

Package ‘cytometree’

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Type Package

Title Automated Cytometry Gating and Annotation

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LinkingTo Rcpp, RcppArmadillo

Description Given the hypothesis of a bi-modal distribution of cells for each marker, the algorithm constructs a binary tree, the nodes of which are subpopulations of cells. At each node, observed cells and markers are modeled by both a family of normal distributions and a family of bi-modal normal mixture distributions. Splitting is done according to a normalized difference of AIC between the two families.

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LazyData true

Depends R (>= 3.1.0), Rcpp (>= 0.12.11)

Imports ggplot2, graphics, igraph, mclust, RColorBrewer, robustbase, stats, grDevices, cowplot

RoxygenNote 6.1.0

BugReports <https://github.com/sistm/Cytometree/issues>

Encoding UTF-8

Suggests knitr, rmarkdown, viridis

VignetteBuilder knitr

NeedsCompilation yes

Repository CRAN

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cytometree-package	<i>Binary tree algorithm for cytometry data analysis.</i>
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Description

The algorithm is based on the construction of a binary tree, the nodes of which are subpopulations of cells. At each node, observed cells and markers are modeled by both a family of normal distributions and a family of bi-modal normal mixture distributions. Splitting is done according to a normalized difference of AIC between the two families. Given the unsupervised nature of the binary tree, some of the available markers may not be used to find the different cell populations present in a given sample. To recover a complete annotation, we defined, as a post processing procedure, an annotation method which allows the user to distinguish two or three expression levels per marker.

Details

Package: cytometree
 Type: Package
 Version: 1.2.0
 Date: 2018-08-28
 License: **LGPL-3**

The main function in this package is [CytomeTree](#).

Author(s)

Chariff Alkhasim, Boris Hejblum — Maintainer: Chariff Alkhasim

References

D. Commenges, C. Alkhassim, B.P. Hejblum, R. Thi'ebaut. cytometree: a Binary tree algorithm for automatic gating in cytometry analysis. Submitted, 2017.

Annotation	<i>Annotates cell populations found using CytomeTree.</i>
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Description

Annotates cell populations found using CytomeTree.

Usage

```
Annotation(CytomeTreeObj, K2markers = NULL, K3markers = NULL,
           plot = TRUE, t = 0.2)
```

Arguments

CytomeTreeObj	An object of class CytomeTree.
K2markers	A vector of class character where the names of the markers for which 2 levels of expression are sought can be specified. Default is NULL i.e. unsupervised.
K3markers	A vector of class character where the names of the markers for which 3 levels of expression are sought can be specified. Default is NULL i.e. unsupervised.
plot	A logical value indicating whether or not to plot the partitioning in 1, 2 or 3 groups for each marker. Default is TRUE.
t	A real positive-or-null number used for comparison with the normalized AIC computed to compare the fits of the marginal distributions obtained by one normal distribution and by a mixture of two or three normal. For markers used in the tree, the algorithm compares the fits obtained by a mixture of two and three normal distributions. Default value is .2. A higher value leads to a smaller number of expression levels per marker.

Details

The algorithm is set to find the partitioning in 1, 2 or 3 groups of cell populations found using CytomeTree. In an unsupervised mode, it minimizes the within-leaves sum of squares of the observed values on each marker and computes the normalized AIC to compare the fits of the marginal distributions obtained by one normal distribution and by a mixture of two or three normal. For markers used in the tree, the algorithm compares the fits obtained by a mixture of two and three normal distributions.

Value

A data.frame containing the annotation of each cell population.

Author(s)

Chariff Alkhassim

bootstrapCI	<i>Bootstrapped Confidence Interval.</i>
-------------	--

Description

Bootstrapped Confidence Interval.

Usage

```
bootstrapCI(stat, n, alpha)
```

Arguments

stat	A numeric vector of statistics for which to compute a confidence interval.
n	An integer giving the number of bootstrap samples.
alpha	A real number comprised in]0, 1[: 1 - desired confidence level.

Author(s)

Chariff Alkhassim

CytomeTree	<i>Binary tree algorithm for cytometry data analysis.</i>
------------	---

Description

Binary tree algorithm for cytometry data analysis.

Usage

```
CytomeTree(M, minleaf = 1, t = 0.1, verbose = TRUE)
```

Arguments

M	A matrix of size n x p containing cytometry measures of n cells on p markers.
minleaf	An integer indicating the minimum number of cells per population. Default is 1.
t	A real positive-or-null number used for comparison with the normalized AIC computed at each node of the tree. A higher value limits the height of the tree.
verbose	A logical controlling if a text progress bar is displayed during the execution of the algorithm. By default is TRUE.

Details

The algorithm is based on the construction of a binary tree, the nodes of which are subpopulations of cells. At each node, observed cells and markers are modeled by both a family of normal distributions and a family of bi-modal normal mixture distributions. Splitting is done according to a normalized difference of AIC between the two families.

Value

An object of class 'cytomeTree' providing a partitioning of the set of n cells.

- `annotation` A data.frame containing the annotation of each cell population underlying the tree pattern.
- `labels` The partitioning of the set of n cells.
- `M` The input matrix.
- `mark_tree` A two level list containing markers used for node splitting.

Author(s)

Chariff Alkhassim, Boris Hejblum

Examples

```
head(DLBCL)

# number of cell event
N <- nrow(DLBCL)

# Cell events
cellevents <- DLBCL[, c("FL1", "FL2", "FL4")]

# Manual partitioning of the set N (from FlowCAP-I)
manual_labels <- DLBCL[, "label"]

# Build the binary tree
Tree <- CytomeTree(cellevents, minleaf = 1, t=.1)

# Retrieve the resulting partition of the set N
Tree_Partition <- Tree$labels

# Plot node distributions
par(mfrow=c(1, 2))
plot_nodes(Tree)

# Choose a node to plot
plot_nodes(Tree, "FL4.1")
```

```

# Plot a graph of the tree
par(mfrow=c(1,1))
plot_graph(Tree,edge.arrow.size=.3, Vcex =.5, vertex.size = 30)

# Run the annotation algorithm
Annot <- Annotation(Tree,plot=FALSE)
Annot$combinations

# Compare to the annotation gotten from the tree
Tree$annotation

# Example of sought phenotypes
# Variable in which sought phenotypes can be entered in the form of matrices.
phenotypes <- list()

# Sought phenotypes:
## FL2+ FL4-.
phenotypes[[1]] <- rbind(c("FL2", 1), c("FL4", 0))

## FL2- FL4+.
phenotypes[[2]] <- rbind(c("FL2", 0), c("FL4", 1))

## FL2+ FL4+.
phenotypes[[3]] <- rbind(c("FL2", 1), c("FL4", 1))

# Retrieve cell populations found using Annotation.
PhenoInfos <- RetrievePops(Annot, phenotypes)
PhenoInfos$phenotypesinfo

# F-measure ignoring cells labeled 0 as in FlowCAP-I.

# Use FmeasureC() in any other case.
FmeasureC_no0(ref>manual_labels, pred=Tree_Partition)

## Not run:

# Scatterplots.
library(ggplot2)

# Ignoring cells labeled 0 as in FlowCAP-I.
rm_zeros <- which(!manual_labels)

# Building the data frame to scatter plot the data.
FL1 <- cellevents[-c(rm_zeros),"FL1"]
FL2 <- cellevents[-c(rm_zeros),"FL2"]
FL4 <- cellevents[-c(rm_zeros),"FL4"]
n <- length(FL1)
Labels <- c(manual_labels[-c(rm_zeros)]%%2+1, Tree_Partition[-c(rm_zeros)])
Labels <- as.factor(Labels)

```

```
method <- as.factor(c(rep("FlowCap-I",n),rep("CytomeTree",n)))

scatter_df <- data.frame("FL2" = FL2, "FL4" = FL4, "labels" = Labels, "method" = method)
p <- ggplot2::ggplot(scatter_df, ggplot2::aes_string(x = "FL2", y = "FL4", colour = "labels")) +
  ggplot2::geom_point(alpha = 1,cex = 1) +
  ggplot2::scale_colour_manual(values = c("green","red","blue")) +
  ggplot2::facet_wrap(~ method) +
  ggplot2::theme_bw() +
  ggplot2::theme(legend.position="bottom")
p

## End(Not run)
```

DLBCL

Diffuse large B-cell lymphoma data set from the FlowCAP-I challenge.

Description

Diffuse large B-cell lymphoma data set from the FlowCAP-I challenge.

Usage

```
data(DLBCL)
```

Format

A data frame with 5524 cell events and 3 markers.

Source

<http://flowcap.flowsite.org/>

FmeasureC

C++ implementation of the F-measure computation

Description

C++ implementation of the F-measure computation

Usage

```
FmeasureC(pred, ref)
```

Arguments

pred	vector of a predicted partition
ref	vector of a reference partition

Author(s)

Boris Hejblum

FmeasureC_no0	<i>C++ implementation of the F-measure computation without the reference class labeled "0"</i>
---------------	--

Description

Aghaeepour in FlowCAP 1 ignore the reference class labeled "0"

Usage

FmeasureC_no0(pred, ref)

Arguments

pred	vector of a predicted partition
ref	vector of a reference partition

Author(s)

Boris Hejblum

HIPC	<i>HIPC T cell data set from HIPC program, patient 12828. The data was analyzed and gated by Stanford.</i>
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Description

HIPC T cell data set from HIPC program, patient 12828. The data was analyzed and gated by Stanford.

Usage

data(HIPC)

Format

A data frame with 33992 cell events and 6 markers.

Source<https://www.immuneprofiling.org/>

plot_cytopop	<i>Plot the cell count for each population using CytomeTree.</i>
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Description

Plot the cell count for each population using CytomeTree.

Usage

```
plot_cytopop(AnnotObj, nbpop = 10, mincount = 1, maxcount = NULL,  
             y_axis = c("abs_count", "prop"))
```

Arguments

AnnotObj	An object of class Annotation.
nbpop	Number indicating the maximum of population plotted. Default is 10
mincount	Number indicating the minimum of cell count for the populations. Default is 1.
maxcount	Number indicating the maximum of cell count for the populations. Default is NULL i.e no maximum selected.
y_axis	a character string either "abs_count" or "prop" indicating whether the absolute cell count or the relative populations proportions should be plotted. Default is "abs_count".

Author(s)

Anthony Devaux, Boris Hejblum

Examples

```
# Run CytomeTree  
data(DLBCL)  
cellevents <- DLBCL[,c("FL1", "FL2", "FL4")]  
Tree <- CytomeTree(cellevents, minleaf = 1, t=.1)  
Annot <- Annotation(Tree,plot=FALSE)  
  
# Plot the cell count  
plot_cytopop(Annot)
```

plot_graph	<i>Plot the binary tree built using CytomeTree.</i>
------------	---

Description

Plot the binary tree built using CytomeTree.

Usage

```
plot_graph(CytomeTreeObj, Ecex = 1, Ecolor = 8, Vcex = 0.8,
           Vcolor = 0, ...)
```

Arguments

CytomeTreeObj	An object of class CytomeTree.
Ecex	Number indicating the amount by which text on the edges should be scaled. Default is 1.
Ecolor	An integer or a string of character to color edges of the graph. Default is 8.
Vcex	Number indicating the amount by which text in the vertices should be scaled. Default is .8.
Vcolor	A vector of class numeric or character to color vertices of the graph. Default is 0.
...	additional arguments to be passed to plot_graph

Author(s)

Chariff Alkhassim

plot_nodes	<i>Plot the distribution of the observed cells at each node of the binary tree built using CytomeTree.</i>
------------	--

Description

Plot the distribution of the observed cells at each node of the binary tree built using CytomeTree.

Usage

```
plot_nodes(CytomeTreeObj, nodes = NULL, nodesPerCol = NULL,
           nodesPerRow = NULL, ...)
```

Arguments

<code>CytomeTreeObj</code>	An object of class <code>CytomeTree</code> .
<code>nodes</code>	A list of character elements containing the name of the nodes for which the distribution is to be plotted. Default is <code>NULL</code> , and plots the distribution for each node.
<code>nodesPerCol</code>	an integer specifying the number of plots to be displayed per column when plotting multiple nodes at once. Default is <code>NULL</code> .
<code>nodesPerRow</code>	an integer specifying the number of plots to be displayed per row when plotting multiple nodes at once. Default is <code>NULL</code> .
<code>...</code>	further arguments to be passed to <code>plot_grid</code> .

Details

if both `nodesPerCol` and `nodesPerRow` are `NULL` then all the nodes are plotted on a single page.

Value

a list of `ggplot2` plot objects, containing each node plot.

Author(s)

Chariff Alkhassim, Boris Hejblum

Examples

```
data(DLBCL)
myct <- CytomeTree(DLBCL[, c("FL1", "FL2", "FL4")], minleaf = 1, t=.1)
plot_nodes(myct)
```

RetrievePops

Retrieve cell populations found using Annotation.

Description

Retrieve cell populations found using Annotation.

Usage

```
RetrievePops(AnnotationObj, phenotypes)
```

Arguments

- AnnotationObj An object of class Annotation.
- phenotypes A list containing at least one element of class matrix describing a sought phenotype. Each matrix should have two columns where the name of a used marker is associated to a value chosen between 0, 1 and 2. 0 translates to -, 1 to + and 2 to ++.

Value

A list of two elements.

- phenotypesinfo A list containing informations about sought populations.
- Mergedleaves The partitioning of the set of n cells with potentially merged leaves.

Author(s)

Chariff Alkassim

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