

Package ‘CytoGLMM’

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Type Package

Title Conditional Differential Analysis for Flow and Mass Cytometry Experiments

Version 1.17.2

Description The CytoGLMM R package implements two multiple regression strategies: A bootstrapped generalized linear model (GLM) and a generalized linear mixed model (GLMM). Most current data analysis tools compare expressions across many computationally discovered cell types. CytoGLMM focuses on just one cell type. Our narrower field of application allows us to define a more specific statistical model with easier to control statistical guarantees. As a result, CytoGLMM finds differential proteins in flow and mass cytometry data while reducing biases arising from marker correlations and safeguarding against false discoveries induced by patient heterogeneity.

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URL <https://christofseiler.github.io/CytoGLMM>,
<https://github.com/ChristofSeiler/CytoGLMM>

BugReports <https://github.com/ChristofSeiler/CytoGLMM/issues>

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Author Christof Seiler [aut, cre] (ORCID:
[<https://orcid.org/0000-0001-8802-3642>](https://orcid.org/0000-0001-8802-3642))

Maintainer Christof Seiler <christof.seiler@maastrichtuniversity.nl>

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cytoflexmix	<i>Logistic mixture regression</i>
-------------	------------------------------------

Description

Logistic mixture regression

Usage

```
cytoflexmix(
  df_samples_subset,
  protein_names,
  condition,
  group = "donor",
  cell_n_min = Inf,
  cell_n_subsample = 0,
```

```
  ks = seq_len(10),
  num_cores = 1
)
```

Arguments

df_samples_subset	Data frame or tibble with proteins counts, cell condition, and group information
protein_names	A vector of column names of protein to use in the analysis
condition	The column name of the condition variable
group	The column name of the group variable
cell_n_min	Remove samples that are below this cell counts threshold
cell_n_subsample	Subsample samples to have this maximum cell count
ks	A vector of cluster sizes
num_cores	Number of computing cores

Value

A list of class `cytglm` containing

flexmixfits	list of <code>flexmix</code> objects
df_samples_subset	possibly subsampled df_samples_subset table
protein_names	input protein names
condition	input condition variable
group	input group names
cell_n_min	input cell_n_min
cell_n_subsample	input cell_n_subsample
ks	input ks
num_cores	input num_cores

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
mix_fit <- CytoGLMM::cytoflexmix(df,
                                    protein_names = protein_names,
                                    condition = "condition",
                                    group = "donor",
                                    ks = 2)
mix_fit
```

cytglm*Fit GLM with bootstrap resampling***Description**

Fit GLM with bootstrap resampling

Usage

```
cytglm(
  df_samples_subset,
  protein_names,
  condition,
  group = "donor",
  covariate_names = NULL,
  cell_n_min = Inf,
  cell_n_subsample = 0,
  num_boot = 100,
  num_cores = 1
)
```

Arguments

<code>df_samples_subset</code>	Data frame or tibble with proteins counts, cell condition, and group information
<code>protein_names</code>	A vector of column names of protein to use in the analysis
<code>condition</code>	The column name of the condition variable
<code>group</code>	The column name of the group variable
<code>covariate_names</code>	The column names of covariates
<code>cell_n_min</code>	Remove samples that are below this cell counts threshold
<code>cell_n_subsample</code>	Subsample samples to have this maximum cell count
<code>num_boot</code>	Number of bootstrap samples
<code>num_cores</code>	Number of computing cores

Value

A list of class <code>cytglm</code> containing	
<code>tb_coef</code>	coefficient table
<code>df_samples_subset</code>	possibly subsampled <code>df_samples_subset</code> table
<code>protein_names</code>	input protein names
<code>condition</code>	input condition variable
<code>group</code>	input group names
<code>covariate_names</code>	input covariates

```

cell_n_min      input cell_n_min
cell_n_subsample      input cell_n_subsample
unpaired      true if unpaired samples were provided as input
num_boot       input num_boot
num_cores      input num_cores
formula_str    formula use in the regression model

```

Examples

```

set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
glm_fit <- CytoGLMM::cytogiM(df,
                                protein_names = protein_names,
                                condition = "condition",
                                group = "donor",
                                num_boot = 10) # in practice >=1000
glm_fit

```

cytogiM

Fit GLMM with method of moments

Description

Fit GLMM with method of moments

Usage

```

cytogiM(
  df_samples_subset,
  protein_names,
  condition,
  group = "donor",
  covariate_names = NULL,
  cell_n_min = Inf,
  cell_n_subsample = 0,
  num_cores = 1
)

```

Arguments

<code>df_samples_subset</code>	Data frame or tibble with proteins counts, cell condition, and group information
<code>protein_names</code>	A vector of column names of protein to use in the analysis
<code>condition</code>	The column name of the condition variable
<code>group</code>	The column name of the group variable
<code>covariate_names</code>	The column names of covariates

```

cell_n_min      Remove samples that are below this cell counts threshold
cell_n_subsample Subsample samples to have this maximum cell count
num_cores       Number of computing cores

```

Value

A list of class `cytoglm` containing

<code>glmmfit</code>	<code>mbest</code> object
<code>df_samples_subset</code>	possibly subsampled <code>df_samples_subset</code> table
<code>protein_names</code>	input protein names
<code>condition</code>	input condition variable
<code>group</code>	input group names
<code>covariate_names</code>	input covariates
<code>cell_n_min</code>	input <code>cell_n_min</code>
<code>cell_n_subsample</code>	input <code>cell_n_subsample</code>
<code>num_cores</code>	input <code>num_cores</code>

Examples

```

set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
glmm_fit <- CytoGLMM::cytoglmm(df,
                                    protein_names = protein_names,
                                    condition = "condition",
                                    group = "donor")
glmm_fit

```

cytogroup	<i>Group-specific fixed effects model</i>
-----------	---

Description

Group-specific fixed effects model

Usage

```

cytogroup(
  df_samples_subset,
  protein_names,
  condition,
  group = "donor",
  cell_n_min = Inf,
  cell_n_subsample = 0
)

```

Arguments

df_samples_subset	Data frame or tibble with proteins counts, cell condition, and group information
protein_names	A vector of column names of protein to use in the analysis
condition	The column name of the condition variable
group	The column name of the group variable
cell_n_min	Remove samples that are below this cell counts threshold
cell_n_subsample	Subsample samples to have this maximum cell count

Value

A list of class `cytoglm` containing

groupfit	<code>glm</code> object
df_samples_subset	possibly subsampled df_samples_subset table
protein_names	input protein names
condition	input condition variable
group	input group names
cell_n_min	input cell_n_min
cell_n_subsample	input cell_n_subsample

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
group_fit <- CytoGLMM::cytogroup(df,
                                    protein_names = protein_names,
                                    condition = "condition",
                                    group = "donor")
group_fit
```

Description

Evaluate parameter stability with respect to gating sheme

Usage

```
cytostab(
  df_samples_subset,
  protein_names,
  condition,
  group = "donor",
  cell_n_min = Inf,
  cell_n_subsample = 0
)
```

Arguments

df_samples_subset	Data frame or tibble with proteins counts, cell condition, and group information
protein_names	A vector of column names of protein to use in the analysis
condition	The column name of the condition variable
group	The column name of the group variable
cell_n_min	Remove samples that are below this cell counts threshold
cell_n_subsample	Subsample samples to have this maximum cell count

Value

A data frame

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
stab <- CytoGLMM::cytostab(df,
                           protein_names = protein_names,
                           condition = "condition",
                           group = "donor")
stab
```

cyto_check

Check if input to cytoxxx function have errors

Description

Check if input to cytoxxx function have errors

Usage

```
cyto_check(cell_n_subsample, cell_n_min, protein_names)
```

Arguments

```
cell_n_subsample      Subsample samples to have this maximum cell count
cell_n_min           A vector of column names of protein to use in the analysis
protein_names        A vector of column names of protein to use in the analysis
```

Value

NULL.

generate_data	<i>Generate dataset for vignettes and simulation studies</i>
---------------	--

Description

Generate dataset for vignettes and simulation studies

Usage

```
generate_data()
```

Value

tibble data frame

Examples

```
set.seed(23)
df <- generate_data()
str(df)
df
```

glmm_moment	<i>Generalized linear mixed model with maximum likelihood</i>
-------------	---

Description

Generalized linear mixed model with maximum likelihood

Usage

```
glmm_moment(
  df_samples,
  protein_names,
  response,
  group = "donor",
  covariate_names = NULL,
  num_cores = 1
)
```

Arguments

<code>df_samples</code>	Data frame or tibble with proteins counts, cell condition, and group information
<code>protein_names</code>	A vector of column names of protein to use in the analysis
<code>response</code>	The column name of the condition variable
<code>group</code>	The column name of the group variable
<code>covariate_names</code>	The column names of covariates
<code>num_cores</code>	Number of computing cores

Value

`mbest` object

<code>is_unpaired</code>	<i>Check if samples match or paired on condition</i>
--------------------------	--

Description

Check if samples match or paired on condition

Usage

```
is_unpaired(df_samples_subset, condition, group)
```

Arguments

<code>df_samples_subset</code>	Data frame or tibble with proteins counts, cell condition, and group information
<code>condition</code>	The column name of the condition variable
<code>group</code>	The column name of the group variable

Value

A boolean

<code>plot.cytoflexmix</code>	<i>Plot all components of mixture regression</i>
-------------------------------	--

Description

Plot all components of mixture regression

Usage

```
## S3 method for class 'cytoflexmix'
plot(x, k = NULL, separate = FALSE, ...)
```

Arguments

x	A cytoflexmix class
k	Number of clusters
separate	create two separate <code>ggplot2</code> objects
...	Other parameters

Value

`ggplot2` object

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
mix_fit <- CytoGLMM::cytoflexmix(df,
                                    protein_names = protein_names,
                                    condition = "condition",
                                    group = "donor",
                                    ks = 2)
plot(mix_fit)
```

plot.cytoglm

Plot bootstraped coefficients

Description

Plot bootstraped coefficients

Usage

```
## S3 method for class 'cytoglm'
plot(x, order = FALSE, separate = FALSE, ...)
```

Arguments

x	A cytoglm class
order	Order the markers according to the mangintute of the coefficients
separate	create two separate <code>ggplot2</code> objects
...	Other parameters

Value

`ggplot2` object

Examples

```

set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
glm_fit <- CytoGLMM::cytoglm(df,
                                protein_names = protein_names,
                                condition = "condition",
                                group = "donor",
                                num_boot = 10) # in practice >=1000
plot(glm_fit)

```

plot.cytoglm

Plot fixed coefficients of random effects model

Description

Plot fixed coefficients of random effects model

Usage

```
## S3 method for class 'cytoglmm'  
plot(x, order = FALSE, separate = FALSE, ...)
```

Arguments

<code>x</code>	A <code>cytoglm</code> class
<code>order</code>	Order the markers according to the magnitude of the coefficients
<code>separate</code>	create two separate <code>ggplot2</code> objects
<code>...</code>	Other parameters

Value

ggplot2 object

Examples

<code>plot.cytogroup</code>	<i>Plot fixed coefficients of group-specific fixed effects model</i>
-----------------------------	--

Description

Plot fixed coefficients of group-specific fixed effects model

Usage

```
## S3 method for class 'cytogroup'
plot(x, order = FALSE, separate = FALSE, ...)
```

Arguments

<code>x</code>	A <code>cytoplmm</code> class
<code>order</code>	Order the markers according to the magnitude of the coefficients
<code>separate</code>	create two separate <code>ggplot2</code> objects
<code>...</code>	Other parameters

Value

`ggplot2` object

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
group_fit <- CytoGLMM::cytogroup(df,
                                    protein_names = protein_names,
                                    condition = "condition",
                                    group = "donor")
plot(group_fit)
```

<code>plot_coeff</code>	<i>Helper function to plot regression coefficient</i>
-------------------------	---

Description

Helper function to plot regression coefficient

Usage

```
plot_coeff(
  tb,
  title_str,
  title_str_right,
  xlab_str,
  redline = 0,
  order = FALSE,
  separate = FALSE
)
```

Arguments

<code>tb</code>	A data frame
<code>title_str</code>	Title string for summary plot
<code>title_str_right</code>	Title for bootstrap sample plot
<code>xlab_str</code>	Label on x-axis
<code>redline</code>	Point on x-axis to draw the red line
<code>order</code>	Order the markers according to the magnitude of the coefficients
<code>separate</code>	Plot both summary and bootstrap samples

Value

[ggplot2](#) object or list of two objects if separate is true

<code>plot_heatmap</code>	<i>Heatmap of median marker expression</i>
---------------------------	--

Description

Heatmap of median marker expression

Usage

```
plot_heatmap(
  df_samples,
  sample_info_names,
  protein_names,
  arrange_by_1,
  arrange_by_2 = "",
  cluster_cols = FALSE,
  fun = median
)
```

Arguments

df_samples	Data frame or tibble with proteins counts, cell condition, and group information
sample_info_names	Column names that contain information about the cell, e.g. donor, condition, file name, or cell type
protein_names	A vector of column names of protein to use in the analysis
arrange_by_1	Column name
arrange_by_2	Column name
cluster_cols	Apply hierarchical cluster to columns
fun	Summary statistics of marker expression

Value

[pheatmap](#) object

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
CytoGLMM::plot_heatmap(df,
                       protein_names = protein_names,
                       sample_info_names = c("donor", "condition"),
                       arrange_by_1 = "condition")
```

plot_lda

LDA on marker expression

Description

LDA on marker expression

Usage

```
plot_lda(
  df_samples,
  protein_names,
  group,
  cor_scaling_factor = 1,
  arrow_color = "black",
  marker_color = "black",
  marker_size = 5
)
```

Arguments

<code>df_samples</code>	Data frame or tibble with proteins counts, cell condition, and group information
<code>protein_names</code>	A vector of column names of protein to use in the analysis
<code>group</code>	The column name of the group variable
<code>cor_scaling_factor</code>	Scaling factor of circle of correlations
<code>arrow_color</code>	Color of correlation circle
<code>marker_color</code>	Colors of marker names
<code>marker_size</code>	Size of marker names

Value

`ggplot2` object

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
df$condition <- rep(c("A", "B", "C", "D"), each = length(df$condition)/4)
CytoGLMM::plot_lda(df,
  protein_names = protein_names,
  group = "condition",
  cor_scaling_factor = 2)
```

`plot_mds`

MDS on median marker expression

Description

MDS on median marker expression

Usage

```
plot_mds(
  df_samples,
  protein_names,
  sample_info_names,
  color,
  sample_label = "")
```

Arguments

<code>df_samples</code>	Data frame or tibble with proteins counts, cell condition, and group information
<code>protein_names</code>	A vector of column names of protein to use in the analysis
<code>sample_info_names</code>	Column names that contain information about the cell, e.g. donor, condition, file name, or cell type
<code>color</code>	Column name
<code>sample_label</code>	Column name

Value

`cowplot` object

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
CytoGLMM::plot_mds(df,
  protein_names = protein_names,
  sample_info_names = c("donor", "condition"),
  color = "condition")
```

`plot_model_selection` *Plot model selection to choose number optimal number of clusters*

Description

Plot model selection to choose number optimal number of clusters

Usage

```
plot_model_selection(fit, k = NULL)
```

Arguments

<code>fit</code>	A <code>cytoflexmix</code> class
<code>k</code>	Number of clusters

Value

`cowplot` object

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
mix_fit <- CytoGLMM::cytoflexmix(df,
  protein_names = protein_names,
  condition = "condition",
  group = "donor",
  ks = 1:2)
plot_model_selection(mix_fit)
```

`plot_prcomp` *Plot PCA of subsampled data using ggplot*

Description

Plot PCA of subsampled data using ggplot

Usage

```
plot_prcomp(
  df_samples,
  protein_names,
  color_var = "treatment",
  subsample_size = 10000,
  repel = TRUE
)
```

Arguments

<code>df_samples</code>	Data frame or tibble with proteins counts, cell condition, and group information
<code>protein_names</code>	A vector of column names of protein to use in the analysis
<code>color_var</code>	A column name
<code>subsample_size</code>	Subsample per color_var variable
<code>repel</code>	Repel labels

Value

`cowplot` object

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
CytoGLMM::plot_prcomp(df,
  protein_names = protein_names,
  color_var = "condition")
```

`print.cytoglm` *Extract and print bootstrap GLM fit*

Description

Extract and print bootstrap GLM fit

Usage

```
## S3 method for class 'cytoglm'
print(x, ...)
```

Arguments

x A cytogl_m class
... Other parameters

Value

NULL.

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
glm_fit <- CytoGLMM::cytoglm(df,
                                protein_names = protein_names,
                                condition = "condition",
                                group = "donor",
                                num_boot = 10) # in practice >=1000

print(glm_fit)
```

`print.cytoglm` *Extract and print GLMM fit*

Description

Extract and print GLMM fit

Usage

```
## S3 method for class 'cytoglmm'  
print(x, ...)
```

Arguments

x A cytoglmm class
... Other parameters

Value

NULL

Examples

remove_samples *Remove samples based on low cell counts*

Description

Remove samples based on low cell counts

Usage

```
remove_samples(df_samples_subset, condition, group, unpaired, cell_n_min)
```

Arguments

<code>df_samples_subset</code>	Data frame or tibble with proteins counts, cell condition, and group information
<code>condition</code>	The column name of the condition variable
<code>group</code>	The column name of the group variable
<code>unpaired</code>	true if unpaired samples were provided as input
<code>cell_n_min</code>	Remove samples that are below this cell counts threshold

Value

NULL.

summary.cytoglm *Extract and calculate p-values of bootstrap GLM fit*

Description

Extract and calculate p-values of bootstrap GLM fit

Usage

```
## S3 method for class 'cytoglm'
summary(object, method = "BH", ...)
```

Arguments

<code>object</code>	A <code>cytoglm</code> class
<code>method</code>	Multiple comparison adjustment method
<code>...</code>	Other parameters

Value

`tibble` data frame

Examples

```
set.seed(23)
df <- generate_data()
protein_names <- names(df)[3:12]
df <- dplyr::mutate_at(df, protein_names, function(x) asinh(x/5))
glm_fit <- CytoGLMM::cytoglm(df,
                                protein_names = protein_names,
                                condition = "condition",
                                group = "donor",
                                num_boot = 10) # in practice >=1000
summary(glm_fit)
```

`summary.cytoglm` Extract and calculate p-values of GLMM fit

Description

Extract and calculate p-values of GLMM fit

Usage

```
## S3 method for class 'cytoglmm'  
summary(object, method = "BH", ...)
```

Arguments

<code>object</code>	A <code>cytoglm</code> class
<code>method</code>	Multiple comparison adjustment method
<code>...</code>	Other parameters

Value

tibble data frame

Examples

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