

# Package ‘RobustCNV’

July 30, 2015

**Type** Package

**Title** A Package for Copy Number Variant Analysis of Targeted Capture Sequencing Data

**Version** 0.0.0.9000

**Date** 2015-06-01

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**Description** A copy number variant tool using robust regression for normalization and DNA-copy for segmentation.

**License** Artistic-2.0

**LazyData** TRUE

**Imports** DNACopy,  
MASS,  
IRanges,

**Suggests** WGCNA,  
knitr

**VignetteBuilder** knitr

## R topics documented:

callSegments . . . . .	2
formatForNexus . . . . .	2
plotGCBias . . . . .	3
plotGenomeIntervalCalls . . . . .	4
plotSampleClusters . . . . .	4
robustNorm . . . . .	5
segment . . . . .	6
summarizeIntervalCalls . . . . .	6
summarizeToGeneCalls . . . . .	7

<b>Index</b>	<b>8</b>
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callSegments	<i>Make calls on segments based on a cutoff calculated from segment SD and a tuning factor.</i>
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### Description

Make calls on segments based on a cutoff calculated from segment SD and a tuning factor.

### Usage

```
callSegments(gsegments, normCoverage, tuning = 0.8)
```

### Arguments

gsegments	a dataframe with columns: Sample,Chromosome,Start,End,NumProbes,Segment,SegmentMean
normCoverage	a vector or dataframe containing normalized coverage
tuning	a parameter for adjusting sensitivity, higher values are less sensitive, more specific.

### Value

A new dataframe with a "SegmentCall" column consisting of (-, 0, +).

### Examples

```
## Not run:
callSegments(gsegments, normCoverage)
callSegments(gsegments, normCoverage, tuning=.2)

## End(Not run)
```

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formatForNexus	<i>Create an output formatted for Nexus compatibility.</i>
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### Description

Create an output formatted for Nexus compatibility.

### Usage

```
formatForNexus(perIntervalCalls)
```

### Arguments

perIntervalCalls	a data.frame containing per-interval calls
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**Value**

A data.frame formatted for Nexus compatibility

**Examples**

```
## Not run:  
formatForNexus(perIntervalCalls)  
  
## End(Not run)
```

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plotGCBias	<i>Plot GC bias plot</i>
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**Description**

Plot GC bias plot

**Usage**

```
plotGCBias(sampleName, pGC, coverage)
```

**Arguments**

sampleName	sample name
pGC	vector of percent GC for each interval
coverage	vector of per-interval coverage

**Value**

A gc-score value which is the sum of abs(predicted values).

**Examples**

```
## Not run:  
plotGCBias(sampleName, pGC, coverage)  
  
## End(Not run)
```

---

plotGenomeIntervalCalls  
*Make genome plots.*

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**Description**

Make genome plots.

**Usage**

```
plotGenomeIntervalCalls(perIntervalCalls, main = "")
```

**Arguments**

perIntervalCalls  
data.frame containing per-Interval calls and other data.

main  
title of plot.

**Value**

nothing.

**Examples**

```
## Not run:  
plotGenomeIntervalCalls(perIntervalCalls)  
  
## End(Not run)
```

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plotSampleClusters     *Make sample-cluster plots.*

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**Description**

Make sample-cluster plots.

**Usage**

```
plotSampleClusters(rawNormalCoverage, rawTumorCoverage)
```

**Arguments**

rawNormalCoverage  
Data.frame containing raw coverage for normal samples

rawTumorCoverage  
Data.frame containing raw coverage for tumor samples

**Value**

nothing.

**Examples**

```
## Not run:
plotSampleClusters(rawNormalCoverage, rawTumorCoverage)

## End(Not run)
```

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robustNorm	<i>Normalize coverage data.</i>
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**Description**

Do a normalization with interval length, gcContent and panel of normals using a robust regression strategy followed by a GC correction using loess.

**Usage**

```
robustNorm(y, x, ilengths, pGC, idx = NA)
```

**Arguments**

y	a vector of coverage values for the sample of interest.
x	a dataframe or matrix with coverage values for normal samples (columns are samples).
ilengths	a vector of interval lengths.
pGC,	the percent GC content for each interval
idx	a vector used to subset the input during model generation only (all intervals are reported)

**Value**

A new vector normalized against the panel of normals, interval lengths, and for GC content

**Examples**

```
## Not run:
robustNorm(y, x, ilengths, pGC)

## End(Not run)
```

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segment	<i>Segment normalized log2 coverage values for one or more samples.</i>
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### Description

Segment normalized log2 coverage values for one or more samples.

### Usage

```
segment(normCoverage, orderedChrom, maploc, idx = NA)
```

### Arguments

normCoverage	a data.frame or matrix with normalized coverage. Columns should be samples.
orderedChrom	a vector of ordered factors indicating the chromosome, must be nrow(norm.coverage) long.
maploc	a vector containing chromosome locus for intervals (end of each interval)
idx	segments can be called using a subset of targeted intervals, idx provides the subset.

### Value

A new dataframe with columns: Sample, Chromosome, Start, End, NumProbes, SegmentMean, SegmentCall

### Examples

```
## Not run:
segment(normalizedTumorCoverage, orderedChrom, maploc)

## End(Not run)
```

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summarizeIntervalCalls	<i>Summarize interval level calls to gene level call. This is an interval function that is used by summarizeToGeneCalls.</i>
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### Description

Summarize interval level calls to gene level call. This is an interval function that is used by summarizeToGeneCalls.

### Usage

```
summarizeIntervalCalls(x)
```

**Arguments**

x                      a vector of calls for a single gene

**Value**

A data.frame containing QC metrics

**Examples**

```
## Not run:  
CNVqc(data)  
  
## End(Not run)
```

---

summarizeToGeneCalls    *Summarize per-IntervalCalls to gene level calls.*

---

**Description**

Summarize per-IntervalCalls to gene level calls.

**Usage**

```
summarizeToGeneCalls(perIntervalCalls)
```

**Arguments**

perIntervalCalls  
                    a vector of calls for a single gene

**Value**

A data.frame containing calls summarized to gene level

**Examples**

```
## Not run:  
CNVqc(data)  
  
## End(Not run)
```

# Index

`callSegments`, [2](#)

`formatForNexus`, [2](#)

`plotGCBias`, [3](#)

`plotGenomeIntervalCalls`, [4](#)

`plotSampleClusters`, [4](#)

`robustNorm`, [5](#)

`segment`, [6](#)

`summarizeIntervalCalls`, [6](#)

`summarizeToGeneCalls`, [7](#)