. Default is NULL.

- **scale.type**
  - Scale type used for

- **grid**
  - logical value indicate showing grid background for track or not.

- **grid.n**
  - integer value indicate horizontal grid line number.

- **grid.background**
  - grid background color.

- **grid.line**
  - grid line color.

- **chr.weight**
  - numeric vectors which sum to <1, the names of vectors has to be matched with seqnames in seqinfo, and you can only specify part of the seqnames, other lengths of chromosomes will be assined proportionally to their seqlengths, for example, you could specify chr1 to be 0.5, so the chr1 will take half of the space and other chromosomes squeezed to take left of the space.

**Value**

A 'Layer'.

Author(s)

Tengfei Yin

Examples

N <- 100
library(GenomicRanges)
## simmulated GRanges
##
gr <- GRanges(seqnames =
    sample(c("chr1", "chr2", "chr3"),
    size = N, replace = TRUE),
IRanges(
    start = sample(1:300, size = N, replace = TRUE),
    width = sample(70:75, size = N, replace = TRUE)),
strand = sample(c("+", "-", "*"), size = N,
    replace = TRUE),
value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
sample = sample(c("Normal", "Tumor"),
    size = N, replace = TRUE),
pair = sample(letters, size = N,
    replace = TRUE))

seqlengths(gr) <- c(400, 500, 700)
values(gr)$to.gr <- gr[sample(1:length(gr), size = length(gr))]

## more formal API

ggplot(gr) + layout_circle(geom = "ideo", fill = "gray70", radius = 7, trackWidth = 3) +
    layout_circle(geom = "bar", radius = 10, trackWidth = 4, aes(fill = score, y = score)) +
    layout_circle(geom = "point", color = "red", radius = 14,
trackWidth = 3, grid = TRUE, aes(y = score)) +
    layout_circle(geom = "link", linked.to = "to.gr", radius = 6, trackWidth = 1)

## more formal API

ggplot(gr) + layout_circle(geom = "ideo", fill = "gray70", radius = 7, trackWidth = 3) +
    layout_circle(geom = "bar", radius = 10, trackWidth = 4, aes(fill = score, y = score)) +
    layout_circle(geom = "point", color = "red", radius = 14,
trackWidth = 3, grid = TRUE, aes(y = score)) +
    layout_circle(geom = "link", linked.to = "to.gr", radius = 6, trackWidth = 1)

layout_karyogram

Create a karyogram layout

Description

Create a karyogram layout.
Usage

```r
## S4 method for signature 'GRanges'
layout_karyogram(data, ..., xlab, ylab, main,
    facets = seqnames ~ ., cytobands = FALSE, geom = "rect",
    stat = NULL, ylim = NULL, rect.height = 10)
```

Arguments

data

da GRanges object, which could contain extra information about cytobands. If you want an accurate genome mapping, please provide seqlengths with this GRanges object, otherwise it will emit a warning and use data space to estimate the chromosome space which is very rough.

... Extra parameters such as aes() or arbitrary color and size.

xlab character vector or expression for x axis label.

ylab character vector or expression for y axis label.

main character vector or expression for plot title.

facets faceting formula to use.

cytobands logical value indicate to show the cytobands or not.

geom The geometric object to use display the data.

stat character vector specifying statistics to use.

ylim limits for y axis, usually the chromosome spaces y limits are from 0 to rect.height, which 10, so if you want to stack some data on top of it, you can set limits to like c(10, 20).

rect.height numeric value indicate half of the rectangle ploting region, used for alignment of multiple layers.

Value

A 'Layer'.

Author(s)

Tengfei Yin

Examples

```r
### R code from vignette source 'karyogram.Rnw'

#############################################################
### code chunk number 1: loading
#############################################################
library(ggbio)
data(hg19IdeogramCyto, package = "biovizBase")
head(hg19IdeogramCyto)
## default pre-set color stored in
```
layout_karyogram

ggetOption("biovizBase")$cytobandColor

########################################################################
### code chunk number 2: default
########################################################################
autoplot(hg19IdeogramCyto, layout = "karyogram", cytobands = TRUE)

########################################################################
### code chunk number 3: change-order
########################################################################
library(GenomicRanges)
hg19 <- keepSeqlevels(hg19IdeogramCyto, paste0("chr", c(1:22, "X", "Y")))
head(hg19)
autoplot(hg19, layout = "karyogram", cytobands = TRUE)

########################################################################
### code chunk number 4: cyto-normal
########################################################################
library(GenomicRanges)
## it's a 'ideogram'
biovizBase::isIdeogram(hg19)
## set to FALSE
autoplot(hg19, layout = "karyogram", cytobands = FALSE, aes(fill = gieStain)) +
  scale_fill_giemsa()

########################################################################
### code chunk number 5: load-RNAediting
########################################################################
data(darned_hg19_subset500, package = "biovizBase")
dn <- darned_hg19_subset500
head(dn)
## add seqlengths
## we have seqlengths information in another data set
data(hg19Ideogram, package = "biovizBase")
seqlengths(dn) <- seqlengths(hg19Ideogram)[names(seqlengths(dn))]
## now we have seqlengths
head(dn)
## then we change order
dn <- keepSeqlevels(dn, paste0("chr", c(1:22, "X")))
autoplot(dn, layout = "karyogram")
## this equivalent to
## autoplot(seqinfo(dn))

########################################################################
### code chunk number 6: load-RNAediting-color
########################################################################
## since default is geom rectangle, even though it's looks like segment
## we still use both fill/color to map colors
autoplot(dn, layout = "karyogram", aes(color = exReg, fill = exReg))

###################################################
### code chunk number 7: load-RNAediting-color-NA
###################################################
## since default is geom rectangle, even though it's looks like segment
## we still use both fill/color to map colors
autoplot(dn, layout = "karyogram", aes(color = exReg, fill = exReg)) +
  scale_color_discrete(na.value = "brown")

###################################################
### code chunk number 8: load-RNAediting-color-fake
###################################################
dn2 <- dn
seqlengths(dn2) <- rep(max(seqlengths(dn2)), length(seqlengths(dn2)) )
autoplot(dn2, layout = "karyogram", aes(color = exReg, fill = exReg))

###################################################
### code chunk number 9: plotKaryogram (eval = FALSE)
###################################################
## plotKaryogram(dn)
## plotKaryogram(dn, aes(color = exReg, fill = exReg))

###################################################
### code chunk number 10: low-default
###################################################
## plot ideogram
p <- ggplot(hg19) + layout_karyogram(cytobands = TRUE)
p
## equivalent autoplot(hg19, layout = "karyogram", cytobands = TRUE)

###################################################
### code chunk number 11: low-default-addon
###################################################
p <- p + layout_karyogram(dn, geom = "rect", ylim = c(11, 21), color = "red")
## commented line below won't work
## the cytoband fill color has been used already.
## p <- p + layout_karyogram(dn, aes(fill = exReg, color = exReg), geom = "rect")
p
###################################################
### code chunk number 12: edit-space
###################################################
## plot chromosome space
p <- autoplot(seqinfo(dn))
## make sure you pass rect as geom
## otherwise you just get background

```r
p <- p + layout_karyogram(dn, aes(fill = exReg, color = exReg), geom = "rect")
values(dn)$pvalue <- rnorm(length(dn))
p + layout_karyogram(dn, aes(x = start, y = pvalue), ylim = c(10, 30), geom = "line", color = "red")
p
```

--------------------------------------------------------------------------------
### code chunk number 13: sessionInfo
--------------------------------------------------------------------------------

```r
sessionInfo()
```

---

**Plot**

### Plot class

**Description**

genealize a graphic object to a Plot object.

**Usage**

```r
## S4 method for signature 'gg'
Plot(x)
## S4 method for signature 'trellis'
Plot(x, mutable = FALSE)
## S4 method for signature 'GGbio'
Plot(x)
## S4 method for signature 'Ideogram'
Plot(x)
```

**Arguments**

- `x` object of gg, GGbio, trellis, Ideogram.
- `mutable` whether a plot repsonse to + method or not.

**Value**

A Plot object.

**Author(s)**

Tengfei Yin
plotFragLength

Plot estimated fragment length for paired-end RNA-seq data

Description

Plot estimated fragment length for paired-end RNA-seq data against single reduced data model.

Usage

```r
## S4 method for signature 'character,GRanges'
plotFragLength(data, model,
               gap.ratio = 0.0025,
               geom = c("segment", "point", "line"),
               type = c("normal", "cut"),
               heights = c(400, 100),
               annotation = TRUE)
```

Arguments

- **data**: A character indicate the bam file.
- **model**: A reduced model to compute estimated fragment length. please see details.
- **gap.ratio**: When type is set to "cut", it will provide a compact view, which cut the common gaps in a certain ratio.
- **geom**: One or all three geoms could be drawn at the same time. y value of "point" and "line" indicate the estimated fragment length. and if geom is set to "segment", the segment is from the left most position to paired right most position, should be equal to "isize".
- **type**: "normal" return a uncut view, loose but the coordinate is true genomic coordinates. "cut" cut the view in a compact way.
- **heights**: Numeric vector indicate the heights of tracks.
- **annotation**: A logical value. TRUE shows model, and FALSE shows only fragment length with labels.

Details

We use a easy way to define this estimated fragment length, we collect all paired reads and model, reduce model first, then find common gaps, remove common gaps between paired-end reads, and compute the new estimated fragment length.

Value

A ggplot object when annotation = FALSE and a frame grob if annotation = TRUE

Author(s)

Tengfei Yin
Examples

```r
## Not run:
data(genesymbol)
bamfile <- system.file("extdata", "SRR027894subRBM17.bam", package="biovizBase")
library(TxDb.Hsapiens.UCSC.hg19.knownGene)
txdb <- TxDb.Hsapiens.UCSC.hg19.knownGene
model <- exonsBy(txdb, by = "tx")
model.new <- subsetByOverlaps(model, genesymbol["RBM17"])
exons.rbm17 <- subsetByOverlaps(exons(txdb), genesymbol["RBM17"])
exons.new <- reduce(exons.rbm17)
plotFragLength(bamfile, exons.new, geom = "line")
plotFragLength(bamfile, exons.new, geom = c("point","segment"))
plotFragLength(bamfile, exons.new, geom = c("point","segment"), annotation = FALSE)
plotFragLength(bamfile, exons.new, geom = c("point","segment"), type = "cut",
gap.ratio = 0.001)
## End(Not run)
```

plotGrandLinear  
**Manhattan for GWAS**

Description

A Manhattan plot is a special scatter plot used to visualize data with a large number of data points, with a distribution of some higher-magnitude values. For example, in genome-wide association studies (GWAS). Here we mainly focus on GWAS Manhattan plots. X-axis is genomic coordinates and Y-axis is the negative logarithm of the associated P-value for each single nucleotide polymorphism. So higher the value, more stronger the association they are.

Usage

```r
plotGrandLinear(obj, ..., facets, space.skip = 0.01, geom = NULL,
cutoff = NULL, cutoff.color = "red", cutoff.size = 1,
legend = FALSE, xlim, ylim, xlab, ylab, main,
highlight.gr = NULL, highlight.name = NULL,
highlight.col = "red", highlight.label = TRUE,
highlight.label.size = 5, highlight.label.offset =
0.05, highlight.label.col = "black", spaceline = FALSE)
```

Arguments

- **obj**  
  GRanges object which contains extra p value, before users pass this object, they need to make sure the p value has been changed to -log10(p).
- **...**  
  extra arguments passed. such as color, size, alpha.
- **facets**  
  facets formula, such as group ~ .
space.skip numeric value for skip ratio, between chromosome spaces. Default is 0.01.
geom geometric object, default is "point".
cutoff A numeric vector which used as cutoff for Manhattan plot.
cutoff.color A character specifying the color used for cutoff. Default is "red".
cutoff.size A numeric value which used as cutoff line size.
legend A logical value indicate whether to show legend or not. Default is FALSE which disabled the legend.
xlim limits for x scale.
ylim limits for y scale.
xlab Label for xscale.
ylab Label foryscale.
main title.
highlight.gr a GRanges object, this will highlight overlapped region with provided intervals.
highlight.name if NULL, using rownames of GRanges object provided by argument highlight.gr, otherwise use character to indicate column used as labeled names.
highlight.col highlight colors.
highlight.label logical value, label the highlighted region or not.
highlight.label.size highlight label size.
highlight.label.offset highlight label offset.
highlight.label.col highlight label color.
spaceline show line between chromosomes.

Details

Please use seqlengths of the object and space.skip arguments to control the layout of the coordiant genome transformation.
aes(y = ...) is required.
aes(color = ) is used to mapping to data variables, if just pass "color" without aes(), then will recycle the color to represent each chromosomes. please see the example below.

Value

Return a ggplot object.

Author(s)

Tengfei Yin
Examples

```r
## load
library(ggbio)
data(hg19IdeogramCyto, package = "biovizBase")
data(hg19Ideogram, package = "biovizBase")
library(GenomicRanges)

## simul_gr
library(biovizBase)
gr <- GRanges(rep(c("chr1", "chr2"), each = 5),
               IRanges(start = rep(seq(1, 100, length = 5), times = 2),
                       width = 50))
autoplot(gr)

## coord:genome
autoplot(gr, coord = "genome")
gr.t <- transformToGenome(gr)
head(gr.t)

## is
is_coord_genome(gr.t)
metadata(gr.t)$coord

## simul_snp
chrs <- as.character(levels(seqnames(hg19IdeogramCyto)))
seqlths <- seqlengths(hg19Ideogram)[chrs]
set.seed(1)
nchr <- length(chrs)
nsnps <- 100
gr.snp <- GRanges(rep(chrs, each = nsnps),
                  IRanges(start =
                      do.call(c, lapply(chrs, function(chr){
                        N <- seqlths[chr]
                        runif(nsnps, 1, N)
                      })),
                       width = 1),
                  SNP = sapply(1:(nchr*nsnps), function(x) paste("rs",x,sep='')),
                  pvalue = -log10(runif(nchr*nsnps)),
                  group = sample(c("Normal", "Tumor"), size = nchr*nsnps,
                      replace = TRUE))

## shorter
seqlengths(gr.snp)
nms <- seqnames(seqinfo(gr.snp))
nms.new <- gsub("chr", "", nms)
names(nms.new) <- nms
gr.snp <- renameSeqlevels(gr.snp, nms.new)
seqlengths(gr.snp)
```
## unorder
```r
autoplot(gr.snp, coord = "genome", geom = "point", aes(y = pvalue), space.skip = 0.01)
```

## sort
```r
gr.snp <- keepSeqlevels(gr.snp, c(1:22, "X", "Y"))
autoplot(gr.snp, coord = "genome", geom = "point", aes(y = pvalue), space.skip = 0.01)
```

## with_seql
```r
names(seqlths) <- gsub("chr", ",", names(seqlths))
seqlengths(gr.snp) <- seqlths[names(seqlengths(gr.snp))]
autoplot(gr.snp, coord = "genome", geom = "point", aes(y = pvalue), space.skip = 0.01)
```

## line
```r
autoplot(gr.snp, coord = "genome", geom = "line", aes(y = pvalue, group = seqnames,
          color = seqnames))
```

## plotGrandLinear
```r
plotGrandLinear(gr.snp, aes(y = pvalue))
```

## morecolor
```r
plotGrandLinear(gr.snp, aes(y = pvalue, color = seqnames))
plotGrandLinear(gr.snp, aes(y = pvalue), color = c("green", "deepskyblue"))
plotGrandLinear(gr.snp, aes(y = pvalue), color = c("green", "deepskyblue", "red"))
plotGrandLinear(gr.snp, aes(y = pvalue), color = "red")
```

## cutoff
```r
plotGrandLinear(gr.snp, aes(y = pvalue), cutoff = 3, cutoff.color = "blue", cutoff.size = 4)
```

## cutoff-low
```r
plotGrandLinear(gr.snp, aes(y = pvalue)) + geom_hline(yintercept = 3, color = "blue", size = 4)
```

## longer
```r
## let's make a long name
nms <- seqnames(seqinfo(gr.snp))
nms.new <- paste("chr0000", nms, sep = ",")
names(nms.new) <- nms
gr.snp <- renameSeqlevels(gr.snp, nms.new)
seqlengths(gr.snp)
```

## rotate
```r
plotGrandLinear(gr.snp, aes(y = pvalue)) + theme(axis.text.x = element_text(angle=-90, hjust=0))
```

## sessionInfo
```r
sessionInfo()
```

---

Plot Ranges Linked with Data
Description
Plot GRanges object structure and linked to a even spaced paralell coordinates plot which representing the data in elementeMetadata.

Usage
```r
## S4 method for signature 'RangedSummarizedExperiment'
plotRangesLinkedToData(data, ..., 
    stat.y = seq_len(ncol(data)), stat.ylab = names(assays(data)[stat.assay]), 
    stat.assay = 1L)

## S4 method for signature 'GenomicRanges.OR_GRangesList'
plotRangesLinkedToData(data, ..., 
    stat.y = seq_len(ncol(mcols(data))), 
    stat.ylab, sig, sig.col = c("black", "red"), 
    stat.coord.trans = coord_trans(), 
    annotation = list(), width.ratio = 0.8, 
    theme.stat = theme_gray(), theme.align = theme_gray(), 
    linetype = 3, heights)
```

Arguments
- **data**: GRanges object with a DataFrame as elementMetadata.
- **...**: Parameters passed to control lines in top plot.
- **stat.y**: integer (variable position starting in DataFrame of data, start from 1) or strings (variable names) which indicate the column names.
- **stat.ylab**: y label for stat track (the top track).
- **stat.assay**: default 1L, element of assays.
- **sig**: a character of element meta data column of logical value, indicates which row is significant and will be shown in link lines and rectangle.
- **sig.col**: colors for significant, valid when you specify "sig" argument, the first color indicates FALSE, non-significant, the second color indicates TRUE.
- **stat.coord.trans**: transformation used for top plot.
- **annotation**: A list of ggplot object.
- **width.ratio**: Control the segment length of statistic layer.
- **theme.stat**: top plot theme.
- **theme.align**: alignment themes.
- **linetype**: linetype
- **heights**: Heights of each track.
Details

Inspired by some graphics produced in some other packages, for example in package DEXseq, the author provides graphics with gene models and linked to an even spaced statistics summary. This is useful because we always plot everything along the genomic coordinates, but genomic features like exons are not evenly distributed, so we could actually treat the statistics associated with exons like categorical data, and show them as “Parallel Coordinates Plots”. This is one special layout which represent the data in a nice manner and also keep the genomic structure information. With ability of tracks, it’s possible to generate such type of a graphic along with other annotations.

The data we want is a normal GRanges object, and make sure the intervals are not overlapped with each other(currently), and you may have multiple columns which store the statistics for multiple samples, then we produce the graphic we introduced above and users could pass other annotation track in the function which will be shown below the main linked track.

The reason you need to pass annotation into the function instead of binding them by tracks later is because binding manually with annotation tracks is tricky and this function doesn’t return a ggplot object.

Value

return a frame grob; side-effect (plotting) if plot=T.

Author(s)

Tengfei Yin

Examples

library(TxDb.Hsapiens.UCSC.hg19.knownGene)
library(ggbio)
data(genesymbol, package = "biovizBase")
txdb <- TxDb.Hsapiens.UCSC.hg19.knownGene
model <- exonsBy(txdb, by = "tx")
model17 <- subsetByOverlaps(model, genesymbol["RBM17"])
exons <- exons(txdb)
exon17 <- subsetByOverlaps(exons, genesymbol["RBM17"])
## reduce to make sure there is no overlap
## just for example
exon.new <- reduce(exon17)
## suppose
values(exon.new)$sample1 <- rnorm(length(exon.new), 10, 3)
values(exon.new)$sample2 <- rnorm(length(exon.new), 10, 10)
values(exon.new)$score <- rnorm(length(exon.new))
values(exon.new)$significant <- sample(c(TRUE, FALSE), size = length(exon.new), replace = TRUE)

plotRangesLinkedToData(exon.new, stat.y = c("sample1", "sample2"))
plotRangesLinkedToData(exon.new, stat.y = 1:2)
plotRangesLinkedToData(exon.new, stat.y = 1:2, size = 3, linetype = 4)
plotRangesLinkedToData(exon.new, stat.y = 1:2, size = 3, linetype = 4,
                       sig = "significant")
plotRangesLinkedToData(exon.new, stat.y = 1:2, size = 3, linetype = 4,
                       sig = "significant", sig.col = c("gray90","red"))
plotSpliceSum

Plot Splice Summary from RNA-seq data

Description

Plot splice summary by simply counting overlaped junction read in weighted way or not.

Usage

```r
## For character,GRangesList
## S4 method for signature 'character,GRangesList'
plotSpliceSum(data, model, ..., weighted = TRUE)

## For character,TxDB
## S4 method for signature 'character,TxDB'
plotSpliceSum(data, model, which, ..., weighted = TRUE)

## For character,EnsDb
## S4 method for signature 'character,EnsDb'
plotSpliceSum(data, model, which, ..., weighted = TRUE)
```

Arguments

data
A character specifying the bam file path of RNA-seq data.

model
A GRangesList which representing different isoforms, a TxDb or an EnsDb object. For the latter cases, users need to pass "which" argument which, for TxDb, is a GRanges object to specify the region and for EnsDb can be a GRanges object, an object extending AnnotationFilter, an AnnotationFilterList combining such filter objects or a filter expression in form of a formula.

which
A GRanges object specifying the region you want to get model from the TxDb object. For EnsDb: can be a GRanges object, an object extending AnnotationFilter, an AnnotationFilterList combining such filter objects or a filter expression in form of a formula.

weighted
If TRUE, weighted by simply add 1/cases matched to each model and if FALSE, simply add 1 to every case.

...
Extra arguments passed to qplot function. such as, offset which control the height of chevron.

Details

Internally we use biovizBase:::spliceSummary for simple counting, but we encourage users to use their own robust way to make slicing summary and store it as GRangesList, then plot the summary by qplot function.

Value

A ggplot object.
plotStackedOverview

Description

Plot stacked overview for genome with or without cytobands. It's a wrapper around layout_karyogram.

Usage

plotStackedOverview(obj, ..., xlab, ylab, main, geom = "rect", cytobands = FALSE, rescale = TRUE, rescale.range = c(0, 10))
plotKaryogram(obj, ..., xlab, ylab, main, geom = "rect", cytobands = FALSE, rescale = TRUE, rescale.range = c(0, 10))

Arguments

obj  a GRanges object, which could contain extra information about cytobands. If it’s missing, will ask user to provide species information and download proper data set from UCSC. If you want an accurate genome mapping, please provide seqlengths with this GRanges object, otherwise it will emit a warning and use data space to estimate the chromosome space which is very rough.
arguments passed to graphic functions to control aesthetics. For example, if you use geom "point", you need to provide "y" in aes(), and if can also pass color, fill, size etc. to control graphics.

xlab label for x
ylab label for y
main title for plot.
geom geom plotted on the stacked layout. Default is "rect", which showing interval data as rectangles. It automatically figures out boundary so you don’t have to provide information in aes, users could specify other supported geom works for data.frame.
cytobands logical value. Default is FALSE. If TRUE, plotting cytobands, this require your data have arbitrary column as name and gieStain. the easiest way is to use getIdeogram to get your data. Notice for this function, when cytobands is TRUE, it will only plot cytobands without overlaying your data. If you really need to overlay extra data on cytobands, please plus layout_karyogram for that purpose.
rescale logical value. Default is TRUE, which rescale your data into the rescale.range, this make sure your data will not be plotted outside the stacked overview box.
rescale.range Numeric range of length 2. Default is (0, 10), because stacked layout draws a white background as chromosome space and this space is of height 10. We hide the y-axis since we don’t need it for stacked overview. Sometime users may want to leave some margin for their data, they can use this arguments to control the rescale.

Details

Stacked overview is just a arbitrary layout for karyogram layout, it use facets seqnaems ~ . as default to stack the genome. For accurate mapping, you need to provide seqlengths information in your GRanges object. Otherwise, data space will be computed for stacked overview chromosome background, this is _NOT_ the actual chromosome space!.

Value

A ggplot object.

Author(s)

Tengfei Yin

Examples

## Not run:
library(biovizBase)
data(hg19IdeogramCyto, package = "biovizBase")
library(GenomicRanges)

# you can also get ideogram by biovizBase::getIdeogram
## make shorter and clean labels
old.chrs <- seqnames(seqinfo(hg19IdeogramCyto))
new.chrs <- gsub("chr", "", old.chrs)
## lst <- as.list(new.chrs)
names(new.chrs) <- old.chrs
new.ideo <- renameSeqlevels(hg19IdeogramCyto, new.chrs)
new.ideo <- keepSeqlevels(new.ideo, c(as.character(1:22) , "X", "Y"))
new.ideo

## sample data
data(darned_hg19_subset500, package = "biovizBase")
idx <- is.na(values(darned_hg19_subset500)$exReg)
values(darned_hg19_subset500)$exReg[idx] <- "unknown"

## you need to add seqlengths for accurate mapping
chrnames <- unique(as.character(seqnames(darned_hg19_subset500)))
data(hg19Ideogram, package = "biovizBase")
seqlengths(darned_hg19_subset500) <- seqlengths(hg19Ideogram)[sort(chrnames)]

dn <- darned_hg19_subset500
values(dn)$score <- rnorm(length(dn))

## plotStackedOverview is a simple wrapper around this functions to
## create a stacked layout
plotStackedOverview(new.ideo, cytobands = TRUE)
plotStackedOverview(dn)
plotStackedOverview(dn, aes(color = exReg, fill = exReg))
## this will did the trick for you to rescale the space
plotStackedOverview(dn, aes(x = midpoint, y = score), geom = "line")
plotStackedOverview(dn, aes(x = midpoint, y = score), geom = "line", rescale.range = c(4, 6))
## no rescale
plotStackedOverview(dn, aes(x = midpoint, y = score), geom = "line", rescale = FALSE,
  xlab = "xlab", ylab = "ylab", main = "main") + ylab("ylab")

## no object? will ask you for species and query the data on the fly
plotStackedOverview()
plotStackedOverview(cytobands = TRUE)

## End(Not run)

### rescale ggplot object

#### Description

Rescale a numeric vector or ggplot object, could be used for static zoom-in in ggbio.
rescale

Usage

## S4 method for signature 'numeric'
rescale(x, to = c(0, 1),
       from = range(x, na.rm = TRUE))

## S4 method for signature 'ggplot'
rescale(x, xlim, ylim, sx = 1, sy = 1)
## S4 method for signature 'gg'
rescale(x, xlim, ylim, sx = 1, sy = 1)

Arguments

x A numeric object or ggplot object to be rescaled.

to For numeric object. it's a vector of two numeric values, specifying the range to be rescale.

from Range of x.

xlim For ggplot object. This specify the new limits on x-scale.

ylim For ggplot object. This specify the new limits on y-scale.

sx Scale fold for x-scale. Default is 1, no change.

sy Scale fold for y-scale. Default is 1, no change.

Details

When x is numeric value, it's just call scales::rescale, please refer to the manual page to check more details. If x is ggplot object, it first try to estimate current x limits and y limits of the ggplot object, then rescale based on those information.

Value

Return the object of the same class as x after rescaling.

Author(s)

Tengfei Yin

Examples

library(ggbio)
head(mtcars)
range(mtcars$mpg)
p <- qplot(data = mtcars, x = mpg, y = disp, geom = "point")
p.new <- rescale(p, xlim = c(20, 25))
p.new
**scale_fill_fold_change**

*scale color for fold change values*

---

**Description**

In biology, lots of data are scaled to value around 0, and people like to show them as blue-white-red scale color, where negative value are blue, 0 is white and positive value is red, and they are scaled for continuous variables.

**Usage**

```r
scale_fill_fold_change()
```

**Value**

a list.

**Author(s)**

Tengfei Yin

**Examples**

```r
p1 <- autoplot(volcano - 150)
p1
p1 + scale_fill_fold_change()
```

---

**scale_fill_giemsa**

*scale filled color to customized giemsa color.*

---

**Description**

scale filled color to customized giemsa color.

**Usage**

```r
scale_fill_giemsa(fill = getOption("biovizBase")$cytobandColor)
```

**Arguments**

- `fill` a character vector to indicate colors, and names of vector mapped to gieStain name.

**Value**

a list.
scale_x_sequnit

Author(s)

Tengfei Yin

Examples

getOption("biovizBase")$cytobandColor
library(biovizBase)
data(hg19IdeogramCyto)
p1 <- autoplot(hg19IdeogramCyto, layout = "karyogram", aes(fill =
gieStain))
p1
p1 + scale_fill_giemsa()

scale_x_sequnit  scale x by unit

Description

scale x by unit 'Mb','kb', 'bp'.

Usage

scale_x_sequnit(unit = c("Mb", "kb", "bp"), append = NULL)

Arguments

unit  unit to scale x. Default is Mb.
append  default NULL. If pass a character, it disalbe unit and arbitrarily append a text
        behind the original x scale numbers.

Value

'position_c'

Author(s)

Tengfei Yin

Examples

library(ggplot2)
p <- qplot(x = seq(1, to = 10000, length.out = 40), y = rnorm(40), geom = "point")
## default mb
p + scale_x_sequnit()
p + scale_x_sequnit("kb")
p + scale_x_sequnit("bp")
stat_aggregate

Generates summaries on the specified windows

Description

Generates summaries on the specified windows

Usage

### S4 method for signature 'GRanges'

```r
stat_aggregate(data, ..., xlab, ylab, main, by, FUN,
   maxgap=-1L, minoverlap=0L,
   type=c("any", "start", "end", "within", "equal"),
   select=c("all", "first", "last", "arbitrary"),
   y = NULL, window = NULL, facets = NULL,
   method = c("mean", "median","max",
     "min", "sum", "count", "identity"),
   geom = NULL)
```

Arguments

- `data` A GRanges or data.frame object.
- `...` Arguments passed to plot function. such as aes() and color.
- `xlab` Label for x
- `ylab` Label for y
- `main` Title for plot.
- `by` An object with 'start', 'end', and 'width' methods. Passed to aggreagate.
- `FUN` The function, found via 'match.fun', to be applied to each window of 'x'. Passed to aggreagate.
- `maxgap`, `minoverlap`, `type` Used in the internal call to findOverlaps() to detect overlaps. See ?findOverlaps in the IRanges package for a description of these arguments.
- `select` It passed to findOverlaps.
- `y` Passed to findOverlaps.
- `window`, `facets` Passed to findOverlaps.
- `method` The function, found via 'match.fun', to be applied to each window of 'x'. Passed to aggreagate.
- `geom` Passed to findOverlaps.

When select is "all" (the default), the results are returned as a Hits object. When select is "first", "last", or "arbitrary" the results are returned as an integer vector of length query containing the first, last, or arbitrary overlapping interval in subject, with NA indicating intervals that did not overlap any intervals in subject.

If select is "all", a Hits object is returned. For all other select the return value depends on the drop argument. When select != "all" && !drop, an IntegerList is returned, where each element of the result corresponds to a space in query. When select != "all" && drop, an integer vector is returned containing indices that are offset to align with the unlisted query.
stat_aggregate

- **y**: A character indicating the variable column for which aggregation is taken on, same as `aes(y = )`.
- **window**: Integer value indicating window size.
- **facets**: Faceting formula to use.
- **method**: Customized method for aggregating, if `FUN` is not provided.
- **geom**: The geometric object to use to display the data.

**Value**

A `Layer`.

**Author(s)**

Tengfei Yin

**Examples**

```r
library(GenomicRanges)
set.seed(1)
N <- 1000
## simmulated GRanges
gr <- GRanges(seqnames = sample(c("chr1", "chr2", "chr3"), size = N, replace = TRUE),
             IRanges(start = sample(1:300, size = N, replace = TRUE),
                     width = sample(70:75, size = N, replace = TRUE),
                     strand = sample(c("+", "-", "*"), size = N,
                                     replace = TRUE),
                     value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
                     sample = sample(c("Normal", "Tumor"), size = N, replace = TRUE),
                     pair = sample(letters, size = N, replace = TRUE))

ggplot(gr) + stat_aggregate(aes(y = value))
## or
ggplot(gr) + stat_aggregate(y = "value")
## Not run:
## no hits
ggplot(gr) + stat_aggregate(aes(y = value), select = "first", type = "within")
## End(Not run)
```

```r
ggplot(gr) + stat_aggregate(window = 30, aes(y = value), fill = "gray40", geom = "bar")
```

```r
ggplot(gr) + stat_aggregate(window = 100, fill = "gray40", aes(y = value),
                           method = "max", geom = "bar")
```
ggplot(gr) + stat_aggregate(aes(y = value), geom = "boxplot")

## now facets need to take place inside stat_* geom_* for an accurate computation

ggplot(gr) + stat_aggregate(aes(y = value), geom = "boxplot", window = 30,
facets = sample ~ seqnames)

## FIXME:
## autoplot(gr, stat = "aggregate", aes(y = value), window = 36)
## autoplot(gr, stat = "aggregate", geom = "boxplot", aes(y = value), window = 36)

stat_bin

### Binning method

**Description**

Binning method especially for Rle and RleList, for data.frame it's just calling ggplot2::stat_bin.

**Usage**

```
## S4 method for signature 'ANY'
stat_bin(data, ...)
```

```
## S4 method for signature 'Rle'
stat_bin(data, ..., binwidth, nbin = 30,
xlab, ylab, main, geom = c("bar", "heatmap"),
type = c("viewSums","viewMins",
         "viewMaxs", "viewMeans"))
```

```
## S4 method for signature 'RleList'
stat_bin(data, ..., binwidth, nbin = 30,
xlab, ylab, main,
indName = "sample",
geom = c("bar", "heatmap"),
type = c("viewSums","viewMins",
         "viewMaxs", "viewMeans"))
```

**Arguments**

- `data` Typically a data.frame or Rle or RleList object.
- `...` arguments passed to aesthetics mapping.
- `binwidth` width of the bins.
- `nbin` number of bins.
- `xlab` x label.
- `ylab` y label.
- `main` title.
- `indName` when faceted by a RleList, name used for labeling faceted factor. Default is 'sample'.
stat_bin

geom  geometric types.
type  statistical summary method used within bins, shown as bar height or heatmap colors.

Value

a ggplot object.

Author(s)

Tengfei Yin

Examples

library(IRanges)
lambda <- c(rep(0.001, 4500), seq(0.001, 10, length = 500),
           seq(10, 0.001, length = 500))
xVector <- rpois(1e4, lambda)
xRle <- Rle(xVector)
xRleList <- RleList(xRle, 2L * xRle)

ggplot() + stat_bin(xRle)
ggplot(xRle) + stat_bin()
ggplot(xRle) + stat_bin(nbin = 100)
ggplot(xRle) + stat_bin(binwidth = 200)

p1 <- ggplot(xRle) + stat_bin(type = "viewMeans")
p2 <- ggplot(xRle) + stat_bin(type = "viewSums")
## y scale are different.
tracks(viewMeans = p1, viewSums = p2)

ggplot(xRle) + stat_bin(geom = "heatmap")
ggplot(xRle) + stat_bin(nbin = 100, geom = "heatmap")
ggplot(xRle) + stat_bin(binwidth = 200, geom = "heatmap")

## for RleList
ggplot(xRleList) + stat_bin()
ggplot(xRleList) + stat_bin(nbin = 100)
ggplot(xRleList) + stat_bin(binwidth = 200)

p1 <- ggplot(xRleList) + stat_bin(type = "viewMeans")
p2 <- ggplot(xRleList) + stat_bin(type = "viewSums")
## y scale are different.
tracks(viewMeans = p1, viewSums = p2)

ggplot(xRleList) + stat_bin(geom = "heatmap")
ggplot(xRleList) + stat_bin(nbin = 100, geom = "heatmap")
ggplot(xRleList) + stat_bin(binwidth = 200, geom = "heatmap")
Calculate coverage.

Usage

# for GRanges
## S4 method for signature 'GRanges'
stat_coverage(data, ..., xlim, xlab, ylab, main,
              facets = NULL, geom = NULL)

# for GRangesList
## S4 method for signature 'GRangesList'
stat_coverage(data, ..., xlim, xlab, ylab, main,
              facets = NULL, geom = NULL)

# for Bamfile
## S4 method for signature 'BamFile'
stat_coverage(data, ..., maxBinSize = 2^14,
              xlim, which, xlab, ylab, main, facets = NULL, geom = NULL,
              method = c("estimate", "raw"),
              space.skip = 0.1, coord = c("linear", "genome"))

Arguments

data A GRanges or data.frame object.

... Extra parameters such as aes() passed to geom_rect, geom_alignment, or geom_segment.

xlim Limits for x.

xlab Label for x

ylab Label for y

main Title for plot.

facets Faceting formula to use.

geom The geometric object to use display the data.

maxBinSize maxBinSize.

method ‘estimate’ for parsing estimated coverage(fast), ‘raw’ is slow and parse the accurate coverage.

which GRanges which defines region to subset the results.

space.skip used for coordinate genome, skip between chromosomes.

coord coordinate system.
**Value**

A `Layer`.

**Author(s)**

Tengfei Yin

**Examples**

```r
library(ggbio)
library(GenomicRanges)

set.seed(1)
N <- 1000

gr <- GRanges(seqnames = sample(c("chr1", "chr2", "chr3"), size = N, replace = TRUE),
IRanges(start = sample(1:300, size = N, replace = TRUE),
width = sample(70:75, size = N, replace = TRUE),
strand = sample(c("+", "-", "x"), size = N, replace = TRUE),
value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
sample = sample(c("Normal", "Tumor"), size = N, replace = TRUE),
pair = sample(letters, size = N, replace = TRUE))

ggplot(gr) + stat_coverage()

ggplot() + stat_coverage(gr)

ggplot(gr) + stat_coverage(geom = "point")

ggplot() + stat_coverage(geom = "area")

ggplot(gr) + stat_coverage(aes(y = ..coverage..), geom = "bar")

ggplot(gr) + stat_coverage(aes(y = ..coverage..)) + geom_point()

## for bam file
## TBD
```

---

**stat_gene**

*Calculate gene structure*

**Description**

Calculate gene structure.
Usage

## S4 method for signature 'TxDb'
stat_gene(data, ...)

Arguments

data A GRanges or data.frame object.

... Extra parameters such as aes() passed to geom_alignment.

Value

A 'Layer'.

Author(s)

Tengfei Yin

See Also

geom_alignment

Examples

## Not run:
## loading package
## Deprecated
library(TxDb.Hsapiens.UCSC.hg19.knownGene)
data(genesymbol, package = "biovizBase")
txdb <- TxDb.Hsapiens.UCSC.hg19.knownGene

## made a track comparing full/reduce stat.
p1 <- ggplot() + geom_alignment(txdb, which = genesymbol["RBM17"])
p1 <- ggplot() + stat_gene(txdb, which = genesymbol["RBM17"])
## or
p1 <- ggplot(txdb) + stat_gene(which = genesymbol["RBM17"])
p1 <- ggplot(txdb) + stat_gene(which = genesymbol["RBM17"])
## ggplot(txdb) + geom_alignment(which = genesymbol["RBM17"])
## ggplot(txdb) + geom_alignment(which = genesymbol["RBM17"])
tracks(full = p1, reduce = p2, heights = c(3, 1))

## change y labels
ggplot(txdb) + stat_gene(which = genesymbol["RBM17"], names.expr = "tx_id:::gene_id")

## End(Not run)
Transform the data to a data.frame and for multiple geoms.

Description

Transform the data to a suitable data.frame and then one could use multiple geom or even stat to re-plot the data.

Usage

## S4 method for signature 'ANY'
stat_identity(data, ...)

## S4 method for signature 'GRanges'
stat_identity(data, ..., geom = NULL)

## S4 method for signature 'Rle'
stat_identity(data, ..., xlab, ylab, main, geom = NULL)

## S4 method for signature 'RleList'
stat_identity(data, ..., xlab, ylab, main,
              geom = NULL, indName = "sample")

Arguments

data        Typically a GRanges or data.frame object.
...         Extra parameters such as aes() passed to geom_rect, geom_alignment, or geom_segment.
geom        The geometric object to use display the data.
xlab        x label.
ylab        y label.
main        title of graphic..
indName     sample name.

Value

A 'Layer'.

Author(s)

Tengfei Yin
Examples

```r
## load
set.seed(1)
N <- 50

require(GenomicRanges)
## simul
## simmulated GRanges
## geom_point_start
gr <- GRanges(seqnames = sample(c("chr1", "chr2", "chr3"), size = N, replace = TRUE),
              IRanges(
                start = sample(1:300, size = N, replace = TRUE),
                width = sample(70:75, size = N, replace = TRUE),
                strand = sample(c("+", "-", "*"), size = N, replace = TRUE),
                value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
                sample = sample(c("Normal", "Tumor"), size = N, replace = TRUE),
                pair = sample(letters, size = N, replace = TRUE))

## or more formal
ggplot(gr) + stat_identity(aes(x = start, y = value), geom = "point")

## geom_point_midpoint
ggplot(gr) + stat_identity(aes(x = midpoint, y = value), geom = "point")

## geom_rect_all
ggplot(gr) + stat_identity(aes(xmin = start, xmax = end,
                               ymin = value - 0.5, ymax = value + 0.5),
                           geom = "rect")

## geom_rect_y
ggplot(gr) + stat_identity(aes(y = value), geom = "rect")

## geom_line
ggplot(gr) + stat_identity(aes(x = start, y = value), geom = "line")

## geom_segment
ggplot(gr) + stat_identity(aes(y = value), geom = "segment")

## Rle/RleList
library(IRanges)
lambda <- c(rep(0.001, 4500), seq(0.001, 10, length = 500),
            seq(10, 0.001, length = 500))
xVector <- rpois(1e4, lambda)
xRle <- Rle(xVector)
```
**stat_mismatch**

```r
xRleList <- RleList(xRle, 2L * xRle)

ggplot(xRle) + stat_identity(geom = "point")
ggplot(xRleList) + stat_identity(geom = "point")
```

---

**Description**

Calculate mismatch summary

**Usage**

```r
## for GRanges
## S4 method for signature 'GRanges'
stat_mismatch(data, ..., bsgenome, 
  xlab, ylab, main, 
  geom = c("segment", "bar"),
  show.coverage = TRUE)

## for BamFile
## S4 method for signature 'BamFile'
stat_mismatch(data, ..., bsgenome, which, 
  xlab, ylab, main, 
  geom = c("segment", "bar"),
  show.coverage = TRUE)
```

**Arguments**

- `data`     A GRanges or BamFile object.
- `...` Extra parameters such as aes() passed to geom_rect, geom_alignment, or geom_segment.
- `bsgenome` BSgenome object.
- `which` GRanges object to subset the data.
- `xlab` Label for x
- `ylab` Label for y
- `main` Title for plot.
- `geom` The geometric object to use display the data.
- `show.coverage` whether to show coverage as background or not.

**Value**

A 'Layer'.

**Author(s)**

Tengfei Yin
stat_reduce

Reduce GRanges, IRanges or TxDb object.

Usage

```r
## S4 method for signature 'GRanges'
stat_reduce(data, ...,
    xlab, ylab, main,
    drop.empty.ranges = FALSE,
    min.gapwidth = 1L,
    facets = NULL, geom = NULL)

## S4 method for signature 'IRanges'
stat_reduce(data, ...,
    xlab, ylab, main,
    drop.empty.ranges = FALSE,
    min.gapwidth = 1L,
    with.inframe.attrib=FALSE,
    facets = NULL, geom = NULL)

## S4 method for signature 'TxDbOREnsDb'
stat_reduce(data, ...)
```

Arguments

- `data` GRanges, IRanges or TxDb object.
- `...` passed to aesthetics mapping.
- `xlab` x label.
- `ylab` y label.
- `main` title.
- `drop.empty.ranges` pass to `reduce` function.
- `min.gapwidth` pass to `reduce` function.
- `with.inframe.attrib` pass to `reduce` function.
- `facets` pass to `reduce` function.
- `geom` geometric type.

Value

a ggplot object.
stat_slice

Author(s)
Tengfei Yin

See Also
reduce.

Examples

```r
set.seed(1)
N <- 1000
library(GenomicRanges)

gr <- GRanges(seqnames =
  sample(c("chr1", "chr2", "chr3"),
  size = N, replace = TRUE),
IRanges(
  start = sample(1:300, size = N, replace = TRUE),
  width = sample(70:75, size = N, replace = TRUE)),
  strand = sample(c("+", "-", "*"), size = N, replace = TRUE),
  value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
  sample = sample(c("Normal", "Tumor"),
  size = N, replace = TRUE),
  pair = sample(letters, size = N, replace = TRUE))

ggplot(gr) + stat_reduce()
autoplot(gr, stat = "reduce")
strand(gr) <- "*"

library(TxDb.Hsapiens.UCSC.hg19.knownGene)
data(genesymbol, package = "biovizBase")
txdb <- TxDb.Hsapiens.UCSC.hg19.knownGene
## made a track comparing full/reduce stat.
ggplot(txdb) + stat_reduce(which = genesymbol["RBM17"])
```

stat_slice

Slice Rle/RleList to view them as bar or heatmap.

Description

Slice Rle/RleList to different view by set lower or other parameters, then view summary for all those viewed region.
Usage

```r
## S4 method for signature 'Rle'
stat_slice(data, ...,
  xlab, ylab, main,
  na.rm = FALSE,
  geom = NULL,
  lower=-Inf, upper=Inf,
  includeLower=TRUE, includeUpper=TRUE,
  rangesOnly = FALSE,
  type = c("viewSums","viewMins",
           "viewMaxs", "viewMeans"))

## S4 method for signature 'RleList'
stat_slice(data, ...,
  xlab, ylab, main,
  indName = "sample",
  na.rm = FALSE,
  geom = NULL,
  lower=-Inf, upper=Inf,
  includeLower=TRUE, includeUpper=TRUE,
  rangesOnly = FALSE,
  type = c("viewSums","viewMins",
           "viewMaxs", "viewMeans"))
```

Arguments

data a data.frame or Rle or RleList object.
...
arguments passed to aesthetics mapping.
xlab x label.
ylab y label.
main title.
indName when faceted by a RleList, name used for labeling faceted factor. Default is 'sample'.
geom geometric types.
type statistical summary method used within bins, shown as bar height or heatmap colors.
na.rm logical value, default FALSE, passed to function like viewMaxs for statistical summary computation.
lower passed to slice.
upper passed to slice.
includeLower passed to slice.
includeUpper passed to slice.
rangesOnly passed to slice.
stat_stepping

Calculate stepping levels

Value

a ggplot object.

Author(s)

Tengfei Yin

See Also

slice

Examples

library(IRanges)
lambda <- c(rep(0.001, 4500), seq(0.001, 10, length = 500),
           seq(10, 0.001, length = 500))
xVector <- rpois(1e4, lambda)
xRle <- Rle(xVector)
xRleList <- RleList(xRle, 2L * xRle)

ggplot(xRle) + stat_slice(lower = 5)
ggplot(xRle) + stat_slice(lower = 5, geom = "bar")
ggplot(xRle) + stat_slice(lower = 5, geom = "heatmap")

p1 <- ggplot(xRle) + stat_slice(type = "viewMeans", lower = 5,
                                geom = "bar")
p2 <- ggplot(xRle) + stat_slice(type = "viewSums", lower = 5,
                                geom = "bar")
## y scale are different.
tracks(viewMeans = p1, viewSums = p2)

ggplot(xRleList) + stat_slice(lower = 5)
ggplot(xRleList) + stat_slice(lower = 5, geom = "bar")
ggplot(xRleList) + stat_slice(lower = 5, geom = "heatmap")

p1 <- ggplot(xRleList) + stat_slice(type = "viewMeans", lower = 5,
                                geom = "bar")
p2 <- ggplot(xRleList) + stat_slice(type = "viewSums", lower = 5,
                                geom = "bar")
## y scale are different.
tracks(viewMeans = p1, viewSums = p2)
Usage

```r
## S4 method for signature 'GRanges'
stat_stepping(data, ..., xlab, ylab, main,
              facets = NULL,
              geom = c("rect", "alignment", "segment"))
```

Arguments

- `data`: A `GRanges` or `data.frame` object.
- `...`: Extra parameters such as `aes()` passed to `geom_rect`, `geom_alignment`, or `geom_segment`.
- `xlab`: Label for x
- `ylab`: Label for y
- `main`: Title for plot.
- `facets`: Faceting formula to use.
- `geom`: The geometric object used to display the data. For 'stepping', could be one of 'rect', 'alignment', 'segment'.

Value

A `Layer`.

Author(s)

Tengfei Yin

Examples

```r
set.seed(1)
N <- 50
require(GenomicRanges)
## simmulated GRanges
gr <- GRanges(sequences =
              sample(c("chr1", "chr2", "chr3"),
                     size = N, replace = TRUE),
              IRanges(  
                        start = sample(1:300, size = N, replace = TRUE),
                        width = sample(70:75, size = N, replace = TRUE),
                        strand = sample(c("+", "-", "x"), size = N,
                                        replace = TRUE),
                        value = rnorm(N, 10, 3),
                        score = rnorm(N, 100, 30),
                        sample = sample(c("Normal", "Tumor"),
                                        size = N, replace = TRUE),
                        pair = sample(letters, size = N,
                                      replace = TRUE)))
```
## default
ggplot(gr) + stat_stepping()
## or
ggplot() + stat_stepping(gr)

## facet_aes
ggplot(gr) + stat_stepping(aes(color = strand, fill = strand),
   facets = sample ~ seqnames)

## geom_segment
## ggplot(gr) + stat_stepping(aes(color = strand),
##   geom = "segment", xlab = "Genomic coord", ylab = "y", main = "hello")

## geom_alignment
## ggplot(gr) + stat_stepping(geom = "alignment")
## geom_alignment_group
## ggplot(gr) + stat_stepping(aes(group = pair), geom = "alignment")

---

### stat_table

**Tabulate a GRanges object**

**Description**

Tabulate a GRanges object

**Usage**

```r
## S4 method for signature 'GRanges'
stat_table(data, ..., xlab, ylab, main,
geom = NULL, stat = NULL)
## S4 method for signature 'GRangesList'
stat_table(data, ..., xlab, ylab, main,
facets = NULL, geom = NULL)
```

**Arguments**

- `data`:
  A GRanges or data.frame object.
- `...`:
  Extra parameters such as aes() passed to geom_rect, geom_alignment, or geom_segment.
- `xlab`:
  Label for x.
- `ylab`:
  Label for y.
- `main`:
  Title for plot.
- `facets`:
  Faceting formula to use.
- `geom`:
  The geometric object to use display the data.
- `stat`:
  The geometric object to use display the data.
Value

A 'Layer'.

Author(s)

Tengfei Yin

Examples

```r
## load
set.seed(1)
N <- 100
require(ggbio)
require(GenomicRanges)
## simul
## ======================================================================
## simmulated GRanges
## ======================================================================
gr <- GRanges(seqnames =
  sample(c("chr1", "chr2", "chr3"),
  size = N, replace = TRUE),
IRanges(
  start = sample(1:300, size = N, replace = TRUE),
  width = sample(70:75, size = N, replace = TRUE)),
strand = sample(c("+", "-", "x"), size = N,
  replace = TRUE),
value = rnorm(N, 10, 3), score = rnorm(N, 100, 30),
sample = sample(c("Normal", "Tumor"),
  size = N, replace = TRUE),
pair = sample(letters, size = N,
  replace = TRUE))
gr <- c(gr[seqnames(gr) == "chr1"][
sample(1:10, size = 1e4, replace = TRUE)], gr)
## default
ggplot(gr) + stat_table()
ggplot(gr) + stat_table(geom = "segment", aes(y = ..score.., color = ..score..))
```

theme

theme in ggbio

Description

Theme defined in ggbio for plot or tracks.
theme

Usage

theme_null()
theme_noexpand()
theme_alignment(ylabel = FALSE, base_size = 12, base_family = "", axis = TRUE, border = TRUE, grid = TRUE)
theme_pack_panels(strip.bg = FALSE, strip.text.y = TRUE)
theme_clear(grid.y = FALSE, grid.x.minor = FALSE, grid.x.major = FALSE,
panel.background.fill = "white", panel.border.color = NA,
axis.ticks.x = FALSE, axis.ticks.y = TRUE, grid.color = "gray95",
axis.line.color = "gray80")
theme_tracks_sunset(bg = "#fffed", alpha = 1, ...)
theme_genome()

Arguments

alpha alpha blending from 0(transparent) to 1(solid).
axis logical value, show axis or not.
axis.line.color color for axis line.
axis.ticks.x show x ticks or not.
axis.ticks.y show y ticks or not.
base_family family for font.
base_size size for font.
bg background color for tracks.
border logical value, show border or not.
grid logical value, show background grid or not.
grid.color grid line color.
grid.x.major show x major grid line or not.
grid.x.minor show x minor grid line or not.
grid.y show y grid or not.
panel.background.fill panel background fill color.
panel.border.color panel border color.
strip.bg if strip background is removed.
strip.text.y if strip text is removed.
ylabel logical value. Show labels or not.
... passed to theme_clear.

details

Themes speciall designed for tracks, are named following naming schema theme_tracks_*
Value

Return a theme.

Author(s)

Tengfei Yin

Examples

```r
## load
library(ggbiogram)
p <- qplot(data = mtcars, x = mpg, y = wt, facets = cyl ~ .)
p + theme_null()
p + theme_clear()
p + theme_pack_panels()
p + theme_alignment()
p1 <- qplot(data = mtcars, x = mpg, y = wt)
tracks(p1 = p, p2 = p1)
tracks(p1 = p, p2 = p1) + theme_tracks_sunset()
```

---

**Tracked**

*Tracked class*

Description

Create a tracked object, designed for tracks function.

Usage

```r
Tracked(mutable = TRUE, fixed = FALSE, labeled = TRUE,
hasAxis = FALSE, bgColor = "white", height = unit(1, "null"))
```

Arguments

- **mutable** logical value, default TRUE. To control whether a track is updatable by applying + on it.
- **fixed** logical value, default FALSE. To control whether the scale response to a xlim change or not.
- **labeled** logical value, default TRUE. To control whether to label it all not.
- **hasAxis** logical value, default FALSE. To control whether to show axis for that track or not.
- **bgColor** character to control background color of a track.
- **height** unit, to control track height.

Value

a Tracked object.
tracks

Author(s)

Tengfei Yin

tracks

Tracks for genomic graphics

Description

tracks is a convenient constructor for binding graphics as tracks. You don’t have to worry about adjusting different graphics, tracks did that for you. It’s NOT just limited to bind genomic tracks, you can use this function to bind any tracks with the same definition of x axis, for example, sets of time series plots you made.

Tracks view is most common way to viewing genome features and annotation data and widely used by most genome browsers. Our assumption is that, most graphics you made with ggbio or by yourself using ggplot2, are almost always sitting on the genomic coordinates or the same x axis. And to compare annotation information along with genome features, we need to align those plots on exactly the same x axis in order to form your hypothesis. This function leaves you the flexibility to construct each tracks separately with worrying your alignments later.

Usage

tracks(..., heights, xlim, xlab = NULL, main = NULL, title = NULL, theme = NULL, track.plot.color = NULL, track.bg.color = NULL, main.height = unit(1.5, "lines"), scale.height = unit(1, "lines"), xlab.height = unit(1.5, "lines"), padding = unit(-1, "lines"), label.bg.color = "white", label.bg.fill = "gray80", label.text.color = "black", label.text.cex = 1, label.text.angle = 90, label.width = unit(2.5, "lines"))

Arguments

... plots of class ggplot, generated from ggplot2 or ggbio.
heights numeric vector of the same length of passed graphic object to indicate the ratio of each track.
xlim limits on x. could be IRanges, GRanges, numeric value
xlab label for x axis.
main title for the tracks.
title title for the tracks, alias like main.
theme

theme object used for building tracks, this will set to default, which could be reseted later.

track.plot.color

Vector of characters of length 1 or the same length of passed plots, background color for each track, default is white.

track.bg.color

background color for the whole tracks.

main.height

unit. Height to control the title track height.

scale.height

unit. Height to control the scale track height.

xlab.height

unit. Height to control the xlab track height.

padding

single numeric value or unit, if numeric value, the unit would be "lines" by default.

label.bg.color

track labeling background rectangle border color.

label.bg.fill

track labeling background fill color.

label.text.color

track labeling text color.

label.text.cex

track labeling text size.

label.text.angle

angle to rotate the track labels.

label.width

track labeling size.

Details

tracks did following modification for passed plots.

- remove x-axis, ticks, xlab and tile for each track and add scales at bottom. We suppose a new xlab and title would be provided by the tracks function for the whole tracks, but we still keep individual’s y axis.
- align x-scale limits to make sure every plots sitting on exactly the same x scale.
- squeezing plots together to some extent.
- labeling tracks if names are provided, please check utilities section about labeled method.
- return a track object. This would allow many features introduced in this manual.

Value

A Tracks object.

Track class

constructor tracks will return a Tracks object, which has following slots.

- grobs: a ggplotGrobList object contains a list of ggplot object, which is our passed graphics.
- backup: a backup of all the slots for holding the original tracks, so users could edit it and reset it back at any time later, and backup method will reset the backedup copy.
- ylim: y limits for each plot.
labeled vector of logical value indicates whether a track is labeled or not, for labeled attributes please check utilities section.

mutable vector of logical value indicates whether a track is mutable for theme editing or not, for mutable attributes please check utilities section.

hasAxis vector of logical value indicates whether a track has axis or not, for hasAxis attributes please check utilities section.

heights, xlim, xlab, main, title, theme, fixed, track.plot.color, track.bg.color, main.height, scale.height, xlab.height, padding, label.bg.color, label.bg.fill, label.text.color, label.text.cex, label.text.angle, label.width those slots are described in arguments section for constructor.

Utilities

Please check examples for usage.

summary(object) summary information about tracks object.

fixed(x), fixed(x) <- value x is the ggplot object, this controls if a track has a fixed x scale or not, if the fixed attributes is TRUE, then when you pass this plot to a tracks, this plot won’t be re-aligned with other tracks and will keep the original x-axis, this allow you to pass some plot like ideogram. fixed function will return a logical value

labeled(x), labeled(x) <- value x is the ggplot object, if you pass named graphics into tracks, it will create the labels on the left for you. Several ways supported to name it. You can pass a list of graphics with names. Or you can use tracks('name1' = p1, 'name 2' = p2, ...) with quotes for complicated words or simply tracks(part1 = p1, part = p2, ...).

mutable(x), mutable(x) <- value x is the ggplot object, this controls whether a plot in the tracks mutable to theme changing or not, when you use + method for Tracks object, add-on edit will only be applied to the the mutable plots.

bgColor(x), bgColor(x) <- value x is the ggplot object, this change the background color for single plot shown in the tracks.

xlim(x), xlim(x) <- value when x is the numeric value, it calls ggplot2::coord_cartesian(xlim = ...) method, we doesn’t use ggplot2::xlim() for the reason it will cut data outside the range, and we believe the best behavior would be zoom-in/out like most browser. when x is IRanges, GRanges, it get the range and passed to ggplot2::coord_cartesian function. when x is Tracks object, xlim(x) will return x limits for that tracks. xlim(x) <- value replace method only works for Tracks object. value could be numeric, IRanges, GRanges object. This will change the x limits associated with tracks.

+ xlim(obj)obj is the numeric range, or IRanges, GRanges object.
+ coord_cartesian(): please read manual in ggplot2, this controls both xlim an ylim, only accept numerical range.

+ The most nice features about Tracks object is the one inherited from ggplot2’s components additive features, with + method you can use any theme object and utilities in ggplot2 package, to add them on a Tracks object, for example, if x is our Tracks object, x + theme would apply theme to any plots in the tracks except those are immutable.

as(x, "grob") Coerces a Tracks object to a grob for embedding in a larger figure.
Backup and reset

reset(obj)  obj is the Tracks object, this reset the tracks back to original or backuped version.

backup(obj)  obj is the Tracks object, this clear previous backup and use current setting for a new backup.

Author(s)

Tengfei Yin

See Also

align.plots

Examples

```r
## make a simulated time series data set
df1 <- data.frame(time = 1:100, score = sin((1:100)/20)*10)
p1 <- qplot(data = df1, x = time, y = score, geom = "line")
df2 <- data.frame(time = 30:120, score = sin((30:120)/20)*10, value = rnorm(120-30 + 1))
p2 <- ggplot(data = df2, aes(x = time, y = score)) +
  geom_line() + geom_point(size = 4, aes(color = value))
## check p2
p1
## check p2
p2

## binding
tracks(p1, p2)

## or
kts <- tracks(p1, p2)
kts

## combine
c(kts, kts)
kts + kts

cbind(kts, kts)
rbind(kts, kts)  ## different with c()!
library(grid)
x <- as(kts, "grob")
grid.draw(cbind(x, x))

## labeling: default labeling a named graphic
## simply pass a name with it
tracks(time1 = p1, time2 = p2)
## or pass a named list with it
lst <- list(time1 = p1, time2 = p2)
tracks(lst)
```
## more complicated case please use quotes
tracks(time1 = p1, "second time" = p2)

## set heights
tracks(time1 = p1, time2 = p2, heights = c(1, 3))

## if you want to disable label arbitrarily
## default label is always TRUE
labeled(p2)
labeled(p2) <- FALSE

## set labeled to FALSE, remove label even the plot has a name
tracks(time1 = p1, time2 = p2)
labeled(p2) <- TRUE

## fix a plot, not synchronize with other plots
p3 <- p1
## default is always FALSE
fixed(p3)
## set to TRUE
fixed(p3) <- TRUE
fixed(p3)

tracks(time1 = p1, time2 = p2, "time3(fixed)" = p3)

fixed(p3) <- FALSE
## otherwise you could run

## control axis
hasAxis(p1)
hasAxis(p1) <- TRUE
# ready for weird looking
tracks(time1 = p1, time2 = p2)
# set it back
hasAxis(p1) <- FALSE

## mutable
mutable(p1)
tracks(time1 = p1, time2 = p2) + theme_bw()
mutable(p1) <- FALSE
# mutable for "+" method
tracks(time1 = p1, time2 = p2) + theme_bw()
mutable(p1) <- TRUE

## bgColor
bgColor(p1)
tracks(time1 = p1, time2 = p2)
bgColor(p1) <- "brown"
# mutable for "+" method
tracks(time1 = p1, time2 = p2)
# set it back
bgColor(p1) <- "white"

## apply a theme to each track
## store it with tracks
## apply a pre-defined theme for tracks!
tracks(time1 = p1, time2 = p2) + theme_tracks_sunset()
tracks(p1, p2) + theme_tracks_sunset()

## change limits
tracks(time1 = p1, time2 = p2) + xlim(c(1, 40))
tracks(time1 = p1, time2 = p2) + xlim(1, 40)
tracks(time1 = p1, time2 = p2) + coord_cartesian(xlim = c(1, 40))
# change y
tracks(time1 = p1, time2 = p2) + xlim(1, 40) + ylim(0, 10)
library(GenomicRanges)
gr <- GRanges("chr", IRanges(1, 40))
## change limits
tracks(time1 = p1, time2 = p2) + xlim(gr)
tracks(time1 = p1, time2 = p2) + xlim(ranges(gr))
tracks(time1 = p1, time2 = p2) + xlim(1, 40)

## xlab, title
tracks(time1 = p1, time2 = p2, xlab = "time")
tracks(time1 = p1, time2 = p2, main = "title")
tracks(time1 = p1, time2 = p2, title = "title")
tracks(time1 = p1, time2 = p2, xlab = "time", title = "title") + theme_tracks_sunset()

## backup and restore
tracks(time1 = p1, time2 = p2)
tracks(time1 = p1, time2 = p2) + xlim(1, 40)
reset(tks)
tks <- tks + xlim(1, 40)
```
tks
  tks <- backup(tks)
  tks <- tks + theme_bw()
  tks
  reset(tks)

  ## padding (need to be fixed for more delicate control)
  tracks(time1 = p1, time2 = p2, padding = 2)

  ## track color
  tracks(time1 = p1, time2 = p2, track.bg.color = "yellow")
  tracks(time1 = p1, time2 = p2, track.plot.color = c("yellow", "brown"))
```

---

**zoom**  
*Simple navigation for ggbio object.*

**Description**  
A set of simple navigation API apply to ggbio object, let you move along the genome and zoom in/out.

**Usage**

```r
zoom(fac = 1/2)
zoom_in(fac = 1/2)
zoom_out(fac = 2)
nextView(unit = c("view", "gene", "exon", "utr"))
prevView(unit = c("view", "gene", "exon", "utr"))
```

**Arguments**

- **fac**  
  numeric value to indicate zoom factor, multiple of current view width. If it’s smaller than 1, then it’s zoom-in operation; if it’s bigger than 1, then it’s zoom-out operation.

- **unit**  
  only support `view` unit now.

**Details**

`zoom_in` and `zoom_out` are just simple wrapper around `zoom` function.  
For more convenient, gene features based jumpting we will support it in the future.

**Value**

A special class of navigation.

**Author(s)**

Tengfei Yin
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