

# Package ‘snpEnrichment’

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**Type** Package

**Title** SNPs Enrichment Analysis

**Version** 1.7.0

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**Description** Implements classes and methods for large scale SNP enrichment analysis (e.g. SNPs associated with genes expression in a GWAS signal).

**License** GPL (>= 2)

**Depends** R (>= 3.0.0), methods

**Imports** parallel, snpStats, grid, ggplot2, grDevices, graphics, stats, utils

**URL** <https://github.com/mcanouil/snpEnrichment>

**Encoding** UTF-8

**BugReports** <https://github.com/mcanouil/snpEnrichment/issues>

**NeedsCompilation** no

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**Repository** CRAN

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## R topics documented:

snpEnrichment-package . . . . .	2
Chromosome-class . . . . .	4
compareEnrichment . . . . .	6
Enrichment-class . . . . .	8
EnrichSNP-class . . . . .	10
excludeSNP . . . . .	11
GC . . . . .	12
getEnrichSNP . . . . .	13

initFiles	14
is.chromosome	15
is.enrichment	16
mclapply2	17
plot-methods	18
print-methods	19
readEnrichment	20
reSample	22
toyEnrichment-dataset	24
transcript-dataset	24
writeLD	25

<b>Index</b>	<b>27</b>
--------------	-----------

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snpEnrichment-package ~ Overview: SNPs enrichment analysis ~

---

## Description

Implements classes and methods for large-scale SNP enrichment analysis (e.g. SNPs associated with genes expression in a GWAS signal).

## Details

```

Package:    snpEnrichment
Title:      SNPs enrichment analysis
Author:     Mickael Canouil
Contributor: Loic Yengo
Maintainer: Mickael Canouil
License:    GPL (>= 2)
Depends:    R (>= 3.0.0), methods
Suggests:   grid, ggplot2
Imports:    parallel, snpStats
URL:        https://github.com/mcanouil/snpEnrichment
Encoding:   UTF-8

```

## Note

Internal data management in 'snpEnrichment' use RefSNP (rs) IDs.

## Author(s)

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)  
Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```
#####  
### 1. Prepare data  
## Not run: snpInfoDir <- system.file("extdata/snpInfo",  
                                     package = "snpEnrichment")  
signalFile <- system.file("extdata/Signal/toySignal.txt",  
                          package = "snpEnrichment")  
  
initFiles(pattern = "Chrom", snpInfoDir, signalFile, mc.cores = 1)  
  
writeLD(pattern = "Chrom", snpInfoDir, signalFile,  
        ldDir = NULL, ldThresh = 0.8, depth = 1000,  
        mc.cores = 1)  
## End(Not run)  
  
#####  
### 2. Read data  
## Not run: snpListDir <- system.file("extdata/List",  
                                     package = "snpEnrichment")  
data(transcript)  
transcriptFile <- transcript  
  
toyData <- readEnrichment(pattern = "Chrom", signalFile,  
                          transcriptFile, snpListDir,  
                          snpInfoDir, distThresh = 1000,  
                          sigThresh = 0.05, LD = TRUE,  
                          ldDir = NULL, mc.cores = 1)  
toyData  
## End(Not run)  
  
#####  
### 3. Compute results  
## Not run: reSample(object = toyData,  
                    nSample = 10,  
                    empiricPvalue = TRUE,  
                    MAFpool = c(0.05, 0.10, 0.2, 0.3, 0.4, 0.5),  
                    mc.cores = 1,  
                    onlyGenome = TRUE)  
## End(Not run)  
  
#####
```

```

### 4. Further analysis: Exclude SNP from original list.
## Not run: excludeFile <- c(
  "rs4376885", "rs17690928", "rs6460708", "rs13061537", "rs11769827",
  "rs12717054", "rs2907627", "rs1380109", "rs7024214", "rs7711972",
  "rs9658282", "rs11750720", "rs1793268", "rs774568", "rs6921786",
  "rs1699031", "rs6994771", "rs16926670", "rs465612", "rs3012084",
  "rs354850", "rs12803455", "rs13384873", "rs4364668", "rs8181047",
  "rs2179993", "rs12049335", "rs6079926", "rs2175144", "rs11564427",
  "rs7786389", "rs7005565", "rs17423335", "rs12474102", "rs191314",
  "rs10513168", "rs1711437", "rs1992620", "rs283115", "rs10754563",
  "rs10851727", "rs2173191", "rs7661353", "rs1342113", "rs7042073",
  "rs1567445", "rs10120375", "rs550060", "rs3761218", "rs4512977"
)
# OR
excludeFile <- system.file("extdata/Exclude/toyExclude.txt",
  package = "snpEnrichment")

toyData_exclude <- excludeSNP(toyData, excludeFile, mc.cores = 1)

# Warning: compareEnrichment is in development!!
compareResults <- compareEnrichment(object.x = toyData,
  object.y = toyData_exclude,
  pattern = "Chrom",
  nSample = 10,
  empiricPvalue = TRUE,
  mc.cores = 1,
  onlyGenome = TRUE)

## End(Not run)

#####
### 5. Watch results
## Not run: show(toyData)
print(toyData)
head(getEnrichSNP(toyData, type = "xSNP"))

show(toyData_exclude)
print(toyData_exclude)
head(getEnrichSNP(toyData_exclude, type = "eSNP"))
## End(Not run)

```

---

Chromosome-class

*Class* [Chromosome](#)


---

## Description

This class is defined to summarize the enrichment analysis about a chromosome.

## Objects from the Class

`chromosome` is defined to build an object of class `Chromosome` in order to compute an enrichment analysis. A `Chromosome` object contains the original data, a list of SNPs, some results and resampling data.

When created, an `Chromosome` object is "empty". `readEnrichment` initializes a `Chromosome` object with value from PLINK computation and user's files. In this step, only the fields "Data", "LD", "SNP" are filled. `reSample` fills the fields: Table, EnrichmentRatio, Z, PValue and Resampling of a `Chromosome`.

Note that if `reSample` is executed on an `Chromosome` every new resampling is added to the original ones, pre-existing statistics are erased and computed again with the new resampling set.

## Slots

**Data** [data.frame]: a data.frame with 6 columns ("SNP", "PVALUE", "CHR", "MAF", "eSNP", "xSNP"). Where "eSNP" and "xSNP" are logical columns defining the lists of SNPs (extended or not).

**LD** [data.frame]: a data.frame which contains LD informations between SNPs (computed with `writeLD` or PLINK).

**eSNP, xSNP** [SNP]: contain a `EnrichSNP` object (whith slots: SNP, Table, EnrichmentRatio, Z, PValue and Resampling) for a list of SNPs (eSNP) and an extended one (xSNP).

## Extends

Class `EnrichSNP`, directly.

## Methods

**chromosome(Data, LD, eSNP, xSNP)**: Generate and initialize a new `Chromosome` object.

**object["slotName" ]**: Get the value of the field slotName.

**object["slotName" <-value:]**: Set value to the field slotName.

**show(object)**: Return the formatted values of `Chromosome` object.

## Note

`Chromosome` object is not intended to be used alone on this version. It is a part of the `Enrichment` object.

## Author(s)

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)  
 Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
 Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
 Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```
Data <- structure(
  list(
    SNP = c("rs4970420", "rs3766178",
            "rs10910030", "rs10910036",
            "rs2234167", "rs6661861"),
    PVALUE = c(0.9244, 0.167, 0.01177, 0.4267, 0.9728, 0.4063),
    CHR = c(1, 1, 1, 1, 1, 1),
    POS = c(1106473, 1478180, 2035684, 2183754, 2494330, 3043121),
    MAF = c(0.2149, 0.3117, 0.374, 0.3753, 0.1565, 0.06101),
    eSNP = c(0, 1, 1, 0, 0, 0),
    xSNP = c(0, 1, 1, 0, 0, 0)
  ),
  .Names = c("SNP", "PVALUE", "CHR", "POS", "MAF", "eSNP", "xSNP"),
  row.names = c("rs4970420", "rs3766178",
                "rs10910030", "rs10910036",
                "rs2234167", "rs6661861"),
  class = "data.frame")

toyChr <- chromosome(Data = Data)
show(toyChr)
toyChr

toyChr <- chromosome()
toyChr["Data"] <- Data
toyChr

is.chromosome(toyChr) # TRUE
```

---

compareEnrichment      *Compare enrichment analysis between two SNPs list*

---

**Description**

Compare the enrichment analysis between two set of SNPs. [compareEnrichment](#) compare two [Enrichment](#) objects.

**Usage**

```
compareEnrichment(object.x, object.y, pattern = "Chrom",
                  nSample = 100, empiricPvalue = TRUE,
                  mc.cores = 1, onlyGenome = TRUE)
```

## Arguments

object.x, object.y	[Enrichment]: an <a href="#">Enrichment</a> object fully filled (e.g. <a href="#">readEnrichment</a> ).
pattern	[character]: character string containing a expression to be matched with all chromosomes files (e.g."Chrom" for files which start by "Chrom" followed by the chromosome number).
nSample	[numeric]: the number of resampling done by <a href="#">reSample</a> for p-values computation (minimum is 100).
empiricPvalue	[logical]: empiricPvalue=TRUE (default) compute PValue based on the null distribution (resampling). If empiricPvalue=TRUE, the empirical p-values are computed instead.
mc.cores	[numeric]: the number of cores to use (default is mc.cores=1), i.e. at most how many child processes will be run simultaneously. Must be at least one, and parallelization requires at least two cores.
onlyGenome	[logical]: onlyGenome=TRUE (default) compute resampling step for all chromosomes.

## Value

Return a list of three elements:

object.xy	<a href="#">Enrichment</a> object from the comparison between object.x and object.y.
object.x	<a href="#">Enrichment</a> object passed in object.x with resampling data.
object.y	<a href="#">Enrichment</a> object passed in object.y with resampling data.

## Note

Still in development.

## Author(s)

Mickael Canouil <mickael.canouil@good.ibl.fr>

## See Also

Overview : [snpEnrichment-package](#)  
Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

## Examples

```
## Not run: data(toyEnrichment)

reSample(object = toyEnrichment,
         nSample = 10,
         empiricPvalue = TRUE,
```

```

MAFpool = c(0.05, 0.10, 0.2, 0.3, 0.4, 0.5),
mc.cores = 1,
onlyGenome = TRUE)

excludeFile <- c(
  "rs7897180", "rs4725479", "rs315404", "rs17390391", "rs1650670",
  "rs6783390", "rs1642009", "rs4756586", "rs11995037", "rs4977345",
  "rs13136448", "rs4233536", "rs11151079", "rs2299657", "rs4833930",
  "rs1384", "rs7168184", "rs6909895", "rs7972667", "rs2293229",
  "rs918216", "rs6040608", "rs2817715", "rs13233541", "rs4486743",
  "rs2127806", "rs10912854", "rs1869052", "rs9853549", "rs448658",
  "rs2451583", "rs17483288", "rs10962314", "rs9612059", "rs1384182",
  "rs8049208", "rs12215176", "rs2980996", "rs1736976", "rs8089268",
  "rs10832329", "rs12446540", "rs7676237", "rs869922", "rs16823426",
  "rs1374393", "rs13268781", "rs11134505", "rs7325241", "rs7520109"
)
# OR
excludeFile <- system.file("extdata/Exclude/toyExclude.txt",
  package = "snpEnrichment")

toyEnrichment_exclude <- excludeSNP(toyEnrichment, excludeFile, mc.cores = 1)

compareResults <- compareEnrichment(object.x = toyEnrichment,
  object.y = toyEnrichment_exclude,
  pattern = "Chrom",
  nSample = 10,
  empiricPvalue = FALSE,
  mc.cores = 1,
  onlyGenome = TRUE)

## End(Not run)

```

---

Enrichment-class      *Class* [Enrichment](#)

---

## Description

This class is defined to summarize the enrichment analysis on each chromosomes and the whole genome.

## Objects from the Class

[enrichment](#) is defined to build an object of class [Enrichment](#) in order to compute an enrichment analysis. [Enrichment](#) is the object containing the results for all [Chromosome](#) object and for the whole genome.

When an [Enrichment](#) object is created, it contains a list of SNPs (e.g. eSNPs). All the others slots are "empty". After [reSample](#) is ran on an [Enrichment](#) object, the slots: [Table](#), [EnrichmentRatio](#), [Z](#), [PValue](#) and [Resampling](#) are filled.

Note that if [reSample](#) is executed on an [Enrichment](#) every new resampling is added to the original ones, pre-existing statistics are erased and computed again with the new resampling set.



## Slots

**Loss** [data.frame]: a four columns data.frame: "Rows", "Unique", "Intersect.Ref.Signal" and "CIS". This slot gives information on data losses.

**Call** [list]: each parameters used for the reading or resampling step are stored in this slot.

**eSNP, xSNP** [SNP]: contain a [EnrichSNP](#) object (whith slots: SNP, Table, EnrichmentRatio, Z, PValue and Resampling) for a list of SNPs (eSNP) and an extended one (xSNP).

**Chromosomes** [list(Chromosome)]: a list of 22 [Chromosome](#) objects.

## Extends

Class [Chromosome](#), directly.

Class [EnrichSNP](#), directly.

## Methods

**enrichment(Loss, Call, eSNP, xSNP, Chromosomes)**: Generate and initialize a new [Enrichment](#) object.

**object["slotName" ]**: Get the value of the field slotName.

**object["slotName" <-value:]** Set value to the field slotName.

**show(object)**: Return the formatted values of [Enrichment](#) object.

**print(object, what, type)**: Return a summary table of an [Enrichment](#) object.

**reSample(object, nSample, MAFpool, mc.cores, onlyGenome)**: Compute P-Values based upon a resampling of SNPs (eSNP and xSNP) and update the [Enrichment](#) object.

**excludeSNP(object, excludeFile, mc.cores)**: Excludes SNPs given in excludeFile from the original list of eSNPs (xSNPs). Then a new enrichment analysis is computed.

**reset(object, "slotName")**: Reset the field slotName.

**plot(object, what, type, ggplot, pvalue)**: Plot p-values or Z-statistics convergence.

**getEnrichSNP(object, type)**: Return eSNP/xSNP which are enriched as a data.frame.

## Author(s)

Mickael Canouil <mickael.canouil@good.ibl.fr>

## See Also

Overview : [snpEnrichment-package](#)

Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)

Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#), [enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)

Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```

data(toyEnrichment)
toyEnrich <- enrichment()
show(toyEnrich)

toyEnrich["Loss"] <- toyEnrichment["Loss"]
toyEnrich["Loss"]

toyEnrich <- enrichment(Loss = toyEnrichment["Loss"],
                        eSNP = toyEnrichment["eSNP"])
toyEnrich <- enrichment(Loss = toyEnrichment["Loss"])

## Not run: reSample(object = toyEnrichment,
                    nSample = 10,
                    empiricPvalue = TRUE,
                    MAFpool = c(0.05, 0.10, 0.2, 0.3, 0.4, 0.5),
                    mc.cores = 1,
                    onlyGenome = TRUE)
print(toyEnrichment)

excludeFile <- c(
"rs7897180", "rs4725479", "rs315404", "rs17390391", "rs1650670",
"rs6783390", "rs1642009", "rs4756586", "rs11995037", "rs4977345",
"rs13136448", "rs4233536", "rs11151079", "rs2299657", "rs4833930",
"rs1384", "rs7168184", "rs6909895", "rs7972667", "rs2293229",
"rs918216", "rs6040608", "rs2817715", "rs13233541", "rs4486743",
"rs2127806", "rs10912854", "rs1869052", "rs9853549", "rs448658",
"rs2451583", "rs17483288", "rs10962314", "rs9612059", "rs1384182",
"rs8049208", "rs12215176", "rs2980996", "rs1736976", "rs8089268",
"rs10832329", "rs12446540", "rs7676237", "rs869922", "rs16823426",
"rs1374393", "rs13268781", "rs11134505", "rs7325241", "rs7520109"
)

toyEnrichment_exclude <- excludeSNP(toyEnrichment, excludeFile, mc.cores = 1)
print(toyEnrichment_exclude)
## End(Not run)

```

---

EnrichSNP-class

Class "EnrichSNP"

---

**Description**

This class is defined to summarize the enrichment analysis. It's a part of [Chromosome](#) and [Enrichment](#) classes.

**Slots**

**List** [vector(character)]: a list of SNPs used to compute enrichment (e.g. eSNP or xSNP).

**Table** [matrix]: Contingency table with SNPs (columns) and P-Values from signal (rows).

**EnrichmentRatio** [numeric]: Enrichment Ratio is computed on the contingency table (Table slot).

**Z** [numeric]: A statistic computed from `EnrichmentRatio` and resampling results.

**PValue** [numeric]: P-Value associated with the statistic Z.

**Resampling** [matrix]: A matrix with by row, the contingency table and the odds ratio for each resampling.

## Methods

`object["slotName" : ]` Get the value of the field `slotName`.

`object["slotName" <-value: ]` Set value to the field `slotName`.

`show(object)`: Return the formatted values of `EnrichSNP` object.

## Note

`EnrichSNP` object is not intended to be use directly by user. It is a part of the `Enrichment` and `Chromosome` object.

## Author(s)

Mickael Canouil <mickael.canouil@good.ibl.fr>

## See Also

Overview : [snpEnrichment-package](#)

Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)

Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#), [enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)

Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

---

excludeSNP

*Exclude SNPs from Enrichment analysis*

---

## Description

Remove a specify set of SNPs and compute a new enrichment analysis.

## Usage

```
excludeSNP(object, excludeFile, mc.cores = 1)
```

## Arguments

`object` [Enrichment]: an `Enrichment` object filled by `reSample`.

`excludeFile` [vector(character)]: a list of SNPs to remove from a previous enrichment analysis. A path to a file which the first column are the SNPs.

`mc.cores` [numeric]: the number of cores to use (default is `mc.cores=1`), i.e. at most how many child processes will be run simultaneously. Must be at least one, and parallelization requires at least two cores.

**Value**

Return the object given in argument where lists of SNPs are updated by removing SNPs in excludeFile.

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)

Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)

Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#), [enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)

Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```
## Not run: data(toyEnrichment)
excludeFile <- c(
  "rs4376885", "rs17690928", "rs6460708", "rs13061537", "rs11769827",
  "rs12717054", "rs2907627", "rs1380109", "rs7024214", "rs7711972",
  "rs9658282", "rs11750720", "rs1793268", "rs774568", "rs6921786",
  "rs1699031", "rs6994771", "rs16926670", "rs465612", "rs3012084",
  "rs354850", "rs12803455", "rs13384873", "rs4364668", "rs8181047",
  "rs2179993", "rs12049335", "rs6079926", "rs2175144", "rs11564427",
  "rs7786389", "rs7005565", "rs17423335", "rs12474102", "rs191314",
  "rs10513168", "rs1711437", "rs1992620", "rs283115", "rs10754563",
  "rs10851727", "rs2173191", "rs7661353", "rs1342113", "rs7042073",
  "rs1567445", "rs10120375", "rs550060", "rs3761218", "rs4512977"
)
# OR
excludeFile <- system.file("extdata/Exclude/toyExclude.txt",
  package = "snpEnrichment")

toyEnrichment_exclude <- excludeSNP(toyEnrichment, excludeFile, mc.cores = 1)
toyEnrichment_exclude
## End(Not run)
```

---

GC

*Full Garbage Collection*


---

**Description**

GC performs garbage collection until free memory indicators show no change.

**Usage**

```
GC(verbose = getOption("verbose"), reset=FALSE)
```

**Arguments**

verbose	[logical]: if TRUE, the garbage collection prints statistics about cons cells and the space allocated for vectors.
reset	[logical]: if TRUE the values for maximum space used are reset to the current values.

**Value**

GC returns a matrix with rows "Ncells" (`_cons cells_`), usually 28 bytes each on 32-bit systems and 56 bytes on 64-bit systems, and "Vcells" (`_vector cells_`, 8 bytes each), and columns "used" and "gc trigger", each also interpreted in megabytes (rounded up to the next 0.1Mb).

If maxima have been set for either "Ncells" or "Vcells", a fifth column is printed giving the current limits in Mb (with NA denoting no limit).

The final two columns show the maximum space used since the last call to `GC(reset=TRUE)` (or since R started).

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

The R Internals manual.

Memory on R's memory management, and `gctorture` if you are an R developer.

`reg.finalizer` for actions to happen at garbage collection.

**Examples**

```
GC() # - do it now
x <- integer(100000); for(i in 1:18) x <- c(x,i)
GC(TRUE)
GC(reset=TRUE)
```

---

getEnrichSNP

*Get all eSNP/xSNP which are enriched*

---

**Description**

`getEnrichSNP` get all eSNP/xSNP in a `Enrichment` object which are significant in the signal according to `sigThresh` defined in `readEnrichment`.

**Usage**

```
## S4 method for signature 'Enrichment'
getEnrichSNP(object, type = "eSNP")
```

**Arguments**

object            [Enrichment]: an object of class [Enrichment](#).  
 type             [character]: extract eSNP or xSNP data.

**Value**

Return a `data.frame` with eSNP/xSNP which are enriched in signal given to `signalFile` in function [readEnrichment](#).

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)  
 Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
 Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
 Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```
## Not run: data(toyEnrichment)
eSNPenriched <- getEnrichSNP(object = toyEnrichment, type = "eSNP")
head(eSNPenriched)
## End(Not run)
```

---

initFiles

*Initialize files for enrichment analysis*

---

**Description**

[initFiles](#) create several files needed to run [readEnrichment](#). ".frq" and ".signal" are created with PLINK. LD computation can be run with [writeLD](#) or with PLINK.

**Usage**

```
initFiles(pattern = "Chrom", snpInfoDir, signalFile, mc.cores = 1)
```

**Arguments**

pattern            [character]: character string containing a expression to be matched with all chromosomes files (e.g."Chrom" for files which start by "Chrom" followed by the chromosome number).  
 snpInfoDir        [character]: character string naming a directory containing the reference data in a PLINK format (\*.bed, \*.bim and \*.fam).

signalFile	[character]: the name of the signal file which the data are to be read from (2 columns: "SNP" and "PVALUE"). Each row of the table appears as one line of the file. If it does not contain an <code>_absolute_</code> path, the file name is <code>_relative_</code> to the current working directory, <code>getwd</code> . The fields separator character have to be a space " " or a tabulation "\t".
mc.cores	[numeric]: the number of cores to use (default is <code>mc.cores=1</code> ), i.e. at most how many child processes will be run simultaneously. Must be at least one, and parallelization requires at least two cores.

**Value**

This function writes several files, in the temporary directory (defined in `R_SESSION_TMPDIR`), nothing else is returned. These files are used to build an [Enrichment](#) object by [readEnrichment](#) in order to compute enrichment analysis ([reSample](#)).

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)  
 Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
 Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
 Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```
## Not run: snpInfoDir <- system.file("extdata/snpInfo",
                                   package = "snpEnrichment")
signalFile <- system.file("extdata/Signal/toySignal.txt",
                         package = "snpEnrichment")
initFiles(pattern = "Chrom",
          snpInfoDir,
          signalFile,
          mc.cores = 1)
## End(Not run)
```

---

is.chromosome

*Is an Chromosome object*

---

**Description**

'is.chromosome' returns 'TRUE' if 'x' is an [Chromosome](#) object and 'FALSE' otherwise.

**Usage**

```
is.chromosome(object)
```

**Arguments**

object            [ANY]: object to be tested.

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)  
 Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
 Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
 Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```
a <- chromosome()
c <- chromosome()
is.chromosome(list())            # FALSE
is.chromosome(1)                # FALSE
is.chromosome(a)                # TRUE
is.chromosome(c(a, c))         # TRUE TRUE
is.chromosome(list(a, b = "char")) # TRUE FALSE
is.chromosome(c(a, b = list(12, c))) # TRUE FALSE TRUE
```

---

is.enrichment	<i>Is an Enrichment object</i>
---------------	--------------------------------

---

**Description**

'is.enrichment' returns 'TRUE' if 'x' is an [Enrichment](#) object and 'FALSE' otherwise.

**Usage**

```
is.enrichment(object)
```

**Arguments**

object            [ANY]: object to be tested.

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>



**See Also**

Overview : [snpEnrichment-package](#)  
 Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
 Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
 Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```
a <- enrichment()
c <- enrichment()
is.enrichment(list())           # FALSE
is.enrichment(1)               # FALSE
is.enrichment(a)               # TRUE
is.enrichment(c(a, c))         # TRUE TRUE
is.enrichment(list(a, b = "char")) # TRUE FALSE
is.enrichment(c(a, b = list(12, c))) # TRUE FALSE TRUE
```

---

mclapply2

*Parallel Versions of lapply with cores and memory control*


---

**Description**

[mclapply2](#) is a [mclapply](#) modification from [parallel](#) package, to avoid a memory overload. The maximum number of cores is computed depending on the amount of memory used by the parent process and the amount of free memory available on the machine. Note: This number is underestimated.

**Usage**

```
mclapply2(X, FUN, ..., mc.preschedule = TRUE, mc.set.seed = TRUE,
          mc.silent = FALSE, mc.cores = getOption("mc.cores", 2L),
          mc.cleanup = TRUE, mc.allow.recursive = TRUE)
```

**Arguments**

X	a vector (atomic or list) or an expressions vector. Other objects (including classed objects) will be coerced by <a href="#">as.list</a> .
FUN	the function to be applied to ( <a href="#">mclapply</a> ) each element of X or ( <a href="#">mcmapply</a> ) in parallel to ....
...	For <a href="#">mclapply</a> , optional arguments to FUN.
mc.preschedule	if set to TRUE then the computation is first divided to (at most) as many jobs are there are cores and then the jobs are started, each job possibly covering more than one value. If set to FALSE then one job is forked for each value of X. The former is better for short computations or large number of values in X, the latter is better for jobs that have high variance of completion time and not too many values of X compared to <code>mc.cores</code> .

<code>mc.set.seed</code>	See <code>mcpParallel</code> .
<code>mc.silent</code>	if set to TRUE then all output on 'stdout' will be suppressed for all parallel processes forked ('stderr' is not affected).
<code>mc.cores</code>	The number of cores to use, i.e. at most how many child processes will be run simultaneously. The option is initialized from environment variable MC_CORES if set. Must be at least one, and parallelization requires at least two cores.
<code>mc.cleanup</code>	if set to TRUE then all children that have been forked by this function will be killed (by sending SIGTERM) before this function returns. Under normal circumstances <code>mclapply</code> waits for the children to deliver results, so this option usually has only effect when <code>mclapply</code> is interrupted. If set to FALSE then child processes are collected, but not forcefully terminated. As a special case this argument can be set to the number of the signal that should be used to kill the children instead of SIGTERM.
<code>mc.allow.recursive</code>	Unless true, calling <code>mclapply</code> in a child process will use the child and not fork again.

**Value**

A list of the same length as `X` and named by `X`.

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

`mclapply` from package `parallel`.

---

plot-methods

*Plot method (S4) for [Enrichment](#) object*

---

**Description**

`plot` is a generic function for plotting of R objects. The function invokes particular methods which depend on the class of the first argument.

**Usage**

```
## S4 method for signature 'Enrichment'
plot(x, what = "Genome", type = c("eSNP", "xSNP"),
      ggplot = FALSE, pvalue = TRUE, ...)
```

**Arguments**

x	[Enrichment]: an object of class <a href="#">Enrichment</a> which the Z statistics or p-values have to be drawn.
what	[character or vector(numeric)]: default what="Genome") plot Z statistics or p-values for genome only (what must be: "All", "Genome" or numeric vector).
type	[vector(character)]: plot the selected analysis for "eSNP" and/or "xSNP".
ggplot	[logical]: use ggplot (default ggplot=FALSE) instead of classic plot method.
pvalue	[logical]: if TRUE, p-value convergense is plotted. Otherwise, Z statistic is plotted.
...	[any]: Arguments to be passed to methods, such as graphical parameters (see par)

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)  
 Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
 Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
 Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```
## Not run: data(toyEnrichment)
reSample(toyEnrichment, 10)
plot(toyEnrichment)
## End(Not run)
```

---

print-methods                      *Print method (S4)*

---

**Description**

[print](#) is a generic function used to print results.

**Usage**

```
## S4 method for signature 'Enrichment'
print(x, what = "Genome", type = c("eSNP", "xSNP"))

## S4 method for signature 'Chromosome'
print(x, type = c("eSNP", "xSNP"))
```

## Arguments

x	[Enrichment or Chromosome]: an object of class <a href="#">Enrichment</a> or <a href="#">Chromosome</a> .
what	[character or numeric]: what="Genome" (default) to print results as a matrix. what could be "All", "Genome" or a numeric from 1 to 22 (numeric vector is allowed).
type	[character]: select if results for "eSNP" and/or "xSNP" should be print.

## Value

Return a matrix for classes [Enrichment](#) and [Chromosome](#).

## Author(s)

Mickael Canouil <mickael.canouil@good.ibl.fr>

## See Also

Overview : [snpEnrichment-package](#)  
Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

## Examples

```
data(toyEnrichment)
print(toyEnrichment, "All", type = "eSNP")
print(toyEnrichment, "Genome")
print(toyEnrichment, 1)
```

---

readEnrichment	<i>Read and create EnrichmentRatio object</i>
----------------	---

---

## Description

Read files created by [initFiles](#) and create an [Enrichment](#) object.

## Usage

```
readEnrichment(pattern = "Chrom", signalFile,
               transcriptFile = "FALSE", snpListDir,
               snpInfoDir, distThresh = 1000,
               sigThresh = 0.05, LD = FALSE, ldDir = NULL,
               mc.cores = 1)
```

**Arguments**

pattern	[character]: character string containing a expression to be matched with all chromosomes files (e.g."Chrom" for files which start by "Chrom" followed by the chromosome number).
signalFile	[character]: the name of the signal file which the data are to be read from (2 columns: "SNP" and "PVALUE"). Each row of the table appears as one line of the file. If it does not contain an <code>_absolute_</code> path, the file name is <code>_relative_</code> to the current working directory, <code>getwd</code> . The fields separator character have to be a space " " or a tabulation "\t".
transcriptFile	[character or data.frame]: character string naming a file or a data.frame with four columns: Chromosome, transcript's name, Starting and Ending positions. <code>data(transcript)</code> can be use as parameters. Default is FALSE.
snpListDir	[character]: character string naming a directory containing a list of SNPs for one or several chromosomes. <code>snpListDir</code> can be a single file with at least two columns: chromosome and rs name.
snpInfoDir	[character]: character string naming a directory containing the reference data in a PLINK format (*.bed, *.bim and *.fam).
distThresh	[numeric]: maximal distance (kb) between SNP and gene. <code>distThresh</code> is used if <code>transcriptFile</code> is set.
sigThresh	[numeric]: statistical threshold for signal (e.g. <code>sigThresh = 0.05</code> for a given GWAS signal) used to compute an Enrichment Ratio.
LD	[logical]: LD=TRUE (default is FALSE) read LD compute with <code>writeLD</code> function or with PLINK. Note that, this setting can increase the computation's time, depending on number of SNPs in the signal file.
ldDir	[character]: character string naming a directory where the linkage disequilibrium files should be read (default <code>ldDir=NULL</code> is in temporary directory). LD files can be the LD output from plink.
mc.cores	[numeric]: the number of cores to use (default is <code>mc.cores=1</code> ), i.e. at most how many child processes will be run simultaneously. Must be at least one, and parallelization requires at least two cores.

**Value**

Return an object of class `Enrichment` partly filled.

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)  
 Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
 Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#), [enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
 Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

## Examples

```
## Not run: snpListDir <- system.file("extdata/List",
                                     package = "snpEnrichment")
signalFile <- system.file("extdata/Signal/toySignal.txt",
                          package = "snpEnrichment")
snpInfoDir <- system.file("extdata/snpInfo", package = "snpEnrichment")
data(transcript)
transcriptFile <- transcript

initFiles(pattern = "Chrom", snpInfoDir, signalFile, mc.cores = 1)
toyData <- readEnrichment(pattern = "Chrom",
                          signalFile,
                          transcriptFile,
                          snpListDir,
                          snpInfoDir,
                          distThresh = 1000,
                          sigThresh = 0.05,
                          LD = FALSE,
                          ldDir = NULL,
                          mc.cores = 1)

toyData
## End(Not run)
```

---

reSample

---

*Compute enrichment analysis on an [Enrichment](#) object*


---

## Description

After [initFiles](#) and [readEnrichment](#) has been run. [reSample](#) computes a statistic value and a p-value for each chromosomes and for the whole genome.

## Usage

```
## S4 method for signature 'Enrichment'
reSample(object, nSample = 100,
          empiricPvalue = TRUE,
          MAFpool = c(0.05, 0.10, 0.2, 0.3, 0.4, 0.5),
          mc.cores = 1, onlyGenome = TRUE)

## S4 method for signature 'Chromosome'
reSample(object, nSample = 100,
          empiricPvalue = TRUE, sigThresh = 0.05,
          MAFpool = c(0.05, 0.10, 0.2, 0.3, 0.4, 0.5),
          mc.cores = 1)
```

## Arguments

object	[Enrichment or Chromosome]: an object to be updated. It is intended, an object returned by the <a href="#">readEnrichment</a> function.
nSample	[numeric]: the number of resampling done by <a href="#">reSample</a> for p-values computation (minimum is 100).
empiricPvalue	[logical]: <code>empiricPvalue=TRUE</code> (default) compute PValue based on the null distribution (resampling). If <code>empiricPvalue=TRUE</code> , the empirical p-values are computed instead.
sigThresh	[numeric]: statistical threshold for signal (e.g. <code>sigThresh = 0.05</code> for a given GWAS signal) used to compute an Enrichment Ratio.
MAFpool	[vector(numeric)]: either a numeric vector giving the breaks points of intervals into which SNP's MAF (Minor Allele Frequency) is to be split.
mc.cores	[numeric]: the number of cores to use (default is <code>mc.cores=1</code> ), i.e. at most how many child processes will be run simultaneously. Must be at least one, and parallelization requires at least two cores.
onlyGenome	[logical]: <code>onlyGenome=TRUE</code> (default) compute resampling step for all chromosomes.

## Value

Return the object given in argument, updated by the resampling results.

## Author(s)

Mickael Canouil <mickael.canouil@good.ibl.fr>

## See Also

Overview : [snpEnrichment-package](#)

Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)

Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#), [enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)

Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

## Examples

```
## Not run: data(toyEnrichment)
reSample(object = toyEnrichment,
          nSample = 10,
          empiricPvalue = TRUE,
          MAFpool = c(0.05, 0.10, 0.2, 0.3, 0.4, 0.5),
          onlyGenome = TRUE)
toyEnrichment
## End(Not run)
```

---

toyEnrichment-dataset *Toy dataset with SNP data*

---

### Description

This data set gives an [Enrichment](#) object after, [initFiles](#) and [readEnrichment](#) is ran. Compute LD for all SNPs in `snpListDir` files two by two. Genome Build 37.3 (hg19).

### Usage

```
data(toyEnrichment)
toyEnrichment
```

### Format

See class [Enrichment](#) for details about the format.

### Author(s)

Mickael Canouil <mickael.canouil@good.ibl.fr>

### See Also

Overview : [snpEnrichment-package](#)  
Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

---

transcript-dataset *Transcript information in order to check the CIS status for SNPs*

---

### Description

This dataset is used by [readEnrichment](#) and [compareEnrichment](#) in order to check the CIS status for each SNP of signal. Genome Build 37.3 (hg19).

### Usage

```
data(transcript)
transcript
```

### Format

See class [readEnrichment](#) and [compareEnrichment](#) for details about how to use this dataset.



**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)

Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)

Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)

Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

---

 writeLD

*Linkage Disequilibrium (LD) computation with PLINK*

---

**Description**

`writeLD` write a '.ld' file for each chromosomes which contains the LD ( $r^2$ ).

**Usage**

```
writeLD(pattern = "Chrom", snpInfoDir, signalFile,
        ldDir = NULL, ldThresh = 0.8, depth = 1000,
        mc.cores = 1)
```

**Arguments**

pattern	[character]: character string containing a expression to be matched with all chromosomes files (e.g."Chrom" for files which start by "Chrom" followed by the chromosome number).
snpInfoDir	[character]: character string naming a directory containing the reference data in a PLINK format (*.bed, *.bim and *.fam).
signalFile	[character]: the name of the signal file which the data are to be read from (2 columns: "SNP" and "PVALUE"). Each row of the table appears as one line of the file. If it does not contain an <code>_absolute_</code> path, the file name is <code>_relative_</code> to the current working directory, <code>getwd</code> . The fields separator character have to be a space " " or a tabulation "\t".
ldDir	[character]: character string naming a directory where the linkage Disequilibrium files should be written (default <code>ldDir=NULL</code> is in temporary directory).
ldThresh	[numeric]: threshold value for LD calculation.
depth	[numeric]: this parameter is mandatory and controls the maximum lag between SNPs considered.
mc.cores	[numeric]: the number of cores to use (default is <code>mc.cores=1</code> ), i.e. at most how many child processes will be run simultaneously. Must be at least one, and parallelization requires at least two cores.

**Value**

One ".ld" file per chromosome is returned by `writeLD` in `snpInfoDir` directory.

**Note**

The LD computation can take a long time depending on number of SNPs in `signalFile`. It is recommended to save LD results in a directory (`ldDir`) which is not a temporary directory.

**Author(s)**

Mickael Canouil <mickael.canouil@good.ibl.fr>

**See Also**

Overview : [snpEnrichment-package](#)  
Classes : [Enrichment](#), [Chromosome](#), [EnrichSNP](#)  
Methods : [plot](#), [reSample](#), [getEnrichSNP](#), [excludeSNP](#), [compareEnrichment](#),  
[enrichment](#), [is.enrichment](#), [chromosome](#), [is.chromosome](#)  
Functions : [initFiles](#), [writeLD](#), [readEnrichment](#)

**Examples**

```
## Not run: signalFile <- system.file("extdata/Signal/toySignal.txt",  
                                   package = "snpEnrichment")  
snpInfoDir <- system.file("extdata/snpInfo",  
                          package = "snpEnrichment")  
writeLD(pattern = "Chrom", snpInfoDir, signalFile,  
        ldDir = NULL, ldThresh = 0.8, mc.cores = 1)  
## End(Not run)
```

# Index

- \*Topic **Enrichment**
    - compareEnrichment, 6
    - snpEnrichment-package, 2
    - writeLD, 25
  - \*Topic **GC**
    - GC, 12
  - \*Topic **chromosome**
    - Chromosome-class, 4
    - is.chromosome, 15
  - \*Topic **classes**
    - Chromosome-class, 4
    - Enrichment-class, 8
    - EnrichSNP-class, 10
  - \*Topic **class**
    - Chromosome-class, 4
    - Enrichment-class, 8
    - EnrichSNP-class, 10
  - \*Topic **compareEnrichment**
    - compareEnrichment, 6
  - \*Topic **core**
    - mclapply2, 17
  - \*Topic **datasets**
    - toyEnrichment-dataset, 24
    - transcript-dataset, 24
  - \*Topic **enrichSNP**
    - EnrichSNP-class, 10
  - \*Topic **enrichment**
    - Enrichment-class, 8
    - is.enrichment, 16
  - \*Topic **excludeSNP**
    - excludeSNP, 11
  - \*Topic **garbage**
    - GC, 12
  - \*Topic **getEnrichSNP**
    - getEnrichSNP, 13
  - \*Topic **initFiles**
    - initFiles, 14
    - readEnrichment, 20
    - writeLD, 25
  - \*Topic **initialize**
    - initFiles, 14
    - readEnrichment, 20
  - \*Topic **is.chromosome**
    - is.chromosome, 15
  - \*Topic **is.enrichment**
    - is.enrichment, 16
  - \*Topic **is**
    - is.chromosome, 15
    - is.enrichment, 16
  - \*Topic **mclapply2**
    - mclapply2, 17
  - \*Topic **methods**
    - excludeSNP, 11
    - getEnrichSNP, 13
    - plot-methods, 18
    - print-methods, 19
    - reSample, 22
  - \*Topic **package**
    - snpEnrichment-package, 2
  - \*Topic **parallel**
    - mclapply2, 17
  - \*Topic **plot**
    - plot-methods, 18
  - \*Topic **print**
    - print-methods, 19
  - \*Topic **reSample**
    - reSample, 22
  - \*Topic **readEnrichment**
    - readEnrichment, 20
  - \*Topic **snpEnrichment**
    - snpEnrichment-package, 2
    - toyEnrichment-dataset, 24
    - transcript-dataset, 24
  - \*Topic **toyEnrichment**
    - toyEnrichment-dataset, 24
  - \*Topic **writeLD**
    - readEnrichment, 20
- [, Chromosome, ANY, ANY-method

- (Chromosome-class), 4
- [,Chromosome-method (Chromosome-class), 4
- [,EnrichSNP,ANY,ANY,ANY-method (EnrichSNP-class), 10
- [,EnrichSNP-method (EnrichSNP-class), 10
- [,Enrichment,ANY,ANY,ANY-method (Enrichment-class), 8
- [,Enrichment-method (Enrichment-class), 8
- [<-,Chromosome,ANY,ANY,ANY-method (Chromosome-class), 4
- [<-,Chromosome-method (Chromosome-class), 4
- [<-,EnrichSNP,ANY,ANY,ANY-method (EnrichSNP-class), 10
- [<-,EnrichSNP-method (EnrichSNP-class), 10
- [<-,Enrichment,ANY,ANY,ANY-method (Enrichment-class), 8
- [<-,Enrichment-method (Enrichment-class), 8
- as.list, 17
- Chromosome, 3–12, 14–17, 19–21, 23–26
- Chromosome (Chromosome-class), 4
- chromosome, 3, 5–7, 9, 11, 12, 14–17, 19–21, 23–26
- chromosome (Chromosome-class), 4
- chromosome,ANY-method (Chromosome-class), 4
- Chromosome-class, 4
- chromosome-methods (Chromosome-class), 4
- compareEnrichment, 3, 6, 6, 7, 9, 11, 12, 14–17, 19–21, 23–26
- compareEnrichment,ANY-method (compareEnrichment), 6
- compareEnrichment,Enrichment,Enrichment,ANY-method (compareEnrichment), 6
- compareEnrichment-methods (compareEnrichment), 6
- Enrichment, 3, 5–26
- Enrichment (Enrichment-class), 8
- enrichment, 3, 6–9, 11, 12, 14–17, 19–21, 23–26
- enrichment (Enrichment-class), 8
- enrichment,ANY-method (Enrichment-class), 8
- Enrichment-class, 8
- enrichment-methods (Enrichment-class), 8
- EnrichSNP, 3, 5–7, 9–12, 14–17, 19–21, 23–26
- EnrichSNP (EnrichSNP-class), 10
- EnrichSNP-class, 10
- excludeSNP, 3, 6, 7, 9, 11, 11, 12, 14–17, 19–21, 23–26
- excludeSNP,ANY-method (excludeSNP), 11
- excludeSNP,Enrichment-method (excludeSNP), 11
- excludeSNP-methods (excludeSNP), 11
- GC, 12, 12, 13
- getEnrichSNP, 3, 6, 7, 9, 11–13, 13, 14–17, 19–21, 23–26
- getEnrichSNP,ANY-method (getEnrichSNP), 13
- getEnrichSNP,Enrichment-method (getEnrichSNP), 13
- getEnrichSNP-methods (getEnrichSNP), 13
- initFiles, 3, 6, 7, 9, 11, 12, 14, 14, 15–17, 19–26
- is.chromosome, 3, 6, 7, 9, 11, 12, 14, 15, 15, 16, 17, 19–21, 23–26
- is.chromosome,ANY-method (is.chromosome), 15
- is.chromosome-methods (is.chromosome), 15
- is.enrichment, 3, 6, 7, 9, 11, 12, 14–16, 16, 17, 19–21, 23–26
- is.enrichment,ANY-method (is.enrichment), 16
- is.enrichment-methods (is.enrichment), 16
- mclapply2, 17, 17
- plot, 3, 6, 7, 9, 11, 12, 14–21, 23–26
- plot (plot-methods), 18
- plot,Enrichment,ANY-method (plot-methods), 18
- plot,Enrichment-method (plot-methods), 18
- plot-methods, 18
- print, 19
- print (print-methods), 19

print,Chromosome-method  
    (print-methods), 19

print,Enrichment-method  
    (print-methods), 19

print,EnrichSNP-method  
    (EnrichSNP-class), 10

print-methods, 19

  

readEnrichment, 3, 5–7, 9, 11–17, 19, 20, 20,  
    21–26

reSample, 3, 5–9, 11, 12, 14–17, 19–22, 22,  
    23–26

reSample,ANY-method (reSample), 22

reSample,Chromosome-method (reSample),  
    22

reSample,Enrichment-method (reSample),  
    22

reSample-methods (reSample), 22

  

show,Chromosome-method  
    (Chromosome-class), 4

show,Enrichment-method  
    (Enrichment-class), 8

show,EnrichSNP-method  
    (EnrichSNP-class), 10

snpEnrichment (snpEnrichment-package), 2

snpEnrichment-package, 2

  

toyEnrichment (toyEnrichment-dataset),  
    24

toyEnrichment-dataset, 24

transcript (transcript-dataset), 24

transcript-dataset, 24

  

writeLD, 3, 5–7, 9, 11, 12, 14–17, 19–21,  
    23–25, 25, 26