

Package ‘spicyR’

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Type Package

Title Spatial analysis of in situ cytometry data

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Description spicyR provides a series of functions to aid in the analysis of both immunofluorescence and mass cytometry imaging data as well as other assays that can deeply phenotype individual cells and their spatial location.

License GPL (>=2)

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Encoding UTF-8

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Suggests BiocStyle, knitr, rmarkdown

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Accessors	<i>Accessors for SegmentedCells</i>
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Description

Methods to access various components of the ‘SegmentedCells’ object.

Usage

```

cellSummary(x, imageID = NULL, bind = TRUE)

cellSummary(x, imageID = NULL) <- value

cellMarks(x, imageID = NULL, bind = TRUE)

cellMarks(x, imageID = NULL) <- value

cellMorph(x, imageID = NULL, bind = TRUE)

cellMorph(x, imageID = NULL) <- value

imagePheno(x, imageID = NULL, bind = TRUE, expand = FALSE)

imagePheno(x, imageID = NULL) <- value

imageID(x, imageID = NULL)

cellID(x, imageID = NULL)

cellID(x) <- value

```

```
imageCellID(x, imageID = NULL)
imageCellID(x) <- value
cellType(x, imageID = NULL)
cellType(x, imageID = NULL) <- value
filterCells(x, select)
cellAnnotation(x, variable, imageID = NULL)
cellAnnotation(x, variable, imageID = NULL) <- value
```

Arguments

x	A 'SegmentedCells' object.
imageID	A vector of imageIDs to specifically extract.
bind	When false outputs a list of DataFrames split by imageID
expand	Used to expand the phenotype information from per image to per cell.
value	The relevant information used to replace.
select	A logical vector of the cells to be kept.
variable	A variable to add or retrieve from cellSummary.

Value

DataFrame or a list of DataFrames

Descriptions

'cellSummary': Retrieves the DataFrame containing 'x' and 'y' coordinates of each cell as well as 'cellID', 'imageID' and 'cellType'. imageID can be used to select specific images and bind=FALSE outputs the information as a list split by imageID.

'cellMorph': Retrieves the DataFrame containing morphology information.

'cellMarks': Retrieves the DataFrame containing intensity of gene or protein markers.

'imagePheno': Retrieves the DataFrame containing the phenotype information for each image. Using expand = TRUE will produce a DataFrame with the number of rows equal to the number of cells.

Examples

```
### Something that resembles cellProfiler data
set.seed(51773)
n = 10
```

```

cells <- data.frame(row.names = seq_len(n))
cells$ObjectNumber <- seq_len(n)
cells$ImageNumber <- rep(1:2,c(n/2,n/2))
cells$AreaShape_Center_X <- runif(n)
cells$AreaShape_Center_Y <- runif(n)
cells$AreaShape_round <- rexp(n)
cells$AreaShape_diameter <- rexp(n, 2)
cells$Intensity_Mean_CD8 <- rexp(n, 10)
cells$Intensity_Mean_CD4 <- rexp(n, 10)

cellExp <- SegmentedCells(cells, cellProfiler = TRUE)

### Cluster cell types
intensities <- cellMarks(cellExp)
kM <- kmeans(intensities,2)
cellType(cellExp) <- paste('cluster',kM$cluster, sep = '')

cellSummary(cellExp, imageID = 1)

```

```

as.data.frame.SegmentedCells
      as.data.frame

```

Description

Function to coerce a SegmentedCells object to a data frame.

Usage

```

## S3 method for class 'SegmentedCells'
as.data.frame(x, ...)

```

Arguments

```

x          A SegmentedCells object.
...       Other arguments.

```

Value

```

A data.frame
## Generate toy data set.seed(51773) x <- round(c(runif(200),runif(200)+1,runif(200)+2,runif(200)+3,
runif(200)+3,runif(200)+2,runif(200)+1,runif(200)),4) y <- round(c(runif(200),runif(200)+1,runif(200)+2,runif(200)+3,
runif(200),runif(200)+1,runif(200)+2,runif(200)+3),4) cellType <- factor(paste('c',rep(rep(c(1:2),rep(200,2)),4),sep
= '')) imageID <- rep(c('s1', 's2'),c(800,800)) cells <- data.frame(x, y, cellType, imageID)
## Store data in SegmentedCells object cellExp <- SegmentedCells(cells, cellTypeString = 'cell-
Type')
## Generate LISA cellsDF <- as.data.frame(cellExp)
NULL

```

diabetesData	<i>Diabetes IMC data</i>
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Description

This is a subset of the Damond et al 2019 imaging mass cytometry dataset. The data contains cells in the pancreatic islets of individuals with early onset diabetes and healthy controls. The object contains single-cell data of 160 images from 8 subjects, with 20 images per subject.

Usage

```
diabetesData
```

Format

```
diabetesData a SegmentedCells object
```

getPairwise	<i>Get statistic from pairwise L curve of a single image.</i>
-------------	---

Description

Get statistic from pairwise L curve of a single image.

Usage

```
getPairwise(  
  cells,  
  from = unique(cellType(cells)),  
  to = unique(cellType(cells)),  
  dist = NULL,  
  window = "convex",  
  window.length,  
  Rs = NULL,  
  sigma = NULL,  
  minLambda = 0.05,  
  fast = TRUE,  
  BPPARAM = BiocParallel::SerialParam()  
)
```

Arguments

<code>cells</code>	A <code>SegmentedCells</code> or data frame that contains at least the variables <code>x</code> and <code>y</code> , giving the location coordinates of each cell, and <code>cellType</code> .
<code>from</code>	The 'from' <code>cellType</code> for generating the L curve.
<code>to</code>	The 'to' <code>cellType</code> for generating the L curve.
<code>dist</code>	The distance at which the statistic is obtained.
<code>window</code>	Should the window around the regions be 'square', 'convex' or 'concave'.
<code>window.length</code>	A tuning parameter for controlling the level of concavity when estimating concave windows.
<code>Rs</code>	A vector of the radii that the measures of association should be calculated.
<code>sigma</code>	A numeric variable used for scaling when fitting inhomogeneous L-curves.
<code>minLambda</code>	Minimum value for density for scaling when fitting inhomogeneous L-curves.
<code>fast</code>	A logical describing whether to use a fast approximation of the inhomogeneous L-curves.
<code>BPPARAM</code>	A <code>BiocParallelParam</code> object.

Value

Statistic from pairwise L curve of a single image.

Examples

```
data("diabetesData")
pairAssoc <- getPairwise(diabetesData)
```

`plot, SegmentedCells, ANY-method`

A basic plot for SegmentedCells object

Description

This function generates a basic x-y plot of the location coordinates and `cellType` data.

Usage

```
## S4 method for signature 'SegmentedCells,ANY'
plot(x, imageID = NULL)
```

Arguments

<code>x</code>	A <code>SegmentedCells</code> object.
<code>imageID</code>	The image that should be plotted.

Value

A ggplot object.

usage

```
'plot(x, imageID = NULL)'
```

Examples

```
### Something that resembles cellProfiler data

set.seed(51773)

n = 10

cells <- data.frame(row.names = seq_len(n))
cells$ObjectNumber <- seq_len(n)
cells$ImageNumber <- rep(1:2,c(n/2,n/2))
cells$AreaShape_Center_X <- runif(n)
cells$AreaShape_Center_Y <- runif(n)
cells$AreaShape_round <- rexp(n)
cells$AreaShape_diameter <- rexp(n, 2)
cells$Intensity_Mean_CD8 <- rexp(n, 10)
cells$Intensity_Mean_CD4 <- rexp(n, 10)

cellExp <- SegmentedCells(cells, cellProfiler = TRUE)

### Cluster cell types
markers <- cellMarks(cellExp)
kM <- kmeans(markers,2)
cellType(cellExp) <- paste('cluster',kM$cluster, sep = '')

#plot(cellExp, imageID=1)
```

SegmentedCells-class *The SegmentedCells class*

Description

The SegmentedCells S4 class is for storing data from segmented imaging cytometry and spatial omics data. It extends DataFrame and defines methods that take advantage of DataFrame nesting to represent elements of cell-based experiments with spatial orientation that are commonly encountered. This object is able to store information on a cell's spatial location, cellType, morphology, intensity of gene/protein markers as well as image level phenotype information.

Usage

```
SegmentedCells(
  cellData,
  cellProfiler = FALSE,
  spatialCoords = c("x", "y"),
  cellTypeString = "cellType",
  intensityString = "intensity_",
  morphologyString = "morphology_",
  phenotypeString = "phenotype_",
  cellIDString = "cellID",
  cellAnnotations = NULL,
  imageCellIDString = "imageCellID",
  imageIDString = "imageID"
)
```

Arguments

cellData	A data frame that contains at least the columns x and y giving the location coordinates of each cell.
cellProfiler	A logical indicating that cellData is in a format similar to what cellProfiler outputs.
spatialCoords	The column names corresponding to spatial coordinates. eg. x, y, z...
cellTypeString	The name of the column that contains cell type calls.
intensityString	A string which can be used to identify the columns which contain marker intensities. (This needs to be extended to take the column names themselves.)
morphologyString	A string which can be used to identify the columns which contains morphology information.
phenotypeString	A string which can be used to identify the columns which contains phenotype information.
cellIDString	The column name for cellID.
cellAnnotations	A vector of variables that provide additional annotation of a cell.
imageCellIDString	The column name for imageCellID.
imageIDString	The column name for imageIDString.

Value

A SegmentedCells object

Examples

```
### Something that resembles cellProfiler data

set.seed(51773)

n = 10

cells <- data.frame(row.names = seq_len(n))
cells$ObjectNumber <- seq_len(n)
cells$ImageNumber <- rep(seq_len(2),c(n/2,n/2))
cells$AreaShape_Center_X <- runif(n)
cells$AreaShape_Center_Y <- runif(n)
cells$AreaShape_round <- rexp(n)
cells$AreaShape_diameter <- rexp(n, 2)
cells$Intensity_Mean_CD8 <- rexp(n, 10)
cells$Intensity_Mean_CD4 <- rexp(n, 10)

cellExp <- SegmentedCells(cells, cellProfiler = TRUE)

### Cluster cell types
intensities <- cellMarks(cellExp)
kM <- kmeans(intensities,2)
cellType(cellExp) <- paste('cluster',kM$cluster, sep = '')
cellSummary(cellExp)
```

show-SegmentedCells *Show SegmentedCells*

Description

This outputs critical information about a SegmentedCells.

Arguments

object A SegmentedCells.

Value

Information of the number of images, cells, intensities, morphologies and phenotypes.

usage

‘show(object)’

Examples

```

### Something that resembles cellProfiler data

set.seed(51773)

n = 10

cells <- data.frame(row.names = seq_len(n))
cells$ObjectNumber <- seq_len(n)
cells$ImageNumber <- rep(1:2,c(n/2,n/2))
cells$AreaShape_Center_X <- runif(n)
cells$AreaShape_Center_Y <- runif(n)
cells$AreaShape_round <- rexp(n)
cells$AreaShape_diameter <- rexp(n, 2)
cells$Intensity_Mean_CD8 <- rexp(n, 10)
cells$Intensity_Mean_CD4 <- rexp(n, 10)

cellExp <- SegmentedCells(cells, cellProfiler = TRUE)

### Cluster cell types
markers <- cellMarks(cellExp)
kM <- kmeans(markers,2)
cellType(cellExp) <- paste('cluster',kM$cluster, sep = '')

cellExp

```

signifPlot

Plots result of signifPlot.

Description

Plots result of signifPlot.

Usage

```

signifPlot(
  results,
  fdr = FALSE,
  breaks = c(-5, 5, 0.5),
  colors = c("blue", "white", "red"),
  marksToPlot = NULL
)

```

Arguments

results	Data frame obtained from spicy.
fdr	TRUE if FDR correction is used.

breaks	Vector of 3 numbers giving breaks used in pheatmap. The first number is the minimum, the second is the maximum, the third is the number of breaks.
colors	Vector of colours to use in pheatmap.
marksToPlot	Vector of marks to include in pheatmap.

Value

a pheatmap object

Examples

```
data(spicyTest)
signifPlot(spicyTest, breaks=c(-3, 3, 0.5))
```

SpicyResults-class *Performs spatial tests on spatial cytometry data.*

Description

Performs spatial tests on spatial cytometry data.

Usage

```
spicy(
  cells,
  condition = NULL,
  subject = NULL,
  covariates = NULL,
  from = NULL,
  to = NULL,
  dist = NULL,
  integrate = TRUE,
  nsim = NULL,
  verbose = TRUE,
  weights = TRUE,
  window = "convex",
  window.length = NULL,
  BPPARAM = BiocParallel::SerialParam(),
  sigma = NULL,
  Rs = NULL,
  minLambda = 0.05,
  fast = TRUE,
  ...
)
```

Arguments

cells	A SegmentedCells or data frame that contains at least the variables x and y, giving the location coordinates of each cell, and cellType.
condition	Vector of conditions to be tested corresponding to each image if cells is a data frame.
subject	Vector of subject IDs corresponding to each image if cells is a data frame.
covariates	Vector of covariate names that should be included in the mixed effects model as fixed effects.
from	vector of cell types which you would like to compare to the to vector
to	vector of cell types which you would like to compare to the from vector
dist	The distance at which the statistic is obtained.
integrate	Should the statistic be the integral from 0 to dist, or the value of the L curve at dist.
nsim	Number of simulations to perform. If empty, the p-value from lmerTest is used.
verbose	logical indicating whether to output messages.
weights	logical indicating whether to include weights based on cell counts.
window	Should the window around the regions be 'square', 'convex' or 'concave'.
window.length	A tuning parameter for controlling the level of concavity when estimating concave windows.
BPPARAM	A BiocParallelParam object.
sigma	A numeric variable used for scaling when fitting inhomogeneous L-curves.
Rs	A vector of the radii that the measures of association should be calculated.
minLambda	Minimum value for density for scaling when fitting inhomogeneous L-curves.
fast	A logical describing whether to use a fast approximation of the inhomogeneous L-curves.
...	Other options to pass to bootstrap.

Value

Data frame of p-values.

Examples

```
data("diabetesData")

# Test with random effect for patient on only one pairwise combination of cell types.
spicy(diabetesData, condition = "stage", subject = "case",
      from = "Tc", to = "Th")

# Test all pairwise combination of cell types without random effect of patient.
#spicyTest <- spicy(diabetesData, condition = "stage", subject = "case")

# Test all pairwise combination of cell types with random effect of patient.
#spicy(diabetesData, condition = "condition", subject = "subject")
```

```
# Test all pairwise combination of cell types with random effect of patient using  
# a bootstrap to calculate significance.  
#spicy(diabetesData, condition = "stage", subject = "case", nsim = 10000)
```

spicyTest

Results from spicy for diabetesData

Description

Results from the call: `spicyTest <- spicy(diabetesData, condition = "condition", subject = "subject")`

Usage

```
spicyTest
```

Format

spicyTest a spicy object

spicyTestBootstrap

Results from spicy with bootstrap for diabetesData

Description

Results from the call: `spicyTestBootstrap <- spicy(diabetesData, condition = "condition", subject = "subject", nsim = 199)`

Usage

```
spicyTestBootstrap
```

Format

spicyTestBootstrap a spicy object

`topPairs`*A table of the significant results from spicy tests*

Description

A table of the significant results from spicy tests

Usage

```
topPairs(x, coef = 1, n = 10, adj = "fdr", cutoff = NULL)
```

Arguments

<code>x</code>	The output from spicy.
<code>coef</code>	Which coefficient to list.
<code>n</code>	Extract the top n most significant pairs.
<code>adj</code>	Which p-value adjustment method to use, argument for <code>p.adjust()</code> .
<code>cutoff</code>	A p-value threshold to extract significant pairs.

Value

A `data.frame`

Examples

```
data(spicyTest)
topPairs(spicyTest)
```

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