

Package ‘r3Cseq’

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Title Analysis of Chromosome Conformation Capture and Next-generation Sequencing (3C-seq)

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Depends GenomicRanges, Rsamtools, rtracklayer, VGAM, qvalue

Imports methods, GenomeInfoDb, IRanges, Biostrings, data.table, sqldf, RColorBrewer

Suggests BSgenome.Mmusculus.UCSC.mm9.masked,
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Description

This package is used for the analysis of long-range chromatin interactions from 3C-seq assay.

License GPL-3

URL <http://r3cseq.genereg.net>, <https://github.com/supatt-lab/r3Cseq/>

Collate AllClasses.R AllGenerics.R Export.R FunctionInCommon.R
FunctionsForBatchAnalysis.R RestrictionEnzymeFunctions.R
FunctionsForNoReplicationAnalysis.R Report.R Visualize3Cseq.R
Annotation.R

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calculateBatchRPM	<i>calculate read per million (RPM) for replicates analysis</i>
-------------------	---

Description

Normalize 3C-Seq data by transforming raw reads to read per million per each region for replication analysis

Usage

```
calculateBatchRPM(object,normalized_method=c("powerlawFittedRPM","normalRPM"))
```

Arguments

object	r3CseqInBatch object
normalized_method	character. method of normalization (default=powerlawFittedRPM)

Author(s)

S. Thongjuea

See Also

[calculateRPM](#), [expRPM](#) [contrRPM](#)

Examples

```
#See the vignette
```

calculateRPM	<i>calculate read per million (RPM)</i>
--------------	---

Description

Normalize 3C-Seq data by transforming raw reads to read per million per each region

Usage

```
calculateRPM(object,normalized_method=c("powerlawFittedRPM","normalRPM"))
```

Arguments

object	r3Cseq object
normalized_method	character. method of normalization (default=powerlawFittedRPM)

Author(s)

S. Thongjuea

See Also

[contrRPM](#), [expRPM](#), [calculateBatchRPM](#)

Examples

#See the vignette

`contrCoverage`

This method has been removed.

Description

This method has been removed.

`contrInteractionRegions`

get interaction regions from the control

Description

get all identified interaction regions from the control

Usage

`contrInteractionRegions(object)`

Arguments

`object` r3Cseq or r3CseqInBatch object

Value

The candidate interaction regions show in the IRange object

Author(s)

S. Thongjuea

See Also

[expInteractionRegions](#), [getInteractions](#)

Examples

#See the vignette

contrRawData*Accessors for the 'contrRawData' slot of a r3Cseq object.*

Description

The 'contrRawData' slot of hold the raw aligned reads data in the GRanges object.

Usage

```
## S4 method for signature 'r3Cseq'  
contrRawData(object)  
## S4 replacement method for signature 'r3Cseq'  
contrRawData(object) <- value
```

Arguments

object	r3Cseq object
value	a GRanges object of aligned reads

Author(s)

S. Thongjuea

See Also

[expRawData](#)

Examples

```
#See the vignette
```

contrReadCount*get read count per region for the control*

Description

get the read count per region for the control

Usage

```
contrReadCount(object)
```

Arguments

object	r3Cseq object
--------	---------------

Author(s)

S. Thongjuea

See Also

[expReadCount](#), [getReadCountPerRestrictionFragment](#)

Examples

#See the vignette

contrRPM

get read per million (RPM) for the control

Description

get the normalized 3C-seq data (RPM) for the control

Usage

contrRPM(object)

Arguments

object r3Cseq or r3CseqInBatch object

Author(s)

S. Thongjuea

See Also

[calculateRPM](#), [expRPM](#)

Examples

#See the vignette

enzymeDb

Rebase The Restriction Enzyme Database

Description

The database includes all restriction enzyme information from the REBASE database.

References

<http://rebase.neb.com/rebase/rebase.html>

expCoverage	<i>This method has been removed.</i>
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Description

This method has been removed.

expInteractionRegions	<i>get interaction regions from the experiment</i>
-----------------------	--

Description

get identified interaction regions from the experiment

Usage

`expInteractionRegions(object)`

Arguments

object r3Cseq or r3CseqInBatch object

Value

The candidate interaction regions show in the IRange object

Author(s)

S. Thongjuea

See Also

[getInteractions](#), [contrInteractionRegions](#)

Examples

#See the vignette

`export3Cseq2bedGraph` *export interaction regions to the 'bedGraph' format*

Description

export interaction regions from RagedData to the bedGraph format, which suitable for uploading to the UCSC genome browser

Usage

```
export3Cseq2bedGraph(object,datatype=c("rpm","read_count"))
```

Arguments

object	r3Cseq object, The object might contain the interaction regions generated by function getInteractions
datatype	read_count : read count per restriction fragment rpm : normalized read per million per restriction fragment

Value

The text file in 'bedGraph' format

Author(s)

S. Thongjuea

See Also

[exportInteractions2text](#)

Examples

```
#See the vignette
```

`export3CseqRawReads2bedGraph`

export the interaction signal from the raw reads to the 'bedGraph' format

Description

export interaction regions signal to the bedGraph format, which suitable for uploading to the UCSC genome browser

Usage

```
export3CseqRawReads2bedGraph(object)
```

Arguments

object r3Cseq object

Value

The text file in 'bedGraph' format

Author(s)

S. Thongjuea

See Also

[exportInteractions2text](#), [export3Cseq2bedGraph](#),

Examples

#See the vignette

exportBatchInteractions2text

export identified interaction regions to the tab separated format for replicates analysis

Description

export interaction regions from RagedData to the tab separated format for replicates analysis

Usage

`exportBatchInteractions2text(object)`

Arguments

object r3CseqInBatch object

Value

The text file in the tab separated format

Author(s)

S. Thongjuea

See Also

[export3Cseq2bedGraph](#), [exportInteractions2text](#)

Examples

#See the vignette

`exportInteractions2text`

export identified interaction regions to the tab separated format

Description

export interaction regions from RagedData to the tab separated format

Usage

`exportInteractions2text(object)`

Arguments

`object` r3Cseq object

Value

The text file in the tab separated format

Author(s)

S. Thongjuea

See Also

[export3Cseq2bedGraph](#)

Examples

#See the vignette

`expRawData`

Accessors for the 'expRawData' slot of a r3Cseq object.

Description

The 'expRawData' slot of hold the raw aligned reads data in the GRanges object.

Usage

```
## S4 method for signature 'r3Cseq'
expRawData(object)
## S4 replacement method for signature 'r3Cseq'
expRawData(object) <- value
```

Arguments

<code>object</code>	r3Cseq object
<code>value</code>	a GRanges object of aligned reads

Author(s)

S. Thongjuea

See Also

[expRawData](#)

Examples

#See the vignette

`expReadCount`

get read count per region for the experiment

Description

get the read count per region for the experiment

Usage

`expReadCount(object)`

Arguments

`object` r3Cseq

Author(s)

S. Thongjuea

See Also

[contrReadCount](#), [getReadCountPerRestrictionFragment](#)

Examples

#See the vignette

expRPM	<i>get read per million (RPM) for the experiment</i>
--------	--

Description

get the normalized 3C-seq data (RPM) for the experiment

Usage

```
expRPM(object)
```

Arguments

object	r3Cseq or r3CseqInBatch
--------	-------------------------

Author(s)

S. Thongjuea

See Also

[calculateRPM](#), [contrRPM](#)

Examples

```
#See the vignette
```

generate3CseqReport	<i>generate reports for analysis results from r3Cseq</i>
---------------------	--

Description

generate reports for analysis results from r3Cseq, the report contains all plots in one pdf file and a text separated out put file.

Usage

```
generate3CseqReport(obj)
```

Arguments

obj	r3Cseq or r3CseqInBatch object
-----	--------------------------------

Value

The text file in the tab separated format and the pdf file of all plots

Author(s)

S. Thongjuea

See Also

[exportInteractions2text](#) [plotOverviewInteractions](#), [plotInteractionsPerChromosome](#), [plotInteractionNearViewpoint](#)

Examples

```
#See the vignette
```

getBatchInteractions *calculate z-score, assign p-value and q-value for each interaction region for replicates data sets*

Description

Calculate z-score, assign p-value and q-value to each interaction regions for replicates data sets

Usage

```
getBatchInteractions(object,method=c("union","intersection"),smoothing.parameter=0.1,fdr=0.05)
```

Arguments

object	r3Cseq object
method	character. The method for combining biological replicates for 3C-Seq analysis (default = "union")
smoothing.parameter	A level at which cubic smoothing spline for the spar (see vsmooth.spline) input parameter. Must be in (0.06,0.4] (default=0.1)
fdr	A level at which to control the FDR. Must be in (0,1] (default=0.05)

Value

The interaction regions show in the RangedData

Author(s)

S. Thongjuea

See Also

[getInteractions](#) [vsmooth.spline](#)

Examples

```
#See the vignette
```

`getBatchRawReads` *Get aligned reads from the replicates BAM files*

Description

Reading in the input BAM files from the 3C-Seq replicates analysis and then save files as the local GRanged object .rData files

Usage

```
getBatchRawReads(object)
```

Arguments

object	r3CseqInBatch object
--------	----------------------

Value

The GRangedData represents the aligned reads from the BAM file

Author(s)

S. Thongjuea

See Also

[getRawReads](#),

Examples

```
#See the vignette
```

`getBatchReadCountPerRestrictionFragment`
count reads for replicates analysis

Description

Counts the number of reads from 3C-Seq data per each restriction fragment for replicates analysis

Usage

```
getBatchReadCountPerRestrictionFragment(object, getReadsMethod = c("wholeReads", "adjacentFragment",  

  nFragmentExcludedReadsNearViewpoint=2))
```

Arguments

object	r3CseqInBatch object
getReadsMethod	character. To count all reads found in the particular restriction fragment uses wholeReads option. To count reads found around the edge of restriction fragment both 5'utr and 3'utr uses adjacentFragmentEndsReads option (default=wholeReads)
nFragmentExcludedReadsNearViewpoint	Numeric. The number of excluded fragments around the viewpoint, reads found in these fragments will be removed from the analysis (default=2)

Value

The RangedData represents the number of reads per each restriction fragment

Author(s)

S. Thongjuea

See Also

[getReadCountPerWindow](#), [getReadCountPerRestrictionFragment](#)

Examples

```
#See the vignette
```

`getBatchReadCountPerWindow`

count reads per window size for replicates analysis

Description

Counts the number of reads from 3C-Seq data per each window size for replicates analysis

Usage

```
getBatchReadCountPerWindow(object,WindowSize=5e3,nFragmentExcludedReadsNearViewpoint=2,mode=c("n
```

Arguments

object	r3CseqInBatch object
windowSize	Numeric. non-overlapping window size for counting reads (default=5e3)
nFragmentExcludedReadsNearViewpoint	Numeric. The number of excluded fragments around the viewpoint, reads found in these fragments will be removed from the analysis (default=2)
mode	character. The window-based modes analysis (default="non-overlapping")

Value

The RangedData represents the number of reads per each window size

Author(s)

S. Thongjuea

See Also

[getReadCountPerRestrictionFragment](#), [getBatchReadCountPerRestrictionFragment](#), [getReadCountPerWindow](#),

Examples

```
#See the vignette
```

`getContrInteractionsInRefseq`

identified significant interaction regions for RefSeq genes

Description

Get a list of genes that contain strong interaction signals in the control

Usage

```
getContrInteractionsInRefseq(obj, cutoff.qvalue=0.05, expanded_upstream=50e3, expanded_downstream=10e3)
```

Arguments

`obj` obj is r3Cseq or r3CseqInBatch object
`cutoff.qvalue` Numeric. The cutoff q-value (default=0.05)
`expanded_upstream` Numeric. The expanded distance from the upstream of a gene start (default=50e3)
`expanded_downstream` Numeric. The expanded distance from the downstream of a gene end (default =10e3)

Value

List of identified genes, which contain strong interaction signals

Author(s)

S. Thongjuea

See Also

[getContrInteractionsInRefseq](#)

Examples

```
# See the vignette
```

getCoverage	<i>This method has been removed.</i>
-------------	--------------------------------------

Description

This method has been removed.

getExpInteractionsInRefseq	<i>identified significant interaction regions for RefSeq genes</i>
----------------------------	--

Description

Get a list of genes that contain strong interaction signals in the experiment

Usage

```
getExpInteractionsInRefseq(obj, cutoff.qvalue=0.05, expanded_upstream=50e3, expanded_downstream=10e3)
```

Arguments

obj	obj is r3Cseq or r3CseqInBatch object
cutoff.qvalue	Numeric. The cutoff q-value (default=0.05)
expanded_upstream	Numeric. The expanded distance from the upstream of a gene start (default=50e3)
expanded_downstream	Numeric. The expanded distance from the downstream of a gene end (default =10e3)

Value

List of identified genes, which contain strong interaction signals

Author(s)

S. Thongjuea

See Also

[getContrInteractionsInRefseq](#)

Examples

```
# See the vignette
```

<code>getInteractions</code>	<i>calculate z-score, assign p-value and q-value for each interaction region</i>
------------------------------	--

Description

Calculate z-score, assign p-value and q-value to each interaction regions

Usage

```
getInteractions(object,smoothing.parameter=0.1,fdr=0.05)
```

Arguments

<code>object</code>	r3Cseq object
<code>smoothing.parameter</code>	A level at which cubic smoothing spline for the spar (see <code>vsmooth.spline</code>) input parameter. Must be in (0.06,0.4] (default=0.1)
<code>fdr</code>	A level at which to control the FDR. Must be in (0,1] (default=0.05)

Value

The interaction regions show in the RangedData

Author(s)

S. Thongjuea

See Also

[getBatchInteractions](#) [vsmooth.spline](#)

Examples

```
#See the vignette
```

<code>getRawReads</code>	<i>Get aligned reads from the BAM file</i>
--------------------------	--

Description

Reading in the input BAM file and then store it in the GRanged object

Usage

```
getRawReads(object)
```

Arguments

<code>object</code>	r3Cseq object
---------------------	---------------

Value

The GRangedData represents the aligned reads from the BAM file

Author(s)

S. Thongjuea

See Also

[getBatchRawReads](#),

Examples

```
#See the vignette
```

getReadCountPerRestrictionFragment
count reads per restriction fragment

Description

Counts the number of reads from 3C-Seq data per each restriction fragment

Usage

```
getReadCountPerRestrictionFragment(object, getReadsMethod = c("wholeReads", "adjacentFragmentEndsReads"),  
nFragmentExcludedReadsNearViewpoint=2)
```

Arguments

object r3Cseq object
getReadsMethod character. To count all reads found in the particular restriction fragment uses wholeReads option. To count reads found around the edge of restriction fragment both 5'utr and 3'utr uses adjacentFragmentEndsReads option (default=wholeReads)
nFragmentExcludedReadsNearViewpoint Numeric. The number of excluded fragments around the viewpoint, reads found in these fragments will be removed from the analysis (default=2)

Value

The RangedData represents the number of reads per each restriction fragment

Author(s)

S. Thongjuea

See Also

[getReadCountPerWindow](#), [getBatchReadCountPerRestrictionFragment](#)

Examples

```
#See the vignette
```

`getReadCountPerWindow` *count reads per window size*

Description

Counts the number of reads from 3C-Seq data per each window size

Usage

```
getReadCountPerWindow(object,WindowSize=5e3,nFragmentExcludedReadsNearViewpoint=2,mode=c("non-overlapping","overlapping"))
```

Arguments

<code>object</code>	r3Cseq object
<code>WindowSize</code>	Numeric. non-overlapping window size for counting reads (default=5e3)
<code>nFragmentExcludedReadsNearViewpoint</code>	Numeric. The number of excluded fragments around the viewpoint, reads found in these fragments will be removed from the analysis (default=2)
<code>mode</code>	character. The window-based modes analysis (default="non-overlapping")

Value

The RangedData represents the number of reads per each window size

Author(s)

S. Thongjuea

See Also

[getReadCountPerRestrictionFragment](#),

Examples

```
#See the vignette
```

`getViewpoint` *get the viewpoint of 3C-seq data*

Description

The viewpoint is the bait of 3C method, which can be a promoter region of an interested gene, an enhancer, and a transcription factor binding region.

Usage

```
getViewpoint(obj)
```

Arguments

obj r3Cseq or r3CseqInBatch object

Value

The viewpoint shows in the IRanges

Author(s)

S. Thongjuea

Examples

#See the vignette

hg18refGene

hg18's refGenes

Description

The human (hg18) reference genes from UCSC

hg19refGene

hg19's refGenes

Description

The human (hg19) reference genes from UCSC

mm10refGene

mm10's refGenes

Description

The mouse (mm10) reference genes from UCSC

mm9refGene

mm9's refGenes

Description

The mouse (mm9) reference genes from UCSC

Myb_prom_FB

Myb_prom_FB a data set for the example of r3Cseq analysis**Description**

The example aligned reads generated by 3C-Seq protocol from fetal brain. The promoter region of the Myb's gene was selected as the viewpoint. This data was transformed from aligned reads shown in the BAM file to GRanged object by using Rsamtools.

Myb_prom_FL

Myb_prom_FL a data set for the example of r3Cseq analysis**Description**

The example aligned reads generated by 3C-Seq protocol from fetal liver. The promoter region of the Myb's gene was selected as the viewpoint. This data was transformed from aligned reads shown in the BAM file to GRanged object by using Rsamtools.

plot3Cecdf

*This method has been removed.***Description**

This method has been removed.

plotDomainogramNearViewpoint

*Plot domainogram of interaction regions near the viewpoint***Description**

Plot domainogram of interaction regions near the viewpoint

Usage

```
plotDomainogramNearViewpoint(object,smoothing.parameter=0.1,distance=5e5,maximum_window=25e3,view="experiment")
```

Arguments

- | | |
|---------------------|--|
| object | r3Cseq or r3CseqInBatch object |
| smoothing.parameter | A level at which cubic smoothing spline for the spar (see vsmooth.spline) input parameter. Must be in (0.06,0.4] (default=0.1) |
| distance | Numeric. The distance relative to the viewpoint (default=5e5) |
| maximum_window | Numeric. The maximum windowing (default=25e3). We normally compute the interaction regions per window starting from 2Kb to maximum window (default=25kb) to make the interaction matrix for visualizing the domainogram. |
| view | character. The selected view of data (default="experiment") |

Value

Plots of domainogram for interaction regions close to the viewpoint

Author(s)

S. Thongjuea

See Also

[plotOverviewInteractions](#), [plotInteractionsPerChromosome](#), [plotInteractionsNearViewpoint](#)

Examples

```
# See the vignette
```

```
plotInteractionsNearViewpoint
```

Plot identified interaction regions near the viewpoint

Description

Plot identified interaction regions near the viewpoint

Usage

```
plotInteractionsNearViewpoint(obj, distance=5e5, log2fc_cutoff=1, yLim=0)
```

Arguments

obj	obj is r3Cseq or r3CseqInBatch object
distance	Numeric. The distance relative to the viewpoint (default=5e5)
log2fc_cutoff	Numeric. The log2 cutoff ratio between the experiment and control (default=1)
yLim	Numeric. The limited height of y-axis (default=0)

Value

Plots of identified interaction regions close to the viewpoint

Author(s)

S. Thongjuea

See Also

[plotOverviewInteractions](#), [plotInteractionsPerChromosome](#), [plotDomainogramNearViewpoint](#)

Examples

```
# See the vignette
```

plotInteractionsPerChromosome

Plot interaction regions per each chromosome of interest

Description

Plot the distribution of interaction regions per each chromosome

Usage

```
plotInteractionsPerChromosome(obj, chromosomeName)
```

Arguments

- obj obj is r3Cseq or r3CseqInBatch object.
- chromosomeName Character. The input chromosome name (e.g. "chr1")

Value

Plots of interaction regions per chromosome.

Author(s)

S. Thongjuea

See Also

[plotInteractionsNearViewpoint](#), [plotOverviewInteractions](#), [plotDomainogramNearViewpoint](#)

Examples

```
# See the vignette
```

plotOverviewInteractions

Plot overview of identified interaction regions for genome-wide

Description

Plot the distribution of identified interaction regions across genome

Usage

```
plotOverviewInteractions(obj, cutoff.qvalue=0.05)
```

Arguments

- obj obj is r3Cseq or r3CseqInBatch object
- cutoff.qvalue Numeric. The cutoff q-value (default=0.05)

Value

Plots of identified 3C-Seq interaction regions genome-wide

Author(s)

S. Thongjuea

See Also

[plotInteractionsNearViewpoint](#), [plotInteractionsPerChromosome](#), [plotDomainogramNearViewpoint](#)

Examples

```
# See the vignette
```

r3Cseq-class

r3Cseq objects

Description

The r3Cseq class is the extended class from r3CseqCommon class. It is a general container for storing and manipulating a set of input parameters, RangeData of interactions regions from r3Cseq analysis , and the raw reads GRanged data of the genome-wide interaction signal generated by next-generation sequencing.

Extends

Class r3CseqCommon, directly.

Slots

organismName Object of class "character" the version of particular assembly genome from UCSC (e.g. mm9, hg18, hg19) . The package supports three genome assemblies consisting of mouse (mm9), and human (hg18, hg19).

restrictionEnzyme Object of class "character" this is the primary restriction enzyme name using in 3C-Seq experiment

viewpoint_chromosome Object of class "character" chromosome name of where is the viewpoint located eg. chr10, chrX etc.

viewpoint_primer_forward Object of class "character" the forward primer DNA sequences for the viewpoint amplification

viewpoint_primer_reverse Object of class "character" the reverse primer DNA sequences for the viewpoint amplification

expReadCount Object of class "RangedData" the read count in experiment

contrReadCount Object of class "RangedData" the read count in control

expRPM Object of class "RangedData" the normalized read read per million in experiment

contrRPM Object of class "RangedData" the normalized read read per million in control

expInteractionRegions Object of class "RangedData" the identified interaction regions in experiment

`contrInteractionRegions` Object of class "RangedData" the identified interaction regions in control
`isControlInvolved` Object of class "logical" the logical to ask whether the control is involved in the analysis or not
`alignedReadsBamExpFile` Object of class "character" the file name of experiment in BAM format
`alignedReadsBamContrFile` Object of class "character" the file name of control in BAM format
`expLabel` Object of class "character" the experiment name
`contrLabel` Object of class "character" the control name
`expLibrarySize` Object of class "integer" the library size of experiment
`contrLibrarySize` Object of class "integer" the library size of control
`expReadLength` Object of class "integer" the read length of experiment
`contrReadLength` Object of class "integer" the read length of experiment
`expRawData` Object of class "GRanges" the raw reads found in experiment
`contrRawData` Object of class "GRanges" the raw reads found in control

Author(s)

S. Thongjuea

See Also

[r3CseqCommon](#), [r3CseqInBatch](#)

Examples

See the vignette

r3CseqCommon-class r3CseqCommon objects

Description

The `r3CseqCommon` class is a general container for storing and manipulating a set of input parameters, `RangeData` of interactions regions from `r3Cseq` analysis. It is a root class for `r3Cseq` and `r3CseqInBatch` classes.

Slots

`organismName` Object of class "character" the version of particular assembly genome from UCSC (e.g. mm9, hg18, hg19). The package supports three genome assemblies consisting of mouse (mm9), and human (hg18, hg19).
`restrictionEnzyme` Object of class "character" this is the primary restriction enzyme name using in 3C-Seq experiment
`viewpoint_chromosome` Object of class "character" chromosome name of where is the viewpoint located eg. chr10, chrX etc.

```

viewpoint_primer_forward Object of class "character" the forward primer DNA sequences
for the viewpoint amplification

viewpoint_primer_reverse Object of class "character" the reverse primer DNA sequences for
the viewpoint amplification

expReadCount Object of class "RangedData" the read count in experiment

contrReadCount Object of class "RangedData" the read count in control

expRPM Object of class "RangedData" the normalized read read per million in experiment

contrRPM Object of class "RangedData" the normalized read read per million in control

expInteractionRegions Object of class "RangedData" the identified interaction regions in ex-
periment

contrInteractionRegions Object of class "RangedData" the identified interaction regions in
control

isControlInvolved Object of class "logical" the logical to ask whether the control is involved
in the analysis or not

```

Author(s)

S. Thongjuea

See Also

[r3Cseq](#), [r3CseqInBatch](#)

Examples

```
# See the vignette
```

r3CseqInBatch-class *r3CseqInBatch objects*

Description

The r3CseqInBatch class is the extended class from r3CseqCommon class. It is a general container for storing and manipulating a set of input parameters, RangeData of interactions regions from r3Cseq analysis for replicates data sets.

Extends

Class r3CseqCommon, directly.

Slots

```

organismName Object of class "character" the version of particular assembly genome from UCSC
(e.g. mm9, hg18, hg19) . The package supports three genome assemblies consisting of mouse
(mm9), and human (hg18, hg19).

restrictionEnzyme Object of class "character" this is the primary restriction enzyme name
using in 3C-Seq experiment

viewpoint_chromosome Object of class "character" chromosome name of where is the view-
point located eg. chr10, chrX etc.

```

`viewpoint_primer_forward` Object of class "character" the forward primer DNA sequences for the viewpoint amplification
`viewpoint_primer_reverse` Object of class "character" the reverse primer DNA sequences for the viewpoint amplification
`expReadCount` Object of class "RangedData" the read count in experiment
`contrReadCount` Object of class "RangedData" the read count in control
`expRPM` Object of class "RangedData" the normalized read read per million in experiment
`contrRPM` Object of class "RangedData" the normalized read read per million in control
`expInteractionRegions` Object of class "RangedData" the identified interaction regions in experiment
`contrInteractionRegions` Object of class "RangedData" the identified interaction regions in control
`isControlInvolved` Object of class "logical" the logical to ask whether the control is involved in the analysis or not
`bamFilesDirectory` Object of class "character" the path name of directory that contains BAM files
`BamExpFiles` Object of class "vector" the file names of BAM files in the experiment
`BamContrFiles` Object of class "vector" the file names of BAM files in the control
`expBatchLabel` Object of class "vector" the labeled experiment names
`contrBatchLabel` Object of class "vector" the labeled control names
`readCountTable` Object of class "RangedData" the read count table
`RPMsTable` Object of class "RangedData" the normalized read per million table
`expBatchLibrarySize` Object of class "vector" the library size of each experiment
`contrBatchLibrarySize` Object of class "vector" the library size of each control
`expBatchReadLength` Object of class "vector" the read length of experiments
`contrBatchReadLength` Object of class "vector" the read length of controls

Author(s)

S. Thongjuea

See Also

[r3CseqCommon](#), [r3CseqInBatch](#)

Examples

```
# See the vignette
```

Description

The rat (rn5) reference genes from UCSC

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