

Package ‘coMET’

April 10, 2023

Type Package

Title coMET: visualisation of regional epigenome-wide association scan (EWAS) results and DNA co-methylation patterns

Version 1.30.0

Date 2022-10-26

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Description Visualisation of EWAS results in a genomic region. In addition to phenotype-association P-values, coMET also generates plots of co-methylation patterns and provides a series of annotation tracks. It can be used to other omic-wide association scans as long as the data can be translated to genomic level and for any species.

Depends R (>= 4.1.0), grid, utils, biomaRt, Gviz, psych

Suggests BiocStyle, knitr, RUnit, BiocGenerics, showtext

Imports hash,grDevices, gridExtra, rtracklayer, IRanges, S4Vectors, GenomicRanges, stats, corrplot

License GPL (>= 2)

URL <http://epigen.kcl.ac.uk/comet>

biocViews Software, DifferentialMethylation, Visualization, Sequencing, Genetics, FunctionalGenomics, Microarray, MethylationArray, MethylSeq, ChIPSeq, DNASEq, RiboSeq, RNASeq, ExomeSeq, DNAMethylation, GenomeWideAssociation, MotifAnnotation

VignetteBuilder knitr

NeedsCompilation no

Repository Bioconductor

git_url <https://git.bioconductor.org/packages/coMET>

git_branch RELEASE_3_16

git_last_commit c75985c

git_last_commit_date 2022-11-01

Date/Publication 2023-04-10

R topics documented:

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coMET-package	<i>visualisation of regional epigenome-wide association scan (EWAS) results and DNA co-methylation patterns (and also for other omic-WAS)</i>
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Description

coMET is an R package for visualising EWAS results in a genomic region. Along with phenotype-association plots, coMET also generates plots of co-methylation patterns and provides a series of annotation tracks. The software is designed for epigenetic data, but can also be applied to genomic and functional genomic datasets (other omic-WAS results) in any species.

Details

Package: coMET
 Type: Package
 Version: 1.11.5
 Date: 2018-04-16
 License: GPL (>=2)

coMET is an R package that can generate regional plots of EWAS results, DNA co-methylation patterns, and genomic information. A coMET figure includes 3 panels with a plot of P-values from EWAS, customized annotation tracks, and a triangle heatmap plot which demonstrates the correlation structure of DNA methylation at the CpG sites in the genomic region. Plots are created as PDF or EPS files.

Author(s)

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Maintainer: Tiphaine Martin <tiphaine.martin@mssm.edu>

Website: <http://www.epigen.kcl.ac.uk/comet>

References

Martin, T.C, Yet, I, Tsai, P-C, Bell, J.T., coMET: visualisation of regional epigenome-wide association scan results and DNA co-methylation patterns, BMC bioinformatics, 2015.

Examples

```

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
configfile <- file.path(extdata, "config_cyp1b1_zoom_4comet.txt")
myinfofile <- file.path(extdata, "cyp1b1_infofile.txt")
myexpressfile <- file.path(extdata, "cyp1b1_infofile_exprGene_region.txt")
mycorrelation <- file.path(extdata, "cyp1b1_res37_rawMatrix.txt")

chrom <- "chr2"
start <- 38290160
end <- 38303219
gen <- "hg38"

if(interactive()){
  genetrack <- genes_ENSEMBL(gen, chrom, start, end, showId=TRUE)
  snptrack <- snpBiomart_ENSEMBL(gen, chrom, start, end,
    dataset="hsapiens_snp_som", showId=FALSE)
  strutrack <- structureBiomart_ENSEMBL(gen, chrom, start, end,
    strand, dataset="hsapiens_structvar_som")
  clinVariant <- ClinVarMain_UCSC(gen, chrom, start, end)
  clinCNV <- ClinVarCnv_UCSC(gen, chrom, start, end)
  gwastrack <- GWAScatalog_UCSC(gen, chrom, start, end)
  geneRtrack <- GeneReviews_UCSC(gen, chrom, start, end)
  listgviz <- list(genetrack, snptrack, strutrack, clinVariant,
    clinCNV, gwastrack, geneRtrack)
  comet(config.file=configfile, mydata.file=myinfofile, mydata.type="file",
    cormatrix.file=mycorrelation, cormatrix.type="listfile",
    mydata.large.file=myexpressfile, mydata.large.type="listfile",
    tracks.gviz=listgviz, verbose=FALSE, print.image=FALSE, disp.pvalueplot=FALSE)
} else {
  data(geneENSEMBLtrack)
  data(snpBiomarttrack)
  data(ISCATrack)
  data(strucBiomarttrack)
  data(ClinVarCnvTrack)
  data(clinVarMaintrack)
  data(GWASTrack)
  data(GeneReviewTrack)

  listgviz <- list(genetrack, snptrack, strutrack, clinVariant,
    clinCNV, gwastrack, geneRtrack)
  comet(config.file=configfile, mydata.file=myinfofile, mydata.type="listfile",
    cormatrix.file=mycorrelation, cormatrix.type="listfile",
    mydata.large.file=myexpressfile, mydata.large.type="listfile",
    tracks.gviz=listgviz,
    verbose=FALSE, print.image=FALSE, disp.pvalueplot=TRUE)
}

```

 bindingMotifsBiomart_ENSEMBL

Creates a binding motif track from ENSEMBL

Description

Creates a binding motif track from ENSEMBL using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
bindingMotifsBiomart_ENSEMBL(gen, chr, start, end, featureDisplay="all",
  datasetEnsembl = NULL, title="Binding Motifs ENSEMBL")
```

Arguments

gen	The name of the genome. Currently only handles human data from either the previous version, GRCh37 (also known as hg19) or the current version, GRCh38 (also known as hg38).
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
featureDisplay	A vector of regulatory features to be displayed, such as Egr1. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. featureDisplay <- "CTCF"), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. featureDisplay <- c("Egr1","CTCF")). Finally, visualisation all features in the genomic region, achieved by using the word "all" (e.g. featureDisplay <- "all"), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
datasetEnsembl	Allows the user to manually set which data set is used if required.
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to ENSEMBLregulation binding motif biomart

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 10000
end <- 50000
featureDisplay <- "CTCF"

if(interactive()){
  bindMotifsBiomartTrackSingle<-bindingMotifsBiomart_ENSEMBL(gen,chr,start,
  end,featureDisplay)
  plotTracks(bindMotifsBiomartTrackSingle, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
} else {
  data(bindMotifsBiomartTrackSingle)
  plotTracks(bindMotifsBiomartTrackSingle, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
}

#####

library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 10000
end <- 50000
featureDisplay <- c("CTCF","Egr1")

if(interactive()){
  bindMotifsBiomartTrackMultiple<-bindingMotifsBiomart_ENSEMBL(gen,chr,start,end,featureDisplay)
  plotTracks(bindMotifsBiomartTrackMultiple, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
} else {
  data(bindMotifsBiomartTrackMultiple)
  plotTracks(bindMotifsBiomartTrackMultiple, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
}

#####

library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 10000
end <- 50000
featureDisplay <- "all"
```

```

if(interactive()){
  bindMotifsBiomartTrackAll<-bindingMotifsBiomart_ENSEMBL(gen,chr,start,end,featureDisplay)
  plotTracks(bindMotifsBiomartTrackAll, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
} else {
  data(bindMotifsBiomartTrackAll)
  plotTracks(bindMotifsBiomartTrackAll, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
}

```

ChIPTF_ENCODE

Creates a TF motif track from ENCODE

Description

Creates a track of TF motifs from ENCODE using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```

ChIPTF_ENCODE(gen="hg19", chr, start, end, bedFilePath,
  featureDisplay='all', motifColorFile, type_stacking='dense',
  showId=FALSE,just_group="above", title="TF motifs ENCODE")

```

Arguments

gen	the name of the genome. Default value=hg19
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
bedFilePath	The path of the BED file from Kheradpour and Kellis, 2014.
featureDisplay	A vector of regulatory features to be displayed, such as Predicted heterochomatin. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. featureDisplay <- "Predicted heterochomatin"), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. featureDisplay <- c("Predicted low activity","Predicted heterochomatin")). Finally, visualison all features in the genomic region, achived by using the word "all" (e.g. featureDisplay <- "all"), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
motifColorFile	The path of the BED file with 2 columns (the first for motif name and the second for the color in hex format without \# in the beginning) with a header.
type_stacking	Object of class"character", the stacking type of overlapping items on the final plot.One in c(hide, dense, squish, pack,full). More information cf the option "stacking" in Gviz

showId	logical. say if we write the name of group
just_group	position. say where we write the name of group (choice in c("above","right","left"))
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to BindingMotifsBiomart binding motif biomart

Examples

```
library("Gviz")
gen <- "hg19"
chr<-"chr1"
start <- 1000
end <- 329000

if(interactive()){
  extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
  bedFilePath <- file.path(extdata, "ENCODE/motifs1000_matches_ENCODE.txt")
  motif_color <- file.path(extdata, "ENCODE/TFmotifs_colors.csv")
  chipTFtrack <- ChIPTF_ENCODE(gen,chr,start, end, bedFilePath,
  featureDisplay=c("AHR::ARNT::HIF1A_1","AIRE_1","AIRE_2","AHR::ARNT_1"),
  motif_color,type_stacking="squish",showId=TRUE)
  plotTracks(chipTFtrack, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
} else {
  data(chipTFtrack)
  plotTracks(chipTFtrack, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
}
```

chromatinHMMAll_UCSC *Creating multiple chromHMM tracks from the UCSC genome browser*

Description

Create multiple chromHMM Broad tracks by connecting to the UCSC genome browser using the Gviz bioconductor package

Usage

```
chromatinHMMAll_UCSC(gen, chr, start, end, mySession, color='coMET',
  pattern = NULL, table.name = NULL)
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in region of interest (the smallest value)
end	the last position in region of interest (the biggest value)
mySession	the object session from the function browserSession of rtracklayer
color	the colour scheme used for plots. By default this is set to 'coMET' to allow easy identification of different elements. The colour scheme set by UCSC can also be used. Consult userguide for table of colours.
pattern	the pattern of the track to visualise
table.name	the name of the table from the track

Value

list of AnnotationTrack objects of GViz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=wg

See Also

[chromatinHMMOne_UCSC](#)

Examples

```
library("Gviz")
library(rtracklayer)
gen <- "hg19"
chr <- "chr2"
start <- 38290160
end <- 38313219
if(interactive()){
  BROWSER.SESSION="UCSC"
  mySession <- browserSession(BROWSER.SESSION)
  genome(mySession) <- gen
  track.name="Broad ChromHMM"
  tabletrack<-ucscTables(gen, track=track.name)
```

```

table.name<-tablestrack[1]
PATTERN.REGULATION<-"GM12878"

chromhmmPattern<-chromatinHMMAll_UCSC(gen,chr,start,end,mySession,
color='coMET',PATTERN.REGULATION)
plotTracks(chromhmmPattern, from = start, to =end,
fontfamily="sans",fontfamily.title="sans")

chromhmmNoPattern<-chromatinHMMAll_UCSC(gen,chr,start,end,
mySession,color='coMET')
plotTracks(chromhmmNoPattern, from = start, to =end,
fontfamily="sans",fontfamily.title="sans")
} else {

data(chromhmmPattern)
plotTracks(chromhmmPattern, from = start, to =end,
fontfamily="sans",fontfamily.title="sans")

data(chromhmmNoPattern)
plotTracks(chromhmmNoPattern, from = start, to =end,
fontfamily="sans",fontfamily.title="sans")
}

```

chromatinHMMOne_UCSC *Creating one chromHMM track from the UCSC genome browser*

Description

Create one track of only one type of chromHMM Broad element from the UCSC genome browser using the Gviz bioconductor package

Usage

```
chromatinHMMOne_UCSC(gen, chr, start, end, mySession, color="coMET",
title="ENCODE/Broad chromHMM", table.name = NULL)
```

Arguments

gen	the name of the genome. Data is not currently available for GRCh38 (hg38).
chr	the chromosome of interest
start	the first position in region of interest (the smallest value)
end	the last position in region of interest (the biggest value)
mySession	the object session from the function browserSession of rtracklayer
color	the color scheme used for plots. By default this is set to 'coMET' to allow easy indentification of differnent elements. The color scheme set by UCSC can also be used. Consult userguide for table of colors.
title	Name of tracks
table.name	the name of the table from the track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtDrFAy6dn&c=chr6&g=wg

See Also

[chromatinHMMAll_UCSC](#)

Examples

```
library("Gviz")
library("rtracklayer")
gen <- "hg19"
chr <- "chr2"
start <- 38290160
end <- 38303219
color <- "coMET"

if(interactive()) {
  BROWSER.SESSION="UCSC"
  mySession <- browserSession(BROWSER.SESSION)
  genome(mySession) <- gen
  track.name="Broad ChromHMM"
  tablestrack<-ucscTables(gen, track.name)
  table.name<-tablestrack[1]
  chromhmmtrackone<-chromatinHMMOne_UCSC(gen,chr,start,end
  ,mySession,color="coMET",table.name)
  plotTracks(chromhmmtrackone, from = start, to =end,
fontfamily="sans",fontfamily.title="sans")
}else {
  data(chromhmmtrackone)
  plotTracks(chromhmmtrackone, from = start, to =end,
fontfamily="sans",fontfamily.title="sans")
}
```

chromHMM_RoadMap *Creates a ChromHMM track from a file of RoadMap*

Description

Creates a ChromHMM track from a file of RoadMap using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
chromHMM_RoadMap(gen="hg19",chr, start, end, bedFilePath,
featureDisplay = 'all', colorcase='roadmap15',
title=" chromHMM RoadMap")
```

Arguments

gen	the name of the genome. Default value=hg19
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
bedFilePath	The file path to the .BED file containing the data to be visualised
featureDisplay	A vector of features to be displayed, such as 1_TssA. Spelling and capitalisation of features must be identical to those in the user guide (in the 'State & Acronym' column). There are three possibilities. First, the visualisation of only one feature (e.g. featureDisplay <- "1_TssA"), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. featureDisplay <- c("1_TssA","2_TssAFlnk")). Finally, visualisation all features in the genomic region, achieved by using the word "all" (e.g. featureDisplay <- "all"), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
colorcase	the type of colors used to visualise different elements contained in ROADmap data with 15-,18-,25- states. choice between roadmap15, roadmap18, comet18, roadmap25 and comet25.
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin
Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to RoadMap Epigenome

Examples

```
library("Gviz")
chr <- "chr1"
start <- 4500000
end <- 4600000
featureDisplay <- "7_Enh"

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
bedFilePath <- file.path(extdata, "RoadMap/E063_15_coreMarks_mnemonics.bed")

if(interactive()){
  chromHMM_RoadMapSingle <- chromHMM_RoadMap(gen="hg19",chr,start, end,
  bedFilePath, featureDisplay = featureDisplay, colorcase='roadmap15' )
  plotTracks(chromHMM_RoadMapSingle, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
} else {
  data(chromHMM_RoadMapSingle)
  plotTracks(chromHMM_RoadMapSingle, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
}

#####

library("Gviz")
chr <- "chr22"
start <- 38291000
end <- 38301200
featureDisplay <- c("7_Enh","13_ReprPC")

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
bedFilePath <- file.path(extdata, "RoadMap/E063_15_coreMarks_mnemonics.bed")

if(interactive()){
  chromHMM_RoadMapMultiple <- chromHMM_RoadMap(gen="hg19",chr,start, end,
  bedFilePath, featureDisplay = featureDisplay, colorcase='roadmap15' )
  plotTracks(chromHMM_RoadMapMultiple, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
} else {
  data(chromHMM_RoadMapMultiple)
  plotTracks(chromHMM_RoadMapMultiple, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
}

#####

library("Gviz")
chr <- "chr22"
```

```
start <- 38291000
end <- 38301200
featureDisplay <- "all"

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
bedFilePath <- file.path(extdata, "RoadMap/E063_15_coreMarks_mnemonics.bed")

if(interactive()){
  chromHMM_RoadMapAll <- chromHMM_RoadMap(gen="hg19",chr,start, end,
    bedFilePath, featureDisplay = featureDisplay, colorcase='roadmap15' )
  plotTracks(chromHMM_RoadMapAll, from = start, to = end,
    fontfamily="sans", fontfamily.title="sans")
} else {
  data(chromHMM_RoadMapAll)
  plotTracks(chromHMM_RoadMapAll, from = start, to = end,
    fontfamily="sans", fontfamily.title="sans")
}
```

chrUCSC2ENSEMBL	<i>Removing "chr" to the chromosome number from UCSC to transform it to ENSEMBL chromosome format</i>
-----------------	---

Description

Removing "chr" at the beginning of the chromosome number

Usage

```
chrUCSC2ENSEMBL(chr)
```

Arguments

chr the chromosome number in UCSC format

Value

the number of chromosome at ENSEMBL format

Author(s)

Tiphaine Martin

Examples

```
chr<-"chr7"
chrUCSC2ENSEMBL(chr)
```

ClinVarCnv_UCSC	<i>Create one track of the genomic positions of variants from the ClinVar database (CNV only)</i>
-----------------	---

Description

Create one track of the genomic positions of variants from the ClinVar database (CNV only, Variants excluded) using the Gviz bioconductor package

Usage

```
ClinVarCnv_UCSC(gen, chr, start, end, title="ClinVar Variants", showId = FALSE)
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in region of interest (the smallest value)
end	the last position in region of interest (the biggest value)
title	The name of the annotation track
showId	Show the ID of the genetic elements

Value

An Ucsctrack object of Gviz

Author(s)

Tiphaine Martin

References

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtDrFAy6dn&c=chr6&g=clin
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[snpLocations_UCSC](#), [structureBiomart_ENSEMBL](#), [snpBiomart_ENSEMBL](#), [CoreilCNV_UCSC](#), [COSMIC_UCSC](#), [ClinVarMain_UCSC](#)

Examples

```

library("Gviz")
chrom <- "chr2"
start <- 38290160
end <- 38303219
gen <- "hg38"
if(interactive()){
  clinCNV<-ClinVarCnv_UCSC(gen,chrom,start,end)
  plotTracks(clinCNV, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}else {
  data(ClinVarCnvTrack)
  plotTracks(clinCNV, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}

```

ClinVarMain_UCSC	<i>Create one track of the genomic positions of variants from the ClinVar database (variants only)</i>
------------------	--

Description

Create one track of the genomic positions of variants from the ClinVar database (Variants only, CNV excluded) using the Gviz bioconductor package

Usage

```
ClinVarMain_UCSC(gen, chr, start, end, title="ClinVar Variants", showId=FALSE)
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in region of interest (the smallest value)
end	the last position in region of interest (the biggest value)
title	The name of the annotation track
showId	Show the ID of the genetic elements

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=clin
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[snpLocations_UCSC](#), [structureBiomart_ENSEMBL](#), [snpBiomart_ENSEMBL](#), [Coreil1CNV_UCSC](#), [COSMIC_UCSC](#), [ClinVarCnv_UCSC](#)

Examples

```
library("Gviz")
gen <- "hg38"
chrom <- "chr2"
start <- 100000
end <- 1000000

if(interactive()) {
  clinVariant<-ClinVarMain_UCSC(gen,chrom,start,end)
  plotTracks(clinVariant, from = start, to =end,
    fontfamily="sans",fontfamily.title="sans")
}else{
  data(clinVarMaintrack)
  plotTracks(clinVariant, from = start, to =end,
    fontfamily="sans",fontfamily.title="sans")
}
```

col2HSV

col2HSV: converts a color to HSV in hexadecimal notation

Description

col2HSV converts an R color (or a set of colors) into an HSV color model, and then returns the color names in hexadecimal notation

Usage

```
col2HSV(color)
```

Arguments

color an R color name or a color in hexadecimal notation

Value

A character vector with the color(s) name(s) in hexadecimal notation

Author(s)

Gaston Sanchez

Examples

```
# convert 'tomato'
col2HSV("tomato")
```

 comet

Visualize EWAS results in a genomic region of interest

Description

coMET is an R-based package to visualize EWAS (epigenome-wide association scans) results in a genomic region of interest. The main feature of coMET is to plot the the significance level of EWAS results in the selected region, along with correlation in DNA methylation values between CpG sites in the region. The coMET package generates plots of phenotype-association, co-methylation patterns, and a series of annotation tracks.

Usage

```
comet(mydata.file = NULL, mydata.format = "site", mydata.type = "file",
      mydata.large.file = NULL, mydata.large.format = "site",
      mydata.large.type = "listfile", cormatrix.file = NULL,
      cormatrix.method = "spearman", cormatrix.format = "raw",
      cormatrix.color.scheme = "bluewhitered", cormatrix.conf.level=0.05,
      cormatrix.sig.level= 1, cormatrix.adjust="none",
      cormatrix.type = "listfile", mydata.ref = NULL,
      start = NULL, end = NULL, zoom = FALSE, lab.Y = "log",
      pval.threshold = 1e-05, pval.threshold.2 = 0, disp.pval.threshold = 1,
      disp.association = FALSE, disp.association.large = FALSE,
      disp.region = FALSE, disp.region.large = FALSE,
      disp.beta.association = FALSE, disp.beta.association.large = FALSE, factor.beta = 0.3,
      symbols = "circle-fill",
      symbols.large = NA, sample.labels = NULL, sample.labels.large = NULL,
      use.colors = TRUE, disp.color.ref = TRUE, color.list = NULL, color.list.large = NULL,
      disp.mydata = TRUE, biofeat.user.file = NULL, biofeat.user.type = NULL,
      biofeat.user.type.plot = NULL,
      genome = "hg19", dataset.gene = "hsapiens_gene_ensembl",
      tracks.gviz = NULL,
      disp.mydata.names = TRUE, disp.color.bar = TRUE, disp.phys.dist = TRUE,
      disp.legend = TRUE, disp.marker.lines = TRUE, disp.cormatrixmap = TRUE,
      disp.pvalueplot = TRUE, disp.type = "symbol", disp.mult.lab.X = FALSE,
      disp.connecting.lines = TRUE, palette.file = NULL, image.title = NULL,
      image.name = "coMET", image.type = NULL, image.size = 3.5,
      fontsize.gviz=5, font.factor = 1,
      symbol.factor = NULL, print.image = TRUE, connecting.lines.factor = 1.5,
```

```
connecting.lines.adj = 0.01, connecting.lines.vert.adj = -1,
connecting.lines.flex = 0, config.file = NULL, verbose = FALSE)
```

Arguments

- `mydata.file` Name of the info file describing the coMET parameters
- `mydata.format` Format of the input data in `mydata.file`. There are 4 different options: `site`, `region`, `site_asso`, `region_asso`.
- `mydata.type` Format of `mydata.file`. There are 2 different options: `FILE` or `MATRIX`.
- `mydata.large.file`
Name of additional info files describing the coMET parameters. File names should be comma-separated. It is optional, but if you add some, they need to be file(s) in tabular format with a header. Additional info file can be a list of CpG sites with/without Beta value (DNA methylation level) or direction sign. If it is a site file then it is mandatory to have the 4 columns as shown below with headers in the same order. Beta can be the 5th column(optional) and it can be either a numeric value (positive or negative values) or only direction sign ("+", "-"). The number of columns and their types are defined but the option `mydata.large.format`.
- `mydata.large.format`
Format of additional data to be visualised in the p-value plot. Format should be comma-separated. There are 4 different options for each file: `site`, `region`, `site_asso`, `region_asso`.
- `mydata.large.type`
Format of `mydata.large.file`. There are 2 different options: `listfile` or `listdataframe`.
- `cormatrix.file` Name of the raw data file or the pre-computed correlation matrix file. It is mandatory and has to be a file in tabular format with an header.
- `cormatrix.method`
Options for calculating the correlation matrix: `spearman`, `pearson` and `kendall`
- `cormatrix.format`
Format of the input `cormatrix.file`. There are two options: `raw file` (raw if CpG sites are by column and samples by row or `raw_rev` if CpG site are by row and samples by column) and `pre-computed correlation matrix` (`cormatrix`)
- `cormatrix.color.scheme`
Color scheme options: `heat`, `bluewhitered`, `cm`, `topo`, `gray`, `bluetored`
- `cormatrix.conf.level`
Alpha level for the confidence interval. Default value= 0.05. CI will be the $\alpha/2$ lower and upper values.
- `cormatrix.sig.level`
Significant level to visualise the correlation. If the correlation has a pvalue under the significant level, the correlation will be colored in "goshwhite", else the color is related to the correlation level and the color scheme chosen. Default value =1.
- `cormatrix.adjust`
indicates which adjustment for multiple tests should be used. "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none". Default value="none"
- `cormatrix.type` Format of `cormatrix.file`. There are 2 different options: `listfile` or `listdataframe`.

<code>mydata.ref</code>	The name of the referenceomic feature (e.g. CpG-site) listed in <code>mydata.file</code>
<code>start</code>	The first nucleotide position to be visualised. It could be bigger or smaller than the first position of our list of omic features.
<code>end</code>	the last nucleotide position to be visualised. It has to be bigger than the value in the option <code>start</code> , but it could be smaller or bigger than the last position of our list of omic features.
<code>zoom</code>	Default=False
<code>lab.Y</code>	Scale of the y-axis. Options: log or ln
<code>pval.threshold</code>	Significance threshold to be displayed as a red dashed line
<code>pval.threshold.2</code>	the second significance threshold to be displayed as a orange dashed line
<code>disp.pval.threshold</code>	Display only the findings that pass the value put in <code>disp.pval.threshold</code>
<code>disp.association</code>	This logical option works only if <code>mydata.file</code> contains the effect direction (<code>mydata.format=site_asso</code> or <code>region_asso</code>). The value can be TRUE or FALSE: if FALSE (default), for each point of data in the p-value plot, the color of symbol is the color of co-methylation pattern between the point and the reference site; if TRUE, the effect direction is shown. If the association is positive, the color is the one defined with the option <code>color.list</code> . On the other hand, if the association is negative, the color is the opposed color.
<code>disp.association.large</code>	This logical option works only if <code>mydata.large.file</code> contains the effect direction (<code>mydata.large.format=site_asso</code> or <code>region_asso</code>). The value can be TRUE or FALSE: if FALSE (default), for each point of data in the p-value plot, the color of symbol is the color of co-methylation pattern between the point and the reference site; if TRUE, the effect direction is shown. If the association is positive, the color is the one defined with the option <code>color.list.large</code> . On the other hand, if the association is negative, the color is the opposed color.
<code>disp.region</code>	This logical option works only if <code>mydata.file</code> contains regions (<code>mydata.format=region</code> or <code>region_asso</code>). The value can be TRUE or FALSE (default). If TRUE, the genomic element will be shown by a continuous line with the color of the element, in addition to the symbol at the center of the region. If FALSE, only the symbol is shown.
<code>disp.region.large</code>	This logical option works only if <code>mydata.large.file</code> contains regions (<code>mydata.large.format=region</code> or <code>region_asso</code>). The value can be TRUE or FALSE (default). If TRUE, the genomic element will be shown by a continuous line with the color of the element, in addition to the symbol at the center of the region. If FALSE, only the symbol is shown.
<code>disp.beta.association</code>	This logical option works only if <code>mydata.file</code> contains the effect direction (<code>mydata.format=site_asso</code> or <code>region_asso</code>). The value can be TRUE or FALSE: if FALSE (default), for each point of data in the p-value plot, the size of symbol is the default size of symbole; if TRUE, the effect direction is shown.

<code>disp.beta.association.large</code>	This logical option works only if <code>mydata.large.file</code> contains the effect direction (<code>mydata.large.format=site_asso</code> or <code>region_asso</code>). The value can be TRUE or FALSE: if FALSE (default), for each point of data in the p-value plot, the size of symbol is the default size of symbol; if TRUE, the effect direction is shown.
<code>factor.beta</code>	Factor to visualise the size of beta. Default value = 0.3.
<code>symbols</code>	The symbol shown in the p-value plot. Options: circle, square, diamond, triangle. symbols can be filled by appending <code>-fill</code> , e.g. <code>square-fill</code> . Example: <code>circle,diamond-fill,triangle</code>
<code>symbols.large</code>	The symbol to visualise the data defined in <code>mydata.large.file</code> . Options: circle, square, diamond, triangle; symbols can either be filled or not filled by appending <code>-fill</code> e.s., <code>square-fill</code> . Example: <code>circle,diamond-fill,triangle</code>
<code>sample.labels</code>	Labels for the sample described in <code>mydata.file</code> to include in the legend
<code>sample.labels.large</code>	Labels for the sample described in <code>mydata.large.file</code> to include in the legend
<code>use.colors</code>	Use the colors defined or use the grey color scheme
<code>disp.color.ref</code>	Logical option TRUE or FALSE (TRUE default). if TRUE, the connection line related to the reference probe is in purple, if FALSE if the connection line related to the reference probe stay black.
<code>color.list</code>	List of colors for displaying the P-value symbols related to the data in <code>mydata.file</code>
<code>color.list.large</code>	List of colors for displaying the P-value symbols related to the data in <code>mydata.large.file</code>
<code>disp.mydata</code>	logical option TRUE or FALSE. TRUE (default). If TRUE, the P-value plot is shown; if FALSE the plot will be defined by Gviz
<code>biofeat.user.file</code>	Name of data file to visualise in the tracks. File names should be comma-separated.
<code>biofeat.user.type</code>	Track type, where multiple tracks can be shown (comma-separated): <code>DataTrack</code> , <code>AnnotationTrack</code> , <code>GeneregionTrack</code> .
<code>biofeat.user.type.plot</code>	Format of the plot if the data are shown with the Gviz's function called <code>DataTrack</code> (comma-separated)
<code>genome</code>	The human genome reference file. e.g. "hg19" for Human genome 19 (NCBI 37), "grch37" (GRCh37), "grch38" (GRCh38)
<code>dataset.gene</code>	The gene names from ENSEMBL. e.g. <code>hsapiens_gene</code>
<code>tracks.gviz</code>	list of tracks created by Gviz.
<code>disp.mydata.names</code>	logical option TRUE or FALSE. If True (default), the names of the CpG sites are displayed.
<code>disp.color.bar</code>	Color legend for the correlation matrix (range -1 to 1). Default: blue-white-red

<code>disp.phys.dist</code>	logical option (TRUE or FALSE). TRUE (default). Display the bp distance on the plots
<code>disp.legend</code>	logical option TRUE or FALSE. TRUE (default) Display the sample labels and corresponding symbols on the lower right side
<code>disp.marker.lines</code>	logical option TRUE or FALSE. TRUE (default), if FALSE the red line for <code>pval.threshold</code> is not shown
<code>disp.cormatrixmap</code>	logical option TRUE or FALSE. TRUE (default), if FALSE correlation matrix is not shown
<code>disp.pvalueplot</code>	logical option (TRUE or FALSE). TRUE (default), if FALSE the pvalue plot is not shown
<code>disp.type</code>	Default: symbol
<code>disp.mult.lab.X</code>	logical option TRUE or FALSE. FALSE (default). Display evenly spaced X-axis labels; up to 5 labels are shown.
<code>disp.connecting.lines</code>	logical option TRUE or FALSE. TRUE (default) displays connecting lines between p-value plot and correlation matrix
<code>palette.file</code>	File that contains color scheme for the heatmap. Colors are hexadecimal HTML color codes; one color per line; if you do not want to use this option, use the color defined by the option <code>cormatrix.color.scheme</code>
<code>image.title</code>	Title of the plot
<code>image.name</code>	The path and the name of the plot file without extension. The extension will be added by coMET depending on the option <code>image.type</code> .
<code>image.type</code>	Options: pdf or eps
<code>image.size</code>	Default: 3.5 inches. Possible sizes : 3.5 or 7
<code>fontsize.gviz</code>	Font size of writing in annotation track. Default value =5
<code>font.factor</code>	Font size of the sample labels. Range: 0-1
<code>symbol.factor</code>	Size of the symbols. Range: 0-1
<code>print.image</code>	Print image in file or not.
<code>connecting.lines.factor</code>	Length of the connecting lines. Range: 0-2
<code>connecting.lines.adj</code>	Position of the connecting lines horizontally. Negative values shift the connecting lines to the left and positive values shift the lines to the right. Range: (-1;1) option -1 means no connecting lines.
<code>connecting.lines.vert.adj</code>	Position of the connecting lines vertically. Can be used to vertically adjust the position of the connecting lines in relation to the CpG-site names. Negative value shift the connecting lines down. Range: (-0.5 - 0), option -1 mean the default value related to the plot size (-0.5 for 3.5 plot size; -0.7 for 7.5 plot size)

connecting.lines.flex	Adjusts the spread of the connecting lines. Range: 0-2
config.file	Configuration file contains the values of these options instead of defining these by command line. It is a file where each line is one option. The name of option and its value are separated by "=". If there are multiple values such as for the option list.tracks or the options for additional data, you need to separated them by a "comma" and not extra space. (i.e. list.tracks=geneENSEMBL,CGI,ChromHMM,DNAse,RegENSEMBL)
verbose	logical option TRUE or FALSE. TRUE (default). If TRUE, shows comments.

Details

The function is limited to visualize 120 omic features.

Value

Create a plot in pdf or eps format depending to some options

Author(s)

Tiphaine Martin

References

<http://epigen.kcl.ac.uk/comet/>

See Also

[comet.web](#), [comet.list](#)

Examples

```
extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
configfile <- file.path(extdata, "config_cyp1b1_zoom_4comet.txt")
myinfofile <- file.path(extdata, "cyp1b1_infofile.txt")
myexpressfile <- file.path(extdata, "cyp1b1_infofile_exprGene_region.txt")
mycorrelation <- file.path(extdata, "cyp1b1_res37_rawMatrix.txt")

chrom <- "chr2"
start <- 38290160
end <- 38303219
gen <- "hg38"

if(interactive()){
  cat("interactive")
  genetrack <- genes_ENSEMBL(gen, chrom, start, end, showId=TRUE)
  snptrack <- snpBiomart_ENSEMBL(gen, chrom, start, end,
    dataset="hsapiens_snp_som", showId=FALSE)
  strutrack <- structureBiomart_ENSEMBL(gen, chrom, start, end,
    strand, dataset="hsapiens_structvar_som")
  clinVariant <- ClinVarMain_UCSC(gen, chrom, start, end)
  clinCNV <- ClinVarCnv_UCSC(gen, chrom, start, end)
```

```

gwastrack <-GWAScatalog_UCSC(gen,chrom,start,end)
geneRtrack <-GeneReviews_UCSC(gen,chrom,start,end)
listgviz <- list(genetrack,snptrack,strutrack,clinVariant,
                clinCNV,gwastrack,geneRtrack)
comet(config.file=configfile, mydata.file=myinfofile, mydata.type="file",
      cormatrix.file=mycorrelation, cormatrix.type="listfile",
      mydata.large.file=myexpressfile, mydata.large.type="listfile",
      tracks.gviz=listgviz, verbose=FALSE, print.image=FALSE,disp.pvalueplot=FALSE)
} else {
  cat("Non interactive")
  data(geneENSEMBLtrack)
  data(snpBiomarttrack)
  data(ISCATrack)
  data(strucBiomarttrack)
  data(ClinVarCnvTrack)
  data(clinVarMaintrack)
  data(GWASTrack)
  data(GeneReviewTrack)
  listgviz <- list(genetrack,snptrack,strutrack,clinVariant,
                  clinCNV,gwastrack,geneRtrack)
  comet(config.file=configfile, mydata.file=myinfofile, mydata.type="file",
        cormatrix.file=mycorrelation, cormatrix.type="listfile",
        mydata.large.file=myexpressfile, mydata.large.type="listfile",
        tracks.gviz=listgviz, verbose=FALSE, print.image=FALSE,disp.pvalueplot=FALSE)
}

```

comet.list

List the correlations between omic features

Description

coMET is an R-based package to visualize EWAS (epigenome-wide association scans) results in a genomic region of interest. The main feature of coMET is to plot the the significance level of EWAS results in the selected region, along with correlation in DNA methylation values between CpG sites in the region. The coMET package generates plots of phenotype-association, co-methylation patterns, and a series of annotation tracks. In addition, the function comet.list gives the list of correlations between omic features

Usage

```

comet.list(cormatrix.file = NULL, cormatrix.method = "spearman", cormatrix.format = "raw",
          cormatrix.conf.level=0.05, cormatrix.sig.level= 1, cormatrix.adjust="none",
          cormatrix.type = "listdataframe", cormatrix.output="cormatrix_list",
          config.file = NULL, verbose = FALSE)

```

Arguments

cormatrix.file Name of the raw data file or the pre-computed correlation matrix file. It is mandatory and has to be a file in tabular format with an header.

cormatrix.method	Options for calculating the correlation matrix: spearman, pearson and kendall. Default value= spearman
cormatrix.format	Format of the input cormatrix.file. TThere are two options: raw file (raw if CpG sites are by column and samples by row or raw_rev if CpG site are by row and samples by column) and pre-computed correlation matrix (cormatrix)
cormatrix.conf.level	Alpha level for the confidence interval. Default value= 0.05. CI will be the alpha/2 lower and upper values.
cormatrix.sig.level	Significant level to visualise the correlation. If the correlation has a pvalue below the significant level, the correlation will be colored in "goshwhite", else the color is related to the correlation level and the color scheme choosen.Default value =1.
cormatrix.adjust	indicates which adjustment for multiple tests should be used. "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none".Default value="none"
cormatrix.type	Format of cormatrix.file. There are 2 different options: listfile or listdataframe.
cormatrix.output	The path and the name of the output file without the extension
config.file	Configuration file contains the values of these options instead of defining these by command line. It is a file where each line is one option. The name of option and its value are separated by "=".
verbose	logical option TRUE or FALSE. TRUE (default). If TRUE, shows comments.

Value

Create a list of correlation between omic features

Author(s)

Tiphaine Martin

References

<http://epigen.kcl.ac.uk/comet/>

See Also

[comet.web](#), [comet](#)

Examples

```
extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
mycorrelation <- file.path(extdata, "cyp1b1_res37_rawMatrix.txt")
myoutput <- file.path(extdata, "cyp1b1_res37_cormatrix_list_BH05.txt")
```

```
comet.list(cormatrix.file=mycorrelation,cormatrix.method = "spearman",
          cormatrix.format= "raw", cormatrix.conf.level=0.05,
          cormatrix.sig.level= 0.05, cormatrix.adjust="BH",
          cormatrix.type = "listfile", cormatrix.output=myoutput,
          verbose=FALSE)
```

comet.web	<i>Visualize EWAS results in a genomic region of interest with predefined annotation tracks</i>
-----------	---

Description

coMET is an R-based package to visualize EWAS (epigenome-wide association scans) results in a genomic region of interest. The main feature of coMET is to plot the the significance level of EWAS results in the selected region, along with correlation in DNA methylation values between CpG sites in the region. The coMET package generates plots of phenotype-association, co-methylation patterns, and a series of annotation tracks.

Usage

```
comet.web(mydata.file = NULL, mydata.format = c("site", "region",
"site_asso", "region_asso"),
          mydata.large.file = NULL,
          mydata.large.format = c("site", "region", "site_asso", "region_asso"),
          cormatrix.file = NULL, cormatrix.method = c("spearman", "pearson", "kendall"),
          cormatrix.format = c("cormatrix", "raw", "raw_rev"),
          cormatrix.color.scheme = "heat", cormatrix.conf.level=0.05,
          cormatrix.sig.level= 1, cormatrix.adjust="none",mydata.ref = NULL,
          genome="hg19", start = NULL, end = NULL, zoom = FALSE, lab.Y = "log",
          pval.threshold = 1e-07, pval.threshold.2 = 0, disp.pval.threshold = 1,
          disp.association= FALSE, disp.association.large = FALSE,
          disp.beta.association = "FALSE", disp.beta.association.large = "FALSE",
          factor.beta = 0.3,
          disp.region = FALSE, disp.region.large = FALSE, symbols = "circle-fill",
          symbols.large = NA, sample.labels = NULL, sample.labels.large = NULL,
          use.colors = TRUE, disp.color.ref = TRUE, color.list = NULL,
          color.list.large = NULL, biofeat.user.file = NULL,
          biofeat.user.type = c("GeneRegion", "Annotation", "Data"),
          biofeat.user.type.plot = NULL,
          list.tracks = "geneENSEMBL,CGI,ChromHMM,DNAse,RegENSEMBL,SNP",
          pattern.regulation = "GM12878",
          image.title = NULL, image.name = "coMET", image.type = c("pdf", "eps"),
          image.size = 3.5, fontsize.gviz=5, font.factor = 1,
          print.image = FALSE, config.file = NULL, verbose = FALSE)
```

Arguments

- .
- Name of the info file describing the coMET parameters. It is mandatory and has to be a file in tabular format with a header. Info file can be a list of CpG sites with/without Beta value (DNA methylation level) or direction sign. If it is a site file then it is mandatory to have the 4 columns as shown below with headers in the same order. Beta can be the 5th column(optional) and it can be either a numeric value (positive or negative values) or only direction sign ("+", "-"). The number of columns and their types are defined but the option mydata.format.
- mydata.format** Format of the input data in mydata.file. There are 4 different options: site, region, site_asso, region_asso.
- mydata.large.file** Name of additional info files describing the coMET parameters. File names should be comma-separated. It is optional, but if you add some, they need to be file(s) in tabular format with a header. Additional info file can be a list of CpG sites with/without Beta value (DNA methylation level) or direction sign. If it is a site file then it is mandatory to have the 4 columns as shown below with headers in the same order. Beta can be the 5th column(optional) and it can be either a numeric value (positive or negative values) or only direction sign ("+", "-"). The number of columns and their types are defined but the option mydata.large.format.
- mydata.large.format** Format of additional data to be visualised in the p-value plot. Format should be comma-separated. There are 4 different options for each file: site, region, site_asso, region_asso.
- cormatrix.file** Name of the raw data file or the pre-computed correlation matrix file. It is mandatory and has to be a file in tabular format with an header.
- cormatrix.method** A character string indicating which correlation coefficient is to be used for the test. One of "pearson", "kendall", or "spearman", can be abbreviated.
- cormatrix.format** A character string indicating which format of the input cormatrix.file is to be used. There are three options: raw file (raw if CpG sites are by column and samples by row or row_rev if CpG site are by row and samples by column) and pre-computed correlation matrix (cormatrix)
- cormatrix.color.scheme** A character string indicating which Color scheme options is to be used: heat, bluewhitered, cm, topo, gray, bluetored
- cormatrix.conf.level** Alpha level for the confidence interval. Default value= 0.05. CI will be the alpha/2 lower and upper values.
- cormatrix.sig.level** Significant level to visualise the correlation. If the correlation has a pvalue under the significant level, the correlation will be colored in "goshwhite", else the color is related to the correlation level and the color scheme choosen. Default value =1.

<code>cormatrix.adjust</code>	indicates which adjustment for multiple tests should be used. "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none".Default value="none"
<code>mydata.ref</code>	The name of the reference omic feature (e.g. CpG-site) listed in mydata.file
<code>genome</code>	The human genome reference file. e.g. "hg19" for Human genome 19 (NCBI 37), "grch37" (GRCh37), "grch38" (GRCh38)
<code>start</code>	The first nucleotide position to be visualised. It could be bigger or smaller than the first position of our list of omic features.
<code>end</code>	the last nucleotide position to be visualised. It has to be bigger than the value in the option start, but it could be smaller or bigger than the last position of our list of omic features.
<code>zoom</code>	logical option TRUE or FALSE. FALSE (default)
<code>lab.Y</code>	Scale of the y-axis. Options: log or ln
<code>pval.threshold</code>	Significance threshold to be displayed as a red dashed line. Default value = 1e-7
<code>pval.threshold.2</code>	the second significance threshold to be displayed as a orange dashed line. Default value= 0 (no printed)
<code>disp.pval.threshold</code>	Display only the findings that pass the value put in disp.pval.threshold
<code>disp.association</code>	This logical option works only if mydata.file contains the effect direction (mydata.format=site_asso or region_asso). The value can be TRUE or FALSE: if FALSE (default), for each point of data in the p-value plot, the color of symbol is the color of co-methylation pattern between the point and the reference site; if TRUE, the effect direction is shown. If the association is positive, the color is the one defined with the option color.list. On the other hand, if the association is negative, the color is the opposed color.
<code>disp.association.large</code>	This logical option works only if mydata.large.file contains the effect direction (MYDATA.large.FORMA=site_asso or region_asso). The value can be TRUE or FALSE: if FALSE (default), for each point of data in the p-value plot, the color of symbol is the color of co-methylation pattern between the point and the reference site; if TRUE, the effect direction is shown. If the association is positive, the color is the one defined with the option color.list.large. On the other hand, if the association is negative, the color is the opposed color.
<code>disp.beta.association</code>	This logical option works only if mydata.file contains the effect direction (mydata.format=site_asso or region_asso). The value can be TRUE or FALSE: if FALSE (default), for each point of data in the p-value plot, the size of symbol is the default size of symbole; if TRUE, the effect direction is shown.
<code>disp.beta.association.large</code>	This logical option works only if mydata.large.file contains the effect direction (mydata.large.format=site_asso or region_asso). The value can be TRUE or FALSE: if FALSE (default), for each point of data in the p-value plot, the size of symbol is ththe default size of symbole; if TRUE, the effect direction is shown.

<code>factor.beta</code>	Factor to visualise the size of beta. Default value = 0.3.
<code>disp.region</code>	This logical option works only if <code>mydata.file</code> contains regions (<code>mydata.format=region</code> or <code>region_asso</code>). The value can be TRUE or FALSE (default). If TRUE, the genomic element will be shown by a continuous line with the color of the element, in addition to the symbol at the center of the region. If FALSE, only the symbol is shown.
<code>disp.region.large</code>	This logical option works only if <code>mydata.large.file</code> contains regions (<code>mydata.large.format=region</code> or <code>region_asso</code>). The value can be TRUE or FALSE (default). If TRUE, the genomic element will be shown by a continuous line with the color of the element, in addition to the symbol at the center of the region. If FALSE, only the symbol is shown.
<code>symbols</code>	The symbol shown in the p-value plot. Options: circle, square, diamond, triangle. symbols can be filled by appending <code>-fill</code> , e.g. <code>square-fill</code> . Example: <code>circle,diamond-fill,triangle</code>
<code>symbols.large</code>	The symbol to visualise the data defined in <code>mydata.large.file</code> . Options: circle, square, diamond, triangle; symbols can either be filled or not filled by appending <code>-fill</code> e.s., <code>square-fill</code> . Example: <code>circle,diamond-fill,triangle</code>
<code>sample.labels</code>	Labels for the sample described in <code>mydata.file</code> to include in the legend
<code>sample.labels.large</code>	Labels for the sample described in <code>mydata.large.file</code> to include in the legend
<code>use.colors</code>	Use the colors defined or use the grey color scheme
<code>disp.color.ref</code>	Logical option TRUE or FALSE (TRUE default). if TRUE, the connection line related to the reference probe is in purple, if FALSE if the connection line related to the reference probe stay black.
<code>color.list</code>	List of colors for displaying the P-value symbols related to the data in <code>mydata.file</code>
<code>color.list.large</code>	List of colors for displaying the P-value symbols related to the data in <code>mydata.large.file</code>
<code>biofeat.user.file</code>	Name of data file to visualise in the tracks. File names should be comma-separated.
<code>biofeat.user.type</code>	Track type, where multiple tracks can be shown (comma-separated): <code>DataTrack</code> , <code>AnnotationTrack</code> , <code>GeneRegionTrack</code> .
<code>biofeat.user.type.plot</code>	Format of the plot if the data are shown with the Gviz's function called <code>DataTrack</code> (comma-separated)
<code>list.tracks</code>	List of annotation tracks to visualise. Options include <code>geneENSEMBL</code> , <code>CGI</code> , <code>ChromHMM</code> , <code>DNase</code> , <code>RegENSEMBL</code> , <code>SNP</code> , <code>transcriptENSEMBL</code> , <code>SNPstoma</code> , <code>SNPstru</code> , <code>SNPstrustoma</code> , <code>BindingMotifENSEMBL</code> , <code>otherRegulatoryENSEMBL</code> , <code>regulatoryEvidenceENSEMBL</code> , <code>regulatoryFeaturesENSEMBL</code> , <code>regulatorySegmeENSEMBL</code> , <code>miRNAENSEMBL</code> , <code>ImprintedtissuesGenes</code> , <code>COSMIC</code> , <code>GAD</code> , <code>ClinVar</code> , <code>GeneReviews</code> , <code>GWAS</code> , <code>ClinVarCNV</code> , <code>GCcontent</code> , <code>genesUCSC</code> , <code>xenogenesUCSC</code> , <code>SegDuplication</code> , <code>RepeatElt</code> .

pattern.regulation	The cell/tissue or the list of cells/tissues to visualise in the regulation region defined by Broad ChromHMM
image.title	Title of the plot
image.name	The path and the name of the plot file without extension. The extension will be added by coMET depending on the option image.type.
image.type	Options: pdf or eps
image.size	Default: 3.5 inches. Possible sizes : 3.5 or 7
fontsize.gviz	Font size of writing in annotation track. Default value =5
font.factor	Font size of the sample labels. Range: 0-1
print.image	Print image in file or not.
config.file	Configuration file contains the values of these options instead of defining these by command line. It is a file where each line is one option. The name of option and its value are separated by "=". If there are multiple values such as for the option list.tracks or the options for additional data, you need to separated them by a "comma" and not extra space. (i.e. list.tracks=geneENSEMBL,CGI,ChromHMM,DNAse,RegENSEMBL)
verbose	logical option TRUE or FALSE. TRUE (default). If TRUE, shows comments.

Details

The function is limited to visualize 120 omic features.

Value

Create a plot in pdf or eps format depending to some options

Author(s)

Tiphaine Martin

References

<http://epigen.kcl.ac.uk/comet/>

See Also

[comet.comet.list](#)

Examples

```
extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
configfile <- file.path(extdata, "config_cyp1b1_zoom_4webserver.txt")
myinfofile <- file.path(extdata, "cyp1b1_infofile.txt")
myexpressfile <- file.path(extdata, "cyp1b1_infofile_exprGene_region.txt")
mycorrelation <- file.path(extdata, "cyp1b1_res37_rawMatrix.txt")

comet.web(config.file=configfile, mydata.file=myinfofile, cormatrix.file=mycorrelation,
           mydata.large.file=myexpressfile, print.image=FALSE, verbose=FALSE)
```

complementary	<i>Complementary or opposite color</i>
---------------	--

Description

Complementary or opposite color scheme is formed by colors that are opposite each other on the color wheel (example: red and green). The high contrast of complementary colors creates a vibrant look that must be managed well so it is not jarring.

Usage

```
complementary(color, plot = TRUE, bg = "white",
              labcol = NULL, cex = 0.8, title = TRUE)
```

Arguments

color	an R color name or color in hexadecimal notation
plot	logical value indicating whether to plot a color wheel with the generated scheme
bg	background color of the plot. Used only when plot=TRUE
labcol	color for the labels (i.e. names of the colors). Used only when plot=TRUE
cex	numeric value indicating the character expansion of the labels
title	logical value indicating whether to display a title in the plot. Used only when plot=TRUE

Details

The complementary color is obtained following a color wheel with 12 colors, each one spaced at 30 degrees from each other. Complementary color schemes are tricky to use in large doses, but work well when you want something to stand out. In addition, complementary colors are really bad for text.

Value

A character vector with the given color and the complementary color in hexadecimal notation

Author(s)

Gaston Sanchez

Examples

```
# complementary color of 'tomato' with no plot
opposite("tomato", plot = FALSE)

# complementary color of 'tomato' with color wheel
opposite("tomato", bg = "gray30")
```

CoreillCNV_UCSC	<i>Create one track of the genomic positions of CNV in chromosomal aberration and inherited disorders from the NIGMS Human Genetic Cell Repository data</i>
-----------------	---

Description

Create one track of the genomic positions of copy-number variants (CNVs) in chromosomal aberration and inherited disorder cell lines from the NIGMS Human Genetic Cell Repository using the Gviz bioconductor package.

Usage

```
CoreillCNV_UCSC(gen, chr, start, end, title="Coriell CNVs", showId=FALSE)
```

Arguments

gen	the name of the genome. Data is not currently available for GRCh38 (hg38).
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	The name of the annotation track
showId	Show the ID of the genetic elements

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtDrFAy6dn&c=chr6&g=coriell

See Also

[snpLocations_UCSC](#), [structureBiomart_ENSEMBL](#), [snpBiomart_ENSEMBL](#), [COSMIC_UCSC](#), [ClinVarMain_UCSC](#), [ClinVarCnv_UCSC](#)

Examples

```

library("Gviz")
gen <- "hg19"
chrom <- "chr2"
start <- 38290160
end <- 38303219

if(interactive()){
  coreilVariant<-CoreillCNV_UCSC(gen,chrom,start,end)
  plotTracks(coreilVariant, from = start, to =end,
    fontfamily="sans",fontfamily.title="sans")
} else {
  data(coreilVarianttrack)
  plotTracks(coreilVariant, from = start, to =end,
    fontfamily="sans",fontfamily.title="sans")
}

```

COSMIC_UCSC

Create one track of the genomic positions of variants from COSMIC [obsolete]

Description

[obsolete] No more possible to extract COSMIC data from UCSC.

Create one track of the genomic positions of variants from COSMIC, the "Catalogue Of Somatic Mutations In Cancer" in extracting data from UCSC and using the Gviz bioconductor package.

Usage

```
COSMIC_UCSC(gen, chr, start, end,title= "COSMIC", showId=FALSE)
```

Arguments

gen	the name of the genome. Data is not currently available for GRCh38 (hg38)
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	The name of the annotation track
showId	Show the ID of the genetic elements

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=cos

See Also

[snpLocations_UCSC](#), [structureBiomart_ENSEMBL](#), [snpBiomart_ENSEMBL](#), [CoreillCNV_UCSC](#), [ClinVarMain_UCSC](#), [ClinVarCnv_UCSC](#),

Examples

```
library("Gviz")
chrom <- "chr2"
start <- 38290160
end <- 38303219
gen <- "hg19"
if(interactive()){
  cosmicVariant<-COSMIC_UCSC(gen,chrom,start,end)
  plotTracks(cosmicVariant, from = start, to =end,
    fontfamily="sans",fontfamily.title="sans")
}else {
  data(cosmicVarianttrack)
  plotTracks(cosmicVariant, from = start, to =end,
    fontfamily="sans",fontfamily.title="sans")
}
```

cpgIslands_UCSC

create track CpG Island from UCSC

Description

create track CpG Island from UCSC using the Gviz bioconductor package

Usage

```
cpgIslands_UCSC(gen, chr, start, end, title="CpG Islands UCSC")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	Name of tracks

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References<http://bioconductor.org/packages/release/bioc/html/Gviz.html>http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=cpg**Examples**

```
library("Gviz")
chrom <- "chr2"
start <- 100000
end <- 1000000
gen <- "hg38"

if(interactive()) {
  cpgIstrack<-cpgIslands_UCSC(gen, chrom, start, end)
  plotTracks(cpgIstrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}else {
  data(cpgIslandtrack)
  plotTracks(cpgIstrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}
```

`dgfootprints_RoadMap` *Creates a track of DNA motif positional bias in digital genomic Footprinting Sites (DGFP) from a file of RoadMap*

Description

Creates a DGFP track from a file of RoadMap using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
dgfootprints_RoadMap(gen="hg19", chr, start, end, bedFilePath,
                    tissueGroupDisplay='Blood & T-cell',showId=FALSE, type_stacking="dense",
                    title= "DGFP RoadMap")
```

Arguments

<code>gen</code>	the name of the genome. Default value=hg19
<code>chr</code>	The chromosome of interest
<code>start</code>	The starting position in the region of interest (the smallest value)

end	The end position in the region of interest (the largest value)
bedFilePath	The file path to the .BED file containing the data to be visualised
tissueGroupDisplay	the group of tissue visualised among list("Neurosp", "Epithelial", "IMR90", "Thymus", "Heart", "Brain", "D & B-cell", "Blood & T-cell"="ES-deriv")
showId	logical. say if we write the name of group
type_stacking	Object of class"character", the stacking type of overlapping items on the final plot. One in c(hide, dense, squish, pack,full). More information cf the option "stacking" in Gviz
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to RoadMap Epigenome

Examples

```
library("Gviz")
chr <- "chr1"
start <- 236728
end <- 238778
gen="hg19"

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
bedFilePath <- file.path(extdata, "RoadMap/CD3-DS17198.hg19_subset.bed")

if(interactive()){
  dgfootprints_RoadMapSingle <- dgfootprints_RoadMap(gen,chr,start, end,
  bedFilePath, tissueGroupDisplay='Blood & T-cell' )
  plotTracks(dgfootprints_RoadMapSingle, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
} else {
  data(dgfootprints_RoadMapSingle)
  plotTracks(dgfootprints_RoadMapSingle, from = start, to = end,
  fontfamily="sans",fontfamily.title="sans")
}
```

DNaseI_FANTOM

Creates a enhancer/promoter track from FANTOM

Description

Creates a track of promoters/enhancers from FANTOM using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
DNaseI_FANTOM(gen="hg19", chr, start, end, bedFilePath,
featureDisplay='enhancer', stacking_type="dense",
title=" DNaseI Fantom")
```

Arguments

gen	the name of the genome. Default value=hg19
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
bedFilePath	The path of the BED file from Kheradpour and Kellis, 2014.
featureDisplay	A vector of regulatory features to be displayed, such as enhancer. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. featureDisplay <- "Predicted heterochomatin"), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. featureDisplay <- c("enhancer","promoter")). Finally, visualisation all features in the genomic region, achieved by using the word "all" (e.g. featureDisplay <- "all"), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
stacking_type	Object of class"character", the stacking type of overlapping items on the final plot.One in c(hide, dense, squish, pack,full). More information cf the option "stacking" in Gviz
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to BindingMotifsBiomart binding motif biomart

Examples

```
library("Gviz")
gen <- "hg19"
chr<- "chr1"
start <- 6000000
end <- 6500000

extdata <- system.file("extdata", package="coMET",mustWork=TRUE)
enhFantomFile <- file.path(extdata,
"/FANTOM/human_permissive_enhancers_phase_1_and_2_example970.bed")

if(interactive()){
  enhFANTOMtrack <- DNaseI_FANTOM(gen,chr,start, end,
  enhFantomFile, featureDisplay='enhancer')
  plotTracks(enhFANTOMtrack, from = start, to = end,
              fontfamily="sans",fontfamily.title="sans")
} else {
  data(enhFANTOMtrack)
  plotTracks(enhFANTOMtrack, from = start, to = end,
              fontfamily="sans",fontfamily.title="sans")
}
```

DNaseI_RoadMap

Creates a promoter/enhancer regions track from a file of RoadMap

Description

Creates a track of promoter/enhancer regions from a file of RoadMap using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
DNaseI_RoadMap(gen="hg19", chr, start, end, bedFilePath,
featureDisplay='promotor',showId=TRUE, type_stacking="dense",
title = "DNaseI RoadMap")
```

Arguments

gen	the name of the genome. Default value=hg19
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)

<code>end</code>	The end position in the region of interest (the largest value)
<code>bedFilePath</code>	The file path to the .BED file containing the data to be visualised
<code>featureDisplay</code>	A vector of features to be displayed, such as 1_TssA. Spelling and capitalisation of features must be identical to those in the user guide (in the 'State & Acronym' column). There are three possibilities. First, the visualisation of only one feature (e.g. <code>featureDisplay <- "1_TssA"</code>), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. <code>featureDisplay <- c("1_TssA","2_TssAFlnk")</code>). Finally, visualise all features in the genomic region, achieved by using the word "all" (e.g. <code>featureDisplay <- "all"</code>), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
<code>showId</code>	Allows to visualise the Id of DNase group.
<code>type_stacking</code>	Object of class "character", the stacking type of overlapping items on the final plot. One in <code>c(hide, dense, squish, pack, full)</code> . More information of the option "stacking" in Gviz
<code>title</code>	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to RoadMap Epigenome

Examples

```
library("Gviz")
chr <- "chr2"
start <- 38300049
end <- 38302592
gen="hg19"

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
bedFilePath <- file.path(extdata, "RoadMap/regions_prom_E063.bed")

if(interactive()){
  DNaseI_RoadMapSingle <- DNaseI_RoadMap(gen,chr,start, end,
    bedFilePath, featureDisplay='promotor' )
  plotTracks(DNaseI_RoadMapSingle, from = start, to = end,
    fontfamily="sans", fontfamily.title="sans")
} else {
  data(DNaseI_RoadMapSingle)
```

```

plotTracks(DNaseI_RoadMapSingle, from = start, to = end,
           fontfamily="sans", fontfamily.title="sans")
}

```

DNase_UCSC

Creation of an UCSC's DNase clusters track - obsolete function

Description

Creation of DNase cluster track from a connection to UCSC genome browser in using the GViz bioconductor package. Obsolete function

Usage

```

DNase_UCSC(gen, chr, start, end, mySession, title="DNA cluster",
           track.name = "DNase Clusters", table.name = NULL)

```

Arguments

gen	the name of the genome. Data is not currently available for GRCh38 (hg38).
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
mySession	the object session from the function browserSession of rtracklayer
title	Name of tracks
track.name	the name of the track DNase_UCSC. "DNase Clusters"(default)
table.name	the name of the table from the track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=wgl

Examples

```

# library("Gviz")
# library("rtracklayer")

# gen <- "hg19"
# chr <- "chr7"
# start <- 38290160
# end <- 38303219
# if(interactive()){
#   BROWSER.SESSION="UCSC"
#   mySession <- browserSession(BROWSER.SESSION)
#   genome(mySession) <- gen
#   track.name="Broad ChromHMM"
#   tabletrack<-tableNames(ucscTableQuery(mySession, track=track.name))
#   table.name<-tabletrack[1]
#   dnasetrack<-DNase_UCSC(gen,chr,start,end,mySession)
#   plotTracks(dnasetrack, from = start, to =end,
#             fontfamily="sans",fontfamily.title="sans")
# }else {
#   data(dnasetrack)
#   plotTracks(dnasetrack, from = start, to =end,
#             fontfamily="sans",fontfamily.title="sans")
# }

```

eQTL

Creates a track from a file for eQTL data

Description

Creates a track from a BED file for eQTL data using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
eQTL(gen,chr, start, end, bedFilePath, featureDisplay, showId=FALSE,
type_stacking="squish",just_group="above", title="eQTL" )
```

Arguments

gen	the name of the genome.
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
bedFilePath	The file path to the .BED file containing the data to be visualised

<code>featureDisplay</code>	A vector of eQTL features to be displayed, such as SNP. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. <code>featureDisplay <- "CpG"</code>), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. <code>featureDisplay <- c("SNP","CpG")</code>). Finally, visualison all features in the genomic region, achived by using the word "all" (e.g. <code>featureDisplay <- "all"</code>), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
<code>showId</code>	Allows to visualise the Id of eQTL group.
<code>type_stacking</code>	Object of class"character", the stacking type of overlapping items on the final plot.One in <code>c(hide, dense, squish, pack,full)</code> . More information cf the option "stacking" in Gviz
<code>just_group</code>	position. say where we write the name of group (choice in <code>c("above","right","left")</code>)
<code>title</code>	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to ENSEMBLregulation binding motif biomart

Examples

```
library("Gviz")
chr <- "chr15"
start <- 74889136
end <- 75018200
featureDisplay <- "SNP"
gen="hg19"

extdata <- system.file("extdata", package="coMET",mustWork=TRUE)
bedFilePath <- file.path(extdata, "eQTL.bed")

if(interactive()){
  eQTLTrackSingle <- eQTL(gen,chr,start, end, bedFilePath,
    featureDisplay = featureDisplay )
  plotTracks(eQTLTrackSingle, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
} else {
  data(eQTLTrackSingle)
```

```

    plotTracks(eQTLTrackSingle, from = start, to = end,
              fontfamily="sans",fontfamily.title="sans")
}

#####

library("Gviz")
chr <- "chr15"
start <- 74889136
end <- 75018200
featureDisplay <- c("SNP","mRNA_pheno")
gen="hg19"

extdata <- system.file("extdata", package="coMET",mustWork=TRUE)
bedFilePath <- file.path(extdata, "eQTL.bed")

if(interactive()){
  eQTLTrackMultiple <- eQTL(gen,chr,start, end, bedFilePath, featureDisplay = featureDisplay )
  plotTracks(eQTLTrackMultiple, from = start, to = end,
            fontfamily="sans",fontfamily.title="sans")
} else {
  data(eQTLTrackMultiple)
  plotTracks(eQTLTrackMultiple, from = start, to = end,
            fontfamily="sans",fontfamily.title="sans")
}

#####

library("Gviz")
chr <- "chr15"
start <- 74889136
end <- 75018200
featureDisplay <- "all"
gen="hg19"

extdata <- system.file("extdata", package="coMET",mustWork=TRUE)
bedFilePath <- file.path(extdata, "eQTL.bed")

if(interactive()){
  eQTLTrackAll <- eQTL(gen,chr,start, end, bedFilePath, featureDisplay = featureDisplay )
  plotTracks(eQTLTrackAll, from = start, to = end,
            fontfamily="sans",fontfamily.title="sans")
} else {
  data(eQTLTrackAll)
  plotTracks(eQTLTrackAll, from = start, to = end,
            fontfamily="sans",fontfamily.title="sans")
}

```

Description

Creates a track of eQTL from GTE_x using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
eQTL_GTEx(gen="hg19",chr,start, end, bedFilePath, featureDisplay = 'all',
showId=FALSE, type_stacking="squish",just_group="above",title="eQTL GTEx")
```

Arguments

gen	the name of the genome. Default value=hg19
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
bedFilePath	The path of the BED file from Kheradpour and Kellis, 2014.
featureDisplay	A vector of regulatory features to be displayed, such as Predicted heterochromatin. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. featureDisplay <- "Predicted heterochromatin"), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. featureDisplay <- c("Predicted low activity","Predicted heterochromatin")). Finally, visualisation all features in the genomic region, achieved by using the word "all" (e.g. featureDisplay <- "all"), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
showId	logical. say if we write the name of group
type_stacking	Object of class"character", the stacking type of overlapping items on the final plot.One in c(hide, dense, squish, pack,full). More information of the option "stacking" in Gviz
just_group	position. say where we write the name of group (choice in c("above","right","left"))
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to BindingMotifsBiomart binding motif biomart

Examples

```

library("Gviz")
gen <- "hg19"
chr<-"chr3"
start <- 132423172
end <- 132442807
extdata <- system.file("extdata", package="coMET",mustWork=TRUE)
bedFilePath <- file.path(extdata, "GTEX/eQTL_Uterus_Analysis_extract100.snpgenes")

if(interactive()){
  eGTexTrackall <- eQTL_GTEEx(gen,chr,start, end, bedFilePath,
    featureDisplay="all", showId=TRUE,just_group="left")
  plotTracks(eGTexTrackall, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
} else {
  data(eGTexTrackall)
  plotTracks(eGTexTrackall, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
}

if(interactive()){
  eGTexTrackSNP <- eQTL_GTEEx(gen,chr,start, end, bedFilePath,
    featureDisplay="SNP", showId=TRUE,just_group="left")
  plotTracks(eGTexTrackSNP, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
} else {
  data(eGTexTrackSNP)
  plotTracks(eGTexTrackSNP, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
}

```

GAD_UCSC

Create one track of the genomic positions of variants from the Genetic Association Database (GAD)

Description

Create one track of the genomic positions of variants from the Genetic Association Database (GAD) (archive of human genetic association studies of complex diseases and disorders) using the Gviz bioconductor package

Usage

```
GAD_UCSC(gen, chr, start, end,title="GAD", showId=FALSE)
```

Arguments

gen	the name of the genome. Data is not currently available for GRCh38 (hg38).
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	The name of the annotation track
showId	Show the ID of the genetic elements

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=gad

See Also

[ISCA_UCSC](#), [GWAScatalog_UCSC](#), [knownGenes_UCSC](#), [genesName_ENSEMBL](#), [GeneReviews_UCSC](#), [genes_ENSEMBL](#), [xenorefGenes_UCSC](#), [transcript_ENSEMBL](#),

Examples

```
library("Gviz")
gen2 <- "hg19"
chrom2 <- "chr2"
start2 <- 38290160
end2 <- 38303219

if(interactive()) {
  gadtrack<-GAD_UCSC(gen=gen2 ,chr=chrom2 ,start=start2 ,end=end2)
  plotTracks(gadtrack, from = start2, to =end2,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(gadtrack)
  plotTracks(gadtrack, from = start2, to =end2,
             fontfamily="sans",fontfamily.title="sans")
}
```

gcContent_UCSC	<i>Create one track of GC content from UCSC</i>
----------------	---

Description

Create a track of GC content from UCSC using the Gviz bioconductor package

Usage

```
gcContent_UCSC(gen, chr, start, end, title="GC Percent")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	Name of tracks

Value

A UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtDrFAy6dn&c=chr6&g=gc5

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr7"
start <- 38290160
end <- 38303219

if(interactive()){
  gctrack<-gcContent_UCSC(gen,chr,start,end)
  plotTracks(gctrack,from= start, to=end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(gctrack)
  plotTracks(gctrack,from= start, to=end,
```

```
fontfamily="sans",fontfamily.title="sans")
}
```

GeneReviews_UCSC *Create one track of the genomic positions of variants from GeneReviews*

Description

Create one track of the genomic positions of variants from GeneReviews using the Gviz bioconductor package

Usage

```
GeneReviews_UCSC(gen, chr, start, end,title="GeneReviews", showId=FALSE)
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	The name of the annotation track
showId	Show the ID of the genetic elements

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=gen

See Also

[ISCA_UCSC](#), [GWAScatalog_UCSC](#), [knownGenes_UCSC](#), [genesName_ENSEMBL](#), [GAD_UCSC](#), [genes_ENSEMBL](#), [xenorefGenes_UCSC](#), [transcript_ENSEMBL](#),

Examples

```
library("Gviz")
gen <- "hg38"
chrom <- "chr2"
start <- 100000000
end <- 100000000
if(interactive()){
  geneRtrack <-GeneReviews_UCSC(gen,chrom,start,end,showId=TRUE)
  plotTracks(geneRtrack, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(GeneReviewTrack)
  plotTracks(geneRtrack, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}
```

genesName_ENSEMBL	<i>Obtain the genes names in the genomic regions of interest from ENSEMBL</i>
-------------------	---

Description

Obtain the genes names in the genomic regions of interest from ENSEMBL

Usage

```
genesName_ENSEMBL(gen, chr, start, end, dataset)
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
dataset	Name of the database to select genes

Details

Can be null

Value

List of name of genes found in this region of interest.

Author(s)

Tiphaine Martin

References

go to ENSEMBL

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[ISCA_UCSC](#), [GWAScatalog_UCSC](#), [knownGenes_UCSC](#), [GeneReviews_UCSC](#), [GAD_UCSC](#), [genes_ENSEMBL](#), [xenorefGenes_UCSC](#), [transcript_ENSEMBL](#),

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr7"
start <- 38290160
end <- 38303219

if(interactive()){
  dataset<- "hsapiens_gene_ensembl"
  geneNameEnsembl<- genesName_ENSEMBL(gen,chr,start,end,dataset)
  geneNameEnsembl
} else {
  data(geneNameEnsembl)
  geneNameEnsembl
}
```

genes_ENSEMBL

Create one track of the genes in the genomic regions of interest from EMSEMBL

Description

Create one track of the genes in the genomic regions of interest from EMSEMBL using the Gviz bioconductor package

Usage

```
genes_ENSEMBL(gen, chr, start, end, showId=FALSE,title="genes (ENSEMBL)")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
showId	Show the ID of the genetic elements
title	Name of tracks

Value

A BiomartGeneRegionTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=ens

See Also

[ISCA_UCSC](#), [GWAScatalog_UCSC](#), [knownGenes_UCSC](#), [genesName_ENSEMBL](#), [GeneReviews_UCSC](#), [GAD_UCSC](#), [xenorefGenes_UCSC](#), [transcript_ENSEMBL](#),

Examples

```
library("Gviz")
gen <- "hg19"
chrom <- "chr2"
start <- 38290160
end <- 38303219
if(interactive()) {
  genetrack <- genes_ENSEMBL(gen, chrom, start, end, showId=TRUE)
  plotTracks(genetrack, from = start, to = end,
    fontfamily="sans", fontfamily.title="sans")
} else {
  data(geneENSEMBLtrack)
  plotTracks(genetrack, from = start, to = end,
    fontfamily="sans", fontfamily.title="sans")
}
```

GWAScatalog_UCSC

Create one track of the genomic positions of variants from the GWAS catalog

Description

Create one track of the genomic positions of variants from the NHGRI Catalog of Published Genome-Wide Association Studies using the Gviz bioconductor package

Usage

```
GWAScatalog_UCSC(gen, chr, start, end, title="GWAS Catalog", showId=FALSE)
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	The name of the annotation track
showId	Show the ID of the genetic elements

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjtrdrFAy6dn&c=chr6&g=gwa
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[ISCA_UCSC](#), [knownGenes_UCSC](#), [genesName_ENSEMBL](#), [GeneReviews_UCSC](#), [GAD_UCSC](#), [genes_ENSEMBL](#), [xenorefGenes_UCSC](#), [transcript_ENSEMBL](#),

Examples

```
library("Gviz")
gen <- "hg38"
chrom <- "chr2"
start <- 10000
end <- 100000

if(interactive()) {
  gwastrack <- GWAScatalog_UCSC(gen,chrom,start,end)
  plotTracks(gwastrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(GWASTrack)
  plotTracks(gwastrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}
```

HiCdata2matrix	<i>Creates a HiC matrix from a file (Rao et al., 2014)</i>
----------------	--

Description

Creates a HiC matrix from Rao et al.,2014.

Usage

```
HiCdata2matrix( chr, start, end, bedFilePath)
```

Arguments

chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
bedFilePath	The path of the BED file from Kheradpour and Kellis, 2014.

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to BindingMotifsBiomart binding motif biomart

Examples

```
library("corrplot")
gen <- "hg19"
chr<-"chr1"
start <- 5000000
end <- 9000000

extdata <- system.file("extdata", package="coMET",mustWork=TRUE)
bedFilePath <- file.path(extdata, "HiC/chr1_1mb.RAWobserved")

if(interactive()){
  matrix_HiC_Rao <- HiCdata2matrix(chr,start, end, bedFilePath)
  cor_matrix_HiC <- cor(matrix_HiC_Rao)
  diag(cor_matrix_HiC)<-1
  corrplot(cor_matrix_HiC, method = "circle")
} else {
```

```

data(matrix_HiC_Rao)
cor_matrix_HiC <- cor(matrix_HiC_Rao)
diag(cor_matrix_HiC)<-1
corrplot(cor_matrix_HiC, method = "circle")
}

```

HistoneAll_UCSC	<i>Create multiple tracks of histone modifications from the UCSC genome browser</i>
-----------------	---

Description

Create multiple tracks of histone modifications from the UCSC genome browser (ENCODE/Broad) using the Gviz bioconductor package

Usage

```

HistoneAll_UCSC(gen, chr, start, end, mySession, pattern = NULL,
               track.name = "Broad Histone", table.name = NULL)

```

Arguments

gen	the name of the genome. Data is not currently available for GRCh38 (hg38).
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
mySession	the object session from the function browserSession of rtracklayer
pattern	The cell type
track.name	the name of the track, for example: "Broad Histone"
table.name	the name of the table from the track

Value

A list of AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=wg
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[HistoneOne_UCSC](#),

Examples

```
library("Gviz")
library("rtracklayer")
gen <- "hg19"
chr <- "chr2"
start <- 38290160
end <- 38313219

if(interactive()){
  BROWSER.SESSION="UCSC"
  mySession <- browserSession(BROWSER.SESSION)
  genome(mySession) <- gen
  pattern1 <- "GM12878"

  histonalltrack<-HistoneAll_UCSC(gen,chr,start,end,mySession,
  pattern=pattern1,track.name="Broad Histone")
  plotTracks(histonalltrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(histonalltrack)
  plotTracks(histonalltrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}
```

HistoneOne_UCSC

Create one track of one histone modification profile from the UCSC genome browser

Description

Create one track of one histone modification profile from the UCSC genome browser (ENCODE/Broad) using the Gviz bioconductor package

Usage

```
HistoneOne_UCSC(gen, chr, start, end, mySession, title="Broad Histone",
track.name = "Broad Histone", table.name = NULL)
```

Arguments

gen	the name of the genome. Data is not currently available for GRCh38 (hg38).
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)

<code>mySession</code>	the object session from the function <code>browserSession</code> of <code>rtracklayer</code>
<code>title</code>	Name of tracks
<code>track.name</code>	the name of the track, for example: "Broad Histone"
<code>table.name</code>	the name of the table from the track

Value

An `AnnotationTrack` object of `Gviz`

Author(s)

Tiphaine Martin

References

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=wg
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[HistoneAll_UCSC](#)

Examples

```
library("Gviz")
library("rtracklayer")
gen <- "hg19"
chr <- "chr2"
start <- 38290160
end <- 38303219

if(interactive()) {
  BROWSER.SESSION="UCSC"
  mySession <- browserSession(BROWSER.SESSION)
  genome(mySession) <- gen
  histoneonetrack<-HistoneOne_UCSC(gen,chr,start,end,mySession)
  plotTracks(histoneonetrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(histoneonetrack)
  plotTracks(histoneonetrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}
```

imprintedGenes_GTEx *Creates a imprinted genes track from GTEx*

Description

Creates a track of imprinted genes from GTEx using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
imprintedGenes_GTEx(gen="hg19", chr,start, end, tissues="all",  
classification="all",showId=FALSE, title="Imprinted genes GTEx")
```

Arguments

gen	the name of the genome. Default value=hg19
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
tissues	list of tissues among 33 tissues in GTEx
classification	list of classification from 5 types (biallelic, consistent with biallelic, consistent with imprinting, imprinted, NC)
showId	logical. say if we write the name of group
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>
Got to BindingMotifsBiomart binding motif biomart

Examples

```
library("Gviz")  
gen<-"hg19"  
chr<- "chr6"  
start <- 144251437  
end <- 144330541
```

```

if(interactive()){
  allIGtrack <- imprintedGenes_GTEEx(gen,chr,start, end,
    tissues="all", classification="imprinted",showId=TRUE)
  allimprintedIGtrack <- imprintedGenes_GTEEx(gen,chr,start, end,
    tissues="all", classification="imprinted",showId=TRUE)
  StomachIGtrack <-imprintedGenes_GTEEx(gen,chr,start, end,
    tissues="Stomach", classification="all",showId=TRUE)
  PancreasIGtrack <- imprintedGenes_GTEEx(gen,chr,start, end,
    tissues="Pancreas", classification="all",showId=TRUE)
  PancreasimprintedIGtrack <- imprintedGenes_GTEEx(gen,chr,start, end,
    tissues="Pancreas", classification="biallelic",showId=TRUE)

  imprintinglist <- list(allIGtrack,allimprintedIGtrack,
    StomachIGtrack,PancreasIGtrack,PancreasimprintedIGtrack)

  plotTracks(imprintinglist, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")

} else {

  data(allIGtrack)
  data(allimprintedIGtrack)
  data(StomachIGtrack)
  data(PancreasIGtrack)
  data(PancreasimprintedIGtrack)

  imprintinglist <- list(allIGtrack,allimprintedIGtrack,
    StomachIGtrack,PancreasIGtrack,PancreasimprintedIGtrack)

  plotTracks(imprintinglist, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
}

```

interestGenes_ENSEMBL *Create one track of the genes in the genomic regions of interest from EMSEMBL*

Description

Create one track of the genes in the genomic regions of interest from EMSEMBL using the Gviz bioconductor package

Usage

```
interestGenes_ENSEMBL(gen, chr, start, end, interestfeatures,interestcolor,
  showId=FALSE,title="genes (ENSEMBL)")
```

Arguments

<code>gen</code>	the name of the genome
<code>chr</code>	the chromosome of interest
<code>start</code>	the first position in the region of interest (the smallest value)
<code>end</code>	the last position in the region of interest (the largest value)
<code>interestfeatures</code>	A data frame with 3 columns: start of features, end of features, and type of features
<code>interestcolor</code>	A list with the color for each new features defined
<code>showId</code>	Show the ID of the genetic elements
<code>title</code>	Name of tracks

Value

A `BiomartGeneRegionTrack` object of `Gviz`

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtDrFAy6dn&c=chr6&g=ens

See Also

[ISCA_UCSC](#), [GWAScatalog_UCSC](#), [knownGenes_UCSC](#), [genesName_ENSEMBL](#), [GeneReviews_UCSC](#), [GAD_UCSC](#), [xenorefGenes_UCSC](#), [transcript_ENSEMBL](#),

Examples

```
library("Gviz")
gen <- "hg19"
chr <- "chr15"
start <- 75011669
end <- 75019876
interestfeatures <- rbind(c("75011883", "75013394", "bad"), c("75013932", "75014410", "good"))
interestcolor <- list("bad"="red", "good"="green")

if(interactive()) {
  interestgenesENSMBLtrack<-interestGenes_ENSEMBL(gen,chr,start,end,
  interestfeatures,interestcolor,showId=TRUE)
  plotTracks(interestgenesENSMBLtrack, from = start, to =end,
             fontfamily="sans", fontfamily.title="sans")
} else {
  data(interestgenesENSMBLtrack)
  plotTracks(interestgenesENSMBLtrack, from = start, to =end,
```

```
fontfamily="sans",fontfamily.title="sans")
}
```

```
interestTranscript_ENSEMBL
```

Create a track of transcripts from ENSEMBL

Description

Create a track to visualize different transcripts from ENSEMBL using the Gviz bioconductor package

Usage

```
interestTranscript_ENSEMBL(gen, chr, start, end,interestfeatures,
interestcolor,showId = FALSE, title="transcripts ENSEMBL")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
interestfeatures	A data frame with 3 columns: start of features, end of features, and type of features
interestcolor	A list with the color for each new features defined
showId	Show the ID of the genetic elements
title	Name of tracks

Value

A BiomartGeneRegionTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=ens

See Also

[ISCA_UCSC](#), [GWAScatalog_UCSC](#), [knownGenes_UCSC](#), [genesName_ENSEMBL](#), [GeneReviews_UCSC](#), [GAD_UCSC](#), [genes_ENSEMBL](#), [xenorefGenes_UCSC](#),

Examples

```
library("Gviz")
gen <- "hg19"
chr <- "chr15"
start <- 75011669
end <- 75019876
interestfeatures <- rbind(c("75017782", "75017835", "bad"), c("75013755", "75013844", "good"))
interestcolor <- list("bad"="red", "good"="green")

if(interactive()){
  interesttransENSMBltrack<-interestTranscript_ENSEMBL(gen,chr,start,end,
  interestfeatures,interestcolor,showId=TRUE)
  plotTracks(interesttransENSMBltrack, from=start, to=end,
             fontfamily="sans", fontfamily.title="sans")
} else {
  data(interesttransENSMBltrack)
  plotTracks(interesttransENSMBltrack, from=start, to=end,
             fontfamily="sans", fontfamily.title="sans")
}
```

ISCA_UCSC

Create one track of the genomic positions of variants from ISCA [obselete database]

Description

Create one track of the genomic positions of variants from International Standards for Cytogenomic Arrays (ISCA) Consortium using the Gviz bioconductor package (obselete database, Impossible to access to data from UCSC from September 2015)

Usage

```
ISCA_UCSC(gen, chr, start, end, mySession, table.name, title="ISCA", showId=FALSE)
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
mySession	the object session from the function browserSession of rtracklayer

table.name	A table of ISCAT classifications: iscaBenign, iscaCuratedBenign, iscaCurated-Pathogenic, iscaLikelyBenign, iscaLikelyPathogenic, iscaPathGainCum, iscaPathLossCum, iscaPathogenic, iscaUncertain
title	The name of the annotation track
showId	Show the ID of the genetic elements

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=isca
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[GWScatalog_UCSC](#), [knownGenes_UCSC](#), [genesName_ENSEMBL](#), [GeneReviews_UCSC](#), [GAD_UCSC](#), [genes_ENSEMBL](#), [xenorefGenes_UCSC](#), [transcript_ENSEMBL](#),

Examples

```
# Oboelet function

library("Gviz")
library("rtracklayer")
gen <- "hg19"
chr <- "chr2"
start <- 38292433
end <- 38305492

if(interactive()){
  BROWSER.SESSION="UCSC"
  mySession <- browserSession(BROWSER.SESSION)
  genome(mySession) <- gen
  iscatrack <- ISCA_UCSC(gen,chrom,start,end,mySession,title="ISCA", table="iscaPathogenic")
  plotTracks(iscatrack, from = start, to =end,
             fontfamily="sans", fontfamily.title="sans")
} else {
  data(ISCAtrack_Grch38)
  plotTracks(iscatrack, from = start, to =end,
             fontfamily="sans", fontfamily.title="sans")
}
```

knownGenes_UCSC	<i>Create a track of known genes from the UCSC genome browser</i>
-----------------	---

Description

Create a track of known genes from the UCSC genome browser using the Gviz bioconductor package

Usage

```
knownGenes_UCSC(gen, chr, start, end, title="UCSC known Genes", showId=TRUE)
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	Name of tracks
showId	Show the ID of the genetic elements

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=knownGenes_UCSC
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[ISCA_UCSC](#), [GWAScatalog_UCSC](#), [genesName_ENSEMBL](#), [GeneReviews_UCSC](#), [GAD_UCSC](#), [genes_ENSEMBL](#), [xenorefGenes_UCSC](#), [transcript_ENSEMBL](#),

Examples

```

library("Gviz")
gen <- "hg38"
chr <- "chr7"
start <- 38290160
end <- 38303219

if(interactive()) {
  genesUcsctrack<-knownGenes_UCSC(gen,chr,start,end,showId=TRUE)
  plotTracks(genesUcsctrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}else {
  data(genesUcsctrack)
  plotTracks(genesUcsctrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}

```

metQTL

*Creates a track from a file for metQTL data***Description**

Creates a track from a BED file for metQTL data using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
metQTL(gen, chr, start, end, bedFilePath, featureDisplay, showId=FALSE,
       type_stacking="squish",just_group="above", title="metQTL")
```

Arguments

gen	the name of the genome. Default value=hg19
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
bedFilePath	The file path to the .BED file containing the data to be visualised
featureDisplay	A vector of metQTL features to be displayed, such as SNP. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. featureDisplay <- "CpG"), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. featureDisplay <- c("SNP","CpG")). Finally, visualison all features in the genomic region, achived by using the word "all" (e.g. featureDisplay <- "all"), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.

showId	Allows the visualization of the Id of metQTL group.
type_stacking	Sets the type of stacking used by Gviz for plots. By default this is set to 'squish'. For more information see Gviz user guide.
just_group	position. say where we write the name of group (choice in c("above","right","left"))
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin
Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>
Got to ENSEMBLregulation binding motif biomart

Examples

```
library("Gviz")

gen <- 'hg19'
chr <- "chr15"
start <- 74889136
end <- 75018200
featureDisplay <- "trans_local_metQTL"
type_stacking <- "squish"

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
mqtbedFilePath <- file.path(extdata, "metQTL.bed")

if(interactive()){
  metQTLTrackSingle <- metQTL(gen,chr,start, end,mqtbedFilePath,
    featureDisplay = featureDisplay )
  plotTracks(metQTLTrackSingle, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
} else {
  data(metQTLTrackSingle)
  plotTracks(metQTLTrackSingle, from = start, to = end,
    fontfamily="sans",fontfamily.title="sans")
}

###

library("Gviz")

gen <- 'hg19'
```

```

chr <- "chr15"
start <- 74889136
end <- 75018200

featureDisplay <- c("trans_local_metQTL", "CpG")

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
bedFilePath <- file.path(extdata, "metQTL.bed")

if(interactive()){
  metQTLTrackMultiple <- metQTL(gen,chr,start, end, bedFilePath,
    featureDisplay = featureDisplay )
  plotTracks(metQTLTrackMultiple, from = start, to = end,
    fontfamily="sans", fontfamily.title="sans")
} else {
  data(metQTLTrackMultiple)
  plotTracks(metQTLTrackMultiple, from = start, to = end,
    fontfamily="sans", fontfamily.title="sans")
}

#####

library("Gviz")

gen <- 'hg19'
chr <- "chr15"
start <- 74889136
end <- 75018200

featureDisplay <- "all"

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
bedFilePath <- file.path(extdata, "metQTL.bed")

if(interactive()){
  metQTLTrackAll <- metQTL(gen,chr,start, end, bedFilePath,
    featureDisplay = featureDisplay )
  plotTracks(metQTLTrackAll, from = start, to = end,
    fontfamily="sans", fontfamily.title="sans")
} else {
  data(metQTLTrackAll)
  plotTracks(metQTLTrackAll, from = start, to = end,
    fontfamily="sans", fontfamily.title="sans")
}

```

```
miRNATargetRegionsBiomart_ENSEMBL
```

Creates a track of miRNA target regions from ENSEMBL

Description

Creates a track of miRNA target regions from ENSEMBL using the Gviz bioconductor package.

Usage

```
miRNATargetRegionsBiomart_ENSEMBL(gen, chr, start, end, showId=FALSE,
  datasetEnsembl = "hsapiens_mirna_target_feature",
  title="miRNA Target Regions ENSEMBL")
```

Arguments

gen	The name of the genome. Currently only handles human data from either the previous version, GRCh37 (also known as hg19) or the current version, GRCh38 (also known as hg38).
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
showId	Show the ID of the genetic elements
datasetEnsembl	Allows the user to manually set which data set is used if required. Default=hsapiens_mirna_target_feature
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin
Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>
Got to ENSEMBLregulation binding motif biomart

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 1000000
end <- 2000000

if(interactive()){
  miRNATargetRegionsBiomartTrack<-miRNATargetRegionsBiomart_ENSEMBL(gen,chr,start,end,
    datasetEnsembl = "hsapiens_mirna_target_feature")
  plotTracks(miRNATargetRegionsBiomartTrack, from = start, to = end,
```

```

                                fontfamily="sans",fontfamily.title="sans")
} else {
  data(miRNATargetRegionsBiomartTrack)
  plotTracks(miRNATargetRegionsBiomartTrack, from = start, to = end,
                                fontfamily="sans",fontfamily.title="sans")
}

```

`otherRegulatoryRegionsBiomart_ENSEMBL`

Creates a track of other regulatory regions from ENSEMBL

Description

Creates a track from ENSEMBL of other regulatory regions using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```

otherRegulatoryRegionsBiomart_ENSEMBL(gen, chr, start, end,
featureDisplay = "all",datasetEnsembl = "hsapiens_external_feature",
title="Other Regulatory Regions ENSEMBL")

```

Arguments

<code>gen</code>	The name of the genome. Currently only handles human data from either the previous version, GRCh37 (also known as hg19) or the current version, GRCh38 (also known as hg38).
<code>chr</code>	The chromosome of interest
<code>start</code>	The starting position in the region of interest (the smallest value)
<code>end</code>	The end position in the region of interest (the largest value)
<code>featureDisplay</code>	A vector of regulatory features to be displayed, such as Enhancer. Spelling and capitalisation of features must be identical to those in the user guide. There are two possibilities. First, the visualisation of only one feature (e.g. <code>featureDisplay <- "Enhancer"</code>), only the name of the specific feature is required. Second, visualisation all features in the genomic region, achieved by using the word "all" (e.g. <code>featureDisplay <- "all"</code>), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
<code>datasetEnsembl</code>	Allows the user to manually set which data set is used if required.
<code>title</code>	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin
Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>
Got to ENSEMBLregulation binding motif biomart

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 100000
end <- 5000000
featureDisplay <- "Enhancer"

if(interactive()){
  otherRegulatoryRegionsTrackSingle<-otherRegulatoryRegionsBiomart_ENSEMBL(gen,
  chr,start,end,featureDisplay)
  plotTracks(otherRegulatoryRegionsTrackSingle, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(otherRegulatoryRegionsTrackSingle)
  plotTracks(otherRegulatoryRegionsTrackSingle, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}

#####

library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 100000
end <- 5000000
featureDisplay <- "all"

if(interactive()){
  otherRegulatoryRegionsTrackAll<-otherRegulatoryRegionsBiomart_ENSEMBL(gen,
  chr,start,end,featureDisplay)
  plotTracks(otherRegulatoryRegionsTrackAll, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(otherRegulatoryRegionsTrackAll)
  plotTracks(otherRegulatoryRegionsTrackAll, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}
```

pizza

Pizza color wheel

Description

This function displays a color wheel with specified colors

Usage

```
pizza(colors, bg = "gray95", border = NA,
       init.angle = 105, cex = 0.8, lty = 1, labcol = NULL,
       ...)
```

Arguments

colors	a vector with R color names of colors in hexadecimal notation
bg	background color of the plot. Default "gray95"
border	color of the border separating the pizza slices
init.angle	integer value indicating the start angle (in degrees) for the slices
cex	numeric value indicating the character expansion of the labels
lty	argument passed to <code>polygon</code> which draws each slice
labcol	color for the labels (i.e. names of the colors)
...	graphical parameters (<code>par</code>) can be given as argument to <code>pizza</code>

Details

This function is based on the `pie` function

Author(s)

Gaston Sanchez

Examples

```
# pizza color wheel for rainbow colors
pizza(rainbow(7))

# pizza color wheel for tomato (18 colors)
pizza(setColors("tomato", 18), bg = "gray20", cex = 0.7)
```

psiQTL_GTEEx	<i>Creates a psiQTL track from GTEEx</i>
--------------	--

Description

Creates a track of psiQTL from GTEEx using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
psiQTL_GTEEx(gen,chr,start, end, bedFilePath, featureDisplay = 'all',
showId=FALSE, type_stacking="squish",just_group="above", title="psiQTL GTEEx")
```

Arguments

gen	the name of the genome.
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
bedFilePath	The path of the BED file from Kheradpour and Kellis, 2014.
featureDisplay	A vector of regulatory features to be displayed, such as Predicted heterochomatin. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. featureDisplay <- "Predicted heterochomatin"), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. featureDisplay <- c("Predicted low activity","Predicted heterochomatin")). Finally, visualison all features in the genomic region, achived by using the word "all" (e.g. featureDisplay <- "all"), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
showId	logical. say if we write the name of group
type_stacking	Object of class"character", the stacking type of overlapping items on the final plot.One in c(hide, dense, squish, pack,full). More information cf the option "stacking" in Gviz
just_group	position. say where we write the name of group (choice in c("above","righ","left"))
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to BindingMotifsBiomart binding motif biomart

Examples

```
library("Gviz")
gen <- "hg19"
chr<-"chr13"
start <- 52713837
end <- 52715894
extdata <- system.file("extdata", package="coMET",mustWork=TRUE)
psiQTLFilePath <- file.path(extdata, "/GTEX/psiQTL_Assoc-total.AdiposeTissue.txt")
```

```
if(interactive()){
  psiGTexTrackall<- psiQTL_GTex(gen,chr,start, end, psiQTLFilePath,
    featureDisplay = 'all', showId=TRUE, type_stacking="squish",
    just_group="above" )
  plotTracks(psiGTexTrackall, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(psiGTexTrackall)
  plotTracks(psiGTexTrackall, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}
```

```
if(interactive()){
  psiGTexTrackSNP<- psiQTL_GTex(gen,chr,start, end, psiQTLFilePath,
    featureDisplay = 'SNP', showId=TRUE, type_stacking="squish",
    just_group="left")
  plotTracks(psiGTexTrackSNP, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(psiGTexTrackSNP)
  plotTracks(psiGTexTrackSNP, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}
```

refGenes_UCSC

Create a track of RefSeq genes from the UCSC genome browser

Description

Create a track of RefSeq genes from the UCSC genome browser using the Gviz bioconductor package

Usage

```
refGenes_UCSC(gen, chr, start, end, title="Ref Genes UCSC", track = "refGene",
  IdType="Ref", showId=TRUE)
```

Arguments

gen	The name of the genome
chr	The chromosome of interest
start	The first position in the region of interest (the smallest value)
end	The last position in the region of interest (the largest value)
title	Name of tracks
track	the name of table in UCSC for the group "Genes and Gene Prediction"
IdType	When set to 'ref' shows the gene reference, when set to "name" shows the gene name
showId	Shows the ID or name of the genetic elements

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin
Tom Hardiman

References

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=known
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[ISCA_UCSC](#), [GWAScatalog_UCSC](#), [genesName_ENSEMBL](#), [GeneReviews_UCSC](#), [GAD_UCSC](#), [genes_ENSEMBL](#), [xenorefGenes_UCSC](#), [transcript_ENSEMBL](#), [knownGenes_UCSC](#)

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr7"
start <- 38203219
end <- 38303219
IdType <- "name"
track <- "refGene"

if(interactive()) {
  genesUcsctrack<-refGenes_UCSC(gen,chr,start,end,track,IdType)
```

```
    plotTracks(genesUcsctrack, from = start, to =end)
  }else {
    data(genesUcsctrack)
    plotTracks(genesUcsctrack, from = start, to =end)
  }
```

regulationBiomart_ENSEMBL

Create a regulation track from ENSEMBL

Description

Create a 'Regulation' track from ENSEMBL using the Gviz bioconductor package

Usage

```
regulationBiomart_ENSEMBL(gen, chr, start, end,title="Regulation ENSEMBL")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	Name of tracks

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to ENSEMBLregulation biomart

Examples

```

library("Gviz")
gen <- "hg19"
chr <- "chr7"
start <- 38290160
end <- 38303219

if(interactive()){
  regulationENSEMBLtrack<-regulationBiomart_ENSEMBL(gen,chr,start,end)
  plotTracks(regulationENSEMBLtrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(regulationENSEMBLtrack)
  plotTracks(regulationENSEMBLtrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}

```

```
regulatoryEvidenceBiomart_ENSEMBL
```

Creates a regulatory feature track from ENSEMBL

Description

Creates a regulatory feature track from ENSEMBL using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
regulatoryEvidenceBiomart_ENSEMBL (gen, chr, start, end,
featureDisplay = "all", datasetEnsembl = "hsapiens_annotated_feature",
title="Other Regulatory Regions ENSEMBL")
```

Arguments

gen	The name of the genome. Currently only handles human data from either the previous version, GRCh37 (also known as hg19) or the current version, GRCh38 (also known as hg38).
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
featureDisplay	A vector of regulatory features to be displayed, such as DNase1. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. featureDisplay <- "DNase1"), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. featureDisplay <- c("CTCF","DNase1")). Finally, visualison all features in the

genomic region, achieved by using the word "all" (e.g. `featureDisplay <- "all"`), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.

`datasetEnsembl` Allows the user to manually set which data set is used if required.
`title` The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to ENSEMBLregulation binding motif biomart

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 40000
end <- 50000
featureDisplay <- "H3K27me3"

if(interactive()){
  regulatoryEvidenceBiomartTrackSingle <- regulatoryEvidenceBiomart_ENSEMBL(gen,
  chr,start,end,featureDisplay)
  plotTracks(regulatoryEvidenceBiomartTrackSingle, from = start, to = end,
             fontfamily="sans", fontfamily.title="sans")
} else {
  data(regulatoryEvidenceBiomartTrackSingle)
  plotTracks(regulatoryEvidenceBiomartTrackSingle, from = start, to = end,
             fontfamily="sans", fontfamily.title="sans")
}

#####

library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 40000
end <- 100000
featureDisplay <- c("H3K27me3","H3K36me3")

if(interactive()){
  regulatoryEvidenceBiomartTrackMultiple<-regulatoryEvidenceBiomart_ENSEMBL (gen,
```

```

chr,start,end,featureDisplay)
  plotTracks(regulatoryEvidenceBiomartTrackMultiple, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(regulatoryEvidenceBiomartTrackMultiple)
  plotTracks(regulatoryEvidenceBiomartTrackMultiple, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}

#####

library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 50000
end <- 100000
featureDisplay <- "all"
if(interactive()){
  regulatoryEvidenceBiomartTrackAll<-regulatoryEvidenceBiomart_ENSEMBL (gen,
chr,start,end,featureDisplay)
  plotTracks(regulatoryEvidenceBiomartTrackAll, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(regulatoryEvidenceBiomartTrackAll)
  plotTracks(regulatoryEvidenceBiomartTrackAll, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}

```

```
regulatoryFeaturesBiomart_ENSEMBL
```

Creates a regulatory feature track from ENSEMBL

Description

Creates a regulatory feature track from ENSEMBL using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
regulatoryFeaturesBiomart_ENSEMBL(gen, chr, start, end, featureDisplay = "all",
datasetEnsembl = "hsapiens_regulatory_feature",
title="Regulatory Features ENSEMBL")
```

Arguments

gen	The name of the genome. Currently only handles human data from either the previous version, GRCh37 (also known as hg19) or the current version, GRCh38 (also known as hg38).
-----	--

<code>chr</code>	The chromosome of interest
<code>start</code>	The starting position in the region of interest (the smallest value)
<code>end</code>	The end position in the region of interest (the largest value)
<code>featureDisplay</code>	A vector of regulatory features to be displayed, such as Promoter. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. <code>featureDisplay <- "Promoter"</code>), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. <code>featureDisplay <- c("TF binding site","Promoter")</code>). Finally, visualisation all features in the genomic region, achieved by using the word "all" (e.g. <code>featureDisplay <- "all"</code>), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
<code>datasetEnsembl</code>	Allows the user to manually set which data set is used if required. Default=hsapiens_regulatory_feature
<code>title</code>	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to ENSEMBLregulation binding motif biomart

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 10000
end <- 500000
featureDisplay <- "Enhancer"

if(interactive()){
  regulatoryFeaturesBiomartTrackSingle<-regulatoryFeaturesBiomart_ENSEMBL(gen,
  chr,start,end,featureDisplay)
  plotTracks(regulatoryFeaturesBiomartTrackSingle, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(regulatoryFeaturesBiomartTrackSingle)
  plotTracks(regulatoryFeaturesBiomartTrackSingle, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}
```

```
#####

library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 10000
end <- 100000
featureDisplay <- c("CTCF Binding Site", "Enhancer")

if(interactive()){
  regulatoryFeaturesBiomartTrackMultiple<-regulatoryFeaturesBiomart_ENSEMBL(gen,
  chr,start,end,featureDisplay)
  plotTracks(regulatoryFeaturesBiomartTrackMultiple, from = start, to = end,
              fontfamily="sans",fontfamily.title="sans")
} else {
  data(regulatoryFeaturesBiomartTrackMultiple)
  plotTracks(regulatoryFeaturesBiomartTrackMultiple, from = start, to = end,
              fontfamily="sans",fontfamily.title="sans")
}

#####

library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 10000
end <- 50000
featureDisplay <- "all"
if(interactive()){
  regulatoryFeaturesBiomartTrackAll<-regulatoryFeaturesBiomart_ENSEMBL(gen,
  chr,start,end,featureDisplay)
  plotTracks(regulatoryFeaturesBiomartTrackAll, from = start, to = end,
              fontfamily="sans",fontfamily.title="sans")
} else {
  data(regulatoryFeaturesBiomartTrackAll)
  plotTracks(regulatoryFeaturesBiomartTrackAll, from = start, to = end,
              fontfamily="sans",fontfamily.title="sans")
}

}
```

regulatorySegmentsBiomart_ENSEMBL

Creates a binding motif track from ENSEMBL [obsolete]

Description

[obsolete] Creates a track of regulatory segments from ENSEMBL using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
regulatorySegmentsBiomart_ENSEMBL(gen, chr, start, end, featureDisplay = 'all',
datasetEnsembl = "hsapiens_external_feature",
title="External Regulatory ENSEMBL")
```

Arguments

gen	The name of the genome. Currently only handles human data from either the previous version, GRCh37 (also known as hg19) or the current version, GRCh38 (also known as hg38).
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
featureDisplay	A vector of regulatory features to be displayed, such as Predicted heterochomatin. Spelling and capitalisation of features must be identical to those in the user guide. There are three possibilities. First, the visualisation of only one feature (e.g. featureDisplay <- "Predicted heterochomatin"), only the name of the specific feature is required. Second, visualisation of a set of features, for this a vector of features is required (e.g. featureDisplay <- c("Predicted low activity", "Predicted heterochomatin")). Finally, visualison all features in the genomic region, achived by using the word "all" (e.g. featureDisplay <- "all"), "all" is set by default. You can find the complete list of features and their associated colours in the user guide.
datasetEnsembl	Allows the user to manually set which data set is used if required.Default=hsapiens_segmentation_feature
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

Got to ENSEMBLregulation binding motif biomart

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 10000
end <- 50000
```



```
featureDisplay <- "CTCF enriched"

if(interactive()){
  regulatorySegmentsBiomartTrackSingle<-regulatorySegmentsBiomart_ENSEMBL(gen,
  chr,start,end,featureDisplay)
  plotTracks(regulatorySegmentsBiomartTrackSingle, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(regulatorySegmentsBiomartTrackSingle)
  plotTracks(regulatorySegmentsBiomartTrackSingle, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}

####

library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 10000
end <- 50000
featureDisplay <- c("CTCF enriched","Predicted Promoter Flank")

if(interactive()){
  regulatorySegmentsBiomartTrackMultiple<-regulatorySegmentsBiomart_ENSEMBL(gen,
  chr,start,end,featureDisplay)
  plotTracks(regulatorySegmentsBiomartTrackMultiple, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(regulatorySegmentsBiomartTrackMultiple)
  plotTracks(regulatorySegmentsBiomartTrackMultiple, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}

####

library("Gviz")
gen <- "hg38"
chr <- "chr1"
start <- 10000
end <- 50000
featureDisplay <- "all"
if(interactive()){
  regulatorySegmentsBiomartTrackAll<-regulatorySegmentsBiomart_ENSEMBL(gen,
  chr,start,end,featureDisplay)
  plotTracks(regulatorySegmentsBiomartTrackAll, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(regulatorySegmentsBiomartTrackAll)
  plotTracks(regulatorySegmentsBiomartTrackAll, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}

}
```

repeatMasker_UCSC *Create one track of the genomic positions of regions from repeatMasker_UCSC*

Description

Create one track of the genomic positions of regions from repeatMasker_UCSC using the Gviz bioconductor package

Usage

```
repeatMasker_UCSC(gen, chr, start, end, title="RepeatMasker",
  showId=FALSE, type_stacking="full")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	The name of the annotation track
showId	Show the ID of the genetic elements
type_stacking	the type of stacking data for this track. More information go to Gviz (the option "stacking")

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=rms

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr2"
start <- 38290160
end <- 38303219
```

```

if(interactive()){
  rmtrack <-repeatMasker_UCSC(gen,chr,start,end,showId=TRUE)
  plotTracks(rmtrack, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(repeatMaskerTrack)
  plotTracks(rmtrack, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}

```

segmentalDups_UCSC	<i>Create one track of the genomic positions of regions from segmentalDups_UCSC</i>
--------------------	---

Description

Create one track of the genomic positions of regions from segmentalDups_UCSC using the Gviz bioconductor package

Usage

```
segmentalDups_UCSC(gen, chr, start, end,title="Segmental Dups UCSC")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	The name of the annotation track

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin
Tom Hardiman

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>
http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=rms

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr2"
start <- 100000
end <- 200000

if(interactive()){
  DupTrack <-segmentalDups_UCSC(gen,chr,start,end)
  plotTracks(DupTrack, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(DupTrack)
  plotTracks(DupTrack, from = start, to = end,
             fontfamily="sans",fontfamily.title="sans")
}
```

setColors*Set Colors for a color wheel*

Description

This function set a given number of colors to create a color wheel

Usage

```
setColors(color, num)
```

Arguments

color	an R color name or a color in hexadecimal notation
num	integer value indicating how many colors to be added to the wheel

Value

A character vector with the given color and the set of colors to create a wheel color

Author(s)

Gaston Sanchez

See Also

[col2HSV](#)

Examples

```
# create a color wheel based on 'tomato'
setColors("tomato", 12)

# set 7 colors for '#3D6DCC'
setColors("#3D6DCC", 7)
```

snpBiomart_ENSEMBL *Create a short variation track from ENSEMBL*

Description

Create a 'Short Variation' track from ENSEMBL using the Gviz bioconductor package

Usage

```
snpBiomart_ENSEMBL(gen,chr, start, end, dataset, showId=FALSE, title = "SNPs ENSEMBL")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
dataset	The name of the database. Example "hsapiens_snp_som"
showId	Show the the ID of element or not
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

Go to ENSEMBL Biomart
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[snpLocations_UCSC](#), [structureBiomart_ENSEMBL](#), [COSMIC_UCSC](#), [CoreillCNV_UCSC](#), [ClinVarMain_UCSC](#), [ClinVarCnv_UCSC](#),

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr2"
start <- 38290160
end <- 38303219

if(interactive()){
  snptrack <- snpBiomart_ENSEMBL(gen,chr, start, end,
                                dataset="hsapiens_snp",showId=FALSE)
  plotTracks(snptrack, from=start, to=end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(snpBiomarttrack)
  plotTracks(snptrack, from=start, to=end,
             fontfamily="sans",fontfamily.title="sans")
}
```

snpLocations_UCSC *Create a SNP track from UCSC*

Description

Create a SNP track from UCSC using the Gviz bioconductor package

Usage

```
snpLocations_UCSC(gen, chr, start, end,title= "SNPs UCSC", track="All SNPs(142)")
```

Arguments

gen	the name of the genome. Data is not currently available for GRCh38 (hg38).
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
title	Name of tracks
track	The name of the database. Default "All SNPs(142)"

Value

An UcsTrack object of Gviz

Author(s)

Tiphaine Martin

References

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=snp
<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[snpLocations_UCSC](#), [structureBiomart_ENSEMBL](#), [COSMIC_UCSC](#), [CoreillCNV_UCSC](#), [ClinVarMain_UCSC](#), [ClinVarCnv_UCSC](#),

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr7"
start <- 38290160
end <- 38303219

if(interactive()) {
  snpUCSCtrack<-snpLocations_UCSC(gen,chr,start,end,"All SNPs(142)")
  plotTracks(snpUCSCtrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(snpUCSCtrack)
  plotTracks(snpUCSCtrack, from = start, to =end,
             fontfamily="sans",fontfamily.title="sans")
}
```

structureBiomart_ENSEMBL

Create a structural variation track from ENSEMBL

Description

Create a 'Structural Variation' track from ENSEMBL using the Gviz bioconductor package

Usage

```
structureBiomart_ENSEMBL(gen, chr, start, end, strand, dataset,
showId=FALSE, title = "Structural variation")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)

strand	the strand to extract structure data for
dataset	The name of the database. Example "hsapiens_structvar_som"
showId	Show the the ID of the element
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

Go to ENSEMBL Biomart

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

See Also

[snpLocations_UCSC](#), [snpBiomart_ENSEMBL](#), [COSMIC_UCSC](#), [CoreillCNV_UCSC](#), [ClinVarMain_UCSC](#), [ClinVarCnv_UCSC](#),

Examples

```
library("Gviz")
gen <- "hg38"
chr <- "chr2"
start <- 38290160
end <- 38303219

if(interactive()){
  strutrack <- structureBiomart_ENSEMBL(chr, start, end,
                                       strand, dataset="hsapiens_structvar_som")
  plotTracks(strutrack, from=start, to=end,
             fontfamily="sans", fontfamily.title="sans")
}else {
  data(strucBiomarttrack)
  plotTracks(strutrack, from=start, to=end,
             fontfamily="sans", fontfamily.title="sans")
}
```

TFBS_FANTOM	<i>Creates a TFBS motif track from FANTOM</i>
-------------	---

Description

Creates a track of TFBS motifs from FANTOM using the Gviz bioconductor package. A complete list of features and their associated colours can be found in the user guide.

Usage

```
TFBS_FANTOM(gen, chr, start, end, bedFilePath, title="TF motif FANTOM5")
```

Arguments

gen	the name of the genome.
chr	The chromosome of interest
start	The starting position in the region of interest (the smallest value)
end	The end position in the region of interest (the largest value)
bedFilePath	The path of the BED file from Kheradpour and Kellis, 2014.
title	The name of the annotation track

Value

An AnnotationTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>
 Got to BindingMotifsBiomart binding motif biomart

Examples

```
library("Gviz")
gen <- "hg19"
chr <- "chr1"
start <- 6000000
end <- 6500000

extdata <- system.file("extdata", package="coMET", mustWork=TRUE)
AP1FantomFile <- file.path(extdata, "FANTOM/Fantom_hg19.AP1_MA0099.2.sites_example970.txt")

if(interactive()){
  tfbsFANTOMtrack <- TFBS_FANTOM(gen,chr,start, end, AP1FantomFile)
```

```

plotTracks(tfbsFANTOMtrack, from = start, to = end,
           fontfamily="sans", fontfamily.title="sans")
} else {
  data(tfbsFANTOMtrack)
  plotTracks(tfbsFANTOMtrack, from = start, to = end,
           fontfamily="sans", fontfamily.title="sans")
}

```

transcript_ENSEMBL *Create a track of transcripts from ENSEMBL*

Description

Create a track to visualize different transcripts from ENSEMBL using the Gviz bioconductor package

Usage

```
transcript_ENSEMBL(gen, chr, start, end, showId = FALSE, title="transcripts ENSEMBL")
```

Arguments

gen	the name of the genome
chr	the chromosome of interest
start	the first position in the region of interest (the smallest value)
end	the last position in the region of interest (the largest value)
showId	Show the ID of the genetic elements
title	Name of tracks

Value

A BiomartGeneRegionTrack object of Gviz

Author(s)

Tiphaine Martin

References

<http://bioconductor.org/packages/release/bioc/html/Gviz.html>

http://genome-euro.ucsc.edu/cgi-bin/hgTrackUi?hgsid=202839739_2hYQ1BAOuBMAR620GjrtdrFAy6dn&c=chr6&g=ens

See Also

[ISCA_UCSC](#), [GWAScatalog_UCSC](#), [knownGenes_UCSC](#), [genesName_ENSEMBL](#), [GeneReviews_UCSC](#), [GAD_UCSC](#), [genes_ENSEMBL](#), [xenorefGenes_UCSC](#),

Examples

```
library("Gviz")
gen <- "hg19"
chr <- "chr2"
start <- 32290160
end <- 33303219

if(interactive()){
  transENSMBLtrack<-transcript_ENSEMBL(gen,chr,start,end,showId=TRUE)
  plotTracks(transENSMBLtrack, from=start, to=end,
             fontfamily="sans",fontfamily.title="sans")
} else {
  data(transENSMBLtrack)
  plotTracks(transENSMBLtrack, from=start, to=end,
             fontfamily="sans",fontfamily.title="sans")
}
```

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