

# Package: diffdriver (via r-universe)

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**Type** Package

**Title** Identify Differential Selection

**Version** 0.1.7

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**Description** Tests for context-dependent selection on cancer driver genes using somatic mutation data. The package implements the DiffDriver statistical framework to assess whether the strength of positive selection on mutations in a driver gene is associated with tumor- or individual-level context variables, such as clinical traits, genomic features, or immune microenvironment subtypes. DiffDriver estimates individual- and position-specific background mutation rates, models selection as a deviation from the background rate using functional annotations, and tests context effects through a latent-variable logistic model. It provides utilities for preparing mutation and annotation data, fitting differential-selection models, running gene-level association tests, summarizing candidate genes, and visualizing mutation patterns. The method is described in Zhou et al. (2026) ``Detecting context-dependent selection on cancer driver genes with DiffDriver" <doi:10.64898/2026.04.06.716771>.

**URL** <https://szhaolab.github.io/diffdriver/>,  
<https://github.com/szhaolab/diffdriver>

**License** MIT + file LICENSE

**Encoding** UTF-8

**Imports** Matrix, stats, data.table, brglm, fastTopics, SQUAREM

**Suggests** knitr, rmarkdown, logging, testthat (>= 3.0.0)

**RoxygenNote** 7.2.3

**Config/testthat/edition** 3

**Depends** R (>= 3.5.0)

**VignetteBuilder** knitr

**NeedsCompilation** no

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**Repository** <https://cran.r-universe.dev>

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ddmcode	<i>Add intercept, if have functypecode, then code and move to the front. different from driverMAPS! only allows functypecode =7   8 when functypecode is included in selectvars.</i>
---------	--

---

## Description

Add intercept, if have functypecode, then code and move to the front. different from driverMAPS! only allows functypecode =7 ||8 when functypecode is included in selectvars.

## Usage

```
ddmcode(matrixlist, selectvars, functypecodelevel = NULL)
```

## Arguments

matrixlist	A list of mutation or annotation matrices.
selectvars	Variables selected for the model.
functypecodelevel	Functional annotation code level.

---

ddmodel	<i>diffDriver model</i>
---------	-------------------------

---

## Description

This model is applied on data of a single gene. It will infer effect size for both sample-level variable and positional level functional annotations. We used an EM algorithm to infer parameters.

## Usage

```
ddmodel(mut, e, mr, fe, label, ...)
```

## Arguments

mut	a matrix of mutation status 0 or 1, rows positions, columns are samples.
e	a vector, phenotype of each sample, should match the columns of mut and mr
mr	a matrix, mutation rate of each sample at each mutation (log scale) that is not dependent on sample level factor
fe	a vector, increased mutation rate at each position, depending on e (log scale), should match the rows of mut and mr
label	A vector or factor of length nrow(mut) identifying mutation-position rows to be grouped and aggregated.
...	Additional arguments passed to internal functions.

## Value

A list with the following elements:

**pvalue** The likelihood-ratio test p-value comparing the null model without a context effect to the alternative model with a context effect.

**res.null** A list returned by the null-model fit. It contains loglikelihood, beta0, and alpha.

**res.alt** A list returned by the alternative-model fit. It contains loglikelihood, beta0, and alpha.

## Examples

```
# Synthetic mutation matrix: rows are positions and columns are samples.
mut <- matrix(0, nrow = 6, ncol = 20)

mut[1, 1] <- 1
mut[2, 5] <- 1

mut[c(1, 2), 11] <- 1
mut[c(2, 3), 12] <- 1
mut[c(3, 4), 13] <- 1
mut[c(4, 5), 14] <- 1
```

```

mut[c(5, 6), 15] <- 1
mut[c(1, 6), 16] <- 1
mut[c(1, 3), 17] <- 1
mut[c(2, 4), 18] <- 1
mut[c(3, 5), 19] <- 1
mut[c(4, 6), 20] <- 1

e <- c(rep(0, 10), rep(1, 10))

# Mutation rates and functional effects are on the log scale.
mr <- matrix(log(0.002), nrow = nrow(mut), ncol = ncol(mut))
fe <- rep(log(3), nrow(mut))

# Each row represents a separate mutation position.
label <- factor(seq_len(nrow(mut)))

result <- ddmodel(mut, e, mr, fe, label = label)
result$pvalue

```

---

ddmodel\_binary

*diffDriver model only for binary phenotype.*


---

## Description

This function uses the model as `cmodel.frac`, but generalizes to take more than 1 functional categories. This model is applied on data of a single gene

## Usage

```
ddmodel_binary(mut, e, bmr, fe)
```

## Arguments

<code>mut</code>	a matrix of mutation status 0 or 1
<code>e</code>	a vector, phenotype of each sample, should match the columns of <code>mut</code> and <code>bmr</code>
<code>bmr</code>	a matrix, background mutation rate of each sample at each mutation (log scale)
<code>fe</code>	a vector, increased mutation rate at each mutation, due to functional effect (log scale), should match the rows of <code>mut</code> and <code>bmr</code>

## Value

A list with the following elements:

`pvalue` The likelihood-ratio test p-value comparing the null and alternative binary-phenotype models.

`null.eta0` The fitted null-model parameter.

`alt.eta` The fitted alternative-model parameters for the two phenotype groups.

null.ll The maximized log-likelihood under the null model.  
 alt.ll The maximized log-likelihood under the alternative model.  
 all A named numeric vector containing pvalue, null\_para, e=0, e=1, null\_likelihood, and alt\_likelihood.

## Examples

```

# Synthetic mutation matrix: rows are positions and columns are samples.
mut <- matrix(
  c(
    1, 0, 0, 0, 0, 0, 1, 0,
    0, 0, 0, 1, 0, 0, 0, 0,
    0, 1, 0, 0, 0, 0, 0, 1
  ),
  nrow = 3,
  byrow = TRUE
)
e <- c(0, 0, 0, 0, 1, 1, 1, 1)

# Background mutation rates and functional effects are on the log scale.
bmr <- matrix(log(0.001), nrow = nrow(mut), ncol = ncol(mut))
fe <- log(c(2, 2, 2))

result <- ddmodel_binary(mut, e, bmr, fe)
result$pvalue

```

---

ddmodel\_binary\_simple *diffDriver model only for binary phenotype, assuming bmr the same across samples.*

---

## Description

This function uses the model as `cmodel.frac`, but generalizes to take more than 1 functional categories. This model is applied on data of a single gene. This should give the same results as `ddmodel_binary` defined above.

## Usage

```
ddmodel_binary_simple(mut, e, bmr, fe)
```

## Arguments

mut	a matrix of mutation status 0 or 1
e	a vector, phenotype of each sample, should match the columns of mut and bmr
bmr	a matrix, background mutation rate of each sample at each mutation (log scale), as we assume bmr the same across samples, only the first column will be used.
fe	a vector, increased mutation rate at each mutation, due to functional effect (log scale), should match the rows of mut and bmr

**Value**

A list with the following elements:

`pvalue` The likelihood-ratio test p-value comparing the null and alternative binary-phenotype models.

`null.eta0` The fitted null-model parameter.

`alt.eta` The fitted alternative-model parameters for the two phenotype groups.

`null.ll` The maximized log-likelihood under the null model.

`alt.ll` The maximized log-likelihood under the alternative model.

**Examples**

```
# Synthetic mutation matrix: rows are positions and columns are samples.
mut <- matrix(
  c(
    1, 0, 0, 0, 0, 0, 1, 0,
    0, 0, 0, 1, 0, 0, 0, 0,
    0, 1, 0, 0, 0, 0, 0, 1
  ),
  nrow = 3,
  byrow = TRUE
)
e <- c(0, 0, 0, 0, 1, 1, 1, 1)

# Background mutation rates and functional effects are on the log scale.
bmr <- matrix(log(0.001), nrow = nrow(mut), ncol = ncol(mut))
fe <- log(c(2, 2, 2))

result <- ddmodel_binary_simple(mut, e, bmr, fe)
result$pvalue
```

---

diffdriver

*Run diffDriver with Input Files*

---

**Description**

This function runs `diffDriver`.

**Usage**

```
diffdriver(
  gene,
  mut,
  pheno,
  anno_dir = ".",
  k = 6,
  totalnttype = 96,
```

```

BMRmode = c("signature", "regular"),
output_dir = tempdir(),
output_prefix = "diffdriver_results"
)

```

## Arguments

gene	A vector of genes to be included in the analysis.
mut	A data frame containing all somatic mutations from the cohort. The format is: <b>Chromosome</b> <int> <b>Position</b> <int> <b>Ref</b> <chr> <b>Alt</b> <chr> <b>SampleID</b> <chr> Example: Chromosome Position Ref Alt SampleID 1 19 55653236 C T TCGA-N6-A4VE-01A-11D-A28R-08
pheno	A data frame containing sample phenotypes. The format is: #' <b>SampleID</b> <chr> <b>Phenotype1</b> <dbl> <b>Phenotype2</b> <dbl> ... .. Example: SampleID SmokingCessation BMI TCGA-N5-A4R8-01A-11D-A28R-08 0.5319630 20.0 TCGA-N5-A4RD-01A-11D-A28R-08 0.0448991 24.4
anno_dir	The path to the directory with all the annotation files. Please download from Zenodo. The default is current folder
k	The number of topics used in modeling background mutation rate. The default is 6.
totalnttype	either 9 or 96. Will look for annotation files anno9_ntypexxx_annodata.txt when totalnttype is 9 or anno96_ntypexxx_annodata when totalnttype is 96.
BMRmode	There are two modes to run diffdriver. One is "signature", this will model individual level BMR, this is the default. The second one is "regular", this assumes BMR is the same across individuals, only models position-level difference.
output_dir	The path to the output directory. Defaults to tempdir().
output_prefix	The prefix being added to the output file names.

## Value

A data frame with one row per tested gene. The returned columns include `dd.p`, `mlr.p`, `mlr.v2.p`, `fisher.p`, `binom.p`, and `lr.p`, which are p-values from DiffDriver and comparator methods; the corresponding `.fdr` columns, which contain Benjamini-Hochberg adjusted p-values; and `mut.E1`, `mut.E0`, `E1`, and `E0`, which summarize mutation and sample counts in the two phenotype groups used by the Fisher test.

**Examples**

```

phenof <- system.file(
  "extdata", "example_phenotypes.txt",
  package = "diffdriver"
)
genef <- system.file(
  "extdata", "example_gene.txt",
  package = "diffdriver"
)
mutf <- system.file(
  "extdata", "example_mutations.txt",
  package = "diffdriver"
)

pheno <- read.table(phenof, header = TRUE)
gene <- read.table(genef, header = FALSE)
mut <- read.table(mutf, header = TRUE)

## Not run:
# Download the annotation files and replace this with their directory.
annodir <- "/path/to/annodir96"

result <- diffdriver(
  gene = gene,
  mut = mut,
  pheno = pheno,
  anno_dir = annodir,
  k = 6,
  totalnttype = 96,
  BMRmode = "signature",
  output_dir = tempdir(),
  output_prefix = "diffdriver_example"
)
result

## End(Not run)

```

---

genebinom

*gene level binomial test*

---

**Description**

gene level binomial test

**Usage**

genebinom(mut, e)

**Arguments**

mut                    Mutation data.  
e                        Phenotype or context variable.

**Value**

A list with the following element:

pvalue The p-value from a gene-level binomial test comparing mutation counts between the two phenotype groups.

**Examples**

```
# Synthetic mutation matrix: rows are positions and columns are samples.
mut <- matrix(
  c(
    1, 0, 0, 1, 0, 0,
    0, 1, 0, 0, 1, 0
  ),
  nrow = 2,
  byrow = TRUE
)
e <- c(0, 0, 0, 1, 1, 1)

result <- genebinom(mut, e)
result$pvalue
```

---

genefisher

*gene level fisher's exact test*

---

**Description**

gene level fisher's exact test

**Usage**

```
genefisher(mut, e)
```

**Arguments**

mut                    Mutation data.  
e                        Phenotype or context variable.

**Value**

A list with the following elements:

`pvalue` The p-value from a gene-level Fisher's exact test.

`count` A numeric vector `c(gmutc, gmutn, mu.c, mu.n)`, where `gmutc` and `gmutn` are mutation counts in the two phenotype groups, and `mu.c` and `mu.n` are the corresponding numbers of samples.

**Examples**

```
# Synthetic mutation matrix: rows are positions and columns are samples.
mut <- matrix(
  c(
    1, 0, 0, 1, 0, 0,
    0, 1, 0, 0, 1, 0
  ),
  nrow = 2,
  byrow = TRUE
)
e <- c(0, 0, 0, 1, 1, 1)

result <- genefisher(mut, e)
result$pvalue
result$count
```

---

genelr

*gene level logistic regression*

---

**Description**

gene level logistic regression

**Usage**

```
genelr(mut, e, covariates = rep(1, length(e)))
```

**Arguments**

<code>mut</code>	Mutation data.
<code>e</code>	Phenotype or context variable.
<code>covariates</code>	Optional covariates included in the model.

**Value**

A list with the following elements:

`res` The summary object from the fitted logistic regression model `glm(e ~ dstatus + covariates, family = binomial(link = "logit"))`.

`pvalue` The p-value for the mutation-status coefficient `dstatus`.

**Examples**

```
# Synthetic mutation matrix: rows are positions and columns are samples.
mut <- matrix(
  c(
    1, 0, 0, 1, 0, 0,
    0, 1, 0, 0, 1, 0
  ),
  nrow = 2,
  byrow = TRUE
)
e <- c(0, 0, 0, 1, 1, 1)

result <- genelr(mut, e)
result$pvalue
```

mlr

*gene level multiple linear regression***Description**

gene level multiple linear regression

**Usage**

```
mlr(mut, e, covariates = rep(1, length(e)))
```

**Arguments**

mut	Mutation data.
e	Phenotype or context variable.
covariates	Optional covariates included in the model.

**Value**

A list with the following elements:

res The summary object from the fitted linear model  $\text{lm}(e \sim \text{dstatus} + \text{covariates})$ .

pvalue The p-value for the mutation-status coefficient dstatus.

**Examples**

```
# Synthetic mutation matrix: rows are positions and columns are samples.
mut <- matrix(
  c(
    1, 0, 0, 1, 0, 0,
    0, 1, 0, 0, 1, 0
  ),
  nrow = 2,
```

```
    byrow = TRUE
  )
  e <- c(0, 0, 0, 1, 1, 1)

  result <- mlr(mut, e)
  result$pvalue
```

---

mlr.v2	<i>gene level multiple linear regression, correcting for total number of mutations</i>
--------	--

---

### Description

gene level multiple linear regression, correcting for total number of mutations

### Usage

```
mlr.v2(mut, e, nmut, covariates = 1)
```

### Arguments

mut	Mutation data.
e	Phenotype or context variable.
nmut	Number of mutations.
covariates	Optional covariates included in the model.

### Value

A list with the following elements:

res The summary object from the fitted linear model  $\text{lm}(e \sim \text{dstatus} + \text{nmut} + \text{covariates})$ .

pvalue The p-value for the mutation-status coefficient `dstatus`, after adjusting for total mutation count and covariates.

### Examples

```
# Synthetic mutation matrix: rows are positions and columns are samples.
mut <- matrix(
  c(
    1, 0, 0, 1, 0, 0,
    0, 1, 0, 0, 1, 0
  ),
  nrow = 2,
  byrow = TRUE
)
e <- c(0, 0, 0, 1, 1, 1)

# Synthetic per-sample mutation counts.
```

```
nmut <- c(3, 4, 5, 6, 7, 8)

result <- mlr.v2(mut, e, nmut)
result$pvalue
```

---

 optifix

*optifix. Optimise with fixed parameters*


---

### Description

its like optim, but with fixed parameters.

### Usage

```
optifix(
  par,
  fixed,
  fn,
  gr = NULL,
  ...,
  method = c("Nelder-Mead", "BFGS", "CG", "L-BFGS-B", "SANN"),
  lower = -Inf,
  upper = Inf,
  control = list(),
  hessian = FALSE
)
```

### Arguments

par	Initial values for the parameters to be optimized.
fixed	Logical or index vector indicating fixed parameters.
fn	Function to be minimized.
gr	Optional gradient function.
...	Additional arguments passed to fn and gr.
method	Optimization method.
lower	Lower bounds for parameters.
upper	Upper bounds for parameters.
control	Control parameters passed to optimization.
hessian	Logical; whether to return the Hessian matrix.

## Details

specify a second argument 'fixed', a vector of TRUE/FALSE values. If TRUE, the corresponding parameter in fn() is fixed. Otherwise its variable and optimised over.

The return thing is the return thing from optim() but with a couple of extra bits - a vector of all the parameters and a vector copy of the 'fixed' argument.

Written by Barry Rowlingson <b.rowlingson@lancaster.ac.uk> October 2011

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and must retain the text: "Originally written by Barry Rowlingson" in comments.

---

plot\_mut                      *plot phenotype, mutation and annotation for a gene across samples*

---

## Description

plot phenotype, mutation and annotation for a gene across samples

## Usage

```
plot_mut(  
  gene_name,  
  mut,  
  pheno,  
  totalnttype = 96,  
  anno_dir = ".",  
  output_prefix = "plot",  
  output_dir = tempdir()  
)
```

## Arguments

gene_name	Gene name to plot.
mut	Mutation data.
pheno	Phenotype or context data.
totalnttype	Total number of nucleotide context types.
anno_dir	Directory containing annotation files.
output_prefix	Prefix for output files.
output_dir	Directory for output files. Defaults to tempdir().

## Value

A numeric vector returned by the final barplot() call, giving the coordinates of the plotted bars. The function is primarily called for its side effect of drawing mutation and phenotype summary plots for the selected gene.

**Examples**

```
phenof <- system.file(
  "extdata", "example_phenotypes.txt",
  package = "diffdriver"
)
mutf <- system.file(
  "extdata", "example_mutations.txt",
  package = "diffdriver"
)

pheno <- read.table(phenof, header = TRUE)
mut <- read.table(mutf, header = TRUE)

## Not run:
# Download the annotation files and replace this with their directory.
annodir <- "/path/to/annodir9"

plot_mut(
  gene_name = "PIK3CA",
  mut = mut,
  pheno = pheno,
  totalnttype = 9,
  anno_dir = annodir,
  output_dir = tempdir()
)

## End(Not run)
```

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