

Package ‘ChIPQC’

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Type Package

Title Quality metrics for ChIPseq data

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Maintainer

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Description Quality metrics for ChIPseq data.

biocViews Sequencing, ChIPSeq, QualityControl, ReportWriting

License GPL (>= 3)

LazyLoad yes

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Suggests BiocStyle

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Description

ChIPQC analyzes aligned reads (in .bam format) for ChIP-seq samples and their associated controls, computing a variety of quality control metrics and statistics, and providing reporting and plotting functions to enable assessment of experimental data for further analysis.

Details

Package: ChIPQC
Type: Package
Version: 0.1
Date: 2014-03-01
License: GPL3

ChIPQC primarily uses two object classes: [ChIPQCsample](#), which encapsulates the information about individual samples, and [ChIPQCexperiment](#), which encapsulates information about larger ChIP-seq experiments (consisting of a number of samples). The primary entry point is the constructor function [ChIPQC](#), which takes a description of an entire experiment, constructs objects for all the samples, and computes the quality metrics.

Author(s)

Tom Carroll and Rory Stark

Maintainers: Tom Carroll <tc.infomatics@gmail.com> and Rory Stark <rory.stark@cruk.cam.ac.uk>

References

Frontiers?

See Also

ChIPQC is designed to work closely with the [DiffBind](#) package, which provides functionality for analyzing ChIP-seq experiments, including performing differential binding analysis to identify significantly differentially bound peaks.

averagepeaksignal-methods

Retrieve average peak profiles

Description

Retrieve the average peak profile for a sample or set of samples.

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a matrix of the average peak profiles for all of the samples in a ChIP-seq experiment. Each column represents a sample, and each row a base pair position, centered on peak summits.

`signature(object = "list")` Retrieve a matrix of the average peak profiles for all of the samples in a list of ChIPQC samples. Each column represents a sample, and each row a base pair position, centered on peak summits.

`signature(object = "ChIPQCsample")` Retrieve a vector representing the average peak profile for a sample. Each column represents a basepair position, centered on the peak summits.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
CTCFprofile = averagepeaksignal(QCsample(exampleExp,1))
length(CTCFprofile)
plot(CTCFprofile,type='l',ylab="mean pileup")

allprofiles = averagepeaksignal(exampleExp)
dim(allprofiles)
for(i in 1:ncol(allprofiles)) lines(allprofiles[,i],col=i)
```

ChIPQC *Construct a [ChIPQCexperiment](#) object*

Description

Constructs a new [ChIPQCexperiment](#) object.

Usage

```
ChIPQC(experiment, annotation, chromosomes, samples,
        consensus=FALSE, bCount=FALSE, mapQCth=15, blacklist=NULL,
        profileWin=400, fragmentLength=125, shifts=1:300, ...)
```

Arguments

`experiment` A specification of the ChIP-seq experiment to evaluate. This can either be a dataframe, a filename for a .csv file, or a [DBA-object](#) object as defined in the [DiffBind](#) package. Columns names in sample sheet may include:

- `SampleID`: Identifier string for sample
- `Tissue`: Identifier string for tissue type
- `Factor`: Identifier string for factor
- `Condition`: Identifier string for condition
- `Treatment`: Identifier string for treatment
- `Replicate`: Replicate number of sample
- `bamReads`: file path for bam file containing aligned reads for ChIP sample
- `bamControl`: file path for bam file containing aligned reads for control sample
- `ControlID`: Identifier string for control sample
- `Peaks`: path for file containing peaks for sample. Format determined by `PeakCaller` field or caller parameter
- `PeakCaller`: Identifier string for peak caller used. If `Peaks` is not a bed file, this will determine how the `Peaks` file is parsed. If missing, will use default peak caller specified in caller parameter. Possible values:
 - “raw”: text file file; peak score is in fourth column
 - “bed”: .bed file; peak score is in fifth column
 - “narrow”: default peak.format: narrowPeaks file
 - “macs”: MACS .xls file
 - “swembl”: SWEMBL .peaks file
 - “bayes”: bayesPeak file
 - “fp4”: FindPeaks v4
- `PeakFormat`: string indicating format for peak files; see `PeakCaller` and [dba.peakset](#)
- `ScoreCol`: column in peak files that contains peak scores

| | |
|-------------|---|
| | <ul style="list-style-type: none"> • LowerBetter: logical indicating that lower scores signify better peaks <p>See the documentation for the <code>sampleSheet</code> parameter of <code>dba</code> for details.</p> |
| annotation | <p>Either a character string indicating the genome and version to use as a genomic annotation, or a previously defined annotation (obtained using <code>QCannotation</code> on a previously defined <code>ChIPQCexperiment</code> object.) May be left unspecified, in which case no genomic feature analysis is performed. The following annotation specifiers are supported:</p> <ul style="list-style-type: none"> • “hg19”: Human, version 19 • “hg18”: Human, version 18 • “mm10”: Mouse, version 10 • “mm9” : Mouse, version 19 • “rn4” : Rat, version 4 • “ce6” : C. Elgans, version 6 • “dm3” : D. Melanogaster, version 3 <p>Alternatively, you can construct your own annotation; see the package vignette for more information.</p> |
| chromosomes | <p>Specification of which chromosomes to use for computing QC statistics. If missing, the first chromosome which has a peak is checked. If NULL, all chromosomes will be checked (which may be time-consuming). This can be a character string (e.g. “chr18”) or a vector or list of character strings. If it is an integer or vector of integers, the chromosomes will be checked based on the order that they are listed in a peak set.</p> |
| samples | <p>list of <code>ChIPCSample</code> objects. If present, the sample objects will be taken directly from this list instead of being computed using the <code>ChIPQCsample</code> constructor.</p> |
| consensus | <p>If consensus is a <code>GRanges</code> object, all samples will use this peakset when computing peak-based metrics. If <code>consensus=TRUE</code>, a consensus peakset will be generated and used for all samples, derived by merging overlapping peaks in all provided peaksets, keeping any peaks that overlap in at least two samples. To avoid this behavior, set <code>consensus=FALSE</code>; this will result in only supplied peaksets being used for calculation of peak-based metrics (and no peak-based metric being computed for samples with no peakset specified, such as controls).</p> |
| bCount | <p>if TRUE, the peak scores for all samples will be based on read counts using <code>dba.count</code> using a consensus peakset. If consensus is missing, any samples (such as controls) that are not already associated with a peakset will be associated with the consensus peakset (if consensus is not missing, all samples will be associated with the consensus peakset). Note that the re-counting process may be time-consuming.</p> |
| mapQcth | <p>An integer representing a mapping quality score threshold. Only reads with mapping quality scores above this threshold will be used for some statistics.</p> |
| blacklist | <p>A <code>GRanges</code> object or filename specifying a bed file containing genomic regions that should be excluded from the analysis. If missing and the annotation is “hg19”, a default blacklist, <code>blacklist_hg19</code> derived from the UCSC list, will be used. No blacklist is used if this is set to NULL, or is left missing and the annotation is not “hg19”.</p> |

| | |
|-----------------------------|---|
| <code>profileWin</code> | An integer indicating the width, in base pairs, of the window to be used for peak profiles. Peaks will be centered on their summits, and include half the window size upstream and half downstream of this point. |
| <code>fragmentLength</code> | An integer indicating the expected fragment length of the libraries. Optional, as this value will be computed for each library. |
| <code>shifts</code> | A vector of values to try when computing optimal shift sizes. |
| <code>...</code> | additional parameters passed to <code>dba.count</code> if <code>bCount=TRUE</code> . |

Details

ChIPQC first constructs a new [DBA-object](#) object if one is not provided. Next it computes the annotation if one is not provided. The main loop constructs new [ChIPQCsample](#) objects for each sample (and unique control sample).

Value

A [ChIPQCexperiment](#) object.

Note

ChIPQC uses [bplapply](#) from the [BiocParallel](#) package to build the [ChIPQCsample](#) object in parallel, if supported. Control of the parallelization can be effected using [BiocParallel](#) functions, such as [register](#).

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#), [DiffBind](#)

Examples

```
## Not run: exampleExp = ChIPQC(samples,annotation="hg19")
data(example_QCexperiment)
exampleExp

## Not run: tamoxifen = ChIPQC(samples, ,annotation="hg19", consensus=TRUE, bCounts=T)
data(tamoxifen_QC)
tamoxifen
```

| | |
|-------------|---|
| ChIPQC-data | <i>Example data sets for ChIPQC package, each containing a ChIPQCexperiment object, as well as a pre-compiled blacklist for hg19.</i> |
|-------------|---|

Description

The `tamoxifen_QC` example data set contains a [ChIPQCexperiment](#) object named `tamoxifen`. This data set, based on Ross-Innes et al (2012), includes 11 ER ChIP libraries, and their input controls, divided into tamoxifen responsive and tamoxifen resistant sample groups. Only data for chromosome 18 (`chr18`) are included.

The `example_QCexperiment` data set contains a [ChIPQCexperiment](#) object named `exampleExp`. This data set, derived from ENCODE data, includes 6 ChIP libraries. Only data for chromosome 22 (`chr22`) are included.

`blacklist_hg19` data set includes a [GRanges](#) object named `blacklist_hg19` containing blacklisted regions for the human genome, derived from the UCSC blacklist.

Usage

```
data(tamoxifen_QC)
data(example_QCexperiment)
data(blacklist_hg19)
```

Format

`tamoxifen_QC`: A single [ChIPQCexperiment](#) object named `tamoxifen` is loaded. This object is used for the [ChIPQC-package](#) examples and vignette. This object can also be used with the [DiffBind](#) package (see related data objects [tamoxifen](#)).

`example_QCexperiment`: A single [ChIPQCexperiment](#) object named `exampleExp` is loaded. This object is used for the [ChIPQC-package](#) examples and vignette.

`blacklist_hg19`: A single [GRanges](#) object named `blacklist.hg19` that is used by default when processing hg19 data sets.

Source

Ross-Innes, C. S., Stark, R., Teschendorff, A. E., Holmes, K. A., Ali, H. R., Dunning, M. J., Brown, G. D., Gojis, O., Ellis, I. O., Green, A. R., Ali, S., Chin, S.-F., Palmieri, C., Caldas, C., and Carroll, J. S. (2012). Differential oestrogen receptor binding is associated with clinical outcome in breast cancer. *Nature* 481, 389-393.

Examples

```
data(tamoxifen_QC)
tamoxifen
plotRegi(tamoxifen)
```



```
data(example_QCexperiment)
exampleExp
```

```
data(blacklist_hg19)
blacklist.hg19
```

ChIPQCExperiment-class

ChIPQCExperiment instances

Description

The ChIPQCExperiment class is built around a list of [ChIPQCsample](#) objects, each representing a ChIP or control sample in a ChIP-seq experiment. These objects are created using the [ChIPQC](#) function.

Slots

.Data: Object of class "list": internal

Samples: Object of class "list": List of [ChIPQCsample](#) objects.

DBA: Object of class "DBA": [DBA-object](#) object (from package [DiffBind](#))

annotation: Object of class "list" : annotation data

Extends

Class "[list](#)".

Methods

QCmetadata signature(object = "ChIPQCExperiment"): see [QCmetadata](#).

QCmetrics signature(object = "ChIPQCExperiment"): see [QCmetrics](#).

QCsample signature(object = "ChIPQCExperiment"): see [QCsample](#).

QCcontrol signature(object = "ChIPQCExperiment"): see [QCcontrol](#).

QCannotation signature(object = "ChIPQCExperiment"): see [QCannotation](#).

QCdba signature(object = "ChIPQCExperiment"): see [QCdba](#).

averagepeaksignal signature(object = "ChIPQCExperiment"): see [averagepeaksignal](#).

coveragehistogram signature(object = "ChIPQCExperiment"): see [coveragehistogram](#).

crosscoverage signature(object = "ChIPQCExperiment"): see [crosscoverage](#).

flagtagcounts signature(object = "ChIPQCExperiment"): see [flagtagcounts](#).

fragmentlength signature(object = "ChIPQCExperiment"): see [fragmentlength](#).

FragmentLengthCrossCoverage signature(object = "ChIPQCExperiment"): see [FragmentLengthCrossCoverage](#).

frip signature(object = "ChIPQCExperiment"): see [frip](#).

mapped signature(object = "ChIPQCExperiment"): see [mapped](#).

reads signature(object = "ChIPQCexperiment"): see [reads](#).

duplicates signature(object = "ChIPQCexperiment"): see [duplicates](#).

duplicateRate signature(object = "ChIPQCexperiment"): see [duplicateRate](#).

Normalisedaveragepeaksignal signature(object = "ChIPQCexperiment"): see [Normalisedaveragepeaksignal](#).

peaks signature(object = "ChIPQCexperiment"): see [peaks](#).

readlength signature(object = "ChIPQCexperiment"): see [readlength](#).

ReadLengthCrossCoverage signature(object = "ChIPQCexperiment"): see [ReadLengthCrossCoverage](#).

RelativeCrossCoverage signature(object = "ChIPQCexperiment"): see [RelativeCrossCoverage](#).

rib1 signature(object = "ChIPQCexperiment"): see [rib1](#).

rip signature(object = "ChIPQCexperiment"): see [rip](#).

show signature(object = "ChIPQCexperiment"): see [show](#).

ssd signature(object = "ChIPQCexperiment"): see [ssd](#).

regi signature(object = "ChIPQCexperiment"): see [regi](#).

plotCC signature(object = "ChIPQCexperiment"): see [plotCC](#).

plotCoverageHist signature(object = "ChIPQCexperiment"): see [plotCoverageHist](#).

plotFrib1 signature(object = "ChIPQCexperiment"): see [plotFrib1](#).

plotPeakProfile signature(object = "ChIPQCexperiment"): see [plotPeakProfile](#).

plotRap signature(object = "ChIPQCexperiment"): see [plotRap](#).

plotRegi signature(object = "ChIPQCexperiment"): see [plotRegi](#).

plotCorHeatmap signature(object = "ChIPQCexperiment"): see [plotCorHeatmap](#).

plotPrincomp signature(object = "ChIPQCexperiment"): see [plotPrincomp](#).

ChIPQCreport signature(object = "ChIPQCexperiment"): see [ChIPQCreport](#).

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCsample](#), [DiffBind](#)

Examples

```
## Not run: exampleExp = ChIPQC(samples)
data(example_QCexperiment)
exampleExp

## Not run: tamoxifen = ChIPQC(samples, consensus=TRUE, bCounts=T)
data(tamoxifen_QC)
tamoxifen
```

ChIPQCreport-methods *Generate a summary QC report*

Description

Generate an HTML summary QC report.

Methods

`signature(experiment = "ChIPQCexperiment", facet=TRUE, reportName="ChIPQC", reportFolder="ChIPQCreport")`
 Generates an summary QC report for the experiment in HTML format.

| | |
|---------------------------|---|
| <code>experiment</code> | <code>ChIPQCexperiment</code> object |
| <code>facet</code> | logical indicating whether or not to facet using experimental metadata. |
| <code>reportName</code> | filename of main report file (.html). |
| <code>reportFolder</code> | directory name where plot graphics will be saved |
| <code>facetBy</code> | metadata fields to use for faceting |
| <code>colourBy</code> | metadata field to color by |

`signature(experiment = "list", facet=TRUE, reportName="ChIPQC", reportFolder="ChIPQCreport", facetBy=c())`
 Generates an summary QC report for a list of ChIPQCsample objects in HTML format.

| | |
|---------------------------|---|
| <code>experiment</code> | <code>list</code> object |
| <code>facet</code> | logical indicating whether or not to facet using experimental metadata. |
| <code>reportName</code> | filename of main report file (.html). |
| <code>reportFolder</code> | directory name where plot graphics will be saved |
| <code>facetBy</code> | metadata fields to use for faceting |
| <code>colourBy</code> | metadata field to color by |

`signature(sample = "ChIPQCsample", reportName="ChIPQC", reportFolder="ChIPQCreport",)`
 Generate a summary QC report for a sample in HTML format.

| | |
|---------------------------|--|
| <code>sample</code> | <code>ChIPQCsample</code> object |
| <code>reportName</code> | filename of main report file (.html). |
| <code>reportFolder</code> | directory name where plot graphics will be saved |

Note

ChIPQCreport uses Nozzle.R2 for generating HTML.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
ChIPQCreport(exampleExp, facetBy=c("Tissue", "Factor"))
#report in ChIPQCreport/Example.html

data(tamoxifen_QC)
ChIPQCreport(tamoxifen, facetBy="Tissue", colourBy="Condition")
#report in ChIPQCreport/ChIPQC.html

ChIPQCreport(tamoxifen, facetBy=c("Tissue", "Condition"))
#report in ChIPQCreport/ChIPQC.html
```

ChIPQCsample-class *Class "ChIPQCsample"*

Description

Object containing quality metrics computed for a ChIP-seq (or associated control) sample.

Objects from the Class

Objects can be created using the [ChIPQCsample](#) function.

Constructor Function

`ChIPQCsample(reads, peaks, annotation, chromosomes = NULL, mapQCth = 15, blacklist, profileWin = 400, fragmentLength = 125, shifts = 1:300, runCrossCor = FALSE, verboseT=FALSE)`

- `readscharacter` string filename of .bam file
- `peaksGRanges` object or character string filename of peaks. If present, peak-based metrics will be computed.
- `annotation` Either a character string indicating the genome and version to use as a genomic annotation, or a previously defined annotation (obtained using [QCannotation](#) on a previously defined [ChIPQCexperiment](#) object.) May be left unspecified, in which case no genomic feature analysis is performed. The following annotation specifiers are supported:

| | |
|--------|----------------------|
| "hg20" | Human, version 20 |
| "hg19" | Human, version 19 |
| "hg18" | Human, version 18 |
| "mm10" | Mouse, version 10 |
| "mm9" | Mouse, version 19 |
| "rn4" | Rat, version 4 |
| "ce6" | C. Elgans, version 6 |

"dm3" D. Melanogaster, version 3

Alternatively, you can construct your own annotation; see the package Vignette for more information.

- **chromosomes** Specification of which chromosomes to use for computing QC statistics. If missing, the first chromosome which has a peak is checked. If NULL, all chromosomes will be checked (which may be time-consuming). This can be a character string (e.g. "chr18") or a vector or list of character strings. If it is an integer or vector of integers, the chromosomes will be checked based on the order that they are listed in a peak set.
- **mapQCth** An integer representing a mapping quality score threshold. Only reads with mapping quality scores above this threshold will be used for some statistics.
- **blacklist** A [GRanges](#) object or filename specifying a bed file containing genomic regions that should be excluded from the analysis. If missing and the annotation is "hg19", a default blacklist, [blacklist_hg19](#) derived from the UCSC list, will be used. No blacklist is used if this is set to NULL, or is left missing and the annotation is not "hg19".
- **profileWin** An integer indicating the width, in base pairs, of the window to be used for peak profiles. Peaks will be centered on their summits, and include half the window size upstream and half downstream of this point.
- **fragmentLength** An integer indicating the expected fragment length of the libraries. Optional, as this value will be computed.
- **shifts** A vector of values to try when computing optimal shift sizes.
- **runCrossCor** Compute cross-correlation in addition to cross-coverage. This will take more compute time, and is currently not used in the final report.
- **verboseT** TRUE or FALSE, specifying whether to report progress. Default is TRUE. When set to FALSE ChIPQC does not report any progress until complete.

Slots

AveragePeakSignal: Object of class "list"
 CrossCoverage: Object of class "numeric"
 CrossCorrelation: Object of class "numeric"
 SSD: Object of class "numeric"
 SSDBL: Object of class "numeric"
 CountsInPeaks: Object of class "numeric"
 CountsInBlackList: Object of class "numeric"
 CountsInFeatures: Object of class "list"
 PropInFeatures: Object of class "list"
 CoverageHistogram: Object of class "numeric"
 FlagAndTagCounts: Object of class "numeric"
 readlength: Object of class "numeric"
 seqnames: Object of class "Rle"

ranges: Object of class "IRanges"
strand: Object of class "Rle"
elementMetadata: Object of class "DataFrame"
seqinfo: Object of class "Seqinfo"
metadata: Object of class "list"

Extends

Class "[GRanges](#)"

Methods

averagepeaksignal signature(object = "ChIPQCsample"): see [averagepeaksignal](#).
coveragehistogram signature(object = "ChIPQCsample"): see [coveragehistogram](#).
crosscoverage signature(object = "ChIPQCsample"): see [crosscoverage](#).
flagtagcounts signature(object = "ChIPQCsample"): see [flagtagcounts](#).
fragmentlength signature(object = "ChIPQCsample"): see [fragmentlength](#).
FragmentLengthCrossCoverage signature(object = "ChIPQCsample"): see [FragmentLengthCrossCoverage](#).
frip signature(object = "ChIPQCsample"): see [frip](#).
mapped signature(object = "ChIPQCsample"): see [mapped](#).
reads signature(object = "ChIPQCsample"): see [reads](#).
duplicates signature(object = "ChIPQCsample"): see [duplicates](#).
duplicateRate signature(object = "ChIPQCsample"): see [duplicateRate](#).
Normalisedaveragepeaksignal signature(object = "ChIPQCsample"): see [Normalisedaveragepeaksignal](#).
peaks signature(object = "ChIPQCsample"):see [peaks](#).
readlength signature(object = "ChIPQCsample"): see [readlength](#).
ReadLengthCrossCoverage signature(object = "ChIPQCsample"): see [ReadLengthCrossCoverage](#).
RelativeCrossCoverage signature(object = "ChIPQCsample"):see [RelativeCrossCoverage](#).
ribl signature(object = "ChIPQCsample"): see [ribl](#).
rip signature(object = "ChIPQCsample"): see [rip](#).
show signature(object = "ChIPQCsample"): see [show](#).
ssd signature(object = "ChIPQCsample"): see [ssd](#).
regi signature(object = "ChIPQCsample"): see [regi](#).
plotCC signature(object = "ChIPQCsample"): see [plotCC](#).
plotCoverageHist signature(object = "ChIPQCsample"): see [plotCoverageHist](#).
plotFribl signature(object = "ChIPQCsample"): see [plotFribl](#).
plotPeakProfile signature(object = "ChIPQCsample"): see [plotPeakProfile](#).
plotRap signature(object = "ChIPQCsample"): see [plotRap](#).
plotRegi signature(object = "ChIPQCsample"): see [plotRegi](#).

Author(s)

Thomas Carroll and Rory Stark

References

Carroll TS, Liang Z, Salama R, Stark R and Santiago Id (in press). Impact of artefact removal on ChIP quality metrics in ChIP-seq and ChIP-exo data. *Frontiers in Genetics*.

See Also

[ChIPQC-package](#), [ChIPQCsample](#)

Examples

```
bamFile <- system.file("extdata", "ex1.bam",  
                      package="Rsamtools")  
ex1 <- ChIPQCsample(bamFile,annotation=NULL)  
readlength(ex1)  
fragmentlength(ex1)
```

coveragehistogram-methods

Retrieve histogram data representing densities of coverage pileups

Description

Retrieve histogram data representing densities of coverage pileups.

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a matrix of coverage histogram data for all samples in a ChIP-seq experiment. Each column represents a sample, and each row a pileup height, with the value representing the number of basepair positions that report this pileup height.

`signature(object = "list")` Retrieve a matrix of coverage histogram data for all ChIPQCsamples in a list. Each column represents a sample, and each row a pileup height, with the value representing the number of basepair positions that report this pileup height.

`signature(object = "ChIPQCsample")` Retrieve a vector representing coverage histogram data for a sample. Values represent the number of base pairs positions that report the pileup value. The value in position 1 of the vector the contains the number of examined basepair positions that are overlapped by exactly zero reads, while position 2 shows the number of basepair positions overlapped by exactly one read, etc.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
CTCFcoverage = coveragehistogram(QCsample(exampleExp,1))
length(CTCFcoverage)
plot(log10(CTCFcoverage),type='l',ylab="log10 Density",
      xlab="Pileup")

allcoverages = coveragehistogram(exampleExp)
dim(allcoverages)
for(i in 1:ncol(allcoverages)) lines(log10(allcoverages[,i]),col=i)
```

crosscoverage-methods *Retrieve the cross coverage values for a range of shift sizes*

Description

Retrieves the cross-coverage values for a range of shift sizes.

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a matrix of cross-coverage data for all samples in an ChIP-seq experiment. Each column represents a sample, and each row a shift size, with the value representing the cross-coverage using that size read.

`signature(object = "list")` Retrieve a matrix of cross-coverage data for all samples in a list of ChIPQCsample objects. Each column represents a sample, and each row a shift size, with the value representing the cross-coverage using that size read.

`signature(object = "ChIPQCsample")` Retrieve a vector of cross-coverage data for a sample. Each position in the vector corresponds to a shift size, with the value representing the cross-coverage using that size read.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
CTCFcoverage = crosscoverage(QCsample(exampleExp,1))
length(CTCFcoverage)
plot(CTCFcoverage,type='l',
     ylab="Cross-coverage",
     xlab="Fragment length")

allcoverages = crosscoverage(exampleExp)
dim(allcoverages)
for(i in 1:ncol(allcoverages)) lines(allcoverages[,i],col=i)
```

duplicateRate-methods *Retrieve duplication rates*

Description

Retrieve duplication rates.

Methods

`signature(object = "ChIPQCexperiment", bFiltered)` Retrieve a vector of the duplication rates for each sample in an experiment. A read is considered duplicated if another read maps to the same genomic location; the duplication rate is the number of duplicated reads divided by the total number of reads for a sample. If `bFiltered=TRUE` (or is missing), only reads that pass the mapping quality filter for each sample are included. if `bFiltered=FALSE`, all reads for each sample will be included.

`signature(object = "list", bFiltered)` Retrieve a vector of the duplication rates for each sample in a list of `ChIPQCsample` objects. A read is considered duplicated if another read maps to the same genomic location; the duplication rate is the number of duplicated reads divided by the total number of reads for a sample. If `bFiltered=TRUE` (or is missing), only reads that pass the mapping quality filter for each sample are included. if `bFiltered=FALSE`, all reads for each sample will be included.

`signature(object = "ChIPQCsample", bFiltered)` Retrieve the duplication rate for a sample. A read is considered duplicated if another read maps to the same genomic location; the duplication rate is the number of duplicated reads divided by the total number of reads for the sample. If `bFiltered=TRUE` (or is missing), only reads that pass the mapping quality filter for the sample are included. if `bFiltered=FALSE`, all reads for the sample will be included.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
duplicateRate(exampleExp)
duplicateRate(QCsample(exampleExp,1))
```

duplicates-methods *Retrieve numbers of duplicate reads.*

Description

Retrieve the numbers of duplicate reads.

Methods

signature(object = "ChIPQCexperiment", bFiltered) Retrieve a vector of the numbers of duplicate reads for each sample in an experiment. A read is considered duplicated if another read maps to the same genomic location. If bFiltered=TRUE (or is missing), this will be the number of duplicates that pass the mapping quality filter for each sample. if bFiltered=FALSE, it will be the total number of duplicates for each sample.

signature(object = "list", bFiltered) Retrieve a vector of the numbers of duplicate reads for each sample in a list of ChIPQCsample objects. A read is considered duplicated if another read maps to the same genomic location. If bFiltered=TRUE (or is missing), this will be the number of duplicates that pass the mapping quality filter for each sample. if bFiltered=FALSE, it will be the total number of duplicates for each sample.

signature(object = "ChIPQCsample", bFiltered) Retrieve the number of duplicates for a sample. A read is considered duplicated if another read maps to the same genomic location. If bFiltered=TRUE (or is missing), this will be the number of duplicates that pass the mapping quality filter. if bFiltered=FALSE, it will be the total number of duplicates for the sample.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
duplicates(exampleExp)
duplicates(QCsample(exampleExp,1))
```

flagtagcounts-methods *Retrieve numbers of reads that pass various filters*

Description

Retrieve numbers of reads that pass various filters

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a matrix of counts passing various filters for all the samples in an experiment. Each column represents the counts in a sample.

`signature(object = "list")` Retrieve a matrix of counts passing various filters for all the samples in a list of ChIPQCsample objects. Each column represents the counts in a sample.

`signature(object = "ChIPQCsample")` Retrieve a vector of counts passing various filters. The values are:

| | |
|----------------|---|
| UnMapped | Number of reads that are not mapped (aligned) |
| Mapped | Number of reads that are mapped (aligned) |
| Duplicates | Number of reads that align to exactly the same place as another read |
| MapQPass | Number of reads with a mapping quality score greater than or equal to the specified threshold |
| MapQPassandDup | Number of reads that are mapped (aligned) and not duplicates |

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
flagtagcounts(exampleExp)
flagtagcounts(QCsample(exampleExp,1))
```

fragmentlength-methods

Retrieve the estimated fragment length

Description

Retrieve the estimated fragment length.

Methods

signature(object = "ChIPQCexperiment") Retrieve a vector of estimated fragments sizes, one for each sample in the experiment.

signature(object = "list") Retrieve a vector of estimated fragments sizes, one for each sample in a list of ChIPQCsample objects.

signature(object = "ChIPQCsample", width) Retrieve the estimated fragment length for a sample. If width is missing, the readlength derived from the bam file is used as the read length.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
fragmentlength(exampleExp)
fragmentlength(QCsample(exampleExp,1))
```

frip-methods

Retrieve fraction of reads in peaks

Description

Retrieve the fraction of reads in peaks

Methods

signature(object = "ChIPQCexperiment") Retrieve a vector of values representing the proportion of reads that overlap peaks for each sample in an experiment.

signature(object = "list") Retrieve a vector of values representing the proportion of reads that overlap peaks for each sample in a list of ChIPQCsample objects.

signature(object = "ChIPQCsample") Retrieve a value representing the proportion of reads that overlap the peaks for a sample.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
frip(exampleExp)
frip(QCsample(exampleExp,1))
```

mapped-methods

Retrieve numbers of mapped reads

Description

Retrieve the numbers of mapped reads.

Methods

signature(object = "ChIPQCexperiment") Retrieve a vector of the numbers of mapped (aligned) reads for each sample in an experiment.

signature(object = "list") Retrieve a vector of the numbers of mapped (aligned) reads for each sample in a list of ChIPQCsample objects.

signature(object = "ChIPQCsample") Retrieve the number of mapped (aligned) reads in a sample.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
mapped(exampleExp)
mapped(QCsample(exampleExp,1))
```

Normalisedaveragepeaksignal-methods

Retrieve normalised average peak profiles

Description

Retrieve normalised average peak profiles

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a matrix of normalised average peak signal data for all samples in a ChIP-seq experiment. Each column represents a sample, and each row a base pair position, centered on peak summits.

`signature(object = "list")` Retrieve a matrix of normalised average peak signal data for all samples in a list of ChIPQCsample objects. Each column represents a sample, and each row a base pair position, centered on peak summits.

`signature(object = "ChIPQCsample")` Retrieve a vector representing the normalised average peak profile for a sample. Each column represents a basepair position, centered on the peak summits.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
CTCFprofile = Normalisedaveragepeaksignal(QCsample(exampleExp,1))
length(CTCFprofile)
plot(CTCFprofile,type='l',ylab="normalised mean pileup")

allprofiles = Normalisedaveragepeaksignal(exampleExp)
dim(allprofiles)
for(i in 1:ncol(allprofiles)) lines(allprofiles[,i],col=i)
```

| | |
|---------------|-----------------------|
| peaks-methods | <i>Retrieve peaks</i> |
|---------------|-----------------------|

Description

Retrieve peaks.

Methods

signature(object = "ChIPQCexperiment") Retrieve a [GRangesList](#) of the peaks associated with all the samples in an experiment.

signature(object = "list") Retrieve a [GRangesList](#) of the peaks associated with all the samples in a list of ChIPQCsample objects.

signature(object = "ChIPQCsample") Retrieve a [GRanges](#) object containing the peaks associated with a sample.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
expPeaks = peaks(exampleExp)
length(expPeaks)
peaks(QCsample(exampleExp,1))
```

| | |
|----------------|--------------------------------------|
| plotCC-methods | <i>Generate Cross-Coverage plots</i> |
|----------------|--------------------------------------|

Description

Generate Cross-Coverage plots.

Methods

signature(object = "ChIPQCexperiment", method) Generate cross-coverage plots for all the samples in an experiment.

signature(object = "list", method) Generate cross-coverage plots for list of samples in an experiment.

signature(object = "ChIPQCsample", methods) Generate cross-coverage plots for a sample. Supported methods include:

"Coverage" [default] Coverage plot

Note

plotCC uses ggplot2 for plotting, and returns a ggplot2 plot dataframe.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
plotCC(exampleExp)
plotCC(exampleExp,excludedBox=TRUE)
plotCC(QCsample(exampleExp,1))
plotCC(QCsample(exampleExp)[1:4])
```

plotCorHeatmap-methods

Generate Correlation Heatmap for ChIP samples

Description

Generates correlation heatmap for ChIP samples.

Methods

signature(object = "ChIPQCexperiment", attributes, ...) Generate correlation heatmap, including clustering dendrogram, for all the samples in an experiment that are associated with a peakset.

attributes character string, or vector of character strings, representing metadata field names, for use in labeling
 ... additional parameters passed to [dba.plotHeatmap](#)

Note

plotCorHeatmap uses [dba.plotHeatmap](#) for plotting.

Author(s)

Rory Stark and Thomas Carroll

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [dba.plotHeatmap](#)

Examples

```
data(tamoxifen_QC)
plotCorHeatmap(tamoxifen, attributes=c("Tissue", "Condition", "Replicate"))
```

plotCoverageHist-methods

Generate coverage histogram plot

Description

Generate coverage histogram plot.

Methods

signature(object = "ChIPQCexperiment") Generate coverage histogram plots for all the samples in an experiment.

signature(object = "list") Generate coverage histogram plots for all the samples in a list of ChIPQCsamples.

signature(object = "ChIPQCsample") Generate coverage histogram plots for a sample.

Note

Uses ggplot2 for plotting, and returns a ggplot2 plot dataframe.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
plotCoverageHist(exampleExp, facetBy=c("Tissue", "Factor"))
plotCoverageHist(QCsample(exampleExp, 1))
```

plotFribl-methods *Generate fraction of reads in blacklist plot*

Description

Generate fraction of reads in blacklist plot.

Methods

signature(object = "ChIPQCexperiment", type="barstacked", facet=T, facetBy=c("Tissue", "Factor"), AsPercent=TRUE)
Generate fraction of reads in blacklist plots for all the samples in an experiment.

signature(object = "list", type="barstacked", facet=T, facetBy=c("Sample"), AsPercent=TRUE)
Generate fraction of reads in blacklist plots for all the samples in a list of ChIPQCsample objects..

signature(object = "ChIPQCsample", type="barstacked", AsPercent=TRUE) Generate fraction of reads in blacklist plots for a sample.

Note

plotFribl uses ggplot2 for plotting, and returns a ggplot2 plot dataframe.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
plotFribl(exampleExp)
plotFribl(QCsample(exampleExp, 1))
```

plotFrip-methods *Generate fraction of reads in peaks plot*

Description

Generate fraction of reads in peaks plot.

Methods

signature(object = "ChIPQCexperiment", type="barstacked", facet=T, facetBy=c("Tissue", "Factor"), AsPercent=TRUE)
Generate fraction of reads in peaks plots for all the samples in an experiment.

signature(object = "list", type="barstacked", facet=T, facetBy=c("Sample"), AsPercent=TRUE)
Generate fraction of reads in peaks plots for all the samples in a list of ChIPQCsample objects.

signature(object = "ChIPQCsample", type="barstacked", facet=T, facetBy=c("Tissue", "Factor"), AsPercent=TRUE)
Generate fraction of reads in peaks plots for a sample.

Note

plotFrip uses ggplot2 for plotting, and returns a ggplot2 plot dataframe.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
plotFrip(exampleExp)
plotFrip(QCsample(exampleExp, 1))
```

plotPeakProfile-methods *Generate peak profile plots*

Description

Generate peak profile plots.

Methods

signature(object = "ChIPQCexperiment", method) Generate peak profile plots for all the samples in an experiment.

signature(object = "list", method) Generate peak profile plots for all the samples in a list of ChIPQCsample objects..

signature(object = "ChIPQCsample", method) Generate peak profile plots for a sample.

Note

plotPeakProfile uses ggplot2 for plotting, and returns a ggplot2 plot dataframe.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
plotCoverageHist(exampleExp, facetBy=c("Tissue", "Factor"))
plotCoverageHist(QCsample(exampleExp, 1))
data(tamoxifen_QC)
plotCoverageHist(tamoxifen, facetBy=c("Tissue", "Condition"))
```

plotPrincomp-methods *Generate Principal Components Analysis plot for ChIP samples*

Description

Generate principal components analysis plot for ChIP samples.

Methods

signature(object = "ChIPQCexperiment", attributes, ...) Generate principal components analysis plot, for all the samples in an experiment that are associated with a peakset.

attributes character string, or vector of character strings, representing metadata field names, for use grouping samples by color
 ... additional parameters passed to [dba.plotPCA](#)

Note

plotPrincomp uses [dba.plotPCA](#) for plotting.

Author(s)

Rory Stark and Thomas Carroll

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [dba.plotPCA](#)

Examples

```
data(tamoxifen_QC)
plotPrincomp(tamoxifen,attributes=c("Condition"))
```

| | |
|-----------------|-------------------------------------|
| plotRap-methods | <i>Generate reads in peaks plot</i> |
|-----------------|-------------------------------------|

Description

Generate reads in peaks plot.

Methods

```
signature(object = "ChIPQCexperiment", type="barstacked", facet=T, facetBy=c("Tissue", "Factor"))
  Generate reads in peaks plots for all the samples in an experiment.
```

```
signature(object = "list", type="barstacked", facet=T, facetBy=c("Sample")) Generate
  reads in peaks plots for all the samples in a list of ChIPQCsample objects.
```

```
signature(object = "ChIPQCexperiment", type="barstacked", facet=T, facetBy=c("Tissue", "Factor"))
  Generate reads in peaks plots for all the samples in an experiment.
```

```
signature(object = "ChIPQCsample", type="barstacked", facet=T, facetBy=c("Tissue", "Factor"))
  Generate reads in peaks plots for a sample.
```

Note

plotRap uses ggplot2 for plotting, and returns a ggplot2 plot dataframe.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
plotRap(exampleExp, facetBy=c("Tissue", "Factor"))
plotRap(QCsample(exampleExp, 1))
```

plotRegi-methods *Generate relative enrichment of genomic features plot*

Description

Generate relative enrichment of genomic features plot.

Methods

signature(object = "ChIPQCexperiment", facet=T, facetBy=c("Tissue", "Factor")) Generate relative enrichment of genomic features plots for all the samples in an experiment.

signature(object = "list", facet=T, facetBy=c("Sample")) Generate relative enrichment of genomic features plots for all the samples in a list of ChIPQCsample objects.

signature(object = "ChIPQCsample") Generate relative enrichment of genomic features plots for a sample.

Note

plotRegi uses ggplot2 for plotting, and returns a ggplot2 plot dataframe.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
plotRegi(exampleExp, facetBy=c("Tissue", "Factor"))
plotRegi(QCsample(exampleExp, 1))
data(tamoxifen_QC)
plotRegi(tamoxifen, facetBy=c("Tissue", "Condition"))
```

plotSSD-methods *Generate SSD plot*

Description

Generate SSD metric plot. If blacklists supplied, will generate SSD prior and post blacklisting

Methods

`signature(object = "ChIPQCexperiment", facet=T, facetBy=c("Tissue", "Factor"))` Generate SSD metric plot for all samples in experiment. If blacklists supplied, will generate SSD prior and post blacklisting

`signature(object = "list", facet=T, facetBy=c("Tissue", "Factor"))` Generate SSD metric plot for list of samples in experiment. If blacklists supplied, will generate SSD prior and post blacklisting

`signature(object = "ChIPQCsample")` Generate SSD metric plot for single sample. If blacklists supplied, will generate SSD prior and post blacklisting

Note

`plotSSD` uses `ggplot2` for plotting, and returns a `ggplot2` plot `gg` object.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
plotSSD(exampleExp, facetBy=c("Tissue", "Factor"))
plotSSD(QCsample(exampleExp, 1))
plotSSD(QCsample(exampleExp)[1:3])
data(tamoxifen_QC)
plotSSD(tamoxifen, facetBy=c("Tissue", "Condition"))
```

QCAnnotation-methods *Retrieve an annotation description, or a processed annotation*

Description

Retrieve an annotation description, or a processed annotation, from a [ChIPQCexperiment](#) object.

Methods

`signature(object = "ChIPQCexperiment", bRetrieve=FALSE)` Retrieve the annotation. If `bRetrieve=FALSE` (default), the character string describing the annotation is returned (currently only "hg19" is supported). If `bRetrieve=TRUE`, a processed annotation is returned (in the form of a list). This can be used in subsequent calls to [ChIPQC](#) and/or [ChIPQCsample](#) for efficiency purposes.

Author(s)

Rory Stark and Thomas Carroll

See Also

[ChIPQC-package](#), [ChIPQCsample](#), [ChIPQCexperiment](#)

Examples

```
data(example_QCexperiment)
QCannotation(exampleExp)
```

QCcontrol-methods

Retrieve control objects associated with a sample

Description

Retrieve the [ChIPQCsample](#) objects representing controls, or a specific [ChIPQCsample](#) representing the control for a specific sample, from a [ChIPQCexperiment](#) object.

Methods:

`signature(object = "ChIPQCexperiment", sampleID)` Get the control sample (as a [ChIPQCsample](#) object) associated with a ChIP sample, or, if `sampleID` is missing, a list of all samples used as controls. `sampleID` is a character string or an integer.

Author(s)

Rory Stark and Thomas Carroll

See Also

[ChIPQC-package](#), [ChIPQCsample](#), [ChIPQCexperiment](#)

Examples

```
data(tamoxifen_QC)
controls = QCcontrol(tamoxifen)
length(controls)
names(controls)
controls[[1]]
bt474control = QCcontrol(tamoxifen,"BT4741")
bt474control
```

QCdba-methods

Retrieve the [DBA-object](#) object associated with an experiment

Description

Retrieve the [DBA-object](#) object associated with a [ChIPQCexperiment](#) object.

Methods:

`signature(object = "ChIPQCexperiment")` Retrieves the [DBA-object](#) object associated with a [ChIPQCexperiment](#). This object can be used with [DiffBind](#) functions to further analyse a ChIP-seq experiment, including performing a differential binding analysis.

Author(s)

Rory Stark and Thomas Carroll

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [DiffBind](#), [dba](#)

Examples

```
data(tamoxifen_QC)
tamoxifenDBA = QCdba(tamoxifen)
## library(DiffBind)
## tamoxifenDBA
```

QCmetadata-methods

Retrieve metadata associated with an experiment

Description

Retrieve metadata for a [ChIPQCexperiment](#) object.

Methods:

`signature(object = "ChIPQCexperiment")` Retrieve a data frame containing metadata for all the samples in a ChIP-seq experiment represented by a [ChIPQCexperiment](#) object.

`signature(object = "list")` Retrieve a data frame containing metadata for all the samples in a list of [ChIPQCsample](#) objects.

Author(s)

Rory Stark and Thomas Carroll

See Also

[ChIPQC-package](#), [ChIPQCsample](#), [ChIPQCexperiment](#)

Examples

```
data(tamoxifen_QC)
meta = QCmetadata(tamoxifen)
meta
```

| | |
|-------------------|--|
| QCmetrics-methods | <i>Retrieve consolidated set of QC metrics</i> |
|-------------------|--|

Description

Retrieves a consolidated set of QC metrics.

Methods

`signature(object = "ChIPQCexperiment")` Retrieves a matrix of QC metrics for all the samples in an experiment, with a column of values for each sample.

`signature(object = "list")` Retrieves a matrix of QC metrics for all the samples in a list of ChIPQCsample objects, with a column of values for each sample.

`signature(object = "ChIPQCsample")` Retrieves a vector of QC metrics for a sample.

Author(s)

Rory Stark and Thomas Carroll

See Also

[ChIPQC-package](#), [ChIPQCsample](#), [ChIPQCexperiment](#)

Examples

```
data(example_QCexperiment)
QCmetrics(exampleExp)

data(tamoxifen_QC)
QCmetrics(tamoxifen)

QCmetrics(QCsample(tamoxifen,1))
```

QCsample-methods *Retrieve sample objects associated with an experiment*

Description

Retrieves a list of [ChIPQCsample](#) objects, or one specific [ChIPQCsample](#) object, from a [ChIPQCexperiment](#) object.

Methods

signature(object = "ChIPQCexperiment", sampleID) If sampleID is missing or equal to 0, the full list of [ChIPQCsample](#) objects is returned. If sampleID is an integer n, the [ChIPQCsample](#) object corresponding to the nth sample is returned. If sampleID is a character string, the [ChIPQCsample](#) object corresponding to the sample with that ID is returned.

Author(s)

Rory Stark and Thomas Carroll

See Also

[ChIPQC-package](#), [ChIPQCsample](#), [ChIPQCexperiment](#)

Examples

```
data(example_QCexperiment)
samples = QCsample(exampleExp)
length(samples)
names(samples)
samples$CTCF_1
```

readlength-methods *Retrieve read length values*

Description

Retrieve read length values.

Methods

signature(object = "ChIPQCexperiment") Retrieve a vector of read length values, one for each sample in an experiment.

signature(object = "list") Retrieve a vector of read length values, one for each sample in a list of [ChIPQCsample](#) objects.

signature(object = "ChIPQCsample") Retrieve the read length value for a sample.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
readlength(exampleExp)
readlength(QCsample(exampleExp,1))
```

ReadLengthFragmentLengthCrossCoverage-methods

Retrieve the cross coverage values when extending reads to the optimal fragment length.

Description

Retrieve the cross coverage values when extending reads to the optimal fragment length.

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a vector of cross-coverage values for all samples in a ChIP-seq experiment, when all reads are shifted by the optimal fragment length (the maximum cross-coverage value).

`signature(object = "list")` Retrieve a vector of cross-coverage values for all samples in a list of ChIPQCsample objects, when all reads are shifted by the optimal fragment length (the maximum cross-coverage value).

`signature(object = "ChIPQCsample")` Retrieve the cross-coverage value for a sample, when all reads are shifted by the optimal fragment length (the maximum cross-coverage value).

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
FragmentLengthCrossCoverage(exampleExp)
FragmentLengthCrossCoverage(QCsample(exampleExp,1))
```

ReadLengthReadLengthCrossCoverage-methods

Retrieve the cross coverage values without extending reads

Description

Retrieve the cross coverage values without extending reads.

Methods

signature(object = "ChIPQCexperiment") Retrieve a vector of cross-coverage values for all samples in a ChIP-seq experiment, with no shift.

signature(object = "list") Retrieve a vector of cross-coverage values for all samples in a list of ChIPQCsample objects, with no shift.

signature(object = "ChIPQCsample") Retrieve the cross-coverage value for a sample, with no shift.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
ReadLengthCrossCoverage(exampleExp)
ReadLengthCrossCoverage(QCsample(exampleExp,1))
```

reads-methods

Retrieve numbers of reads

Description

Retrieve the numbers of reads.

Methods

`signature(object = "ChIPQCexperiment", bFiltered)` Retrieve a vector of the numbers of reads for each sample in an experiment. If `bFiltered=TRUE` (or is missing), this will be the number of reads that pass the mapping quality filter for each sample. if `bFiltered=FALSE`, it will be the total number of reads for each sample.

`signature(object = "list", bFiltered)` Retrieve a vector of the numbers of reads for each sample in a list of `ChIPQCsample` objects. If `bFiltered=TRUE` (or is missing), this will be the number of reads that pass the mapping quality filter for each sample. if `bFiltered=FALSE`, it will be the total number of reads for each sample.

`signature(object = "ChIPQCsample", bFiltered)` Retrieve the number of reads for a sample. If `bFiltered=TRUE` (or is missing), this will be the number of reads that pass the mapping quality filter. if `bFiltered=FALSE`, it will be the total number of reads for the sample.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
reads(exampleExp)
reads(QCsample(exampleExp,1))
```

regi-methods

Retrieve genomic profile information

Description

Retrieve genomic profile information in terms of relative enrichment over background genomic distribution.

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a matrix of relative enrichment values for a variety of genomic features. Each column represents the enrichment values for one sample in the experiment.

`signature(object = "list")` Retrieve a matrix of relative enrichment values for a variety of genomic features. Each column represents the enrichment values for one sample in a list of `ChIPQCsample` objects.

`signature(object = "ChIPQCsample")` Retrieve a vector of relative enrichment values for a variety of genomic features for a sample. Relative enrichment is computed as the proportion of reads overlapping a genomic feature type compared to the overall proportion of base pairs in the genome comprising those features. Genomic features include:

| | |
|----------------------|--|
| 3UTRs | 3' UTRs |
| 5UTRs | 5' UTRs |
| Introns | Intronic (non-coding) portions of gene bodies |
| Transcripts | Transcribed regions, including exons |
| Promoters500 | 500bp regions immediately upstream of annotated TSSs |
| Promoters2000to500 | 2500bp regions from 2000bp immediately upstream of annotated TSSs to 500bp downstream |
| Promoters20000to2000 | 22000bp regions from 20000bp immediately upstream of annotated TSSs to 2000bp downstream |

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
genomicprofile = regi(exampleExp)
heatmap(genomicprofile)
regi(QCsample(exampleExp,1))
```

RelativeCrossCoverage-methods

Retrieve the relative cross coverage values for a range of shift sizes

Description

Retrieve the relative cross-coverage values for a range of shift sizes

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a vector of relative cross-coverage values for all samples in a ChIP-seq experiment, computed based on the maximal value (when extending the reads to the optimal fragment length) versus the cross-coverage values using non-extended reads.

`signature(object = "list")` Retrieve a vector of relative cross-coverage values for all samples in a list of ChIPQCsample objects, computed based on the maximal value (when extending the reads to the optimal fragment length) versus the cross-coverage values using non-extended reads.

`signature(object = "ChIPQCsample")` Retrieve the relative cross-coverage value for a sample, computed based on the maximal value (when extending the reads to the optimal fragment length) versus the cross-coverage values using non-extended reads.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
RelativeCrossCoverage(exampleExp)
RelativeCrossCoverage(QCsample(exampleExp,1))
```

ribl-methods

Retrieve numbers of reads overlapping blacklisted regions

Description

Retrieve the numbers of reads overlapping blacklisted regions.

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a vector of the numbers of reads overlapping blacklisted regions for each sample in an experiment.

`signature(object = "list")` Retrieve a vector of the numbers of reads overlapping blacklisted regions for each sample in a list of `ChIPQCsample` objects.

`signature(object = "ChIPQCsample")` Retrieve the number of reads overlapping blacklisted regions in a sample.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
ribl(exampleExp)
ribl(QCsample(exampleExp,1))
```

`rip-methods`*Retrieve numbers of reads overlapping peaks*

Description

Retrieve the numbers of reads overlapping peaks.

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a vector of the numbers of reads overlapping peaks for each sample in an experiment.

`signature(object = "list")` Retrieve a vector of the numbers of reads overlapping peaks for each sample in a list of `ChIPQCsample` objects.

`signature(object = "ChIPQCsample")` Retrieve the number of reads overlapping peaks in a sample.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
rip(exampleExp)
rip(QCsample(exampleExp,1))
```

`ssd-methods`*Retrieve SSD (squared sum of deviations) values of peak coverage*

Description

Retrieve SSD (squared sum of deviations) values of peak coverage density.

Methods

`signature(object = "ChIPQCexperiment")` Retrieve a vector of SSD values, one for each sample in an experiment.

`signature(object = "list")` Retrieve a vector of SSD values, one for each sample in a list of `ChIPQCsample` objects.

`signature(object = "ChIPQCsample")` Retrieve the SSD (squared sum of deviations) for a sample, computed from the standard deviation of the [coveragehistogram](#).

Note

uses the SSD calculation from the chipseq package.

Author(s)

Thomas Carroll and Rory Stark

See Also

[ChIPQC-package](#), [ChIPQCexperiment](#), [ChIPQCsample](#)

Examples

```
data(example_QCexperiment)
ssd(exampleExp)
ssd(QCsample(exampleExp,1))
```

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