Package: DegCre (via r-universe)

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Type Package

Title Probabilistic association of DEGs to CREs from differential data

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Description DegCre generates associations between differentially
expressed genes (DEGs) and cis-regulatory elements (CREs) based
on non-parametric concordance between differential data. The
user provides GRanges of DEG TSS and CRE regions with
differential p-value and optionally log-fold changes and DegCre
returns an annotated Hits object with associations and their
calculated probabilities. Additionally, the package provides
functionality for visualization and conversion to other
formats.

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Encoding UTF-8

RoxygenNote 7.3.1

Depends R (>= 4.3)

URL https://github.com/brianSroberts/DegCre

Imports GenomicRanges, InteractionSet, plotgardener, S4Vectors, stats,
graphics, grDevices, BiocGenerics, GenomeInfoDb, IRanges, utils

biocViews GeneExpression, GeneRegulation, ATACSeq, ChIPSeq, DNaseSeq,
RNASeq

Suggests BSgenome, BSgenome.Hsapiens.UCSC.hg38, org.Hs.eg.db,
BiocStyle, magick, knitr, rmarkdown,
TxDb.Hsapiens.UCSC.hg38.knownGene,
TxDb.Mmusculus.UCSC.mm10.knownGene, testthat (>= 3.0.0)

VignetteBuilder knitr

Config/testthat/edition 3

BugReports <https://github.com/brianSroberts/DegCre/issues>

Repository https://bioc.r-universe.dev

RemoteUrl https://github.com/bioc/DegCre

RemoteRef HEAD

RemoteSha ee9a9ae2ced4973404a7e257db7c3afb2d24f556

Contents

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calcRawAssocProbOR *Calculate Raw Association Probability Odds Ratio (OR)*

Description

Given a DegCre results list, this function calculates the raw association probability odds ratio (OR) for each association.

Usage

```
calcRawAssocProbOR(degCreResList)
```

Arguments

degCreResList   List of DegCre results.

Details

This function calculates the raw association probability odds ratio (OR) for each association in a DegCre analysis output. The OR is calculated relative to the distance bin null association probability, which would happen if all CRE p-values were identical. Thus it is a measure of the increase in association probability due to CRE p-value information content over what would occur by random chance.

Value

A numeric vector of raw association probability odds ratios (OR) for each association.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load test data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                   DegP=subDegGR$pVal,
                                   DegLfc=subDegGR$logFC,
                                   CreGR=subCreGR,
                                   CreP=subCreGR$pVal,
                                   CreLfc=subCreGR$logFC)

#Calculate raw odds ratio.
ORvec <- calcRawAssocProbOR(degCreResListDexNR3C1)
```

```
convertDegCreDataFrame
```
Convert DegCre Results List to DataFrame

Description

Given a DegCre results list, this function converts it into a DataFrame for further analysis and export.

Usage

```
convertDegCreDataFrame(degCreResList, assocAlpha = 0.05)
```

Arguments

degCreResList   List of DegCre results.

assocAlpha   The significance threshold for associations to be included in the output (Default:
                0.05).

Details

This function takes a DegCre results list as input and extracts the significant associations based on
the adjusted p-values assocProbFDR compared to the specified significance threshold assocAlpha.
It then creates a [DataFrame](#) with the genomic coordinates of the significant associations from both
the DegGR and CreGR components of the input list. These are marked as Deg_ or Cre_ with chr,
start, end, and strand. The coordinates are followed by the metadata of the [Hits DataFrame](#)d
by [runDegCre](#). These are then followed by all metadata columns in the input DegGR or CreGR
proceeded by either Deg_ or Cre_ in the colname.

If no associations pass the significance threshold, the function returns NA.

Value

A [DataFrame](#) containing the significant associations that pass the specified significance threshold.
It is roughly in BEDPE format.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load test data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                    DegP=subDegGR$pVal,
                                    DegLfc=subDegGR$logFC,
                                    CreGR=subCreGR,
                                    CreP=subCreGR$pVal,
                                    CreLfc=subCreGR$logFC)
```

```
#Create DataFrame.
outDf <-
  convertDegCreDataFrame(degCreResList=degCreResListDexNR3C1,
                                      assocAlpha = 0.05)

#Write out as text file.
degCreDfFile <- tempfile(pattern="myDegCreResults",fileext=".tsv")

write.table(outDf,file=degCreDfFile[1],sep="\t",row.names=FALSE,quote=FALSE)

unlink(degCreDfFile[1])
```

---

convertdegCreResListToGInteraction

Convert DegCre Results List to GInteractions Object

Description

Given a DegCre results list, this function converts it into a GInteractions object.

Usage

```
convertdegCreResListToGInteraction(degCreResList, assocAlpha = 0.05)
```

Arguments

degCreResList	List of DegCre results.
assocAlpha	The significance threshold for associations to be included in the output (Default: 0.05).

Details

This function takes a DegCre results list as input and extracts the significant associations based on the assocProbFDR compared to the specified significance threshold assocAlpha. It then creates a [GInteractions](#) object metadata columns from the input list.

If no associations pass the significance threshold, the function returns NA' and prints a message.

Value

A [GInteractions](#) object containing the associations that pass the specified significance threshold.

The [GInteractions](#) object has same metadata columns as the [Hits](#) returned from [runDegCre](#) with additional columns. These additional columns are every metadata column in the input DegGR or CreGR proceeded by either Deg_ or Cre_ in the colname.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load test data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                   DegP=subDegGR$pVal,
                                   DegLfc=subDegGR$logFC,
                                   CreGR=subCreGR,
                                   CreP=subCreGR$pVal,
                                   CreLfc=subCreGR$logFC)

#Create GInteractions object.
gInteractions <-
 convertdegCreResListToGInteraction(degCreResList=degCreResListDexNR3C1,
                                    assocAlpha = 0.01)
```

DegCre   *DegCre*

Description

Probabilistic association of DEGs to CREs from differential data.

Author(s)

Maintainer: Brian S. Roberts <brianroberts1976@yahoo.com> (ORCID)

See Also

Useful links:

- https://github.com/brianSroberts/DegCre
- Report bugs at https://github.com/brianSroberts/DegCre/issues

---

degCrePRAUC   *Calculate PR AUC for DegCre results.*

Description

This function calculates the Precision-Recall Area Under the Curve (AUC) from a DegCre results list.

Usage

```
degCrePRAUC(
  degCreResList,
  makePlot = TRUE,
  nShuff = 100,
  alphaVal = degCreResList$alphaVal,
  nThresh = 200
)
```

Arguments

degCreResList	A list of DegCre results.
makePlot	Logical indicating whether to generate a plot of the Precision-Recall curve. (Default: `TRUE`)
nShuff	Integer number of shuffles for no-skill curve. (Default: `100`)
alphaVal	Numeric from 0 to 1 threshold alpha value of DEG significance. (Default: alpha value from `degCreResList`)
nThresh	Integer number of threshold values for the Precision-Recall curve. (Default: `200`)

Details

This function calculates the Precision-Recall curve and AUC based on the provided DegCre results. It also estimates the statistical significance of the AUC by shuffling the associations and calculating AUC for shuffled data. Note that the PR AUCs tend to be small (0.05-0.2). Under the calculation framework, a PR AUC of 1 could only be achieved from DegCre results in which every association involves a significant DEG and has an association probability of 1. This situation will never actually occur but serves as a theoretical optimum for comparison.

Value

Invisibly, a list containing:

actualTprPpvMat A matrix of actual True Positive Rate (TPR) and apparent Positive Predictive Value (PPV).

shuffTprQMat A matrix of shuffled TPR quantiles.

shuffPpvQMat A matrix of shuffled PPV quantiles.

AUC Numeric of the total Area Under the Curve (AUC) for the Precision-Recall curve.

deltaAUC Numeric of the difference in AUC between the actual curve and shuffled curves.

normDeltaAUC Numeric of the normalized difference in AUC.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load sample data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                   DegP=subDegGR$pVal,
                                   DegLfc=subDegGR$logFC,
                                   CreGR=subCreGR,
                                   CreP=subCreGR$pVal,
                                   CreLfc=subCreGR$logFC)

#Plot PR curve.

degCrePRAUC(degCreResList=degCreResListDexNR3C1)

#Get PR results with out plotting.

prAUCList <- degCrePRAUC(degCreResList=degCreResListDexNR3C1,
                         makePlot=FALSE)
```

DexNR3C1   *DegCre input data for examples.*

Description

DegCre input data for examples.

Format

A named list with two slots: DegGR and CreGR.

DegGR [GRanges](#) of RNA-seq data. The coordinates reference TSS sites. It has the following mcols:

    promGeneName [EPDNew](#) promoter names

    GeneSymb Gene symbols

    GeneID Ensembl gene ids

    baseMean baseMean values from [DESeq2](#)

    logFC Log-2 fold-changes from [DESeq2](#)

    pVal P-values from [DESeq2](#)

    pAdj Adjusted p-values from [DESeq2](#)

CreGR [GRanges](#) of differntial CRE data. The coordinates reference signal regions. It has the following mcols:

    logFC Log-2 fold-changes from [csaw](#)

    pVal P-values from [csaw](#)

    pAdj Adjusted p-values from [csaw](#)

Details

This is a list with two slots: DegGR and CreGR. This data was derived from work by McDowell et al. in which they generated RNA-seq and ChIP-seq data by treating A549 cells with dexamethasone at several time points. Specifically this is RNA-seq and NR3C1 ChIP-seq at four hours versus control.

Author(s)

Brian S. Roberts

References

<https://genome.cshlp.org/content/28/9/1272>

---

getAssocDistHits *Get Associations and Distances Between Genomic Regions*

Description

This function finds associations and distances between two sets of genomic regions.

Usage

```
getAssocDistHits(DegGR, CreGR, maxDist = 1e+06)
```

Arguments

`DegGR`	A [GRanges](#) object representing DEG TSSs.
`CreGR`	A [GRanges](#) object representing the CREs.
`maxDist`	Integer value representing the maximum distance allowed for associations. Regions further apart than this threshold will not be associated. (Default: `1e6`)

Details

This function identifies associations between genomic regions from two GenomicRanges objects (DegGR and CreGR) based on their spatial overlap within a specified maximum distance threshold using [findOverlaps](#). It calculates the distances with [distance](#) between associated regions and stores them in the metadata of the [Hits](#) object. Large values `maxDist` will require more computational resources.

Value

A Hits object containing associations and their distances between the genomic regions represented by DegGR and CreGR.

Author(s)

Brian S. Roberts

Examples

```
#Load sample data.
data(DexNR3C1)

# Get hits with association distances.
hits <- getAssocDistHits(DegGR = DexNR3C1$DegGR,
                         CreGR = DexNR3C1$CreGR,
                         maxDist = 1e6)
```

`getDistBinNullAssocProb`

Calculate Null Association Probability for Each Distance Bin

Description

Calculates the null association probability for each distance bin in the DegCre analysis.

Usage

```
getDistBinNullAssocProb(degCreResList)
```

Arguments

degCreResList   A list of DegCre results.

Details

This function takes the results of the DegCre analysis and computes the null association probability for each unique distance bin. The null association probability represents the expected proportion of differentially expressed genes (DEGs) in each distance bin under the null hypothesis.

Value

A matrix with these columns:

binAssocDist Numeric value representing the distance bin (TSS to CRE) in base pairs.

nullAssocProb Numeric value representing the null association probability of the bin.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load example data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                   DegP=subDegGR$pVal,
                                   DegLfc=subDegGR$logFC,
                                   CreGR=subCreGR,
                                   CreP=subCreGR$pVal,
                                   CreLfc=subCreGR$logFC)

# Calculate null association probabilities.
outNullMat <- getDistBinNullAssocProb(degCreResList = degCreResListDexNR3C1)
```

---

getExpectAssocPerDEG   *Get Expected Associations per DEG*

Description

Calculates the expected associations per DEG (Differentially Expressed Gene).

Usage

```
getExpectAssocPerDEG(degCreResList, geneNameColName = NULL, assocAlpha = 0.05)
```

Arguments

degCreResList   A list of DegCre results.

geneNameColName

         Character value of the name of the column in DegGR (that was inputted to [run-DegCre](#)) that contains gene names. If NULL, the function will attempt to automatically find the gene name column. (Default: NULL)

assocAlpha   Numeric value from 0 to 1 specifying the significance threshold for associations. (Default: 0.05)

Details

This function calculates the expected associations per DEG based on DegCre analysis results. It first filters significant associations based on the provided association significance threshold (assocAlpha) and then computes the expected associations per gene. The function returns a [DataFrame](#) with gene-level information, including expected associations, number of associations, and significance thresholds.

Value

A [DataFrame](#) with the all data in the input DegGR with these columns added:

geneName Character values of gene names extracted from geneNameColName column (or column found if geneNameColName = NULL) in DegGR.

expectAssocs Numeric values of the expected associations per gene.

nAssocs Integer value of the number of associations passing assocAlpha per gene.

assocAlpha Numeric value from 0 to 1 of input assocAlpha

degAlpha Numeric value from 0 to 1 of the significance threshold for DEGs. Obtained from degCreResList

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load example data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                   DegP=subDegGR$pVal,
                                   DegLfc=subDegGR$logFC,
                                   CreGR=subCreGR,
                                   CreP=subCreGR$pVal,
                                   CreLfc=subCreGR$logFC)


# Get expected associations per DEG
expectAssocsDf <- getExpectAssocPerDEG(degCreResList = degCreResListDexNR3C1,
                                       geneNameColName = "GeneSymb",
                                       assocAlpha = 0.05)

head(expectAssocsDf)
```

---

optimizeAlphaDegCre   *Run DegCre with DEG alpha optimization.*

Description

Runs DegCre across a set of DEG alpha thesholds to find optimal performance.

Usage

```
optimizeAlphaDegCre(
  DegGR,
  DegP,
  DegLfc = NULL,
  CreGR,
  CreP,
  CreLfc = NULL,
  reqEffectDirConcord = TRUE,
  padjMethod = "bonferroni",
  maxDist = 1e+06,
```

```
  verbose = FALSE,
  smallestTestBinSize = 100,
  fracMinKsMedianThresh = 0.2,
  testedAlphaVals = c(0.005, 0.01, 0.02, 0.03, 0.05, 0.1)
)
```

Arguments

DegGR	A [GRanges](#) object of gene TSSs. Multiple TSSs per gene are allowed.
DegP	A numeric vector of differential expression p-values for genes in DegGR.
DegLfc	A numeric vector of log fold-change values of differential expression for gene in DegGR. Required when reqEffectDirConcord = TRUE. (Default: NULL)
CreGR	A [GRanges](#) object of CRE regions.
CreP	A numeric vector differential signal p-values for regions in CreGR.
CreLfc	A numeric vector log fold-change values of differential signal for regions in CreGR. Required when reqEffectDirConcord = TRUE. (Default: NULL)
reqEffectDirConcord	
	A logical whether to require concordance between the effect direction between DEG and CRE differential values. (Default: NULL)
padjMethod	A character value indicating the method for p-value adjustment. Do not change from default under most circumstances. Can be any method name accepted by p.adjust() (Default: bonferroni)
maxDist	An integer value specifying the maximum distance for probability calculation of TSS to CRE associations. (Default: 1e6)
verbose	A logical indicating whether to print messages of step completion and algorithm results. (Default: NULL)
smallestTestBinSize	
	An integer value specifying the size (number of elements) of the smallest distance bin to be considered in the optimization algorithm. (Default: 100)
fracMinKsMedianThresh	
	A numeric value between 0 and 1 specifying the optimization criterion for the distance bin size algorithm (See Details). (Default: 0.2)
testedAlphaVals	
	A numeric vector of DEG alpha values to test (Default: c(0.005,0.01,0.02,0.03,0.05,0.1)).

Details

This function runs [runDegCre](#) for each value in testedAlphaVals. The performance at each tested alpha is evaluated with [degCrePRAUC](#). which generates a Precision-Recall curve based on the recovery rate of DEGs by associations. Various AUCs are calculated as performance metrics. Using the alpha with the highest value of normDeltaAUC is recommended (see Examples).

Value

A named list containing:

alphaPRMat A matrix of Precision-Recall Area Under the Curve (AUC) values.

degCreResListsByAlpha Named list of DegCre results lists indexed by the `testedAlphaVals`.

The columns of `alphaPRMat` are:

alphaVal Numeric vector of tested DEG alpha value.

AUC Numeric vector of Area under the curve of a Precision-Recall (PR) curve based on associations recovering significant DEGs.

deltaAUC Numeric vector of PR AUC minus the AUC of the no-skill line.

normDeltaAUC Numeric vector of the value of `deltaAUC` divided by one minus the no-skill AUC.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load sample data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

# Run DegCre over range of alpha values:
alphaOptList <- optimizeAlphaDegCre(DegGR = subDegGR,
                          DegP = subDegGR$pVal,
                          DegLfc = subDegGR$logFC,
                          CreGR = subCreGR,
                          CreP = subCreGR$pVal,
                          CreLfc = subCreGR$logFC)

bestAlphaId <- which.max(alphaOptList$alphaPRMat[,4])
bestDegCreResList <- alphaOptList$degCreResListsByAlpha[[bestAlphaId]]
```

---

plotBrowserDegCre   *Make Browser Plots from DegCre Results*

Description

Creates browser plots of specified genomic regions or gene regions based on the provided DegCre analysis results.

Usage

```
plotBrowserDegCre(
  degCreResList,
  assocAlpha = 0.05,
  browserWinPad = 1000,
  geneName = NULL,
  plotRegionGR = NULL,
  CreSignalName = "CRE",
  assembly = "hg38",
  plotWidth = grDevices::dev.size("in")[1],
  plotHeight = grDevices::dev.size("in")[2],
  plotXbegin = 0.9,
  mergeGenePromotersDist = 1000,
  sigPlotMaxY = 4,
  assocColorRange = NULL,
  lowAssocColor = "#88CCEE",
  hiAssocColor = "#CC6677",
  signalColor = "#DDCC77",
  geneLabelFontsize = 8,
  axisFontSize = 6,
  panelTitleFontSize = 7,
  geneNameColName = NULL,
  geneHighlightDf = NULL,
  dePrioritizeSmallRNA = TRUE,
  useLogFC = TRUE,
  creSignalBinRes = 100
)
```

Arguments

degCreResList	List of DegCre results.
assocAlpha	Numeric value from 0 to 1 of significance threshold for associations. (Default: 0.05)
browserWinPad	Numeric value of the padding size (in base pairs) to extend the plotting region. (Default: 1000)
geneName	Character of name of the gene of interest. If specified, the function will plot the region associated with this gene. (Default: NULL)
plotRegionGR	[GRanges](#) of length 1 specifying the region to plot. If provided, geneName is ignored. (Default: NULL)
CreSignalName	Character of name of the differential CRE signal track. For plot labeling purposes only (Default: CRE)
assembly	Character of genome assembly name, e.g., "hg38". Must be one of accepted inputs to assembly argument to [plotGenomeLabel.](#) (Default: hg38)
plotWidth	Numeric value of width of the browser plot in inches. (Default: dev.size("in")[1])
plotHeight	Numeric value of height of the browser plot in inches. (Default: dev.size("in")[2])

plotXbegin	Numeric value of the width of the left margin (where track annotations will be) in inches. (Default: `0.9`)
mergeGenePromotersDist	
	Maximum distance (in base pairs) for merging promoters of the same gene in plot. (Default: 1000)
sigPlotMaxY	Numeric value of maximum value for the CRE differential signal plot (Y-axis). (Default: 4)
assocColorRange	
	Numeric vector of values from 0 to 1 of length 2. These values specify the lower and upper values of DegCre association probabilities for color saturation for arch color. (Default: `NULL`. If `NULL` will be set to 0 and maximum association probability in input data.)
lowAssocColor	Character color for low saturation point of association probabilities in arches plot. (Default: `#88CCEE`)
hiAssocColor	Character color for high saturation point of association probabilities in arches plot. (Default: `#CC6677`)
signalColor	Character color for the CRE differential signal plot. (Default: `#DDCC77`)
geneLabelFontsize	
	Numeric of font size (as implemented in [plotgardener](#)) for gene labels. (Default: 8)
axisFontSize	Numeric of font size for axis labels and tick marks. (Default: 6)
panelTitleFontSize	
	Numeric of font size for panel titles. (Default: 7)
geneNameColName	
	Character of name of the column in DegGR metadata that was inputted to [runDegCre](#) that contains gene names to query by geneName. (Default: `NULL`. If omitted, the column name will be guessed, with warnings if not.)
geneHighlightDf	
	[DataFrame](#) specifying genes to highlight in the plot as accepted by [plotGenes](#) argument `geneHighlights`.
dePrioritizeSmallRNA	
	Logical, indicating whether small RNA genes should be deprioritized in plotting. (Default: `TRUE`)
useLogFC	Logical, indicating whether to use log-fold change values for the CRE differential signal. (Default: `TRUE`)
creSignalBinRes	
	Bin resolution in base pairs for the CRE signal track. Only used for initial calculation and will likely differ from display resolution. (Default: 100)

Details

This function uses [plotgardener](#) functionality to generate browser plots for visualizing DegCre analysis results in specified regions. The user can input genomic regions or gene names. The output plot consists of an arches plot made with [plotPairsArches](#) of DegCre associations colored by association probability. Below is a signal track plot made via [plotSignal](#) of the data in `CreGR` that was inputted

to runDegCre. This plot displays the signed negative log p-value, meaning the $-log_{10}(p_{CRE})$ multiplied by the sign of the the CRE log fold-change. Beneath this panel genomic coordinates via plotGenomeLabel and gene models via plotGenes are displayed.

Value

Invisibly, a named list containing:

plotRegionGR GRanges of the plotted region.

creSignalPlotGR GRanges of the CRE signal (signed negative log CRE p-value) across the plotted region.

assocGinter GInteractions of the DegCre associations in the plotted region.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load example data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                   DegP=subDegGR$pVal,
                                   DegLfc=subDegGR$logFC,
                                   CreGR=subCreGR,
                                   CreP=subCreGR$pVal,
                                   CreLfc=subCreGR$logFC)

#Make browser plot from specified gene name.
browserOuts <- plotBrowserDegCre(degCreResList=degCreResListDexNR3C1,
                                 geneName="ERRFI1",
                                 geneNameColName="GeneSymb",
                                 CreSignalName="NR3C1")
dev.off()

#Make plot of specified region.
zoomGR <- GenomicRanges::GRanges(seqnames="chr1",
                                 ranges=IRanges(start=7900e3,end=8400e3))

zoomedBrowserOuts <- plotBrowserDegCre(degCreResList=degCreResListDexNR3C1,
                                       plotRegionGR=zoomGR,
                                       geneNameColName="GeneSymb",
```

```
                                          CreSignalName="NR3C1")
  dev.off()
```

---

plotDegCreAssocProbVsDist

Plot DegCre Association Probability vs. Binned Genomic Distance

---

Description

Plots the DegCre association probability against binned genomic distance and highlights the quantile range.

Usage

```
plotDegCreAssocProbVsDist(
  degCreResList,
  assocProbFDRThresh = 0.05,
  plotQRange = c(0.25, 0.75),
  hiYLim = NULL,
  loYLim = NULL,
  qRangeFillColor = "#88CCEE",
  nullLineColor = "#CC6677"
)
```

Arguments

degCreResList	A list of DegCre results.
assocProbFDRThresh	
	Numeric value from 0 to 1 specifying the FDR threshold for association probability. (Default: `0.05`)
plotQRange	Numeric vector of quantile range for plotting (e.g., `c(0.25, 0.75)` for interquartile range). (Default: `c(0.25, 0.75)`)
hiYLim	Numeric value specifying the upper limit of the y-axis. (Default: NULL)
loYLim	Numeric value specifying the lower limit of the y-axis. (Default: NULL)
qRangeFillColor	
	Color for filling the quantile range polygon. (Default: #88CCEE)
nullLineColor	Color for the null association probability line. (Default: #CC6677)

Details

This function takes the results of the DegCre analysis, including genomic distances and association probabilities, and creates a plot of association probabilities against binned genomic distances. It highlights the quantile range (e.g., interquartile range) and includes a line for null association probabilities. The top panel shows the number of associations passing `assocProbFDRThresh`. The bottom panel shows the median FDR-passing association probability as a black line, with the specified

quantile range (defaults to interquartile) plotted as `qRangeFillColor` region. The `nullLineColor` colored line is the null association probability, that is the association probability for a bin with uniform CRE p-values.

Value

Invisibly, a matrix with these columns:

binMidDist Numeric value of the midpoint distance of the bin (TSS to CRE) in kb.

q_<`plotQRange[1]` x 100**>** Numeric value of lower bound of the highlight region.

q_50 Numeric value of plotted line.

q_<`plotQRange[2]` x 100**>** Numeric value of upper bound of the highlight region.

nullAssocProb Numeric of null association probability of the bin.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load example data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                   DegP=subDegGR$pVal,
                                   DegLfc=subDegGR$logFC,
                                   CreGR=subCreGR,
                                   CreP=subCreGR$pVal,
                                   CreLfc=subCreGR$logFC)

#Plot association probability versus binned genomic distance.

outProbVsDistMat <-
 plotDegCreAssocProbVsDist(degCreResList=degCreResListDexNR3C1)
```

```
plotDegCreBinHeuristic
```
Plot DegCre Bin Algorithm Statistics

Description

Plots the DegCre distance bin optimization statistic against different bin sizes, highlighting the optimal bin size.

Usage

```
plotDegCreBinHeuristic(degCreResList)
```

Arguments

degCreResList   A list of DegCre results.

Details

This function takes a DegCre results list and plots the bin heuristic statistics against different bin sizes. It also highlights the optimal bin size chosen based on the analysis. The y-axis of the plot is the median KS statistic of all bins versus the global CRE p-value distribution.

Value

Invisibly, the picked optimal bin size.

Author(s)

Brian S. Roberts

See Also

distBinHeuristic for calculating the DEG-CRE bin heuristic.

Examples

```
#Load required packages.
library(GenomicRanges)

#Load example data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
```

```
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                   DegP=subDegGR$pVal,
                                   DegLfc=subDegGR$logFC,
                                   CreGR=subCreGR,
                                   CreP=subCreGR$pVal,
                                   CreLfc=subCreGR$logFC)

#Plot distance bin median KS statistic curve.

plotDegCreBinHeuristic(degCreResList=degCreResListDexNR3C1)
```

---

plotExpectedAssocsPerDeg

Plot Histogram of Expected Associations per DEG

Description

Plots a histogram of the expected number of associations per DEG (Differentially Expressed Gene) based on DegCre analysis.

Usage

```
plotExpectedAssocsPerDeg(
  expectAssocPerDegDf,
  barOutlineColor = "#88CCEE",
  barFillColor = NULL,
  extraText = FALSE
)
```

Arguments

expectAssocPerDegDf

> [DataFrame](#) output of [getExpectAssocPerDEG](#).

barOutlineColor

> Color for the outline of the histogram bars. (Default: #88CCEE)

barFillColor   Fill color for the histogram bars. If `NULL`, it will be derived from `barOutlineColor` with adjusted transparency.

extraText   Logical, indicating whether additional text information (Details) should be added to the plot.

Details

This function generates a histogram of the expected number of associations per DEG and optionally adds additional text information to the plot, such as DEG FDR, association FDR, and the fraction of DEGs with at least one association. Plot displays a dashed line a value indicating the median expected associations per DEG.

Value

Invisibly, the median expected associations per DEG.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load example data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#Generate DegCre results.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                   DegP=subDegGR$pVal,
                                   DegLfc=subDegGR$logFC,
                                   CreGR=subCreGR,
                                   CreP=subCreGR$pVal,
                                   CreLfc=subCreGR$logFC)

# Generate data frame of expected associations per DEG
expectAssocPerDegDf <-
 getExpectAssocPerDEG(degCreResList = degCreResListDexNR3C1,
                      geneNameColName = "GeneSymb",
                      assocAlpha = 0.05)

# Plot histogram of expected associations per DEG
medianExpAssocs <- plotExpectedAssocsPerDeg(expectAssocPerDegDf,
                                            barOutlineColor = "blue",
                                            extraText = TRUE)
```

---

runDegCre   *Generate DegCre associations*

Description

Create DEG to CRE associations from differential data.

Usage

```
runDegCre(
  DegGR,
  DegP,
  DegLfc = NULL,
  CreGR,
  CreP,
  CreLfc = NULL,
  reqEffectDirConcord = TRUE,
  padjMethod = "bonferroni",
  maxDist = 1e+06,
  verbose = TRUE,
  smallestTestBinSize = 100,
  fracMinKsMedianThresh = 0.2,
  alphaVal = 0.01,
  binNOverride = NULL
)
```

Arguments

DegGR	A [GRanges](#) object of gene TSSs. Multiple TSSs per gene are allowed.
DegP	A numeric vector of differential expression p-values for genes in `DegGR`.
DegLfc	A numeric vector of log fold-change values of differential expression for gene in `DegGR`. Required when `reqEffectDirConcord = TRUE`. (Default: NULL)
CreGR	A [GRanges](#) object of CRE regions.
CreP	A numeric vector differential signal p-values for regions in `CreGR`.
CreLfc	A numeric vector log fold-change values of differential signal for regions in `CreGR`. Required when `reqEffectDirConcord = TRUE`. (Default: NULL)
reqEffectDirConcord	
	A logical whether to require concordance between the effect direction between DEG and CRE differential values. (Default: TRUE)
padjMethod	A character value indicating the method for p-value adjustment. Do not change from default under most circumstances. Can be any method name accepted by [p.adjust](#) (Default: bonferroni)
maxDist	An integer value specifying the maximum distance for probability calculation of TSS to CRE associations. (Default: 1e6)
verbose	A logical indicating whether to print messages of step completion and algorithm results. (Default: TRUE)
smallestTestBinSize	
	An integer value specifying the size (number of elements) of the smallest distance bin to be considered in the optimization algorithm. (Default: 100)
fracMinKsMedianThresh	
	A numeric value between 0 and 1 specifying the optimization criterion for the distance bin size algorithm (See Details). (Default: 0.2)

alphaVal   A numeric value between 0 and 1 specifying the alpha value for DEG signifi-
                  cance. (Default: `0.01`)

binNOverride   An integer value specifying the number of elements per distance bin. When
                  specified, overrides distance bin size optimization (Not recommended). (De-
                  fault: NULL)

Details

The DegCre algorithm considers experimental data from a perturbation experiment and produces
associations between cis-regulatory elements (CREs) and differentially expressed genes (DEGs).
The user provides differential expression data such as RNA-seq, and differential regulatory signal
data such as ATAC-seq, DNase Hypersensitivity, and ChIP-seq. For RNA-seq analysis, we suggest
methods such as DESeq2 or edgeR. For the analysis of differential regulatory data we recommend
csaw. As an example experiment, we use data from McDowell et al. (PMID = 30097539) in
which A549 cells were treated with dexamethasone and control. RNA-seq and ChIP-seq data were
collected at various time points.

A complete description of the mathematical basis of the DegCre core algorithms is provided in
DegCre bioRxiv. DegCre takes two inputs. The first is a GRanges of p-values and optionally
log fold-changes associated with DEG TSSs. The second input is a GRanges of differential sig-
nal p-values and optionally log fold-changes for CRE regions. DegCre generates a Hits object of
all associations between DEG TSSs and CREs within maxDist. Associations are then binned by
TSS-to-CRE distance according to an algorithm that balances resolution (many bins with few mem-
bers) versus minimization of the deviance of each bin's CRE p-value distribution from the global
distribution, seleting an optimal bin size.

Next, DegCre applies a non-parametric algorithm to find concordance between and CRE differential
effects within bins and derives an association probability. For all association probabilities involving
one given CRE, the probabilities are adjusted to favor associations across shorter distances. An
FDR of the association probability is then estimated. Results are returned in list containing a Hits
object and both input GRanges.

Value

A named list containing:

degCreHits A Hits object with metadata. The queryHits of degCreHits reference DegGR. The
    subjectHits of degCreHits reference CreGR

binHeurOutputs List of outputs from the distance binning algorithm.

alphaVal Numeric alpha value used for DEG significance threshold.

DegGR GRanges of input DegGR with added metadata columns "pVal", "pAdj",and possibly "logFC"
    if reqEffectDirConcord==TRUE. Will overwrite existing metadata with same colnames.

CreGR GRanges of input CreGR with added metadata columns "pVal", "pAdj",and possibly "logFC"
    if reqEffectDirConcord==TRUE. Will overwrite existing metadata with same colnames.

The degCreHits Hits object metadata has these columns:

assocDist Integer of distance in base pairs between the TSS and CRE for the association.

assocProb Numeric from 0 to 1 of association probability.

assocProbFDR Numeric from 0 to 1 of False discovery rate of the association probability exceeding distance only null.

rawAssocProb Numeric from 0 to 1 of association probability not adjusted for DEG significance or shorter associations involving this CRE.

CreP Numeric of differential p-value of the CRE.

DegP Numeric of differential p-value of the DEG.

DegPadj Numeric of differential adjusted p-value of the DEG.

binAssocDist Integer of the maximum association distance cutoff for the bin containing the association.

numObs Integer number of associations in the distance bin containing the association.

distBinId Integer that uniquely identifies the distance containing the association.

Author(s)

Brian S. Roberts

Examples

```
#Load required packages.
library(GenomicRanges)

#Load sample data.
data(DexNR3C1)

subDegGR <-
 DexNR3C1$DegGR[which(GenomeInfoDb::seqnames(DexNR3C1$DegGR)=="chr1")]
subCreGR <-
 DexNR3C1$CreGR[which(GenomeInfoDb::seqnames(DexNR3C1$CreGR)=="chr1")]

#With defaults.
degCreResListDexNR3C1 <- runDegCre(DegGR=subDegGR,
                                    DegP=subDegGR$pVal,
                                    DegLfc=subDegGR$logFC,
                                    CreGR=subCreGR,
                                    CreP=subCreGR$pVal,
                                    CreLfc=subCreGR$logFC)

#With custom settings.
modDegCreResList <- runDegCre(DegGR=subDegGR,
                             DegP=subDegGR$pVal,
                             CreGR=subCreGR,
                             CreP=subCreGR$pVal,
                             reqEffectDirConcord=FALSE,
                             maxDist=1e5,
                             alphaVal=0.001)
```

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