

# Package ‘SIM’

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**Type** Package

**Title** Integrated Analysis on two human genomic datasets

**Version** 1.81.0

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**Imports** graphics, stats, globaltest, quantsmooth

**Depends** R (>= 3.5), quantreg

**Suggests** biomaRt, RColorBrewer

**Description** Finds associations between two human genomic datasets.

**License** GPL (>= 2)

**biocViews** Microarray, Visualization

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SIM-package

*Statistical Integration of Microarrays*

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## Description

SIM is a statistical model to identify associations between two genomic datasets. Where one is assigned as dependent variable and the other as independent e.g. copy number measurements on several samples versus expression measurements on the same samples. A region of interest can be chosen to run the integrated analysis on either the same region for both dependent and independent datasets or different regions. For each dependent feature a P-value measures the association with the independent data, the contribution of each independent feature is given as Z-scores. The integrated analysis is based on the random-effect model for gene-sets as implemented in [gt](#).

maybe something about annotation?

By default we use `method.adjust = "BY"` (Benjamini-Yekutieli) for multiple testing correction. This method accounts for dependence between measurements and is more conservative than "BH" (Benjamini-Hochberg). For details on the multiple testing correction methods see [p.adjust](#). We have experienced that a rather low stringency cut-off on the BY-values of 20% allows the detection of associations for data with a low number of samples or a low frequency of aberrations. False positives are rarely observed.

Make sure that the array probes are mapped to the same builds of the genome, and that the [chrom.table](#) used by the [integrated.analysis](#) is from the same build as well. See [sim.update.chrom.table](#).

## Details

Package: SIM  
 Type: Package  
 Version: 1.7.1  
 Date: 2010-09-14  
 License: Open

**Author(s)**

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <R.X.Menezes@lumc.nl>

**References**

Menezes RX, Boetzer M, Sieswerda M, van Ommen GJ, Boer JM (2009). Integrated analysis of DNA copy number and gene expression microarray data using gene sets. *BMC Bioinformatics*, **10**, 203-.

Goeman JJ, van de Geer SA, de Kort F, van Houwelingen HC (2004). A global test for groups of genes: testing association with a clinical outcome. *Bioinformatics*, **20**, 93-109.

**See Also**

[assemble.data](#), [integrated.analysis](#), [sim.plot.zscore.heatmap](#), [sim.plot.pvals.on.region](#), [sim.plot.pvals.on.genome](#), [tabulate.pvals](#), [tabulate.top.dep.features](#), [tabulate.top.indep.features](#), [impute.nas.by.surrounding](#), [sim.update.chrom.table](#), [sim.plot.overlapping.indep.dep.features](#), [getoverlappingregions](#)

**Examples**

```
#load the datasets and the samples to run the integrated analysis
data(expr.data)
data(acgh.data)
data(samples)

#assemble the data
assemble.data(dep.data = acgh.data,
              indep.data = expr.data,
              dep.ann = colnames(acgh.data)[1:4],
              indep.ann = colnames(expr.data)[1:4],
              dep.id="ID",
              dep.chr = "CHROMOSOME",
              dep.pos = "STARTPOS",
              dep.symb="Symbol",
              indep.id="ID",
              indep.chr = "CHROMOSOME",
              indep.pos = "STARTPOS",
              indep.symb="Symbol",
              overwrite = TRUE,
              run.name = "chr8q")

#run the integrated analysis
integrated.analysis(samples = samples,
                   input.regions = "8q",
                   zscores=TRUE,
                   run.name = "chr8q")

# use functions to plot the results of the integrated analysis

#plot the P-values along the genome
sim.plot.pvals.on.genome(input.regions = "8q",
                        significance = c(0.2, 0.05),
```

```
        adjust.method = "BY",
        pdf = FALSE,
        run.name = "chr8q")

#plot the P-values along the regions
sim.plot.pvals.on.region(input.regions = "8q",
                        adjust.method="BY",
                        run.name = "chr8q")

#plot the z-scores in an association heatmap
#plot the zscores in a heatmap
sim.plot.zscore.heatmap(input.regions = "8q",
                        method="full",
                        significance=0.2,
                        z.threshold=3,
                        show.names.indep=TRUE,
                        show.names.dep=TRUE,
                        adjust.method = "BY",
                        add.plot = "smooth",
                        smooth.lambda = 2,
                        pdf = FALSE,
                        run.name = "chr8q")

sim.plot.zscore.heatmap(input.regions = "8q",
                        method="full",
                        significance = 0.05,
                        z.threshold = 1,
                        show.names.indep=TRUE,
                        show.names.dep=FALSE,
                        adjust.method = "BY",
                        add.plot = "heatmap",
                        smooth.lambda = 2,
                        pdf = FALSE,
                        run.name = "chr8q")

sim.plot.zscore.heatmap(input.regions = "8q",
                        method="full",
                        significance = 0.05,
                        z.threshold = 1,
                        show.names.indep=TRUE,
                        show.names.dep=TRUE,
                        adjust.method = "BY",
                        add.plot = "none",
                        pdf = FALSE,
                        run.name = "chr8q")

#tabulate the P-values per region (prints to screen)
tabulate.pvals(input.regions = "8q",
              adjust.method="BY",
              bins=c(0.001,0.005,0.01,0.025,0.05,0.075,0.10,0.20,1.0),
              run.name = "chr8q")

table.dep <- tabulate.top.dep.features(input.regions="8q",
```

```

        adjust.method="BY",
        method="full",
        significance=0.05,
        run.name="chr8q")
head(table.dep[["8q"]])

table.indep <- tabulate.top.indep.features(input.regions="8q",
        adjust.method="BY",
        method="full",
        significance= 0.05,
        z.threshold=c(-1, 1),
        run.name="chr8q")
head(table.indep[["8q"]])

sim.plot.overlapping.indep.dep.features(input.regions="8q",
        adjust.method="BY",
        significance=0.1,
        z.threshold= c(-1,1),
        log=TRUE,
        summarize="consecutive",
        pdf=FALSE,
        method="full",
        run.name="chr8q")

```

---

acgh.data

*Array Comparative Genomic Hybridization data*


---

## Description

Copy number data taken from Pollack et al. PNAS. 2002, 99(20): 12963-8.

## Usage

```
data(acgh.data)
```

## Format

A data frame with 99 observations on 45 variables. The first 4 columns are the unique identifier, symbol for the chromosome and start position of the probe the next 41 columns are the copy number measurements of 41 samples.

## Details

A subset of the original data is taken namely all the probes on the long arm of chromosome 8.

## Source

[http://genome-www.stanford.edu/aCGH\\_breast/data.shtml](http://genome-www.stanford.edu/aCGH_breast/data.shtml)

## References

Pollack JR, Sorlie T, Perou CM, Rees CA, Jeffrey SS, Lonning PE, Tibshirani R, Botstein D, Borresen-Dale AL, Brown PO (2002). Microarray analysis reveals a major direct role of DNA copy number alteration in the transcriptional program of human breast tumors. *Proc Natl Acad Sci USA*. **1**, 99(20), 12963-8.

## Examples

```
data(acgh.data)
```

---

assemble.data	<i>Assemble the data to run the integrated analysis</i>
---------------	---

---

## Description

Assembles the dependent and independent data and annotation of the both data sets.

## Usage

```
assemble.data(dep.data,
  indep.data,
  dep.id = "ID",
  dep.chr = "CHROMOSOME",
  dep.pos = "STARTPOS",
  dep.ann = NULL,
  dep.symb,
  indep.id = "ID",
  indep.chr = "CHROMOSOME",
  indep.pos = "STARTPOS",
  indep.ann = NULL,
  indep.symb,
  overwrite = FALSE,
  run.name = "analysis_results")
```

## Arguments

dep.data	The dependent data ( <code>data.frame</code> ), along with annotations. Each row should correspond to one feature. The following columns are expected to exist, and the column names should be inserted in the function. <code>dep.id</code> : A unique identifier. <code>dep.chr</code> : The number of the chromosome (1,2, ..., 22, X, Y). <code>dep.pos</code> : The base pair position, relative to the chromosome. <code>dep.symb</code> : Gene symbol (optional). <code>dep.ann</code> : Annotation can be multiple columns.
indep.data	<code>data.frame</code> The independent data, along with annotations. Each row should correspond to one feature. The following columns are expected to exist, and the column names should be inserted in the function. <code>indep.id</code> : A unique identifier. <code>indep.chr</code> : The number of the chromosome (1,2, ..., 22, X, Y). <code>indep.pos</code> : The base pair position, relative to the chromosome. <code>indep.symb</code> : Gene symbol (optional). <code>indep.ann</code> : Annotation can be multiple columns.

dep.ann	vector with either the names of the columns or the column numbers in the dependent data that contain the annotation.
indep.ann	vector with either the names of the columns or the column numbers in the independent data that contain the annotation.
dep.id	vector with the column name in the dependent data that contains the ID. Will be used in the <a href="#">sim.plot.zscore.heatmap</a> function. Empty ID's will be substituted by NA.
dep.chr	vector with column name in the dependent data that contains the chromosome numbers.
dep.pos	vector with the column name in the dependent data that contains the position on the chromosome in bases.
dep.symb	Optional, either missing or a single vector with the column name in the dependent data that contains the symbols. Will be used in <a href="#">sim.plot.zscore.heatmap</a> as label.
indep.id	vector with the column name in the independent data that contains the ID. Will be used in the <a href="#">sim.plot.zscore.heatmap</a> function. Empty ID's will be substituted by NA.
indep.chr	vector with the column name in the independent data that contains the chromosome numbers.
indep.pos	vector with the column name in the independent data that contains the position on the chromosome in bases.
indep.symb	Optional, either missing or a vector with the column name in the dependent data that contains the Symbols. Will be used in <a href="#">sim.plot.zscore.heatmap</a> as label.
overwrite	logical, indicate when a run.name is already present, the results can be overwritten.
run.name	Name of the analysis. The results will be stored in a folder with this name in the current working directory (use <code>getwd()</code> to print the current working directory). If the missing, the default folder "analysis_results" will be generated.

### Details

Based on the chromosome and probe position an absolute position is calculated according to  $chromosomenumber * 1e9 + probeposition$ . Chromosome column is converted to factor and relevelled according to the levels of the [chrom.table](#), so the only levels allowed are `c(1:22, "X", "Y")`. Currently only human genome support without mitochondrial DNA.

### Value

No values are returned. Instead, the datasets and annotation columns are stored in separate files in the data folder in the directory specified in `run.name`. If `assemble.data` has run successfully, the [integrated.analysis](#) function can be performed.

### Author(s)

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <[R.X.Menezes@lumc.nl](mailto:R.X.Menezes@lumc.nl)>

**See Also**

[SIM, integrated.analysis](#)

**Examples**

```
# Generate datasets and the samples to run the integrated analysis
set.seed(53245)
ngenes <- 100
nsamples <- 100
# generate copy number measurements
x <- matrix(rnorm(n = ngenes*nsamples), nrow = ngenes, ncol = nsamples)
# add mean shift effect for half of the samples, copy gain for 2nd half of the genes
x[ seq_len(ngenes/2), seq_len(nsamples/2)] <- x[ seq_len(ngenes/2), seq_len(nsamples/2)] + 2
# generate gene expression with normal distribution and mean equal to gene copy number
y <- rnorm(n = ngenes*nsamples, mean = matrix(x, nrow = ngenes*nsamples, ncol = 1), sd = 0.8)
y <- matrix(y, nrow = ngenes, ncol = nsamples)
samples <- paste0("S", seq_len(nsamples))
colnames(x) <- colnames(y) <- samples
# Making data objects
acgh.data <- data.frame(ID = paste0("G", seq_len(ngenes)),
                       CHROMOSOME = rep(1, ngenes),
                       STARTPOS = seq_len(ngenes)*12*10^5,
                       Symbol = paste0("Gene", seq_len(ngenes)),
                       x)
expr.data <- data.frame(ID = paste0("G", seq_len(ngenes)),
                       CHROMOSOME = rep(1, ngenes),
                       STARTPOS = seq_len(ngenes)*12*10^5,
                       Symbol = paste0("Gene", seq_len(ngenes)),
                       y)

#assemble the data
assemble.data(dep.data = acgh.data,
              indep.data = expr.data,
              dep.ann = colnames(acgh.data)[1:4],
              indep.ann = colnames(expr.data)[1:4],
              dep.id="ID",
              dep.chr = "CHROMOSOME",
              dep.pos = "STARTPOS",
              dep.symb="Symbol",
              indep.id="ID",
              indep.chr = "CHROMOSOME",
              indep.pos = "STARTPOS",
              indep.symb="Symbol",
              overwrite = TRUE,
              run.name = "chr1p")
```

---

chrom.table

*Table with chromosome information*

---

**Description**

A table indicating the base positions of the beginning and end of chromosome arms and bands. Currently based on the UCSC March 2006/NCBI 36 build of the human genome.

**Usage**

```
data(chrom.table)
```

**Format**

A data frame with 862 observations on the following 6 variables.  
chr, arm, band, start, end, stain

**Details**

Possibly the chrom.table can be update by [sim.update.chrom.table](#). Currently only human genome support without mitochondrial DNA.

**See Also**

[sim.update.chrom.table](#)

**Examples**

```
data(chrom.table)
```

---

expr.data

*Expression data example*

---

**Description**

Expression data taken from Pollack et al. PNAS. 2002, 99(20): 12963-8.

**Usage**

```
data(expr.data)
```

**Format**

A data frame with 99 observations on 45 variables. The first 4 columns are the unique identifier, symbol for the chromosome and start position of the probe the next 41 columns are the expression log-ratios of 41 samples.

**Details**

A subset of the original data is taken namely all the probes on the long arm of chromosome 8.

**Source**

[http://genome-www.stanford.edu/aCGH\\_breast/data.shtml](http://genome-www.stanford.edu/aCGH_breast/data.shtml)

**References**

Pollack JR, Sorlie T, Perou CM, Rees CA, Jeffrey SS, Lonning PE, Tibshirani R, Botstein D, Borresen-Dale AL, Brown PO (2002). Microarray analysis reveals a major direct role of DNA copy number alteration in the transcriptional program of human breast tumors. *Proc Natl Acad Sci USA*. **1**, 99(20), 12963-8.

**Examples**

```
data(expr.data)
```

---

getoverlappingregions *Get the overlapping regions between independent and dependent regions*

---

**Description**

Generates a table with overlapping regions.

**Usage**

```
getoverlappingregions(independent_regions,  
                      dependent_regions,  
                      method = c("union", "overlapping"))
```

**Arguments**

`independent_regions` `data.frame()`. Independent regions found with tab [tabulate.top.indep.features](#).

`dependent_regions` `data.frame()`. Independent regions found with tab [tabulate.top.dep.features](#).

`method` method to estimate the overlapping regions, either "union" or "overlapping". overlapping outputs only the overlapping parts of the overlapping regions. union outputs the whole region. Say independent region = 1-10, dependent region = 5-12. The output is 1-12.

**Details**

Calculates the overlap between two tables.

**Author(s)**

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <[R.X.Menezes@lumc.nl](mailto:R.X.Menezes@lumc.nl)>

**See Also**

[SIM](#), [tabulate.top.dep.features](#), [tabulate.top.indep.features](#), [sim.plot.overlapping.indep.dep.features](#)

**Examples**

```
#no examples yet!
```

---

```
impute.nas.by.surrounding  
    Impute NA's in array-CGH data
```

---

**Description**

Replace an NA by the median of the surrounding features in the same sample.

**Usage**

```
impute.nas.by.surrounding(dataset,  
                           window.size = 5)
```

**Arguments**

dataset	<a href="#">data.frame</a> with the dataset to replace the NA's by the medians of the surrounding features.
window.size	numeric value, specifying of how many features around the NA the median should be taken.

**Details**

This function can be used when the dependent dataset in the [integrated.analysis](#) function is array-CGH data and contains probes that have an NA. To avoid losing data by throwing away the probes with NA's, the `impute.nas.by.surrounding` function can be used which simply takes the median of the probes around an NA. The number of probes used for the imputation is chosen by giving a value for `window.size`. This script takes quite long to run!

**Value**

A `data.frame` is returned, containing the inserted "dataset" all NA replaced with median of the window of size "window.size" around the NA.

**Author(s)**

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <[R.X.Menezes@lumc.nl](mailto:R.X.Menezes@lumc.nl)>

**See Also**

[SIM](#), [assemble.data](#), [integrated.analysis](#)

**Examples**

```
#no examples yet!
```

---

```
integrated.analysis Integrated analysis of dependent and independent microarray data
```

---

**Description**

Runs the Integrated Analysis to test for associations between dependent and independent microarray data on the same set of samples.

**Usage**

```
integrated.analysis(samples,
                    input.regions = "all chrs",
                    input.region.indep = NULL,
                    zscores = FALSE,
                    method = c("full", "smooth", "window", "overlap"),
                    dep.end = 1e5,
                    window = c(1e6, 1e6),
                    smooth.lambda=2,
                    adjust = ~1,
                    run.name = "analysis_results",
                    ...)
```

**Arguments**

<code>samples</code>	vector with either the names of the columns in the dependent and independent data corresponding to the samples, or a numerical vector containing the column numbers to include in the analysis, e.g. 5:10 means columns 5 till 10. Make sure that both datasets have the same number of samples with the same column names!
<code>input.regions</code>	vector indicating the dependent regions to be analyzed. Can be defined in four ways: 1) predefined input region: insert a predefined input region, choices are: "all chrs", "all chrs auto", "all arms", "all arms auto" In the predefined regions "all arms" and "all arms auto" the arms 13p, 14p, 15p, 21p and 22p are left out, because in most studies there are no or few probes in these regions. To include them, just make your own vector of arms. 2) whole chromosome(s): insert a single chromosome or a list of chromosomes as a vector: c(1, 2, 3). 3) chromosome arms: insert a single chromosome arm or a list of chromosome arms like c("1q", "2p", "2q"). 4) subregions of a chromosome: insert a chromosome number followed by the start and end position like "chr1:1-1000000" These regions can also be combined, e.g. c("chr1:1-1000000", "2q", 3). See details for more information.

<code>input.region.indep</code>	indicating the independent region which will be analysed in combination of the dependent region. Only one input region can given using the same format as the dependent input region.
<code>zscores</code>	logical indicates whether the Z-scores are calculated (takes longer time to run). If <code>zscores = FALSE</code> , only P-values are calculated.
<code>method</code>	either one of “full”, “window”, “overlap” or “smooth”. This defines how the data is used for the <code>integrated.analysis</code> . <code>full</code> : the whole dependent data region is taken. <code>window</code> : takes the middle of the dependent probe and does the integration on the independent probes that are within the window given at <code>window-size</code> given by <code>window</code> . <code>overlap</code> : does the integration on the independent probes that are within the start and end of the dependent probes given at <code>dep.end</code> . <code>smooth</code> : does smooth on the dependent probes with smoothing factor given at <code>smooth.lambda</code> , finds the value of smooth for each independent probe and does the integration on them. Only needed when <code>method = "smooth"</code> , default <code>smooth.lambda = 2</code>
<code>dep.end</code>	numeric or character either the name of the column “end” in the dependent data or, when not available, an numeric value which indicates the end deviating from the start. When a numeric value is inserted, the function will do: $start + dep.end = end$ . Only needed when <code>method = "window"</code> or “overlap”.
<code>window</code>	numeric values. Window to search for overlapping independent features per dependent probe. First value is the number of positions to the left from the middle of the probe, the second value is the number of positions to the right from the middle of the probe. Only needed when <code>method = "window"</code> .
<code>smooth.lambda</code>	numeric factor used for smoothing the dependent data. Only needed when <code>method = "smooth"</code> . See <a href="#">quantsmooth</a> for more information. By default the <code>segment = min(nrow(dep.data), 100)</code> .
<code>adjust</code>	formula a formula like <code>~gender</code> , where <code>gender</code> is a vector of the same size as samples. The regression models is correct for the gender effect, see <a href="#">gt</a> .
<code>run.name</code>	character name of the analysis. The results will be stored in a folder with this name in the current working directory (use <code>getwd()</code> to print the current working directory). If missing the default folder “analysis\_results” will be generated.
<code>...</code>	additional arguments for <a href="#">gt</a> e.g. <code>model="logistic"</code> or when <code>permutations &gt; 0</code> the null distribution is estimated using permutations, see <a href="#">gt</a> . See Details.

## Details

The Integrated Analysis is a regression of the independent data on the dependent features. The regression itself is done using the [gt](#), which means that the genes in a region (e.g. a chromosome arm) are tested as a gene set. The individual associations between each dependent and each independent feature are calculated as Z-scores (standardized influences, see [?gt](#)).

This function splits the datasets into separate sets for each region (as specified by the `input.regions`) and runs the analysis for each region separately.

When running the Integrated Analysis for a predefined input region, like “all arms” and “all chrs”, output can be obtained for all input regions, as well as subsets of it. But note that the genomic

unit must be the same: if `integrated.analysis` was run using chromosomes as units, any of the functions and plots must also use chromosomes as units, and not chromosome arms. Similarly, if `integrated.analysis` was run using chromosome arms as units, these units must also be used to produce plots and outputs. For example if the `input.regions = "all arms"` was used, P-value plots (see [sim.plot.pvals.on.region](#)) can be produced by inserting the `input.regions = "all arms"`, but also for instance "1p" or "20q". However, to produce a plot of the whole chromosome, for example chromosome 1, the `integrated` should be re-run with `input.region=1`. The same goes for "all chrs": P-value plots etc. can be produced for chromosome 1,2 and so on... but to produce plots for an arm, the `integrated.analysis` should be re-run for that region. This also goes for subregions of the chromosome like "chr1:1-1000000".

By default the `gt` uses a "linear" model, only when the dependent data is a logical matrix containing TRUE and FALSE a "logistic" model is selected. All other models need `model = ""`, see `gt` for available models.

### Value

No values are returned. Instead, the results of the analysis are stored in the subdirectories of the directory specified in `run.name`. E.g. the z-score matrices are saved in subfolder `method`.

The following functions can be used to visualize the data:

- 1) [sim.plot.zscore.heatmap](#) (only possible when `zscores = TRUE`)
- 2) [sim.plot.pvals.on.region](#)
- 3) [sim.plot.pvals.on.genome](#)
- 4) [sim.plot.overlapping.indep.dep.features](#)

Other functions can be used to tabulate the results:

- 1) [tabulate.pvals](#)
- 2) [tabulate.top.dep.features](#)
- 3) [tabulate.top.indep.features](#) (only possible when `zscores = TRUE`)
- 4) [getoverlappingregions](#) (only possible when `tabulate.top.dep.features` and `tabulate.top.indep.features` were run).

### Author(s)

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <R.X.Menezes@lumc.nl>

### References

- Menezes RX, Boetzer M, Sieswerda M, van Ommen GJ, Boer JM (2009). Integrated analysis of DNA copy number and gene expression microarray data using gene sets. *BMC Bioinformatics*, **10**, 203-.
- Goeman JJ, van de Geer SA, de Kort F, van Houwelingen HC (2004). A global test for groups of genes: testing association with a clinical outcome. *Bioinformatics*, **20**, 93-109.

### See Also

`SIM`, [sim.plot.zscore.heatmap](#), [sim.plot.pvals.on.region](#), [sim.plot.pvals.on.genome](#), [tabulate.pvals](#), [tabulate.top.dep.features](#), [tabulate.top.indep.features](#), [getoverlappingregions](#), [sim.plot.overlapping.indep.dep.features](#), `gt`

**Examples**

```
#first run example(assemble.data)
data(samples)
#perform integrated analysis without Z-scores using the method = "full"
integrated.analysis(samples=samples,
  input.regions="8q",
  zscores=FALSE,
  method="full",
  run.name="chr8q")
```

---

link.metadata	<i>Link a metadata annotation file to expression ID</i>
---------------	---

---

**Description**

Get annotation out of a AnnotationData package and link them to the expression data using the expression probe ID's

**Usage**

```
link.metadata(data = expr.data,
  col.ID.link = 1,
  chr = as.list(hgu133plus2CHR),
  chrloc = as.list(hgu133plus2CHRLOC),
  symbol = as.list(hgu133plus2SYMBOL))
```

**Arguments**

data	data.frame with expression data including an expression probe ID column.
col.ID.link	numeric value, specifying the column of data that contains the ID to link with the poslist.
chr	list specifying the metadata annotation of the chromosome location on the genome.
chrloc	list specifying the metadata annotation of the location of the probe on the chromosome.
symbol	list specifying the metadata annotation of the symbol corresponding to the probe.

**Details**

Often, the annotation for expression array probes lack chromosome position information. Therefore, this function adds the two required columns to run the [integrated.analysis](#): "CHROMOSOME" and "STARTPOS". In addition, the optional column, "Symbol" is added.

**Value**

A data.frame is returned, containing a dataset with annotation columns which can be used for [integrated.analysis](#).

**Author(s)**

Marten Boetzer, Melle Sieswerda, Renee x Menezes <R.X.Menezes@lumc.nl>

**See Also**

[RESOURCERER.annotation.to.ID](#)

**Examples**

```
# first download and install the AnnotationData package for your expression array platform
# for example
## Not run: library(hgu133plus2)
## Not run: expr.data <- link.metadata(data, col.ID.link = 1, chr = as.list(hgu133plus2CHR),
chrloc = as.list(hgu133plus2CHRLoc), symbol = as.list(hgu133plus2SYMBOL))
## End(Not run)
```

---

RESOURCERER.annotation.to.ID

*Link RESOURCERER annotation file to expression ID*

---

**Description**

Get annotation out of the RESOURCERER annotation file and link them to expression data with help of expression ID's

**Usage**

```
RESOURCERER.annotation.to.ID(data, poslist, col.ID.link = 1, col.poslist.link = 1)
```

**Arguments**

data	<a href="#">data.frame</a> with expression data including an expression ID column.
poslist	<a href="#">data.frame</a> containing the RESOURCERER annotation file
col.ID.link	numeric value, specifying the column of data that contains the ID to link with the poslist.
col.poslist.link	numeric value, specifying the column of poslist that contains the ID to link with the data.

**Details**

This function will output the inserted dataset, including the necessary, for [integrated.analysis](#), a nnotation columns: "CHROMOSOME", "STARTPOS"and "Symbol" out of the inserted RESOURCE RER annotation file poslist.

**Value**

A `data.frame` is returned, containing a dataset with annotation columns which can be used for [integrated.analysis](#)

**Author(s)**

Marten Boetzer, Melle Sieswerda, Renee x Menezes <R.X.Menezes@lumc.nl>

**See Also**

[link.metadata](#)

**Examples**

```
# download expression array annotation from RESOURCERER
# ftp://occams.dfc.harvard.edu/pub/bio/tgi/data/Resourcerer
# it may be necessary to remove the first row, which states the genome build used for mapping
## Not run: read.an <- read.delim("affy_U133Plus2.txt", sep="\t", header=T)

# get physical mapping columns
## Not run: expr.data <- RESOURCERER_annotation_to_ID(data = read.expr, poslist = read.an, col.ID.link = 1, col.pos
```

---

samples

*Samples for example data*

---

**Description**

Vector of sample names corresponding to the column headers containing the data in both the copy number (`acgh.data`) and expression (`expr.data`) example datasets.

**Usage**

```
data(samples)
```

**Format**

A character vector.

**Source**

[http://genome-www.stanford.edu/aCGH\\_breast/data.shtml](http://genome-www.stanford.edu/aCGH_breast/data.shtml)

**References**

Pollack JR, Sorlie T, Perou CM, Rees CA, Jeffrey SS, Lonning PE, Tibshirani R, Botstein D, Borresen-Dale AL, Brown PO (2002). Microarray analysis reveals a major direct role of DNA copy number alteration in the transcriptional program of human breast tumors. *Proc Natl Acad Sci USA*. **1**, 99(20), 12963-8.

**Examples**

```
data(samples)
```

---

```
sim.plot.overlapping.indep.dep.features
```

*P-value plot and mean-zscore plots with indication of overlapping features.*

---

**Description**

Generates three plots: The first plot contains the P-values along the region, with the cut-off displayed. The second plot contains the mean-zscores along the region, with the cut-offs displayed. The third plots generates the cytobands of the region.

**Usage**

```
sim.plot.overlapping.indep.dep.features(input.regions,
input.region.indep = NULL,
adjust.method = "BY",
log = FALSE,
significance = 0.2,
max.pow = 5,
z.threshold = c(-3,3),
summarize = c("consecutive", "stretch", "window"),
stretch = 10,
window = 1e6,
percentage = 0.5,
xlim=NULL,
pdf = FALSE,
method = c("full", "smooth", "window", "overlap"),
run.name = "analysis_results", ...)
```

**Arguments**

**input.regions** vector indicating the dependent regions to be analyzed. Can be defined in four ways: 1) predefined input region: insert a predefined input region, choices are: "all chrs", "all chrs auto", "all arms", "all arms auto" In the predefined regions "all arms" and "all arms auto" the arms 13p, 14p, 15p, 21p and 22p are left out, because in most studies there are no or few probes in these regions. To include them, just make your own vector of arms. 2) whole chromosome(s): insert a single chromosome or a list of chromosomes as a vector: c(1, 2, 3). 3) chromosome arms: insert a single chromosome arm or a list of chromosome arms like c("1q", "2p", "2q"). 4) subregions of a chromosome: insert a chromosome number followed by the start and end position like "chr1:1-1000000" These regions can also be combined, e.g. c("chr1:1-1000000", "2q", 3). See [integrated.analysis](#) for more information.

input.region.indep	indicating the independent region which will be analysed in combination of the dependent region. Only one input region can given using the same format as the dependent input region.
adjust.method	Method used to adjust the P-values for multiple testing, see <a href="#">p.adjust</a> . Default is “BY” recommended when copy number is used as dependent data. See <a href="#">SIM</a> for more information about adjusting P-values.
log	logical default log = FALSE, if log = TRUE P-values are plotting on $\log_{10}$ scale.
significance	The threshold for selecting significant P-values.
max.pow	numeric only when log = TRUE scale of the y-axis.
z.threshold	Threshold to display a green or red bar in the color bar on top of the heatmap for independent features with mean z-scores above z.threshold (high positive association) or below -z.threshold (high negative association).
summarize	either one of “consecutive”, “stretch”, “window” which visualizes the subregions according to the selected summarization a) “consecutive” shows the consecutive significant regions b) “stretch” shows a percentage percentage significant stretch of size stretch and c) “window” a percentage percentage significant window of length window
stretch	integer length of stretch, default stretch = 10
window	integer length of window, default window = 1e6
percentage	numeric a number between [0, 1] given the percentage of significance in either the “stretch” or “window”
xlim	c(min, max) scale of the x-axis. Can be used for zooming in on a region.
pdf	logical indicate whether to generate a pdf of the plots in the current working directory or not.
method	this must be the either full, window, overlap or smooth but the data should generated by the same method in <code>integrated.analysis</code> .
run.name	This must be the same a given to <code>integrated.analysis</code>
...	not used in this version

**Details**

details: Cytobands plot adapted from SNPChip

**Value**

No values are returned. The results are stored in a subdirectory of `run.name` as `pdf`.

**Author(s)**

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <[R.X.Menezes@lumc.nl](mailto:R.X.Menezes@lumc.nl)>

**See Also**

[SIM](#), [tabulate.top.dep.features](#), [tabulate.top.indep.features](#), [getoverlappingregions](#)

**Examples**

```
#first run example(assemble.data)
#and example(integrated.analysis)
#overview plot of the dependent and independent features
sim.plot.overlapping.indep.dep.features(input.regions="8q",
                                       adjust.method="BY",
                                       significance=0.1,
                                       z.threshold= c(-1,1),
                                       log=TRUE,
                                       summarize="consecutive",
                                       pdf=FALSE,
                                       method="full",
                                       run.name="chr8q")
```

---

```
sim.plot.pvals.on.genome
```

*Plot the P-values in whole genome overview*

---

**Description**

Generates a plot of the analyzed dependent data probe positions and their significance on all chromosomes.

**Usage**

```
sim.plot.pvals.on.genome(input.regions = "all chrs",
                         significance = c(0.05, 0.20),
                         adjust.method = "BY",
                         method = c("full", "smooth", "window", "overlap"),
                         run.name = "analysis_results",
                         pdf = TRUE,
                         main = "Significantly associated features",
                         ylab = "Chromosomes",
                         ann=par("ann"),
                         ...)
```

**Arguments**

**input.regions** vector indicating the dependent regions to be analyzed. Can be defined in four ways: 1) predefined input region: insert a predefined input region, choices are: "all chrs", "all chrs auto", "all arms", "all arms auto" In the predefined regions "all arms" and "all arms auto" the arms 13p, 14p, 15p, 21p and 22p are left out, because in most studies there are no or few probes in these regions. To include them, just make your own vector of arms. 2) whole

	chromosome(s): insert a single chromosome or a list of chromosomes as a vector: <code>c(1, 2, 3)</code> . 3) chromosome arms: insert a single chromosome arm or a list of chromosome arms like <code>c("1q", "2p", "2q")</code> . 4) subregions of a chromosome: insert a chromosome number followed by the start and end position like <code>"chr1:1-1000000"</code> These regions can also be combined, e.g. <code>c("chr1:1-1000000", "2q", 3)</code> . See <a href="#">integrated.analysis</a> for more information.
significance	Two values that categorize the P-values on the selected chromosomes. These margins are indicated by different colors shown in the legend. These values can be defined, e.g. <code>pval.sig = c(0.3, 0.075)</code>
adjust.method	Method used to adjust the P-values for multiple testing. see <a href="#">p.adjust</a> Default is "BY" recommended when copy number is used as dependent data. See <a href="#">SIM</a> for more information about adjusting P-values.
method	this must be the either full, window, overlap or smooth but the data should generated by the same method in <code>integrated.analysis</code> .
pdf	Boolean. Indicate whether to generate a pdf of the plots in the current working directory or not.
run.name	This must be the same a given to <code>integrated.analysis</code>
main	the usual graphical parameter for the caption of the plot.
ylab	the usual graphical parameter for the y-axis label of the plot.
ann	the usual graphical parameter for annotation of the plot.
...	Arguments to be passed to pdf when pdf = TRUE, see Details.

### Details

Grey vertical lines indicate insignificant probes on top the significant ones are plotted. A purple dot indicates the centromere and a orange line the input region.

Sometimes it is useful to make the genome-plot as A4 landscape-format, add the following parameters to the `sim.plot.pvals.on.genome(..., paper='a4r', width=0, height=0)`

### Value

No values are returned. The results are stored in the folder "pvalue.plots" in directory `run.name` as pdf.

### Author(s)

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <[R.X.Menezes@lumc.nl](mailto:R.X.Menezes@lumc.nl)>

### See Also

[SIM](#), [sim.plot.zscore.heatmap](#), [sim.plot.pvals.on.region](#)

## Examples

```
#first run example(assemble.data)
#and example(integrated.analysis)
#plot the p-values along the genome
sim.plot.pvals.on.genome(input.regions="8q",
                        significance=c(0.05, 0.005),
                        adjust.method="BY",
                        method="full",
                        pdf=FALSE,
                        run.name="chr8q")
```

---

```
sim.plot.pvals.on.region
```

*P-value histograms and P-values along the genome per region*

---

## Description

Generates two plots of the P-values for an analyzed region. The first plot contains the distribution of the raw P-values and ranked plots of the raw and adjusted P-values. The second plot contains the P-values along the genome of analyzed input regions.

## Usage

```
sim.plot.pvals.on.region(input.regions = c("all chrs"),
                        significance = 0.2,
                        adjust.method = "BY",
                        method = c("full", "smooth", "window", "overlap"),
                        run.name = "analysis_results", ...)
```

## Arguments

**input.regions** vector indicating the dependent regions to be analyzed. Can be defined in four ways: 1) predefined input region: insert a predefined input region, choices are: "all chrs", "all chrs auto", "all arms", "all arms auto" In the predefined regions "all arms" and "all arms auto" the arms 13p, 14p, 15p, 21p and 22p are left out, because in most studies there are no or few probes in these regions. To include them, just make your own vector of arms. 2) whole chromosome(s): insert a single chromosome or a list of chromosomes as a vector: c(1, 2, 3). 3) chromosome arms: insert a single chromosome arm or a list of chromosome arms like c("1q", "2p", "2q"). 4) subregions of a chromosome: insert a chromosome number followed by the start and end position like "chr1:1-1000000" These regions can also be combined, e.g. c("chr1:1-1000000", "2q", 3). See [integrated.analysis](#) for more information.

**significance** The threshold for selecting significant P-values.

**adjust.method** Method used to adjust the P-values for multiple testing. see [p.adjust](#) Default is "BY" recommended when copy number is used as dependent data. See [SIM](#) for more information about adjusting P-values.

method	this must be the either full, window, overlap or smooth but the data should generated by the same method in <code>integrated.analysis</code> .
run.name	This must be the same a given to <code>integrated.analysis</code>
...	Arguments to be passed to pdf.

### Details

This function returns a pdf containing the P-value plots. The second plot contains the multiple testing corrected P-values plotted along the chromosome (arm). On the x-axis, the start positions of the dependent features are displayed. On the y-axis, the P-value levels are displayed. Two dotted lines indicate P-value levels 0.2 and 0.1. In general, P-values below 0.2 are said to be “significant”.

### Value

No values are returned. The results are stored in a subdirectory of `run.name` as pdf.

### Author(s)

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <R.X.Menezes@lumc.nl>

### See Also

[SIM](#), [sim.plot.pvals.on.genome](#)

### Examples

```
#first run example(assemble.data)
#and example(integrated.analysis)
#plot the p-values along the regions
sim.plot.pvals.on.region(input.regions="8q",
                        adjust.method="BY",
                        method="full",
                        run.name="chr8q")
```

---

sim.plot.zoom.in      *Zoom in on heatmap*

---

### Description

Zoom in on pervious produced heatmap by [sim.plot.zscore.heatmap](#)

### Usage

```
sim.plot.zoom.in(call)
```

### Arguments

call                    language, function call of [sim.plot.zscore.heatmap](#)

**Details**

[sim.plot.zscore.heatmap](#) returns (invisible) the function call and as attribute the local function environment. So by adjusting the the xlim and ylim a zoom in is created.

**Value**

an additional plot is created, on the previous plot a rectangle indicating the zoomed region.

**Author(s)**

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <R.X.Menezes@lumc.nl>

**See Also**

[SIM](#), [sim.plot.zscore.heatmap](#)

**Examples**

```
#first run example(assemble.data)
#and example(integrated.analysis)

#plot the zscores in a heatmap
sim.plot <- sim.plot.zscore.heatmap(input.regions = "8q",
                                   adjust.method = "BY",
                                   run.name = "chr8q", pdf = FALSE)

#only when runned interactive
if(interactive())
  sim.plot.zoom.in(sim.plot)
```

---

```
sim.plot.zscore.heatmap
```

*Association heatmap from z-scores*

---

**Description**

Produces an association heatmap that shows the association (standardized influence) of each independent feature (expression measurement) with each dependent feature (copy number measurement). A P-value bar on the left indicates test significance. A color bar on top indicates genes with mean z-scores across the significant copy number probes above a set threshold. A summary of the copy number data helps to identify what copy number alterations are present in a region of association with expression. Positive association can mean copy number gain and increased expression, or deletion and decreased expression. The heatmaps can also be used in an exploratory analysis, looking for very local effects of copy number changes (usually small amplifications) on gene expression, that do not lead to a significant test result.

**Usage**

```
sim.plot.zscore.heatmap(input.regions = "all chrs",
  input.region.indep = NULL,
  method = c("full", "smooth", "window", "overlap"),
  adjust = ~1,
  significance = 0.2,
  z.threshold = 3,
  colRamp = colorRampPalette(c("red", "black", "green")),
  add.colRamp = colorRampPalette(c("blue", "black", "yellow"))(7),
  show.names.indep = FALSE,
  show.names.dep = FALSE,
  adjust.method = "BY",
  scale,
  add.scale,
  add.plot = c("smooth", "none", "heatmap"),
  smooth.lambda = 2,
  pdf = TRUE,
  run.name = "analysis_results",...)
```

**Arguments**

- input.regions** vector indicating the dependent regions to be analyzed. Can be defined in four ways: 1) predefined input region: insert a predefined input region, choices are: "all chrs", "all chrs auto", "all arms", "all arms auto" In the predefined regions "all arms" and "all arms auto" the arms 13p, 14p, 15p, 21p and 22p are left out, because in most studies there are no or few probes in these regions. To include them, just make your own vector of arms. 2) whole chromosome(s): insert a single chromosome or a list of chromosomes as a vector: c(1, 2, 3). 3) chromosome arms: insert a single chromosome arm or a list of chromosome arms like c("1q", "2p", "2q"). 4) subregions of a chromosome: insert a chromosome number followed by the start and end position like "chr1:1-1000000" These regions can also be combined, e.g. c("chr1:1-1000000", "2q", 3). See [integrated.analysis](#) for more information.
- input.region.indep** indicating the independent region which will be analysed in combination of the dependent region. Only one input region can given using the same format as the dependent input region.
- method** this must be the either full, window, overlap or smooth but the data should generated by the same method in [integrated.analysis](#).
- adjust** This variable must be a vector with the same length as samples or FALSE. The vector will be transformed to a factor and the levels of this will be coloured according to their subtype. When subtype=FALSE, all the samples will be coloured black.
- significance** The threshold for selecting significant P-values.
- z.threshold** Threshold to display a green or red bar in the color bar on top of the heatmap for independent features with mean z-scores above `z.threshold` (high positive association) or below `-z.threshold` (high negative association).

colRamp	Palette of colors to be used in the heatmap.
add.colRamp	Palette of colors to be used in the added plot.
show.names.indep	logical if set to TRUE, displays the names (indep.id and indep.symb entered in the <a href="#">assemble.data</a> ) of the independent features with mean z-scores above or below the z.threshold in the heatmap.
show.names.dep	logical if set to TRUE, displays the names (dep.id and dep.symb entered in the <a href="#">assemble.data</a> ) of the significant dependent features in the heatmap.
adjust.method	Method used to adjust the P-values for multiple testing, see <a href="#">p.adjust</a> . Default is "BY" recommended when copy number is used as dependent data. See <a href="#">SIM</a> for more information about adjusting P-values.
scale	Vector specifying the color scale in the heatmap. If scale="auto", the maximum and minimum value of all z-scores will be calculated and set as the limits for all analyzed regions. Another option is to define a custom scale, e.g. scale = c(-5,5).
add.scale	Vector specifying the color scale in the left plot near the heatmap. If scale="auto", the maximum and minimum value of all the values will be calculated and set as the limits for all analyzed regions. Another option is to define a custom scale, e.g. scale = c(-5,5).
add.plot	Summary plot of copy number data in left panel. Either "smooth", "heatmap", or "none". The "smooth" plot smoothes the copy number log ratios per sample, see <a href="#">quantsmooth</a> for more details. The "heatmap" method produces an aCGH heatmap where green indicates gain, and red loss. The scale of the aCGH heatmap is automatically set to the min and max of the aCGH measurements of the analyzed regions. Default is plot.method = "none", no additional plot will be drawn.
smooth.lambda	Numeric value, specifying the quantile smoothing parameter for plot.method="smooth". See <a href="#">quantsmooth</a> and references for more information.
pdf	logical; indicate whether to generate a pdf of the plots in the current working directory or not.
run.name	This must be the same a given to <code>integrated.analysis</code>
...	not used in this version

## Details

The `sim.plot.zscore.heatmap` function can only run after the [integrated.analysis](#) is run with `zscores = TRUE`.

The results are returned as a single-page pdf containing an association heatmap of the regions listed in `input.regions`. For high-density arrays large files will be produced, both demanding more memory available from your computer to produce them as well as being heavier to open on screen. To avoid this, analyze chromosome arms as units instead of chromosomes, both here and in `input.regions = "all arms"`.

The heatmap contains the z-scores generated by the function [integrated.analysis](#) with `zscores=TRUE`. The dependent features are plotted from bottom to top, the independent features from left to right. Positive associations are shown in green, negative associations in red (color scale on the right).

At the left side of the heatmap a color bar represents the multiple testing corrected P-values of the probes in the dependent data (copy number), also with a color legend. Depending on which `plot.method` is used, a summary of copy number changes is shown on the left. At the top of the heatmap is a color bar corresponding to the mean z-scores of the independent features (expression data) that are above or below the `z.threshold`. If `show.names.indep` is set to TRUE, labels will be drawn for the probes with mean z-scores greater than `z.threshold` or lower than `-z.threshold` at the bottom of the heatmap. If `show.names.dep` is set to TRUE, labels will be drawn for the significant dependent probes lower than significance to the right of the heatmap.

### Value

No values are returned. The results are stored in a subdirectory of `run.name` as pdf.

### Author(s)

Marten Boetzer, Melle Sieswerda, Renee X. de Menezes <R.X.Menezes@lumc.nl>

### References

Eilers PH, de Menezes RX. 2005 Apr 1, Quantile smoothing of array CGH data. *Bioinformatics*, **21**(7):1146-53.

Wang P, Kim Y, Pollack J, Narasimhan B, Tibshirani R. 2005, A method for calling gains and losses in array CGH data. *Biostatistics*, **6**:45-58.

### See Also

[SIM](#), [tabulate.pvals](#), [tabulate.top.dep.features](#), [tabulate.top.indep.features](#), [sim.plot.overlapping.indep.dep.features](#)

### Examples

```
#first run example(assemble.data)
#and example(integrated.analysis)

#plot the zscores in a heatmap
sim.plot.zscore.heatmap(input.regions = "8q", adjust.method = "BY",
                        run.name = "chr8q", pdf = FALSE)

sim.plot.zscore.heatmap(input.regions = "8q",
                        method="full",
                        significance = 0.05,
                        z.threshold = 1,
                        colRamp = colorRampPalette(c("red", "black",
                                                    "green"))(15),
                        show.names.indep=TRUE,
                        show.names.dep=TRUE,
                        adjust.method = "holm",
                        add.plot = "heatmap",
                        smooth.lambda = 2,
                        pdf = FALSE,
                        run.name = "chr8q")
```

```
sim.plot.zscore.heatmap(input.regions = "8q",
                        method="full",
                        significance = 0.05,
                        z.threshold = 1,
                        show.names.indep = TRUE,
                        show.names.dep = TRUE,
                        add.plot = "none",
                        smooth.lambda = 2,
                        scale = c(-2, 2),
                        pdf = FALSE,
                        run.name = "chr8q")
```

---

sim.update.chrom.table

*Update the chromosome table*

---

### Description

A function to update the genomic positions of chromosome arms. Base locations of the start and end of chromosome arms should be used from the same organism and build of genome as the location provided as annotation with the datasets.

### Usage

```
sim.update.chrom.table(db = "homo_sapiens_core_40_36b")
```

### Arguments

db                    database name

### Details

This functions requires library RMySQL. Currently **SIM** only supports integrated analysis on the human genome without mitochondrial DNA.

### Value

Chromosome table [chrom.table](#).

### Author(s)

Marten Boetzer, Renee X. de Menezes <R.X.Menezes@lumc.nl>

### References

<http://www.ensembl.org/info/data/mysql.html>

### See Also

[SIM](#), [chrom.table](#)

## Examples

```
#you need internet connection for this!
#sim.update.chrom.table(db = "homo_sapiens_core_40_36b")
```

---

tabulate.pvals	<i>Sums significant P-values for the analyzed regions</i>
----------------	---

---

## Description

Generates a data.frame with the significance of P-values in the analyzed regions, dividing them into bins.

## Usage

```
tabulate.pvals(input.regions = "all chrs",
               adjust.method = "BY",
               bins = c(0.001, 0.005, 0.01, 0.025, 0.05, 0.075, 0.1, 0.2, 1),
               significance.idx = 8,
               order.by,
               decreasing = TRUE,
               method = c("full", "smooth", "window", "overlap"),
               run.name = "analysis_results")
```

## Arguments

- input.regions** vector indicating the dependent regions to be analyzed. Can be defined in four ways: 1) predefined input region: insert a predefined input region, choices are: "all chrs", "all chrs auto", "all arms", "all arms auto" In the predefined regions "all arms" and "all arms auto" the arms 13p, 14p, 15p, 21p and 22p are left out, because in most studies there are no or few probes in these regions. To include them, just make your own vector of arms. 2) whole chromosome(s): insert a single chromosome or a list of chromosomes as a vector: c(1, 2, 3). 3) chromosome arms: insert a single chromosome arm or a list of chromosome arms like c("1q", "2p", "2q"). 4) subregions of a chromosome: insert a chromosome number followed by the start and end position like "chr1:1-1000000" These regions can also be combined, e.g. c("chr1:1-1000000", "2q", 3). See [integrated.analysis](#) for more information.
- adjust.method** Method used to adjust the P-values for multiple testing, see [p.adjust](#). Default is "BY" recommended when copy number is used as dependent data. See [SIM](#) for more information about adjusting P-values.
- bins** vector of significance thresholds. This function will calculate the number of features having a P-value lower than the bin.
- significance.idx** Index of "bins" to use when computing the percentage of significant P-values. Defaults to 8 (i.e. the first entry in "bins"), in this case 0.20.



**Arguments**

<code>input.regions</code>	vector indicating the dependent regions to be analyzed. Can be defined in four ways: 1) predefined input region: insert a predefined input region, choices are: “all chrs”, “all chrs auto”, “all arms”, “all arms auto” In the predefined regions “all arms” and “all arms auto” the arms 13p, 14p, 15p, 21p and 22p are left out, because in most studies there are no or few probes in these regions. To include them, just make your own vector of arms. 2) whole chromosome(s): insert a single chromosome or a list of chromosomes as a vector: <code>c(1, 2, 3)</code> . 3) chromosome arms: insert a single chromosome arm or a list of chromosome arms like <code>c("1q", "2p", "2q")</code> . 4) subregions of a chromosome: insert a chromosome number followed by the start and end position like <code>"chr1:1-1000000"</code> These regions can also be combined, e.g. <code>c("chr1:1-1000000", "2q", 3)</code> . See <a href="#">integrated.analysis</a> for more information.
<code>adjust.method</code>	Method used to adjust the P-values for multiple testing, see <a href="#">p.adjust</a> . Default is “BY” recommended when copy number is used as dependent data. See <a href="#">SIM</a> for more information about adjusting P-values.
<code>method</code>	this must be the either one of “full”, “window”, “overlap” or “smooth” but the data should generated by the same method in <a href="#">integrated.analysis</a> .
<code>significance</code>	threshold used to select the significant dependent features. Pvalues below this threshold will be used to estimate regions.
<code>run.name</code>	This must be the same a given to <a href="#">integrated.analysis</a>

**Details**

Output is a .txt file containing a table with sorted integrated analysis P-values of the dependent features. It includes the `ann.dep` columns that were read in the [assemble.data](#) function. Additionally it returns a .txt file containing the significant P-value rich regions. No P-value rich regions are returned when `zscores.diag = "all"`.

**Value**

Returns a list of `data.frame`’s for each input region. Significant P-value rich regions are returned as a `data.frame`. This `data.frame` can be used as an input for [getoverlappingregions](#). Additionally, the results are stored in a subdirectory of `run.name` as `txt`.

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**See Also**

[SIM](#), [tabulate.pvals](#), [tabulate.top.indep.features](#)

**Examples**

```
#first run example(assemble.data)
#and example(integrated.analysis)
#get the top dependent features sorted by p-value
```

```
table.dep <- tabulate.top.dep.features(input.regions="8q",
                                     adjust.method="BY",
                                     method="full",
                                     significance=0.05,
                                     run.name="chr8q")
head(table.dep[["8q"]])
```

---

tabulate.top.indep.features

*Lists the mean z-scores for the independent features*

---

### Description

Lists the mean z-scores for independent features in the analyzed regions, calculated across the significant dependent features. Gives insight in the expression levels most strongly associated with copy number changes.

### Usage

```
tabulate.top.indep.features(input.regions = "all chrs",
                           input.region.indep = NULL,
                           method = c("full", "smooth", "window", "overlap"),
                           adjust.method = "BY",
                           significance = 1,
                           decreasing=TRUE,
                           z.threshold = c(0, 0),
                           run.name = "analysis_results")
```

### Arguments

**input.regions** vector indicating the dependent regions to be analyzed. Can be defined in four ways: 1) predefined input region: insert a predefined input region, choices are: "all chrs", "all chrs auto", "all arms", "all arms auto" In the predefined regions "all arms" and "all arms auto" the arms 13p, 14p, 15p, 21p and 22p are left out, because in most studies there are no or few probes in these regions. To include them, just make your own vector of arms. 2) whole chromosome(s): insert a single chromosome or a list of chromosomes as a vector: c(1, 2, 3). 3) chromosome arms: insert a single chromosome arm or a list of chromosome arms like c("1q", "2p", "2q"). 4) subregions of a chromosome: insert a chromosome number followed by the start and end position like "chr1:1-1000000" These regions can also be combined, e.g. c("chr1:1-1000000", "2q", 3). See [integrated.analysis](#) for more information.

**input.region.indep** fill in

**method** this must be the either one of "full", "window", "overlap" or "smooth" but the data should generated by the same method in [integrated.analysis](#).

adjust.method	Method used to adjust the P-values for multiple testing, see <a href="#">p.adjust</a> . Default is "BY" recommended when copy number is used as dependent data. See <a href="#">SIM</a> for more information about adjusting P-values.
significance	threshold used to select the significant dependent features. Only the z-scores with these features are used to calculate the mean z-scores across the independent features.
decreasing	fill in
z.threshold	fill in
run.name	This must be the same as given to <code>integrated.analysis</code>

### Details

`tabulate.top.indep.features` can only be run after [integrated.analysis](#) with `zscores = TRUE`.

Output is a .txt file containing a table with the mean z-scores of all independent features per analyzed region. It includes the `ann.indep` columns that were read in the [assemble.data](#) function.

Additionally it returns a .txt file containing the significant zscores rich regions.

Depending on the argument "adjust.method", the P-values are first corrected for multiple testing. Next, the z-scores are filtered to include only those entries that correspond to significant (P-value < "significance") dependent features to calculate the mean z-scores.

The dependent table can not be generated for diagonal integrated runs.

### Value

Returns a list of data.frame's for each input region. Significant P-value rich regions are returned as a data.frame. This data.frame can be used as an input for [getoverlappingregions](#). Additionally, the results are stored in a subdirectory of `run.name` as txt.

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### See Also

[SIM](#), [tabulate.pvals](#), [tabulate.top.dep.features](#)

### Examples

```
#first run example(assemble.data)
#and example(integrated.analysis)
table.indep <- tabulate.top.indep.features(input.regions="8q",
                                         adjust.method="BY",
                                         method="full",
                                         significance= 0.05,
                                         z.threshold=c(-1, 1),
                                         run.name="chr8q")
head(table.indep[["8q"]])
```

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