

# Package ‘PlasmaMutationDetector’

January 20, 2025

**Type** Package

**Title** Tumor Mutation Detection in Plasma

**Version** 1.7.2

**Date** 2018-05-16

**Author** Yves Rozenholc, Nicolas Pécuchet, Pierre Laurent-Puig

**Maintainer** Yves Rozenholc <yves.rozenholc@parisdescartes.fr>

**Description** Aims at detecting single nucleotide variation (SNV) and insertion/deletion (INDEL) in circulating tumor DNA (ctDNA), used as a surrogate marker for tumor, at each base position of an Next Generation Sequencing (NGS) analysis. Mutations are assessed by comparing the minor-allele frequency at each position to the measured PER in control samples.

**License** MIT + file LICENSE

**Imports** S4Vectors (>= 0.16.0), Rsamtools (>= 1.30.0), rtracklayer (>= 1.38.0), robustbase (>= 0.92-8), SummarizedExperiment (>= 1.8.0)

**Depends** R (>= 3.4.0), ggplot2 (>= 2.2.0), grid (>= 3.4.0), GenomicRanges (>= 1.30.0), VariantAnnotation (>= 1.24.0)

**Encoding** UTF-8

**RoxygenNote** 6.0.1

**LazyData** true

**NeedsCompilation** no

**Repository** CRAN

**Date/Publication** 2018-06-11 07:43:09 UTC

## Contents

background_error_rate . . . . .	2
BuildCtrlErrorRate . . . . .	3
DetectPlasmaMutation . . . . .	4
hotspot . . . . .	6
LoadBackgroundErrorRate . . . . .	6

MAF_from_BAM . . . . .	7
positions_ranges . . . . .	9
PrepareLibrary . . . . .	9

<b>Index</b>	<b>12</b>
--------------	-----------

---

background\_error\_rate *The package provide the SNV and INDEL PERs computed for the Ion AmpliSeq™ Colon and Lung Cancer Panel v2 from 29 controls in a table available in the data file background\_error\_rate.txt.*

---

## Description

This table contains 9 variables for each genomic position

- chrpos, char, of the form chrN:XXXXXXXXXX defining genomic position
- N0, integer, the coverage in the controls
- E0, integer, the number of errors in the controls
- p.sain, numeric, the ratio E0/N0
- up.sain, numeric, the 95th quantile of the Binomial with parameter N0 and E0/N0
- E0indel, integer, the amount of indel
- indel.p.sain, numeric, the ratio E0indel/N0
- indel.up.sain, numeric, the 95th quantile of the Binomial with parameter N0 and E0indel/N0
- hotspot, char, either 'Non-hotspot' or 'Hotspot' depending if the genomic position is known as hotspot or not.

## Usage

```
data(background_error_rate)
```

## Author(s)

N. Pécuchet, P. Laurent-Puig and Y. Rozenholc

## References

*Analysis of base-position error rate of next-generation sequencing to detect tumor mutations in circulating DNA* N. Pécuchet, Y. Rozenholc, E. Zonta, D. Pietraz, A. Didelot, P. Combe, L. Gibault, J-B. Bachet, V. Taly, E. Fabre, H. Blons, P. Laurent-Puig in *Clinical Chemistry*

## See Also

BuildCtrlErrorRate

---

BuildCtrlErrorRate     *function BuildCtrlErrorRate*

---

### Description

Compute the SNV Position-Error Rates and INDEL Position-Error Rates from control samples (available in the control directory `ctrl.dir`). This function requires MAF files, that will be automatically generated if not present in the specified control folder. SNV PER is computed as the sum in control samples of SNV background counts / sum in control samples of depths where SNV background counts = depth - major allele count. INDEL PER is computed as sum in control samples of INDEL background counts / sum in control samples of depths where INDEL background counts = sum of insertion and deletion counts.

### Usage

```
BuildCtrlErrorRate(ctrl.dir = "Plasma ctrl/", bai.ext = ".bai",
  pos_ranges.file = NULL, hotspot.file = NULL, force = FALSE,
  output.dir = ctrl.dir)
```

### Arguments

<code>ctrl.dir</code>	char, foldername containing the control files (default 'Plasma ctrl/'). The typical folder hierarchy will consist of 'Plasma ctrl/rBAM'
<code>bai.ext</code>	char, filename extension of the bai files (default '.bai')
<code>pos_ranges.file</code>	char, name of the Rdata file containing the three variables <code>pos_ind</code> , <code>pos_snp</code> and <code>pos_ranges</code> as build by the function <code>PrepareLibrary</code> . Default NULL, use the <code>position_ranges.rda</code> provided, used for our analysis.
<code>hotspot.file</code>	char, name of the text file containing a list of the genomic positions of the hotspots (default NULL, read the provide <code>hotspot.txt</code> , see <code>hotspot</code> )
<code>force</code>	boolean, (default FALSE) if TRUE force all computations to all files including already processed ones
<code>output.dir</code>	char, name of the folder to save results (default <code>ctrl.dir</code> ).

### Value

the number of processed files

### Author(s)

N. Pécuchet, P. Laurent-Puig and Y. Rozenholc

### References

*Analysis of base-position error rate of next-generation sequencing to detect tumor mutations in circulating DNA* N. Pécuchet, Y. Rozenholc, E. Zonta, D. Pietraz, A. Didelot, P. Combe, L. Gibault, J-B. Bachet, V. Taly, E. Fabre, H. Blons, P. Laurent-Puig in *Clinical Chemistry*

**Examples**

```
## Not run:
ctrl.dir = system.file("extdata", "4test_only/ctrl/", package = "PlasmaMutationDetector")
if (substr(ctrl.dir,nchar(ctrl.dir),nchar(ctrl.dir))!='/')
  ctrl.dir = paste0(ctrl.dir, '/') # TO RUN UNDER WINDOWS
BuildCtrlErrorRate(ctrl.dir,output.dir=paste0(tempdir(), '/'))

## End(Not run)
```

---

DetectPlasmaMutation *function DetectPlasmaMutation*

---

**Description**

This is the main function of the package that calls mutations by comparing at each genomic position the SNV or INDEL frequencies computed in one tested sample to the SNV or INDEL Position-Error Rates computed from several control samples by a binomial test. An outlier detection is performed among all intra-sample p-values to call a mutation. For users wishing to develop their own analysis for other sequencing panel, it requires recalibrated BAM files control samples to be processed to compute the Position-Error Rates stored in a file specified in `ber.ctrl.file`.

**Usage**

```
DetectPlasmaMutation(patient.dir = "./", patient.name = NULL,
  pos_ranges.file = NULL, ber.ctrl.file = NULL, bai.ext = ".bai",
  n.trim = 8, cov.min = 0, force = FALSE, show.more = FALSE,
  qcutoff.snv = 0.95, qcutoff.indel = 0.99, cutoff.sb.ref = 0.1,
  cutoff.sb.hotspot = 3.1, cutoff.sb.nonhotspot = 2.5,
  hotspot.indel = "chr7:55227950:55249171", output.dir = patient.dir)
```

**Arguments**

<code>patient.dir</code>	char, foldername containing the rBAM folder of the patients. The typical folder hierarchy will consist of 'Plasma/rBAM'
<code>patient.name</code>	char, filename of the patient .bam file(s) (default NULL read all patients in folder <code>patient.dir</code> )
<code>pos_ranges.file</code>	char, name of the Rdata file containing the three variables <code>pos_ind</code> , <code>pos_snp</code> , <code>pos_ranges</code> as build by the function <code>PrepareLibrary</code> . Default NULL, use the <code>position_ranges.rda</code> provides that we used for our analysis.
<code>ber.ctrl.file</code>	char, pathname of the file providing the background error rates obtained from the controls (default NULL use the provided background error rates obtained from our 29 controls). See <code>background_error_rate.txt</code> data and <code>BuildCtrlErrorRate</code> function.
<code>bai.ext</code>	char, filename extension of the bai files (default '.bai')

n.trim	integer, number of base positions trimmed at the ends of each amplicon (default 8)
cov.min	integer, minimal coverage required at each position (default 0)
force	boolean, (default FALSE) if TRUE force all computations to all files including already processed ones
show.more	boolean, (default FALSE show only detected positions) if TRUE additional annotations on result plots are given for non-significant mutations
qcutoff.snv	numeric, proportion of kept base positions ranged by increasing 95th percentile SNV PER in control samples (default 0.95)
qcutoff.indel	numeric, proportion of kept base positions ranged by increasing 95th percentile INDEL PER in control samples (default 0.99)
cutoff.sb.ref	numeric, exclude reference positions without cutoff < strand bias < 1-cutoff (default 0.1) (see Supplementary Materials in References)
cutoff.sb.hotspot	numeric, exclude hotspot positions with Symmetric Odds Ratio test > cutoff (default 3.1) (see Supplementary Materials in References)
cutoff.sb.nonhotspot	numeric, exclude non-hotspot positions with Symmetric Odds Ratio test > cutoff (default 2.5) (see Supplementary Materials in References)
hotspot.indel	char, a vector containing the known positions of hotspot deletion/insertion defined as chrX:start:end (default 'chr7:55227950:55249171')
output.dir	char, name of the folder to save results (default patient.dir).

**Value**

the number of processed patients

**Author(s)**

N. Pécuchet, P. Laurent-Puig and Y. Rozenholc

**References**

*Analysis of base-position error rate of next-generation sequencing to detect tumor mutations in circulating DNA* N. Pécuchet, Y. Rozenholc, E. Zonta, D. Pietraz, A. Didelot, P. Combe, L. Gibault, J-B. Bachet, V. Taly, E. Fabre, H. Blons, P. Laurent-Puig in *Clinical Chemistry*

**Examples**

```
patient.dir=system.file("extdata", "4test_only/case/", package="PlasmaMutationDetector")
if (substr(patient.dir, nchar(patient.dir), nchar(patient.dir)) != '/')
  patient.dir = paste0(patient.dir, '/') # TO RUN UNDER WINDOWS
DetectPlasmaMutation(patient.dir, output.dir=paste0(tempdir(), '/'))
```

---

hotspot	<i>The package provide a list of known hotspot positions located on the amplicons of the Ion AmpliSeq™ Colon and Lung Cancer Panel v2 as a txt file hotspot.txt which contains a vector/variable —named chrpos (first row)— of chars, of the form chrN:XXXXXXXXXX defining genomic positions.</i>
---------	---

---

### Description

The package provide a list of known hotspot positions located on the amplicons of the Ion AmpliSeq™ Colon and Lung Cancer Panel v2 as a txt file hotspot.txt which contains a vector/variable —named chrpos (first row)— of chars, of the form chrN:XXXXXXXXXX defining genomic positions.

### Usage

```
data(hotspot)
```

### Author(s)

N. Pécuchet, P. Laurent-Puig and Y. Rozenholc

### References

*Analysis of base-position error rate of next-generation sequencing to detect tumor mutations in circulating DNA* N. Pécuchet, Y. Rozenholc, E. Zonta, D. Pietraz, A. Didelot, P. Combe, L. Gibault, J-B. Bachet, V. Taly, E. Fabre, H. Blons, P. Laurent-Puig in *Clinical Chemistry*

---

LoadBackgroundErrorRate

*function LoadBackgroundErrorRate*

---

### Description

This function will load the background error rates created from the controls using the function BuildCtrlErrorRate

### Usage

```
LoadBackgroundErrorRate(pos_ranges.file, ber.ctrl.file, n.trim)
```

**Arguments**

<code>pos_ranges.file</code>	char, name of the Rdata file containing the three variables <code>pos_ind</code> , <code>pos_snp</code> , <code>pos_ranges</code> as build by the function <code>PrepareLibrary</code> . Default NULL, use the <code>position_ranges.rda</code> provides that we used for our analysis.
<code>ber.ctrl.file</code>	char, pathname of the file providing the background error rates obtained from the controls (default NULL use the provided background error rates obtained from our 29 controls). See <code>background_error_rate.txt</code> data and <code>BuildCtrlErrorRate</code> function.
<code>n.trim</code>	integer, number of base positions trimmed at the ends of each amplicon (default 8)

**Value**

the adapted background error rate

**Author(s)**

N. Pécuchet, P. Laurent-Puig and Y. Rozenholc

**References**

*Analysis of base-position error rate of next-generation sequencing to detect tumor mutations in circulating DNA* N. Pécuchet, Y. Rozenholc, E. Zonta, D. Pietraz, A. Didelot, P. Combe, L. Gibault, J-B. Bachet, V. Taly, E. Fabre, H. Blons, P. Laurent-Puig in *Clinical Chemistry*

---

MAF_from_BAM	<i>function MAF_from_BAM</i>
--------------	------------------------------

---

**Description**

Read BAM files and create MAF file. BAMfiles are stored in a sub-folder 'rBAM'. MAF files are intermediate files stored in a sub-folder '/BER'. MAF files contain the raw counts of A,T,C,G, insertion, deletion, insertion>2bp, deletion>2bp for strand plus and stand minus. Note : we strongly recommend to externally recalibrate BAM files using tools like GATK.

**Usage**

```
MAF_from_BAM(study.dir = "Plasma/", input.filesnames = NULL,  
  bai.ext = ".bai", pos_ranges.file = NULL, force = FALSE,  
  output.dir = study.dir)
```

**Arguments**

<code>study.dir</code>	char, name of the folder containing the rBAM directory (default 'Plasma/'). The typical folder hierarchy will consist of 'Plasma/rBAM'
<code>input.filesnames</code>	a vector of char (default NULL), the names of the BAM files to process. If NULL all BAM files in the rBAM folder will be processed
<code>bai.ext</code>	char, filename extension of the bai files (default '.bai')
<code>pos_ranges.file</code>	char, name of the Rdata file containing the three variables <code>pos_ind</code> , <code>pos_snp</code> and <code>pos_ranges</code> as build by the function <code>PrepareLibrary</code> . Default NULL, use the <code>position_ranges.rda</code> provided, used for our analysis.
<code>force</code>	boolean, (default FALSE) if TRUE force all computations to all files including already processed ones
<code>output.dir</code>	char, name of the folder to save results (default <code>study.dir</code> ).

**Value**

the path/names of the MAF files

**Author(s)**

N. Pécuchet, P. Laurent-Puig and Y. Rozenholc

**References**

*Analysis of base-position error rate of next-generation sequencing to detect tumor mutations in circulating DNA* N. Pécuchet, Y. Rozenholc, E. Zonta, D. Pietraz, A. Didelot, P. Combe, L. Gibault, J-B. Bachet, V. Taly, E. Fabre, H. Blons, P. Laurent-Puig in *Clinical Chemistry*

**Examples**

```
## Not run:
ctrl.dir = system.file("extdata", "4test_only/ctrl/", package = "PlasmaMutationDetector")
if (substr(ctrl.dir, nchar(ctrl.dir), nchar(ctrl.dir)) != '/')
  ctrl.dir = paste0(ctrl.dir, '/') # TO RUN UNDER WINDOWS
MAF_from_BAM(ctrl.dir, force=TRUE, output.dir=paste0(tempdir(), '/'))

## End(Not run)
```



---

positions_ranges	<i>The package provide the positions and ranges computed for the Ion AmpliSeq™ Colon and Lung Cancer Panel v2 as a Rdata file positions_ranges.rda.</i>
------------------	---

---

## Description

This file contains 4 variables

- pos\_ind, vector of chars, of the form chrN:XXXXXXXXXX defining genomic positions of the Ion AmpliSeq™ Colon and Lung Cancer Panel v2
- pos\_snp, vector of chars, of the form chrN:XXXXXXXXXX defining the known snp genomic positions
- pos\_ranges, GRanges object, describing the 92 amplicons of the Ion AmpliSeq™ Colon and Lung Cancer Panel v2

## Usage

```
data(positions_ranges)
```

## Author(s)

N. Pécuchet, P. Laurent-Puig and Y. Rozenholc

## References

*Analysis of base-position error rate of next-generation sequencing to detect tumor mutations in circulating DNA* N. Pécuchet, Y. Rozenholc, E. Zonta, D. Pietraz, A. Didelot, P. Combe, L. Gibault, J-B. Bachet, V. Taly, E. Fabre, H. Blons, P. Laurent-Puig in *Clinical Chemistry*

## See Also

Prepare\_Library

---

PrepareLibrary	<i>function PrepareLibrary</i>
----------------	--------------------------------

---

## Description

Define the Genomic Ranges and Genomic Positions covered by the AmpliSeq™ Panel to include in the study and define SNP positions to exclude from the study. Trimming amplicon ends is performed if specified. This function is mostly useful if you want to add some SNP positions which are not existing in the positions\_ranges.rda file provided within the package. It is provided to be able to reconstruct positions\_ranges.rda data.

**Usage**

```
PrepareLibrary(info.dir = "Info/", bed.filename = "lungcolonV2.bed.txt",
  snp.filename = "ExAC.r0.3.sites.vep.vcf.gz",
  snp.extra = c("chr2:212812097", "chr4:1807909", "chr7:140481511",
  "chr14:105246474", "chr18:48586344", "chr19:1223055"),
  output.name = "positions_ranges.rda", output.dir = info.dir,
  load.from.broad.insitute = FALSE)
```

**Arguments**

<code>info.dir</code>	char, name of the folder containing the library information files (default 'Info')
<code>bed.filename</code>	char, name of a BED table (tab-delimited) describing the Panel (with first 3 columns: "chr" (ex:chr1), "start position" (ex:115252190), "end position" (ex:115252305), i.e. the Ion AmpliSeq™ Colon and Lung Cancer Research Panel v2 (default 'lungcolonV2.bed.txt' as provided in the inst/extdata/Info folder of the package).
<code>snp.filename</code>	char, name of the vcf file describing known SNP positions, obtained from ftp://ftp.broadinstitute.org/pub/ExAC (default 'ExAC.r0.3.sites.vep.vcf.gz'). It requires a corresponding TBI file to be in the same folder (obtained from ftp://ftp.broadinstitute.org/pub/ExAC_release/release0.3/ExAC.r0.3.sites.vep.vcf.gz).
<code>snp.extra</code>	a vector of char, a vector of extra known snp positions manually curated (ex:"chrN:XXXXXXXXXX")
<code>output.name</code>	char, filename to save pos_ind and pos_snp (default 'positions_ranges.rda')
<code>output.dir</code>	char, directory where to save pos_ind and pos_snp (default info.dir)
<code>load.from.broad.insitute</code>	boolean, if TRUE load snp.filename from Broad Institute ftp server otherwise use the file positions_ranges_broad.rda (default FALSE)

**Value**

Save the following variables in a .rda file defined by `output.name` in the folder defined by `output.dir`:

- `pos_ranges`, a GRanges descriptor of amplicon positions
- `pos_ind`, a vector of char "chrN:XXXXXXXXXX", defining ALL index positions
- `pos_snp`, a vector of char "chrN:XXXXXXXXXX", defining SNP positions

**Author(s)**

N. Pécuchet, P. Laurent-Puig and Y. Rozenholc

**References**

*Analysis of base-position error rate of next-generation sequencing to detect tumor mutations in circulating DNA* N. Pécuchet, Y. Rozenholc, E. Zonta, D. Pietraz, A. Didelot, P. Combe, L. Gibault, J-B. Bachet, V. Taly, E. Fabre, H. Blons, P. Laurent-Puig in *Clinical Chemistry*

**See Also**

`positions_ranges`,

**Examples**

```
bad.pos = "chr7:15478"  
PrepareLibrary(info.dir='./',snp.extra=bad.pos,output.dir=paste0(tempdir(),'/'))
```

# Index

## \* data

- background\_error\_rate, 2
- hotspot, 6
- positions\_ranges, 9

background\_error\_rate, 2  
BuildCtrlErrorRate, 3

DetectPlasmaMutation, 4

hotspot, 6

LoadBackgroundErrorRate, 6

MAF\_from\_BAM, 7

pos\_ind (positions\_ranges), 9  
pos\_ranges (positions\_ranges), 9  
pos\_snp (positions\_ranges), 9  
positions\_ranges, 9  
PrepareLibrary, 9