

# Package ‘QDNAseq’

July 10, 2025

**Version** 1.44.0

**Title** Quantitative DNA Sequencing for Chromosomal Aberrations

**Depends** R (>= 3.1.0)

**Imports** graphics, methods, stats, utils, Biobase (>= 2.18.0), CGHbase (>= 1.18.0), CGHcall (>= 2.18.0), DNACopy (>= 1.32.0), GenomicRanges (>= 1.20), IRanges (>= 2.2), matrixStats (>= 0.60.0), R.utils (>= 2.9.0), Rsamtools (>= 1.20), future.apply (>= 1.8.1)

**Suggests** BiocStyle (>= 1.8.0), BSgenome (>= 1.38.0), digest (>= 0.6.20), GenomeInfoDb (>= 1.6.0), future (>= 1.22.1), parallelly (>= 1.28.1), R.cache (>= 0.13.0), QDNAseq.hg19, QDNAseq.mm10

**Description** Quantitative DNA sequencing for chromosomal aberrations. The genome is divided into non-overlapping fixed-sized bins, number of sequence reads in each counted, adjusted with a simultaneous two-dimensional loess correction for sequence mappability and GC content, and filtered to remove spurious regions in the genome. Downstream steps of segmentation and calling are also implemented via packages DNACopy and CGHcall, respectively.

**biocViews** CopyNumberVariation, DNASEq, Genetics, GenomeAnnotation, Preprocessing, QualityControl, Sequencing

**License** GPL

**URL** <https://github.com/ccagc/QDNAseq>

**BugReports** <https://github.com/ccagc/QDNAseq/issues>

**RoxygenNote** 7.3.2

**git\_url** <https://git.bioconductor.org/packages/QDNAseq>

**git\_branch** RELEASE\_3\_21

**git\_last\_commit** 9c070eb

**git\_last\_commit\_date** 2025-04-15

**Repository** Bioconductor 3.21

**Date/Publication** 2025-07-09

**Author** Ilari Scheinin [aut],  
 Daoud Sie [aut, cre],  
 Henrik Bengtsson [aut],  
 Erik van Dijk [ctb]

**Maintainer** Daoud Sie <d.sie@vumc.nl>

## Contents

QDNaseq-package . . . . .	2
addPhenodata . . . . .	3
applyFilters . . . . .	4
binReadCounts . . . . .	5
callBins . . . . .	7
compareToReference . . . . .	8
correctBins . . . . .	9
createBins . . . . .	10
estimateCorrection . . . . .	12
exportBins . . . . .	13
frequencyPlot . . . . .	14
getBinAnnotations . . . . .	15
highlightFilters . . . . .	16
isobarPlot . . . . .	17
LGG150 . . . . .	18
makeCgh . . . . .	18
noisePlot . . . . .	19
normalizeBins . . . . .	20
normalizeSegmentedBins . . . . .	21
plot . . . . .	22
poolRuns . . . . .	22
QDNaseq-defunct . . . . .	23
QDNaseqCopyNumbers . . . . .	24
QDNaseqReadCounts . . . . .	24
QDNaseqSignals . . . . .	25
segmentBins . . . . .	25
smoothOutlierBins . . . . .	27
<b>Index</b>	<b>28</b>

**Description**

Quantitative DNA sequencing for chromosomal aberrations. The genome is divided into non-overlapping fixed-sized bins, number of sequence reads in each counted, adjusted with a simultaneous two-dimensional loess correction for sequence mappability and GC content, and filtered to remove spurious regions in the genome. Downstream steps of segmentation and calling are also implemented via packages DNACopy and CGHcall, respectively.

**Details**

A package to detect chromosomal aberrations from whole-genome sequencing data. [QDNaseqReadCounts](#) and [QDNaseqCopyNumbers](#) classes are used as the main data structures.

**How to cite this package**

Whenever using this package, please cite: Scheinin I, Sie D, Bengtsson H, van de Wiel MA, Olshen AB, van Thuijl HF, van Essen HF, Eijk PP, Rustenburg F, Meijer GA, Reijneveld JC, Wesseling P, Pinkel D, Albertson DG, Ylstra B (2014). "DNA copy number analysis of fresh and formalin-fixed specimens by shallow whole-genome sequencing with identification and exclusion of problematic regions in the genome assembly." *\_Genome Research\_, \*24\**, 2022-2032.

**License**

This package is licensed under GPL.

**Author(s)**

Ilari Scheinin

---

addPhenodata	<i>Adds phenotype data from a file to a <a href="#">QDNaseqReadCounts</a> or a <a href="#">QDNaseqCopyNumbers</a> object</i>
--------------	--

---

**Description**

Adds phenotype data from a file to a [QDNaseqReadCounts](#) or a [QDNaseqCopyNumbers](#) object.

**Usage**

```
addPhenodata(object, phenofile)
```

**Arguments**

object	A <a href="#">QDNaseqReadCounts</a> or <a href="#">QDNaseqCopyNumbers</a> object.
phenofile	A file name with phenotypic data for samples in object.

**Value**

Returns a [QDNaseqReadCounts](#) or [QDNaseqCopyNumbers](#) object with phenotype data added.

**Author(s)**

Ilari Scheinin

**Examples**

```

data(LGG150)
readCounts <- LGG150
## Not run:
readCounts <- addPhenodata(readCounts, "phenodata.txt")

## End(Not run)

```

---

applyFilters

*Adjusts the filtering on which bins are used*


---

**Description**

Adjusts the filtering on which bins are used.

**Usage**

```

applyFilters(object, residual=TRUE, blacklist=TRUE, mappability=NA, bases=NA,
  chromosomes=c("X", "Y"), verbose=getOption("QDNaseq::verbose", TRUE))

```

**Arguments**

object	A <a href="#">QDNaseqReadCounts</a> object.
residual	Either a <a href="#">logical</a> specifying whether to filter based on loess residuals of the calibration set, or if a <a href="#">numeric</a> , the cutoff as number of standard deviations estimated with <a href="#">madDiff</a> to use for. Default is <a href="#">TRUE</a> , which corresponds to 4.0 standard deviations.
blacklist	Either a <a href="#">logical</a> specifying whether to filter based on overlap with blacklisted regions, or if numeric, the maximum percentage of overlap allowed. Default is <a href="#">TRUE</a> , which corresponds to no overlap allowed (i.e. value of 0).
mappability	A <a href="#">numeric</a> in [0, 100] to specify filtering out bins with mappabilities lower than the number specified. NA (default) or <a href="#">FALSE</a> will not filter based on mappability.
bases	A <a href="#">numeric</a> specifying the minimum percentage of characterized bases (not Ns) in the reference genome sequence. NA (default) or <a href="#">FALSE</a> will not filter based on uncharacterized bases.
chromosomes	A <a href="#">character</a> vector specifying which chromosomes to filter out. Defaults to the sex chromosomes and mitochondria, i.e. <code>c("X", "Y", "MT")</code> .
verbose	If <a href="#">TRUE</a> , verbose messages are produced.

**Value**

Returns a [QDNaseqReadCounts](#) object with updated filtering.

**Author(s)**

Ilari Scheinin

**Examples**

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
```

---

binReadCounts	<i>Calculate binned read counts from a set of BAM files</i>
---------------	---

---

**Description**

Calculate binned read counts from a set of BAM files.

**Usage**

```
binReadCounts(bins, bamfiles=NULL, path=NULL, ext="bam", bamnames=NULL, phenofile=NULL,
  chunkSize=NULL, cache=getOption("QDNaseq::cache", FALSE), force=!cache, isPaired=NA,
  isProperPair=NA, isUnmappedQuery=FALSE, hasUnmappedMate=NA, isMinusStrand=NA,
  isMateMinusStrand=NA, isFirstMateRead=NA, isSecondMateRead=NA, isSecondaryAlignment=NA,
  isNotPassingQualityControls=FALSE, isDuplicate=FALSE, minMapq=37, pairedEnds=NULL,
  verbose=getOption("QDNaseq::verbose", TRUE))
```

**Arguments**

bins	A data.frame or an <a href="#">AnnotatedDataFrame</a> object containing bin annotations.
bamfiles	A character vector of (BAM) file names. If NULL (default), all files with extension ext, are read from directory path.
path	If bamfiles is NULL, directory path to read input files from. Defaults to the current working directory.
ext	File name extension of input files to read, default is "bam".
bamnames	An optional character vector of sample names. Defaults to file names with extension ext removed.
phenofile	An optional character(1) specifying a file name for phenotype data.
chunkSize	An optional integer specifying the chunk size (nt) by which to process the bam file.
cache	Whether to read and write intermediate cache files, which speeds up subsequent analyses of the same files. Requires packages R.cache and digest (both available on CRAN) to be installed. Defaults to getOption("QDNaseq::cache", FALSE).
force	When using the cache, whether to force reading input data from the BAM files even when an intermediate cache file is present.
isPaired	A logical(1) indicating whether unpaired (FALSE), paired (TRUE), or any (NA, default) read should be returned.

isProperPair	A logical(1) indicating whether improperly paired (FALSE), properly paired (TRUE), or any (NA, default) read should be returned. A properly paired read is defined by the alignment algorithm and might, e.g., represent reads aligning to identical reference sequences and with a specified distance.
isUnmappedQuery	A logical(1) indicating whether unmapped (TRUE), mapped (FALSE, default), or any (NA) read should be returned.
hasUnmappedMate	A logical(1) indicating whether reads with mapped (FALSE), unmapped (TRUE), or any (NA, default) mate should be returned.
isMinusStrand	A logical(1) indicating whether reads aligned to the plus (FALSE), minus (TRUE), or any (NA, default) strand should be returned.
isMateMinusStrand	A logical(1) indicating whether mate reads aligned to the plus (FALSE), minus (TRUE), or any (NA, default) strand should be returned.
isFirstMateRead	A logical(1) indicating whether the first mate read should be returned (TRUE) or not (FALSE), or whether mate read number should be ignored (NA, default).
isSecondMateRead	A logical(1) indicating whether the second mate read should be returned (TRUE) or not (FALSE), or whether mate read number should be ignored (NA, default).
isSecondaryAlignment	A logical(1) indicating whether alignments that are primary (FALSE), are not primary (TRUE) or whose primary status does not matter (NA, default) should be returned. A non-primary alignment ("secondary alignment" in the SAM specification) might result when a read aligns to multiple locations. One alignment is designated as primary and has this flag set to FALSE; the remainder, for which this flag is TRUE, are designated by the aligner as secondary.
isNotPassingQualityControls	A logical(1) indicating whether reads passing quality controls (FALSE, default), reads not passing quality controls (TRUE), or any (NA) read should be returned.
isDuplicate	A logical(1) indicating that un-duplicated (FALSE, default), duplicated (TRUE), or any (NA) reads should be returned. 'Duplicated' reads may represent PCR or optical duplicates.
minMapq	If quality scores exists, the minimum quality score required in order to keep a read, otherwise all reads are kept.
pairedEnds	A boolean value or vector specifying whether the BAM files contain paired-end data or not. Only affects the calculation of the expected variance.
verbose	If TRUE, verbose messages are produced.

### Value

Returns a [QDNaseqReadCounts](#) object with assay data element counts containing the binned read counts as non-negative integers.

**Author(s)**

Ilari Scheinin, Daoud Sie

**Examples**

```
## Not run: # read all files from the current directory with names ending in .bam
bins <- getBinAnnotations(15)
readCounts <- binReadCounts(bins)

## End(Not run)
```

---

callBins

*Call aberrations from segmented copy number data*

---

**Description**

Call aberrations from segmented copy number data.

**Usage**

```
callBins(object, organism=c("human", "other"), method=c("CGHcall", "cutoff"),
  cutoffs=log2(c(deletion = 0.5, loss = 1.5, gain = 2.5, amplification = 10)/2), ...)
```

**Arguments**

object	An object of class <code>QDNAseqCopyNumbers</code>
organism	Either “human” or “other”, see manual page for <a href="#">CGHcall</a> for more details. This is only used for chromosome arm information when “prior” is set to “all” or “auto” (and <code>samplesize &gt; 20</code> ). Ignored when method is not “CGHcall”.
method	Calling method to use. Options currently implemented are: “CGHcall” or “cutoff”.
cutoffs	When <code>method=“cutoff”</code> , a numeric vector of (log2-transformed) thresholds to use for calling. At least one positive and one negative value must be provided. The smallest positive value is used as the cutoff for calling gains, and the negative value closest to zero is used as the cutoff for losses. If a second positive value is provided, it is used as the cutoff for amplifications. And if a second negative value is provided, it is used as the cutoff for homozygous deletions.
...	Additional arguments passed to <a href="#">CGHcall</a> .

**Details**

By default, chromosomal aberrations are called with **CGHcall**. It has been developed for the analysis of series of cancer samples, and uses a model that contains both gains and losses. If used on a single sample, or especially only on a subset of chromosomes, or especially on a single non-cancer sample, it may fail, but method “cutoff” can be used instead.

When using method “cutoff”, the default values assume a uniform cell population and correspond to thresholds of (assuming a diploid genome) 0.5, 1.5, 2.5, and 10 copies to distinguish between homozygous deletions, (hemizygous) losses, normal copy number, gains, and amplifications, respectively. When using with cancer samples, these values might require adjustments to account for tumor cell percentage.

### Value

Returns an object of class `QDNaseqCopyNumbers` with calling results added.

### Author(s)

Ilari Scheinin

### See Also

Internally, `CGHcall` and `ExpandCGHcall` of the `CGHcall` package are used when method=“CGHcall”.

### Examples

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
copyNumbersNormalized <- normalizeBins(copyNumbers)
copyNumbersSmooth <- smoothOutlierBins(copyNumbersNormalized)
copyNumbersSegmented <- segmentBins(copyNumbersSmooth)
copyNumbersSegmented <- normalizeSegmentedBins(copyNumbersSegmented)
copyNumbersCalled <- callBins(copyNumbersSegmented)
```

---

`compareToReference`      *Divide binned read counts with those of reference samples*

---

### Description

Divide binned read counts with those of reference samples.

### Usage

```
compareToReference(object, references, force=FALSE)
```

**Arguments**

object	An object of class <code>QDNaseqCopyNumbers</code> .
references	A numeric vector of indexes of the reference sample. Must be the same length as there are samples in object. When <code>NA</code> , the sample will be kept as is. When <code>FALSE</code> , the sample will be removed from the output. As an example, object contains three samples: tumor1, tumor2, and normal2. There is no reference for tumor1, but normal2 is a matched normal sample from the same patient as tumor2. The keep tumor1 as is, but to divide tumor2 with normal2, argument references should be <code>c(NA, 3, FALSE)</code> .
force	Whether to force the operation even when downstream data will be lost.

**Value**

Returns a `QDNaseqCopyNumbers` object with the desired samples divided by the signal of their reference samples.

**Author(s)**

Ilari Scheinin

**Examples**

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
copyNumbersNormalized <- normalizeBins(copyNumbers)
copyNumbersSmooth <- smoothOutlierBins(copyNumbersNormalized)
# Note: the following command will compare the sample to itself, which
# does not really make sense:
tumorVsNormal <- compareToReference(copyNumbersSmooth, c(1))
```

---

correctBins

*Correct binned read counts for GC content and mappability*

---

**Description**

Correct binned read counts for GC content and mappability.

**Usage**

```
correctBins(object, fit=NULL, method="ratio", adjustIncompletes=TRUE, ...)
```

**Arguments**

object	An <a href="#">QDNaseqReadCounts</a> object with counts data.
fit	An optional matrix of values to use for the correction. If NULL (default), assay data fit from object is used. If it is missing, it is generated with a call to <a href="#">estimateCorrection()</a> .
method	A character(1) string specifying the correction method. ratio (default) divides counts with fit. median calculates the median fit, and defines the correction for bins with GC content gc and mappability map as median(fit) - fit(gc, map), which is added to counts. Method none leaves counts untouched.
adjustIncompletes	A boolean(1) specifying whether counts for bins with uncharacterized nucleotides (N's) in their reference genome sequence should be adjusted by dividing them with the percentage of characterized (A, C, G, T) nucleotides. Defaults to <a href="#">TRUE</a> .
...	Additional arguments passed to <a href="#">estimateCorrection()</a> .

**Value**

Returns a [QDNaseqCopyNumbers](#) object with assay data element copynumber.

**Author(s)**

Ilari Scheinin

**See Also**

Internally, [loess](#) is used to fit the regression model.

**Examples**

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
```

---

createBins

*Builds bin annotation data for a particular bin size*

---

**Description**

Builds bin annotation data for a particular bin size.

**Usage**

```
createBins(bsgenome, binSize, ignoreMitochondria=TRUE, excludeSeqnames=NULL,
  verbose=getOption("QDNaseq::verbose", TRUE))
```

**Arguments**

bsgenome	A BSgenome package.
binSize	A <a href="#">numeric</a> scalar specifying the width of the bins in units of kbp (1000 base pairs), e.g. binSize=15 corresponds to 15 kbp bins.
ignoreMitochondria	Whether to ignore the mitochondria, defined as chromosomes named 'chrM', 'chrMT', 'M', or 'MT'.
excludeSeqnames	Character vector of seqnames which should be ignored.
verbose	If <a href="#">TRUE</a> , verbose messages are produced.

**Value**

Returns a [data.frame](#) with columns chromosome, start, end, bases, and gc, which correspond to the chromosome name, positions of the first and last base pair in the bin, the percentage of characterized nucleotides (A, C, G, or T, i.e. non-N), and GC content (percentage of C and G nucleotides of non-N nucleotides).

**Parallel processing**

The [future](#) is used parallelize the following functions:

- `createBins()` - parallelizes binned GC content across chromosomes
- `calculateBlacklist()` - parallelizes overlap counts across bins

**Author(s)**

Ilari Scheinin

**See Also**

[getBinAnnotations\(\)](#).

**Examples**

```
## Not run: # NOTE: These take a very long time to run.
library(BSgenome.Hsapiens.UCSC.hg19)
bins <- createBins(BSgenome.Hsapiens.UCSC.hg19, 15)
bins$mappability <- calculateMappability(bins,
  bigWigFile='/path/to/wgEncodeCrgMapabilityAlign50mer.bigWig',
  bigWigAverageOverBed='/path/to/bigWigAverageOverBed')
bins$blacklist <- calculateBlacklist(bins,
  bedFiles=c('/path/to/wgEncodeDacMapabilityConsensusExcludable.bed',
    '/path/to/wgEncodeDukeMapabilityRegionsExcludable.bed'))
bins$residual <- iterateResiduals(readCountsG1K)

## End(Not run)
```

---

estimateCorrection      *Estimate correction to read counts for GC content and mappability*

---

### Description

Estimate correction to read counts for GC content and mappability.

### Usage

```
estimateCorrection(object, span=0.65, family="symmetric", adjustIncompletes=TRUE,
  maxIter=1, cutoff=4, variables=c("gc", "mappability"), ...)
```

### Arguments

object	An <a href="#">QDNaseqReadCounts</a> object with counts data.
span	For <a href="#">loess</a> , the parameter alpha which controls the degree of smoothing.
family	For <a href="#">loess</a> , if "gaussian" fitting is by least-squares, and if "symmetric" a re-descending M estimator is used with Tukey's biweight function.
adjustIncompletes	A boolean(1) specifying whether counts for bins with uncharacterized nucleotides (N's) in their reference genome sequence should be adjusted by dividing them with the percentage of characterized (A, C, G, T) nucleotides. Defaults to <a href="#">TRUE</a> .
maxIter	An integer(1) specifying the maximum number of iterations to perform, default is 1. If larger, after the first loess fit, bins with median residuals larger than cutoff are removed, and the fitting repeated until the list of bins to use stabilizes or after maxIter iterations.
cutoff	A numeric(1) specifying the number of standard deviations (as estimated with <a href="#">madDiff</a> ) the cutoff for removal of bins with median residuals larger than the cutoff. Not used if maxIter=1 (default).
variables	A character vector specifying which variables to include in the correction. Can be c("gc", "mappability") (the default), "gc", or "mappability".
...	Additional arguments passed to <a href="#">loess</a> .

### Value

Returns a [QDNaseqReadCounts](#) object with the assay data element fit added.

### Parallel processing

This function uses **future** to calculate the QDNaseq model across samples in parallel.

### Author(s)

Ilari Scheinin

**See Also**

Internally, [loess](#) is used to fit the regression model.

**Examples**

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
```

---

exportBins	<i>Exports to a file</i>
------------	--------------------------

---

**Description**

Exports to a file.

**Usage**

```
exportBins(object, file, format=c("tsv", "igv", "bed", "vcf", "seg"),
  type=c("copynumber", "segments", "calls"), filter=TRUE, logTransform=TRUE, digits=3,
  chromosomeReplacements=c(`23` = "X", `24` = "Y", `25` = "MT"), ...)
```

**Arguments**

<code>object</code>	A <a href="#">QDNAseqReadCounts</a> or <a href="#">QDNAseqCopyNumbers</a> object.
<code>file</code>	Filename. For formats that support only one sample per file, such as BED, '%s' can be used as a placeholder for sample name or '%d' for sample number.
<code>format</code>	Format to export in. Currently supported ones are "tsv" (tab separated values), "igv" (Integrative Genomics Viewer), and "bed" (BED file format).
<code>type</code>	Type of data to export, options are "copynumber" (corrected or uncorrected read counts), "segments", or "calls".
<code>filter</code>	If <a href="#">TRUE</a> , bins are filtered, otherwise not.
<code>logTransform</code>	If <a href="#">TRUE</a> (default), exported data will be log2 transformed for format in "tsv", "igv", and "bed". This argument is ignored if type = "calls".
<code>digits</code>	The number of digits to round to. If not <a href="#">numeric</a> , no no rounding is performed.
<code>chromosomeReplacements</code>	A named character vector of chromosome name replacements to be done. Only used when object is of class <a href="#">cghRaw</a> , <a href="#">cghSeg</a> , <a href="#">cghCall</a> , or <a href="#">cghRegions</a> , since these classes store chromosome names as integers, whereas all QDNAseq object types use character vectors. Defaults to c("23"="X", "24"="Y", "25"="MT") for human.
<code>...</code>	Additional arguments passed to <a href="#">write.table</a> .

**Details**

Exports object to a file.

**Value**

Returns the pathnames of the files written.

**Author(s)**

Ilari Scheinin

**Examples**

```
## Not run:
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
copyNumbersNormalized <- normalizeBins(copyNumbers)
copyNumbersSmooth <- smoothOutlierBins(copyNumbersNormalized)
exportBins(copyNumbersSmooth, file="LGG150.igv", format="igv")

## End(Not run)
```

---

frequencyPlot

*Plot copy number aberration frequencies*

---

**Description**

Plot copy number aberration frequencies.

**Usage**

```
frequencyPlot(x, y, ...)
```

**Arguments**

x	A <a href="#">QDNAseqCopyNumbers</a> object with calls data.
y	missing
...	...

**Author(s)**

Ilari Scheinin

**Examples**

```

data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
copyNumbersNormalized <- normalizeBins(copyNumbers)
copyNumbersSmooth <- smoothOutlierBins(copyNumbersNormalized)
copyNumbersSegmented <- segmentBins(copyNumbersSmooth)
copyNumbersSegmented <- normalizeSegmentedBins(copyNumbersSegmented)
copyNumbersCalled <- callBins(copyNumbersSegmented)
frequencyPlot(copyNumbersCalled)

```

---

getBinAnnotations	<i>Gets bin annotation data for a particular bin size</i>
-------------------	---

---

**Description**

Gets bin annotation data for a particular bin size.

**Usage**

```

getBinAnnotations(binSize, genome="hg19", type="SR50",
  path=getOption("QDNaseq:binAnnotationPath", NULL),
  verbose=getOption("QDNaseq:verbose", TRUE))

```

**Arguments**

binSize	A <a href="#">numeric</a> scalar specifying the width of the bins in units of kbp (1000 base pairs), e.g. binSize=15 corresponds to 15 kbp bins.
genome	A <a href="#">character</a> string specify the genome and genome version to be used.
type	A <a href="#">character</a> string specify the experiment type, e.g. "SR50" or "PE100".
path	A <a href="#">character</a> string specifying the path for the bin annotation file to be downloaded. The path can either be on the local file system or a URL online. If <a href="#">NULL</a> (default), then data loaded from an R package named <b>QDNaseq.{{genome}}</b> . The default value can be controlled via R options <code>QDNaseq:binAnnotationPath</code> .
verbose	If <a href="#">TRUE</a> , verbose messages are produced.

**Details**

Gets bin annotation data for a particular bin size.

**Value**

Returns a [AnnotatedDataFrame](#) object.

**Author(s)**

Ilari Scheinin

**See Also**[createBins\(\)](#).**Examples**

```
## Not run:
bins <- getBinAnnotations(15)

## End(Not run)
```

highlightFilters

*Highlights data points in a plotted profile to evaluate filtering***Description**

Highlights data points in a plotted profile to evaluate filtering.

**Usage**

```
highlightFilters(object, col="red", residual=NA, blacklist=NA, mappability=NA, bases=NA,
  type="union", ...)
```

**Arguments**

object	A <a href="#">QDNaseqCopyNumbers</a> object.
col	The color used for highlighting.
residual	Either a <a href="#">logical</a> specifying whether to filter based on loess residuals of the calibration set, or if a <a href="#">numeric</a> , the cutoff as number of standard deviations estimated with <a href="#">madDiff</a> to use for. Default is <a href="#">TRUE</a> , which corresponds to 4.0 standard deviations.
blacklist	Either a <a href="#">logical</a> specifying whether to filter based on overlap with blacklisted regions, or if numeric, the maximum percentage of overlap allowed. Default is <a href="#">TRUE</a> , which corresponds to no overlap allowed (i.e. value of 0).
mappability	A <a href="#">numeric</a> in [0, 100] to specify filtering out bins with mappabilities lower than the number specified. NA (default) or <a href="#">FALSE</a> will not filter based on mappability.
bases	A <a href="#">numeric</a> specifying the minimum percentage of characterized bases (not Ns) in the reference genome sequence. NA (default) or <a href="#">FALSE</a> will not filter based on uncharacterized bases.
type	When specifying multiple filters (residual, blacklist, mappability, bases), whether to highlight their union (default) or intersection.
...	Further arguments to <a href="#">points</a> .

**Author(s)**

Ilari Scheinin

**Examples**

```
data(LGG150)
readCounts <- LGG150
plot(readCounts)
highlightFilters(readCounts, residual=TRUE, blacklist=TRUE)
```

---

isobarPlot

*Plot median read counts as a function of GC content and mappability*

---

**Description**

Plot median read counts as a function of GC content and mappability.

**Usage**

```
isobarPlot(x, y, ...)
```

**Arguments**

x	A <a href="#">QDNaseqReadCounts</a> object.
y	missing
...	...

**Author(s)**

Ilari Scheinin

**Examples**

```
data(LGG150)
readCounts <- LGG150
isobarPlot(readCounts)
```

LGG150

*LGG150 chromosomes 7-10***Description**

An example data set of read counts from chromosomes 7-10 of sample LGG150, contained within a [QDNaseqReadCounts](#) object

**Author(s)**

Ilari Scheinin

makeCgh

*Constructs a 'cghRaw', 'cghSeg', or 'cghCall' object***Description**

Constructs a 'cghRaw', 'cghSeg', or 'cghCall' object.

**Usage**

```
makeCgh(object, filter=TRUE, chromosomeReplacements=c(X = 23, Y = 24, MT = 25), ...)
```

**Arguments**

object	A <a href="#">QDNaseqCopyNumbers</a> object.
filter	If <code>TRUE</code> , bins are filtered, otherwise not.
chromosomeReplacements	A named integer vector of chromosome name replacements to be done. QDNaseq stores chromosome names as characters, but CGHcall expects them to be integers. Defaults to <code>c(X=23, Y=24, MT=25)</code> for human. Value of "auto" will use running numbers in order of appearance in the bin annotations.
...	Not used.

**Value**

Returns a [cghRaw](#) if the object has not been segmented, a [cghSeg](#) if it has been segmented but not called, or [cghCall](#) if it has been called as well.

**Author(s)**

Ilari Scheinin

**Examples**

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
copyNumbersNormalized <- normalizeBins(copyNumbers)
copyNumbersSmooth <- smoothOutlierBins(copyNumbersNormalized)
cgh <- makeCgh(copyNumbersSmooth)
```

---

noisePlot

*Plot noise as a function of sequence depth*

---

**Description**

Plot noise as a function of sequence depth.

**Usage**

```
noisePlot(x, y, ...)
```

**Arguments**

x	A <a href="#">QDNaseqReadCounts</a> object.
y	missing
...	Further arguments to <a href="#">plot()</a> and <a href="#">text</a> .

**Author(s)**

Ilari Scheinin

**Examples**

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
noisePlot(readCountsFiltered)
```

---

normalizeBins	<i>Normalizes binned read counts</i>
---------------	--------------------------------------

---

### Description

Normalizes binned read counts.

### Usage

```
normalizeBins(object, method="median", force=FALSE, verbose=getOption("QDNaseq:verbose", TRUE))
```

### Arguments

object	A <a href="#">QDNaseqCopyNumbers</a> object with copynumber data.
method	A <a href="#">character</a> string specifying the normalization method. Choices are "mean", "median" (default), or "mode". A partial string sufficient to uniquely identify the choice is permitted.
force	Running this function will remove possible segmentation and calling results. When they are present, running requires specifying force is <a href="#">TRUE</a> .
verbose	If <a href="#">TRUE</a> , verbose messages are produced.

### Value

Returns a [QDNaseqCopyNumbers](#) object with the assay data element copynumber scaled with the chosen method.

### Author(s)

Ilari Scheinin

### Examples

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
copyNumbersNormalized <- normalizeBins(copyNumbers)
```

---

`normalizeSegmentedBins`*Normalize segmented bins*

---

**Description**

Normalize segmented bins.

**Usage**

```
normalizeSegmentedBins(object, inter=c(-0.1, 0.1), force=FALSE)
```

**Arguments**

<code>object</code>	An object of class <code>QDNAseqCopyNumbers</code> .
<code>inter</code>	The interval in which the function should search for the normal level.
<code>force</code>	Whether to force execution when it causes removal of downstream calling results.

**Details**

This function recursively searches for the interval containing the most segmented data, decreasing the interval length in each recursion. The recursive search makes the post-segmentation normalization robust against local maxima. This function is particularly useful for profiles for which, after segmentation, the 0-level does not coincide with many segments. It is more or less harmless to other profiles. We advise to keep the search interval (`inter`) small, in particular at the positive (`gain`) side to avoid that the 0-level is set to a common gain level.

**Value**

Returns an object of class `QDNAseqCopyNumbers` with re-normalized data.

**Author(s)**

Ilari Scheinin

**See Also**

Internally, [postsegnormalize](#) of the **CGHcall** package is used.

**Examples**

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
copyNumbersNormalized <- normalizeBins(copyNumbers)
```

```
copyNumbersSmooth <- smoothOutlierBins(copyNumbersNormalized)
copyNumbersSegmented <- segmentBins(copyNumbersSmooth)
copyNumbersSegmented <- normalizeSegmentedBins(copyNumbersSegmented)
```

---

plot *Plot copy number profile*

---

### Description

Plot copy number profile.

### Usage

```
plot(x, y, ...)
```

### Arguments

x	A <a href="#">QDNAseqReadCounts</a> or <a href="#">QDNAseqCopyNumbers</a> object.
y	missing
...	...

### Author(s)

Ilari Scheinin

### Examples

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
plot(copyNumbers)
```

---

poolRuns *Pools binned read counts across samples*

---

### Description

Pools binned read counts across samples.

### Usage

```
poolRuns(object, samples, force=FALSE)
```

**Arguments**

object	A <a href="#">QDNaseqReadCounts</a> or <a href="#">QDNaseqCopyNumbers</a> object.
samples	A character vector of new sample names. Samples with identical names will be pooled together. Must be the same length as there are samples in object.
force	Whether to force the operation even when downstream data will be lost.

**Value**

Returns a [QDNaseqReadCounts](#) or [QDNaseqCopyNumbers](#) object.

**Author(s)**

Ilari Scheinin

**Examples**

```
data(LGG150)
readCounts <- LGG150
# Note: the following command will "pool" data from a single run, which
# does not really make sense:
pooledReadCounts <- poolRuns(readCounts, samples = "LGG150")
```

---

QDNaseq-defunct

*Defunct functions in package 'QDNaseq'*

---

**Description**

These functions are defunct and no longer available.

**Details**

The following functions are defunct; use the replacement indicated below:

- downloadBinAnnotations: [getBinAnnotations](#)

---

QDNaseqCopyNumbers      *Container for QDNaseq read count data*

---

### Description

Container for QDNaseq read count data

### Assay data elements

An object of this class contains the following elements:

`copynumber` (**numeric**) Corrected "count" signals in  $[0, +\infty)$  An object with only this field is created by `correctBins()`.

`segmented` (**numeric**; optional) Segmented data in  $[0, +\infty)$ , added by calling `segmentBins()`.

`calls` (**integer**; optional) Calls as -2=deletion, -1=loss, 0=normal, 1=gain, 2=amplification, added by calling `callBins()`.

`probdlloss` (**numeric**; optional) Probabilities of deletions in  $[0, 1]$ , added by calling `callBins()`.

`problloss` (**numeric**; optional) Probabilities of losses in  $[0, 1]$ , added by calling `callBins()`.

`probnorm` (**numeric**; optional) Probabilities of normal copy number in  $[0, 1]$ , added by calling `callBins()`.

`probgain` (**numeric**; optional) Probabilities of gains in  $[0, 1]$ , added by calling `callBins()`.

`probamp` (**numeric**; optional) Probabilities of amplifications in  $[0, 1]$ , added by calling `callBins()`.

### Missing values

The bin data (assay data) may contain missing values.

### Author(s)

Ilari Scheinin

---

QDNaseqReadCounts      *Container for QDNaseq read count data*

---

### Description

Container for QDNaseq read count data

### Assay data elements

An object of this class contains (a subset) the following elements:

`counts` (**numeric**) Binned read counts as non-negative integers in  $\{0, 1, 2, \dots\}$ . An object with only this field is created by `binReadCounts()`.

`fit` (**numeric**; optional) Loess fit of "count" signals as doubles. Normally, these should all be positive values, but a small number of edge case bins might contain negatives, especially when fitting unfiltered data. This element is added after calling `estimateCorrection()`.

**Missing values**

The bin data (assay data) may contain missing values.

**Author(s)**

Ilari Scheinin

---

QDNAseqSignals	<i>A parent class for containers of QDNAseq data</i>
----------------	--

---

**Description**

A parent class for containers of QDNAseq data

**Author(s)**

Ilari Scheinin

---

segmentBins	<i>Segments normalized copy number data</i>
-------------	---

---

**Description**

Segments normalized copy number data.

**Usage**

```
segmentBins(object, smoothBy=FALSE, alpha=1e-10, undo.splits="sdundo", undo.SD=1,
            force=FALSE, transformFun="log2", ...)
```

**Arguments**

object	An object of class QDNAseqCopyNumbers.
smoothBy	An optional integer value to perform smoothing before segmentation by taking the mean of every smoothBy bins, and then segment those means. Default ( <code>FALSE</code> ) is to perform no smoothing. <code>smoothBy=1L</code> is a special case that will not perform smoothing, but will split the segmentation process by chromosome instead of by sample.
alpha	Significance levels for the test to accept change-points. Default is 1e-10.
undo.splits	A character string specifying how change-points are to be undone, if at all. Default is "sdundo", which undoes splits that are not at least this many SDs apart. Other choices are "prune", which uses a sum of squares criterion, and "none".
undo.SD	The number of SDs between means to keep a split if undo.splits="sdundo". Default is 1.0.

force	Whether to force execution when it causes removal of downstream calling results.
transformFun	A function to transform the data with. This can be the default "log2" for $\log_2(x + .Machine\$double.xmin)$ , "sqrt" for the Anscombe transform of $\sqrt{x * 3/8}$ which stabilizes the variance, "none" for no transformation, or any R function that performs the desired transformation and also its inverse when called with parameter <code>inv=TRUE</code> .
...	Additional arguments passed to <a href="#">segment</a> .

**Value**

Returns an object of class `QDNAseqCopyNumbers` with segmentation results added.

**Numerical reproducibility**

This method make use of random number generation (RNG) via the [segment](#) used internally. Because of this, calling the method with the same input data multiple times will each time give slightly different results. To get numerically reproducible results, the random seed must be fixed, e.g. by using `'set.seed()'` at the top of the script.

**Parallel processing**

This function uses **future** to segment samples in parallel.

**Author(s)**

Ilari Scheinin

**References**

- [1] A.B. Olshen, E.S. Venkatraman (aka Venkatraman E. Seshan), R. Lucito and M. Wigler, *Circular binary segmentation for the analysis of array-based DNA copy number data*, Biostatistics, 2004
- [2] E.S. Venkatraman and A.B. Olshen, *A faster circular binary segmentation algorithm for the analysis of array CGH data*, Bioinformatics, 2007

**See Also**

Internally, [segment](#) of the **DNAcopy** package, which implements the CBS method [1,2], is used to segment the data.

**Examples**

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
copyNumbersNormalized <- normalizeBins(copyNumbers)
copyNumbersSmooth <- smoothOutlierBins(copyNumbersNormalized)
```

```
copyNumbersSegmented <- segmentBins(copyNumbersSmooth)
```

---

smoothOutlierBins      *Smooth outlier bins after normalization*

---

## Description

Smooth outlier bins after normalization.

## Usage

```
smoothOutlierBins(object, logTransform=TRUE, force=FALSE, ...)
```

## Arguments

object	A <a href="#">QDNaseqCopyNumbers</a> object with copynumber data.
logTransform	If <a href="#">TRUE</a> (default), data will be log2-transformed.
force	Running this function will remove possible segmentation and calling results. When they are present, running requires specifying force is <a href="#">TRUE</a> .
...	Additional arguments passed to <a href="#">smooth.CNA</a> .

## Value

Returns a [QDNaseqCopyNumbers](#) object with the values for outliers smoothed. See [smooth.CNA](#) for more details. If [logTransform](#) is [TRUE](#), these signals are log2-transformed prior to smoothing, but afterwards back-transformed..

## Author(s)

Ilari Scheinin

## Examples

```
data(LGG150)
readCounts <- LGG150
readCountsFiltered <- applyFilters(readCounts)
readCountsFiltered <- estimateCorrection(readCountsFiltered)
copyNumbers <- correctBins(readCountsFiltered)
copyNumbersNormalized <- normalizeBins(copyNumbers)
copyNumbersSmooth <- smoothOutlierBins(copyNumbersNormalized)
```

# Index

- \* **IO**
  - addPhenodata, 3
  - binReadCounts, 5
  - exportBins, 13
  - getBinAnnotations, 15
- \* **aplot**
  - highlightFilters, 16
- \* **classes**
  - QDNaseqCopyNumbers, 24
  - QDNaseqReadCounts, 24
  - QDNaseqSignals, 25
- \* **datasets**
  - LGG150, 18
- \* **file**
  - addPhenodata, 3
  - binReadCounts, 5
  - exportBins, 13
- \* **hplot**
  - frequencyPlot, 14
  - isobarPlot, 17
  - noisePlot, 19
  - plot, 22
- \* **loess**
  - correctBins, 9
  - estimateCorrection, 12
- \* **manip**
  - applyFilters, 4
  - callBins, 7
  - compareToReference, 8
  - correctBins, 9
  - estimateCorrection, 12
  - makeCgh, 18
  - normalizeBins, 20
  - normalizeSegmentedBins, 21
  - poolRuns, 22
  - segmentBins, 25
  - smoothOutlierBins, 27
- \* **package**
  - QDNaseq-package, 2
- \* **smooth**
  - segmentBins, 25
- addPhenodata, 3
- AnnotatedDataFrame, 5, 15
- applyFilters, 4
- applyFilters, QDNaseqReadCounts-method (applyFilters), 4
- binReadCounts, 5, 24
- bpend, QDNaseqSignals-method (QDNaseqSignals), 25
- bpstart, QDNaseqSignals-method (QDNaseqSignals), 25
- calculateBlacklist (createBins), 10
- calculateBlacklistByRegions (createBins), 10
- calculateMappability (createBins), 10
- callBins, 7, 24
- callBins, QDNaseqCopyNumbers-method (callBins), 7
- CGHcall, 7, 8
- cghCall, 13, 18
- cghRaw, 13, 18
- cghRegions, 13
- cghSeg, 13, 18
- character, 4, 15, 20
- chromosomes, QDNaseqSignals-method (QDNaseqSignals), 25
- compareToReference, 8
- compareToReference, QDNaseqCopyNumbers, numeric-method (compareToReference), 8
- correctBins, 9, 24
- correctBins, QDNaseqReadCounts-method (correctBins), 9
- createBins, 10, 16
- data.frame, 11
- downloadBinAnnotations (QDNaseq-defunct), 23

- estimateCorrection, [10, 12, 24](#)
- estimateCorrection,QDNAseqReadCounts-method  
(estimateCorrection), [12](#)
- ExpandCGHcall, [8](#)
- exportBins, [13](#)
- exportBins,QDNAseqSignals-method  
(exportBins), [13](#)
- FALSE, [4, 9, 16, 25](#)
- frequencyPlot, [14](#)
- frequencyPlot,QDNAseqCopyNumbers,missing-method  
(frequencyPlot), [14](#)
- getBinAnnotations, [11, 15, 23](#)
- highlightFilters, [16](#)
- highlightFilters,QDNAseqSignals-method  
(highlightFilters), [16](#)
- integer, [6, 24](#)
- isobarPlot, [17](#)
- isobarPlot,QDNAseqReadCounts,missing-method  
(isobarPlot), [17](#)
- iterateResiduals (createBins), [10](#)
- LGG150, [18](#)
- loess, [10, 12, 13](#)
- logical, [4, 16](#)
- madDiff, [4, 12, 16](#)
- makeCgh, [18](#)
- makeCgh,QDNAseqCopyNumbers-method  
(makeCgh), [18](#)
- NA, [9](#)
- noisePlot, [19](#)
- noisePlot,QDNAseqReadCounts,missing-method  
(noisePlot), [19](#)
- normalizeBins, [20](#)
- normalizeBins,QDNAseqCopyNumbers-method  
(normalizeBins), [20](#)
- normalizeSegmentedBins, [21](#)
- normalizeSegmentedBins,QDNAseqCopyNumbers-method  
(normalizeSegmentedBins), [21](#)
- NULL, [15](#)
- numeric, [4, 11, 13, 15, 16, 24](#)
- plot, [19, 22](#)
- plot,QDNAseqSignals,missing-method  
(plot), [22](#)
- points, [16](#)
- poolRuns, [22](#)
- poolRuns,QDNAseqSignals,character-method  
(poolRuns), [22](#)
- postsegnormalize, [21](#)
- QDNAseq (QDNAseq-package), [2](#)
- QDNAseq-defunct, [23](#)
- QDNAseq-package, [2](#)
- QDNAseqCopyNumbers, [3, 8–10, 13, 14, 16, 18,](#)  
[20, 22, 23, 24, 27](#)
- QDNAseqCopyNumbers-class  
(QDNAseqCopyNumbers), [24](#)
- QDNAseqReadCounts, [3, 4, 6, 10, 12, 13,](#)  
[17–19, 22, 23, 24](#)
- QDNAseqReadCounts-class  
(QDNAseqReadCounts), [24](#)
- QDNAseqSignals, [25](#)
- QDNAseqSignals-class (QDNAseqSignals),  
[25](#)
- segment, [26](#)
- segmentBins, [24, 25](#)
- segmentBins,QDNAseqCopyNumbers-method  
(segmentBins), [25](#)
- smooth.CNA, [27](#)
- smoothOutlierBins, [27](#)
- smoothOutlierBins,QDNAseqCopyNumbers-method  
(smoothOutlierBins), [27](#)
- text, [19](#)
- TRUE, [4, 6, 10–13, 15, 16, 18, 20, 27](#)
- write.table, [13](#)