

# Package ‘conumee’

June 17, 2016

**Title** Enhanced copy-number variation analysis using Illumina 450k methylation arrays

**Version** 1.4.2

**Author** Volker Hovestadt, Marc Zapatka

**Maintainer** Volker Hovestadt <conumee@hovestadt.bio>

**Address** Division of Molecular Genetics, German Cancer Research Center (DKFZ), Heidelberg, Germany

**Description** This package contains a set of processing and plotting methods for performing copy-number variation (CNV) analysis using Illumina 450k methylation arrays.

**Imports** methods, stats, DNACopy, rtracklayer, GenomicRanges, IRanges, GenomeInfoDb

**Depends** R (>= 3.0), minfi, IlluminaHumanMethylation450kmanifest, IlluminaHumanMethylation450kanno.ilmn12.hg19

**Suggests** BiocStyle, knitr, rmarkdown, minfiData, CopyNumber450kData, RCurl

**License** GPL (>= 2)

**LazyData** false

**Collate** classes.R annotation.R load.R process.R output.R data.R

**biocViews** CopyNumberVariation, DNAMethylation, MethylationArray, Microarray, Normalization, Preprocessing, QualityControl, Software

**VignetteBuilder** knitr

**NeedsCompilation** no

## R topics documented:

CNV.analysis-class . . . . .	2
CNV.anno-class . . . . .	3
CNV.bin . . . . .	4
CNV.check . . . . .	6

CNV.create_anno	6
CNV.create_bins	7
CNV.data-class	8
CNV.detail	9
CNV.detailplot	10
CNV.detailplot_wrap	12
CNV.fit	13
CNV.genomeplot	14
CNV.load	16
CNV.merge_bins	17
CNV.segment	18
CNV.write	19
detail_regions	20
exclude_regions	21
read.450k.url	21
tbl_ucsc	22
tcgaBRCA.sentrrix2name	22
<b>Index</b>	<b>23</b>

---

CNV.analysis-class	<i>CNV.analysis class</i>
--------------------	---------------------------

---

## Description

CNV analysis data of a single sample is stored in this class

## Usage

```
## S4 method for signature 'CNV.analysis'
show(object)

## S4 method for signature 'CNV.analysis'
names(x)

## S4 replacement method for signature 'CNV.analysis'
names(x) <- value

## S4 method for signature 'CNV.analysis'
coef(object)
```

## Arguments

object	CNV.analysis object
x	CNV.analysis object (defined by show generic).
value	Replacement names.

**Details**

Use `CNV.fit` to create. Modified by `CNV.bin`, `CNV.detail` and `CNV.segment`.

**Value**

CNV.analysis class.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```
# prepare
library(minfiData)
data(MsetEx)
d <- CNV.load(MsetEx)
anno <- CNV.create_anno()

# create object
x <- CNV.fit(query = d['GroupB_1'], ref = d[c('GroupA_1', 'GroupA_2', 'GroupA_3')], anno)

# modify object
x <- CNV.bin(x)
x <- CNV.detail(x)
x <- CNV.segment(x)

# general information
x
show(x)

# coefficients of linear regression
coef(x)

# show or replace sample name
names(x)
names(x) <- 'Sample 1'

# output plots
CNV.genomeplot(x)
CNV.genomeplot(x, chr = 'chr6')
#CNV.detailplot(x, name = 'MYCN')
#CNV.detailplot_wrap(x)
CNV.write(x, what = 'segments')
```

**Description**

Annotations required for CNV analysis are stored in this class.

**Usage**

```
## S4 method for signature 'CNV.anno'  
show(object)
```

**Arguments**

object            CNV.anno object

**Details**

This class does not contain any sample data. Use `CNV.create_anno` to create.

**Value**

CNV.anno class.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```
# create object  
anno <- CNV.create_anno()  
  
# general information  
anno  
show(anno)
```

---

CNV.bin

*CNV.bin*

---

**Description**

Combine single probe intensity values into predefined bins.

**Usage**

```
CNV.bin(object, ...)  
  
## S4 method for signature 'CNV.analysis'  
CNV.bin(object)
```

**Arguments**

object            CNV.analysis object.  
...                Additional parameters (CNV.bin generic, currently not used).

**Details**

The median intensity per bin is calculated. Bins are defined using `CNV.create_anno`. A value by which all probe and bin intensity values are shifted in subsequent analysis steps is calculated by minimizing the median absolute deviation from all bins to zero (ideally shifting the copy-neutral state to 0).

**Value**

CNV.analysis object.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```
# prepare
library(minfiData)
data(MsetEx)
d <- CNV.load(MsetEx)
data(detail_regions)
anno <- CNV.create_anno(detail_regions = detail_regions)

# create object
x <- CNV.fit(query = d['GroupB_1'], ref = d[c('GroupA_1', 'GroupA_2', 'GroupA_3')], anno)

# modify object
x <- CNV.bin(x)
#x <- CNV.detail(x)
#x <- CNV.segment(x)

# general information
x
show(x)

# coefficients of linear regression
coef(x)

# show or replace sample name
names(x)
names(x) <- 'Sample 1'
```

CNV.check

*CNV.check*

---

**Description**

Check intensity values.

**Usage**

```
CNV.check(object)
```

```
## S4 method for signature 'CNV.data'
```

```
CNV.check(object)
```

**Arguments**

object            CNV.data object.

**Details**

This method checks if intensities are positive and not NA. If not, they are set to 1. Warnings are given if intensities are abnormally high or low (> 50000 or < 5000, respectively).

**Value**

CNV.data object.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

---

CNV.create\_anno*CNV.create\_anno*

---

**Description**

Create annotations for CNV analysis.

**Usage**

```
CNV.create_anno(bin_minprobes = 15, bin_minsize = 50000,  
  bin_maxsize = 5e+06, chrXY = FALSE, exclude_regions = NULL,  
  detail_regions = NULL)
```

**Arguments**

bin_minprobes	numeric. Minimum number of probes per bin. Bins are iteratively merged with neighboring bin until minimum number is reached.
bin_minsize	numeric. Minimum size of a bin.
bin_maxsize	numeric. Maximum size of a bin. Merged bins that are larger are filtered out.
chrXY	logical. Should chromosome X and Y be included in the analysis?
exclude_regions	GRanges object or path to bed file containing genomic regions to be excluded.
detail_regions	GRanges object or path to bed file containing genomic regions to be examined in detail.

**Details**

This function collects all annotations required for CNV analysis using 450k arrays. The output CNV.anno object is not editable. Rerun CNV.create\_anno to change parameters.

**Value**

CNV.anno object.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```
# create annotation object
anno <- CNV.create_anno()
anno
```

---

CNV.create_bins	<i>CNV.create_bins</i>
-----------------	------------------------

---

**Description**

Split genome into bins of defined size.

**Usage**

```
CNV.create_bins(hg19.anno, bin_minsize = 50000, hg19.gap, hg19.exclude)
```

**Arguments**

hg19.anno	foo
bin_minsize	foo
hg19.gap	foo
hg19.exclude	foo

**Value**

GRanges object.

---

CNV.data-class	<i>CNV.data class</i>
----------------	-----------------------

---

**Description**

Intensities of one or multiple samples are stored in this class.

**Usage**

```
## S4 method for signature 'CNV.data'  
show(object)  
  
## S4 method for signature 'CNV.data,ANY,ANY'  
x[i]  
  
## S4 method for signature 'CNV.data'  
names(x)  
  
## S4 replacement method for signature 'CNV.data'  
names(x) <- value
```

**Arguments**

object	CNV.data object
x	CNV.data object (defined by Extract generic).
i	index. logical, numeric or character.
value	Replacement names.

**Details**

Use `CNV.load` to create.

**Value**

CNV.data class.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>



**Examples**

```
# create object
library(minfiData)
data(MsetEx)

d <- CNV.load(MsetEx)

# general information
d
show(d)

# show or replace sample names
names(d)
names(d) <- toupper(names(d))

# subset samples
d[1:2]
```

---

CNV.detail

*CNV.detail*

---

**Description**

Combine single probe values within detail regions.

**Usage**

```
CNV.detail(object, ...)
```

## S4 method for signature 'CNV.analysis'

```
CNV.detail(object)
```

**Arguments**

object            CNV.analysis object.

...                Additional parameters (CNV.detail generic, currently not used).

**Details**

The median intensity per detail region is calculated. Detail regions are defined using `CNV.create_anno(detail_bed=)`

**Value**

CNV.analysis object.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```

# prepare
library(minfiData)
data(MsetEx)
d <- CNV.load(MsetEx)
data(detail_regions)
anno <- CNV.create_anno(detail_regions = detail_regions)

# create object
x <- CNV.fit(query = d['GroupB_1'], ref = d[c('GroupA_1', 'GroupA_2', 'GroupA_3')], anno)

# modify object
x <- CNV.bin(x)
x <- CNV.detail(x)
#x <- CNV.segment(x)

# general information
x
show(x)

# coefficients of linear regression
coef(x)

# show or replace sample name
names(x)
names(x) <- 'Sample 1'

```

---

CNV.detailplot

*CNV.detailplot*


---

**Description**

Create CNV plot for detail region.

**Usage**

```
CNV.detailplot(object, ...)
```

```

## S4 method for signature 'CNV.analysis'
CNV.detailplot(object, name, yaxt = "l",
  ylim = c(-1.25, 1.25), set_par = TRUE, cols = c("red", "red",
  "lightgrey", "green", "green"))

```

**Arguments**

object	CNV.analysis object.
...	Additional parameters (CNV.detailplot generic, currently not used).
name	character. Name of detail region to plot.

yaxt	character. Include y-axis? 'l': left, 'r': right, 'n': no. Defaults to 'l'.
ylim	numeric vector. The y limits of the plot. Defaults to c(-1.25, 1.25).
set_par	logical. Use recommended graphical parameters for oma and mar? Defaults to TRUE. Original parameters are restored afterwards.
cols	character vector. Colors to use for plotting intensity levels of bins. Centered around 0. Defaults to c('red', 'red', 'lightgrey', 'green', 'green').

### Details

This method provides the functionality for generating detail regions CNV plots. Probes are shown as dots, bins are shown as lines. See parameters for more information.

### Value

NULL.

### Author(s)

Volker Hovestadt <conumee@hovestadt.bio>

### Examples

```
# prepare
library(minfiData)
data(MsetEx)
d <- CNV.load(MsetEx)
data(detail_regions)
anno <- CNV.create_anno(detail_regions = detail_regions)

# create/modify object
x <- CNV.segment(CNV.detail(CNV.bin(CNV.fit(query = d['GroupB_1'],
  ref = d[c('GroupA_1', 'GroupA_2', 'GroupA_3')], anno))))

# output plots
CNV.genomeplot(x)
CNV.genomeplot(x, chr = 'chr6')
CNV.detailplot(x, name = 'PTEN')
CNV.detailplot_wrap(x)

# output text files
CNV.write(x, what = 'segments')
CNV.write(x, what = 'detail')
CNV.write(x, what = 'bins')
CNV.write(x, what = 'probes')
```

---

CNV.detailplot\_wrap    *CNV.detailplot\_wrap*

---

### Description

Create CNV plot for all detail regions.

### Usage

```
CNV.detailplot_wrap(object, ...)

## S4 method for signature 'CNV.analysis'
CNV.detailplot_wrap(object, set_par = TRUE,
  main = NULL, ...)
```

### Arguments

object	CNV.analysis object.
...	Additional paramters supplied to CNV.detailplot.
set_par	logical. Use recommended graphical parameters for oma and mar? Defaults to TRUE. Original parameters are restored afterwards.
main	character. Title of the plot. Defaults to sample name.

### Details

This method is a wrapper of the CNV.detailplot method to plot all detail regions.

### Value

NULL.

### Author(s)

Volker Hovestadt <conumee@hovestadt.bio>

### Examples

```
# prepare
library(minfiData)
data(MsetEx)
d <- CNV.load(MsetEx)
data(detail_regions)
anno <- CNV.create_anno(detail_regions = detail_regions)

# create/modify object
x <- CNV.segment(CNV.detail(CNV.bin(CNV.fit(query = d['GroupB_1'],
  ref = d[c('GroupA_1', 'GroupA_2', 'GroupA_3')], anno))))
```

```

# output plots
CNV.genomeplot(x)
CNV.genomeplot(x, chr = 'chr6')
CNV.detailplot(x, name = 'PTEN')
CNV.detailplot_wrap(x)

# output text files
CNV.write(x, what = 'segments')
CNV.write(x, what = 'detail')
CNV.write(x, what = 'bins')
CNV.write(x, what = 'probes')

```

---

CNV.fit

*CNV.fit*


---

### Description

Normalize query sample intensities by fitting intensities to reference set using a linear regression model.

### Usage

```
CNV.fit(query, ref, anno, ...)
```

```
## S4 method for signature 'CNV.data,CNV.data,CNV.anno'
CNV.fit(query, ref, anno, name = NULL,
        intercept = TRUE)
```

### Arguments

query	CNV.data object of query sample (single sample).
ref	CNV.data object of reference set.
anno	CNV.anno object. Use CNV.create_anno do create.
...	Additional parameters (CNV.fit generic, currently not used).
name	character. Optional parameter to set query sample name.
intercept	logical. Should intercept be considered? Defaults to TRUE.

### Details

The log<sub>2</sub> ratio of query intensities versus a linear combination of reference set intensities that best reflects query intensities is calculated (as determined by linear regression). The annotations provided to CNV.fit are saved within the returned CNV.analysis object and used for subsequent analysis steps.

### Value

CNV.analysis object.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```
# prepare
library(minfiData)
data(MsetEx)
d <- CNV.load(MsetEx)
data(detail_regions)
anno <- CNV.create_anno(detail_regions = detail_regions)

# create object
x <- CNV.fit(query = d['GroupB_1'], ref = d[c('GroupA_1', 'GroupA_2', 'GroupA_3')], anno)

# modify object
#x <- CNV.bin(x)
#x <- CNV.detail(x)
#x <- CNV.segment(x)

# general information
x
show(x)

# coefficients of linear regression
coef(x)

# show or replace sample name
names(x)
names(x) <- 'Sample 1'
```

---

CNV.genomeplot

*CNV.genomeplot*

---

**Description**

Create CNV plot for the whole genome or chromosomes.

**Usage**

```
CNV.genomeplot(object, ...)

## S4 method for signature 'CNV.analysis'
CNV.genomeplot(object, chr = "all", chrX = TRUE,
  chrY = TRUE, centromere = TRUE, detail = TRUE, main = NULL,
  ylim = c(-1.25, 1.25), set_par = TRUE, cols = c("red", "red",
  "lightgrey", "green", "green"))

## S4 method for signature 'CNV.analysis,ANY'
plot(x, y = NULL, ...)
```

**Arguments**

object	CNV.analysis object.
...	Additional parameters supplied to CNV.genomeplot.
chr	character vector. Which chromosomes to plot. Defaults to 'all'.
chrX	logical. Plot values for chrX? Defaults to TRUE. Set CNV.create_anno(chrXY = FALSE) if chrX and Y should not be included at all.
chrY	logical. Plot values for chrY? Defaults to TRUE.
centromere	logical. Show dashed lines at centromeres? Defaults to TRUE.
detail	logical. If available, include labels of detail regions? Defaults to TRUE.
main	character. Title of the plot. Defaults to sample name.
ylim	numeric vector. The y limits of the plot. Defaults to c(-1.25, 1.25).
set_par	logical. Use recommended graphical parameters for oma and mar? Defaults to TRUE. Original parameters are restored afterwards.
cols	character vector. Colors to use for plotting intensity levels of bins. Centered around 0. Defaults to c('red', 'red', 'lightgrey', 'green', 'green').
x	CNV.analysis object (defined by plot generic).
y	NULL (defined by plot generic).

**Details**

This method provides the functionality for generating CNV plots for the whole genome or defined chromosomes. Bins are shown as dots, segments are shown as lines. See parameters for more information.

**Value**

NULL.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```
# prepare
library(minfiData)
data(MsetEx)
d <- CNV.load(MsetEx)
data(detail_regions)
anno <- CNV.create_anno(detail_regions = detail_regions)

# create/modify object
x <- CNV.segment(CNV.detail(CNV.bin(CNV.fit(query = d['GroupB_1'],
  ref = d[c('GroupA_1', 'GroupA_2', 'GroupA_3')], anno))))

# output plots
```

```

CNV.genomeplot(x)
CNV.genomeplot(x, chr = 'chr6')
CNV.detailplot(x, name = 'PTEN')
CNV.detailplot_wrap(x)

# output text files
CNV.write(x, what = 'segments')
CNV.write(x, what = 'detail')
CNV.write(x, what = 'bins')
CNV.write(x, what = 'probes')

```

---

CNV.load

*CNV.load*


---

### Description

Prepare combined intensities from various input objects.

### Usage

```

CNV.load(input, ...)

## S4 method for signature 'MethylSet'
CNV.load(input, names = NULL)

## S4 method for signature 'data.frame'
CNV.load(input, names = NULL)

## S4 method for signature 'matrix'
CNV.load(input, names = NULL)

## S4 method for signature 'numeric'
CNV.load(input, names = NULL)

```

### Arguments

input	Object of MethylSet class (minfi package), data.frame class, matrix class or numeric class.
...	Additional parameters (CNV.load generic, currently not used).
names	Vector specifying sample names. If not supplied, colnames are used. For MethylSet input, the first column of pData(input) matching 'name' (grep) is used.

### Details

This method gathers combined intensities of the Methylated and Unmethylated signals for all supplied probes. Probe IDs must be supplied as row names or in a separate column named 'ID\_REF' or 'TargetID'. If column names match 'intensity', only those columns are used. Else, if column names match 'signal' or 'methylated', only those columns are used. Otherwise, all columns are used.



**Value**

CNV.data object.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```
library(minfiData)
d <- CNV.load(MsetEx)
d
```

---

CNV.merge\_bins

*CNV.merge\_bins*

---

**Description**

Merge bins containing less than the defined number probes with neighboring bin containing fewer probes.

**Usage**

```
CNV.merge_bins(hg19.anno, hg19.tile, bin_minprobes = 20, hg19.probes,
  bin_maxsize = 5e+06, verbose = FALSE)
```

**Arguments**

hg19.anno	foo
hg19.tile	foo
bin_minprobes	foo
hg19.probes	foo
bin_maxsize	foo
verbose	foo

**Value**

GRanges object.

---

`CNV.segment`*CNV.segment*

---

**Description**

Segment bin values (wrapper of DNACopy package).

**Usage**

```
CNV.segment(object, ...)
```

```
## S4 method for signature 'CNV.analysis'  
CNV.segment(object, alpha = 0.001, nperm = 50000,  
  min.width = 5, undo.splits = "sdundo", undo.SD = 2.2, verbose = 0,  
  ...)
```

**Arguments**

<code>object</code>	CNV.analysis object.
<code>...</code>	Additional parameters supplied to the segment method of the DNACopy package.
<code>alpha</code>	See details. Defaults to 0.001.
<code>nperm</code>	See details. Defaults to 50000.
<code>min.width</code>	See details. Defaults to 5.
<code>undo.splits</code>	See details. Defaults to 'sdundo'.
<code>undo.SD</code>	See details. Defaults to 2.2.
<code>verbose</code>	See details. Defaults to 0.

**Details**

This method is a wrapper of the CNA, segment, segments.summary and segments.p methods of the DNACopy package. Please refer to the respective man pages for more detailed information. The default parameters of CNV.segment override some of the default parameters of segment and are optimized for 450k data CNV analysis.

**Value**

CNV.analysis object.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```

# prepare
library(minfiData)
data(MsetEx)
d <- CNV.load(MsetEx)
data(detail_regions)
anno <- CNV.create_anno(detail_regions = detail_regions)

# create object
x <- CNV.fit(query = d['GroupB_1'], ref = d[c('GroupA_1', 'GroupA_2', 'GroupA_3')], anno)

# modify object
x <- CNV.bin(x)
x <- CNV.detail(x)
x <- CNV.segment(x)

# general information
x
show(x)

# coefficients of linear regression
coef(x)

# show or replace sample name
names(x)
names(x) <- 'Sample 1'

```

---

CNV.write

*CNV.write*


---

**Description**

Output CNV analysis results as table.

**Usage**

```
CNV.write(object, ...)
```

```
## S4 method for signature 'CNV.analysis'
CNV.write(object, file = NULL, what = "segments")
```

**Arguments**

object	CNV.analysis object.
...	Additional parameters (CNV.write generic, currently not used).
file	Path where output file should be written to. Defaults to NULL: No file is written, table is returned as data.frame object.
what	character. This should be (an unambiguous abbreviation of) one of 'probes', 'bins', 'detail' or 'segments'. Defaults to 'segments'.

**Value**

if parameter file is not supplied, the table is returned as a data.frame object.

**Examples**

```
# prepare
library(minfiData)
data(MsetEx)
d <- CNV.load(MsetEx)
data(detail_regions)
anno <- CNV.create_anno(detail_regions = detail_regions)

# create/modify object
x <- CNV.segment(CNV.detail(CNV.bin(CNV.fit(query = d['GroupB_1'],
  ref = d[c('GroupA_1', 'GroupA_2', 'GroupA_3')], anno))))

# output plots
CNV.genomeplot(x)
CNV.genomeplot(x, chr = 'chr6')
CNV.detailplot(x, name = 'PTEN')
CNV.detailplot_wrap(x)

# output text files
CNV.write(x, what = 'segments')
CNV.write(x, what = 'detail')
CNV.write(x, what = 'bins')
CNV.write(x, what = 'probes')
```

---

detail\_regions

*detail\_regions*

---

**Description**

Example of genomic regions to be analyzed in detail (e.g. candidate oncogenes/TSGs).

**Details**

Imported using rtracklayer. Raw data stored in inst/extdata/detail\_regions.bed.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

---

exclude_regions	<i>exclude_regions</i>
-----------------	------------------------

---

**Description**

Example of genomic regions to exclude (e.g. known polymorphic regions).

**Details**

Imported using `rtracklayer`. Raw data stored in `inst/extdata/exclude_regions.bed`.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

---

read.450k.url	<i>read.450k.url</i>
---------------	----------------------

---

**Description**

Read IDAT files from the web.

**Usage**

```
read.450k.url(url = NULL, idat = NULL)
```

**Arguments**

<code>url</code>	URL of the directory in which the IDAT files are located.
<code>idat</code>	Vector of IDAT names. <code>url</code> and <code>idat</code> default to the TCGA example described in the vignette.

**Details**

This method downloads the provided list of IDAT files to a temporary folder (using the `RCurl` package). It then uses the `'read.450k.exp'` method of the `'minfi'` package.

**Value**

`RGChannelSet` object.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

**Examples**

```
RGsetTCGA <- read.450k.url()
```

---

tbl_ucsc	<i>tbl_ucsc</i>
----------	-----------------

---

**Description**

UCSC tables required for creating annotation object.

**Details**

Imported using `rtracklayer::browserSession('UCSC'):chromInfo, gap, cytoBand`.

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

---

tcgaBRCA.sentrrix2name	<i>tcgaBRCA.sentrrix2name</i>
------------------------	-------------------------------

---

**Description**

Named vector for Sentrrix ID to TCGA ID conversion of breast cancer example data (see README).

**Details**

Based on [https://tcga-data.nci.nih.gov/tcgafiles/ftp\\_auth/distro\\_ftpusers/anonymous/tumor/brca/cgcc/](https://tcga-data.nci.nih.gov/tcgafiles/ftp_auth/distro_ftpusers/anonymous/tumor/brca/cgcc/)

**Author(s)**

Volker Hovestadt <conumee@hovestadt.bio>

# Index

[, CNV.data, ANY, ANY-method  
(CNV.data-class), 8

CNV.analysis-class, 2  
CNV.anno-class, 3  
CNV.bin, 4  
CNV.bin, CNV.analysis-method (CNV.bin), 4  
CNV.check, 6  
CNV.check, CNV.data-method (CNV.check), 6  
CNV.create\_anno, 6  
CNV.create\_bins, 7  
CNV.data-class, 8  
CNV.detail, 9  
CNV.detail, CNV.analysis-method  
(CNV.detail), 9  
CNV.detailplot, 10  
CNV.detailplot, CNV.analysis-method  
(CNV.detailplot), 10  
CNV.detailplot\_wrap, 12  
CNV.detailplot\_wrap, CNV.analysis-method  
(CNV.detailplot\_wrap), 12  
CNV.fit, 13  
CNV.fit, CNV.data, CNV.data, CNV.anno-method  
(CNV.fit), 13  
CNV.genomeplot, 14  
CNV.genomeplot, CNV.analysis-method  
(CNV.genomeplot), 14  
CNV.load, 16  
CNV.load, data.frame-method (CNV.load),  
16  
CNV.load, matrix-method (CNV.load), 16  
CNV.load, MethylSet-method (CNV.load), 16  
CNV.load, numeric-method (CNV.load), 16  
CNV.merge\_bins, 17  
CNV.segment, 18  
CNV.segment, CNV.analysis-method  
(CNV.segment), 18  
CNV.write, 19  
CNV.write, CNV.analysis-method  
(CNV.write), 19

coef, CNV.analysis-method  
(CNV.analysis-class), 2

detail\_regions, 20

exclude\_regions, 21

names, CNV.analysis-method  
(CNV.analysis-class), 2  
names, CNV.data-method (CNV.data-class),  
8  
names<-, CNV.analysis-method  
(CNV.analysis-class), 2  
names<-, CNV.data-method  
(CNV.data-class), 8

plot, CNV.analysis, ANY-method  
(CNV.genomeplot), 14

read.450k.url, 21

show, CNV.analysis-method  
(CNV.analysis-class), 2  
show, CNV.anno-method (CNV.anno-class), 3  
show, CNV.data-method (CNV.data-class), 8

tbl\_ucsc, 22  
tcgaBRCA.sentrax2name, 22