

# Package ‘UMAPVizR’

November 16, 2022

**Encoding** UTF-8

**Version** 1.0-0

**Type** Package

**Title** Visualization and statistical analysis of high dimensional cytometry data using manifold representations

## Description

Cytometry data are now classically analyzed using non linear dimensionality reduction approaches, but it is still challenging to easily handle the whole pipeline of computational analyses. UMAPVizR allows the statistical analysis and visualization of high dimensional cytometry data using manifold algorithms and clustering methods. Especially, several key analysis steps are available to perform data importation, manifold generation, cell cluster identification, statistical analyses, cluster visualization, and quality controls of generated results. UMAPVizR can import cell events from FCS or txt file formats using different transformation, down-sampling, and normalization approaches.

Manifold representations can be generated using the UMAP, tSNE or LargeVis algorithms to project cell events into a lower dimensionality space. The identification of cell clusters can be done using multiple clustering algorithms, depending on user's assumptions. The characteristics of cell clusters can be visualized using scatter plot, categorical heatmap of marker expressions, or using parallel coordinates representations. Cell clusters having abundances differently expressed between biological conditions can be identified using several statistical tests.

Statistical results can be visualized using volcano plots or heatmaps.

Unsupervised and supervised analysis approaches can be conducted by users to appreciate the homogeneity/heterogeneity of biological conditions, and to identify cell population biomarkers in a multivariate manner.

Additionally, UMAPVizR provides a workflow for asserting the quality of identified cell clusters using statistical approaches.

**Author** Gwendolyn MARGUERIT, Nicolas TCHITCHEK

**Imports** checkmate,

cluster,  
concaveman,  
cowplot,  
dbscan,  
dendextend,  
diptest,  
FactoMineR,  
flowCore,  
FNN,

ggdendro,  
 ggiraph,  
 ggnewscale,  
 ggplot2,  
 ggpubr,  
 ggrepel,  
 ggridges,  
 Gmedian,  
 gridExtra,  
 gtools,  
 kohonen,  
 MASS,  
 methods,  
 plyr,  
 reshape,  
 reshape2,  
 rstatix,  
 Rtsne,  
 scales,  
 spade,  
 stats,  
 stringr,  
 uwot,  
 viridis

**Suggests** knitr

**License** GPL-3 | file LICENSE

**VignetteBuilder** knitr

**biocViews** Clustering, DataImport, FlowCytometry, Normalization, StatisticalMethod, Software, Visualization

**RoxygenNote** 7.2.1

## R topics documented:

assignMetadata . . . . .	3
Celldata-class . . . . .	4
computeStatistics . . . . .	4
CreateMetaclusters . . . . .	5
export . . . . .	5
generateManifold . . . . .	6
identifyClusters . . . . .	7
import . . . . .	8
performUpsampling . . . . .	9
plot . . . . .	9
plotBoxplot . . . . .	10
plotCellCounts . . . . .	11
plotClustersCounts . . . . .	12
plotCombineHM . . . . .	12
plotCoordinates . . . . .	13
plotDistogram . . . . .	13
plotHmAbundances . . . . .	14

<i>assignMetadata</i>	3
plotHmExpressions . . . . .	15
plotHmStatistics . . . . .	16
plotLDA . . . . .	16
plotManifold . . . . .	17
plotMDS . . . . .	18
plotPCA . . . . .	19
plotPhenoClusters . . . . .	20
plotScatter . . . . .	20
plotVolcano . . . . .	21
print . . . . .	21
QCMarkerNames . . . . .	22
QCMarkerRanges . . . . .	22
QCSmallClusters . . . . .	23
QCUnciformClusters . . . . .	23
renameMarkers . . . . .	24
show . . . . .	25
<b>Index</b>	<b>26</b>

---

<code>assignMetadata</code>	<i>Assigns meta-information about biological samples</i>
-----------------------------	--

---

## Description

This function aims to attach meta-information to each biological sample.

Especially, the biological individual, the biological condition and the time point of each sample can be specified for subsequent analyses.

## Usage

```
assignMetadata(Celldata, metadata)
```

## Arguments

<code>Celldata</code>	a <code>Celldata</code> object
<code>metadata</code>	a <code>data.frame</code> containing contextual information about the biological samples. This <code>data.frame</code> must have 3 columns specifying for each sample the associated individual (column named 'individual'), the biological condition (column named 'condition') and the time point (column named 'timepoint')

## Value

a S4 object of class 'Celldata'

---

Celldata-class	<i>Celldata class definition</i>
----------------	----------------------------------

---

### Description

The Celldata object is a S4 object containing all cytometry expressions.

### Slots

**samples** a character vector containing the names of the biological samples  
**raw.markers** a character vector containing the names of the raw markers  
**matrix.expression.r** a data.frame containing the raw marker expressions of each cell  
**matrix.expression** a data.frame containing the marker expressions of each cell  
**manifold** a data.frame containing the manifold coordinates  
**manifold.params** a list containing the parameters used for manifold creation  
**identify.clusters** a vector containing the identifiers of cell clusters  
**identify.clusters.params** a vector containing the parameters used for the identification of the cell clusters  
**concave.hulls** a data.frame containing the coordinates of the cell cluster of the concave hulls for each cluster  
**matrix.cell.count** a data.frame containing the number of cells associated to each cluster for each sample  
**matrix.abundance** a data.frame containing the percentage of cells associated to each cluster for each sample  
**statistic** a data.frame containing the statistics of cell clusters  
**metadata** a data.frame containing the metadata associated to each sample

---

computeStatistics	<i>Computes differential analysis statistics for cell clusters</i>
-------------------	--

---

### Description

This function aims to compute the statistics of Differentially Abundant Clusters 'DAC'.

DAC correspond to cell clusters having abundances statistically different between two biological conditions. The statistical test used for the comparisons can be defined by users. For each cluster, the p-value, log2 fold-change and effect size relative to the reference condition are computed. Statistical comparison can be performed in a paired and unpaired manner.

### Usage

```
computeStatistics(
  Celldata,
  condition,
  ref.condition,
  test.statistics = c("wilcox.test", "t.test"),
  paired = FALSE
)
```

**Arguments**

Celldata	a Celldata object
condition	a character value providing the name of the condition to be compared
ref.condition	a character value providing the name of reference condition
test.statistics	a character value providing the type of statistical test to use. Possible values are: 'wilcoxon' or 't-test'
paired	a boolean value indicating if a paired or unpaired comparison should be applied

**Value**

a S4 object of class 'Celldata'

---

CreateMetaclusters	<i>Create Metaclusters</i>
--------------------	----------------------------

---

**Description**

This function aims to xxx

**Usage**

```
CreateMetaclusters(Celldata, clusters, metaclusters)
```

**Arguments**

Celldata	a Celldata object
clusters	a character vector containing the identifiers of the clusters to use. By default, all clusters are used
metaclusters	a character value containing the name of metaclusters

**Value**

xx

---

export	<i>Exports cell expression profiles to TSV or FCS files</i>
--------	---

---

**Description**

Exports cell expression profiles from a Celldata object to a tab-separated or FCS files

Cell expression profiles can be exported for a set of given samples and for a set of given cell clusters

**Usage**

```
export(Celldata, filename, clusters = NULL, samples = NULL)
```

```
## S4 method for signature 'Celldata'
```

```
export(Celldata, filename, clusters = NULL, samples = NULL)
```

**Arguments**

Celldata	a Celldata object
filename	a character value providing the name of the output file
clusters	a character vector containing the identifiers of the cell clusters to export. By default, all clusters are extracted
samples	a character vector containing the names of biological samples to export. By default, all samples are extracted

**Value**

none

---

generateManifold	<i>Generates a manifold of cell events</i>
------------------	--

---

**Description**

This function aims to generate a manifold representation for cell events stored in a Celldata object

This function allows the use of several non-linear dimension reduction techniques such as UMAP, t-SNE or LargeVis The whole set of cell markers or specific cell markers can be used during the dimensionality reduction process

**Usage**

```
generateManifold(
  Celldata,
  type = c("UMAP", "tSNE", "lvish"),
  markers = NULL,
  seed = 42,
  verbose = TRUE,
  ...
)
```

**Arguments**

Celldata	a Celldata object
type	a character value specifying the type of manifold to compute. Possible values are: 'UMAP' for Uniform Manifold Approximation and Projection, 'tSNE' for t Stochastic Neighbor Embedding, and 'lvish' for LargeVis
markers	a character vector providing the cell markers to use for the manifold generation
seed	a numeric value providing the random seed to use during stochastic operations
verbose	a boolean value indicating if computational details must be displayed on the console
...	Other arguments passed on to methods

**Value**

a S4 object of class 'Celldata'

---

identifyClusters	<i>Identify cell cluster of having similar marker expressions</i>
------------------	---

---

## Description

This function aims to identify cell clusters, which are groups of cells having similar expressions for selected markers, using different unsupervised clustering methods.

Several clustering method are available such as kmeans, kmedian, clara, DBSCAN, HDBSCAN and SOM. The cell clustering can be performed on the manifold representation or based on marker expression.

## Usage

```
identifyClusters(
  Celldata,
  space = c("manifold", "markers"),
  markers = NULL,
  method = c("kmeans", "kmedian", "clara", "DBSCAN", "SOM"),
  concavity = 2,
  length.threshold = 0,
  seed = 42,
  ...
)
```

## Arguments

Celldata	a Celldata object
space	a character value containing the space of clustering method to use. Possible values are: 'manifold' or 'markers'
markers	a character vector providing the cell markers to use for the manifold generation
method	a character value containing the name of the clustering method to use. Possible values are: 'kmeans', 'kmedian', 'clara', 'DBSCAN' and 'SOM'
concavity	a numeric value providing a relative measure of concavity for the computation of the concave hulls (please refer to the function 'concaveman' of the 'concave-man' package)
length.threshold	a numeric value providing a threshold of the segment length for the computation of the concave hulls (please refer to the function 'concaveman' of the 'concave-man' package)
seed	a numeric value providing the random seed to use during stochastic operations
...	Other arguments passed on to methods

## Details

For each identify cell cluster, the boundaries of cells belonging to this cluster are delineated using a concave hull

## Value

a S4 object of class 'Celldata'

---

import

---

*Imports of cell expression profiles from TSV or FCS files*


---

## Description

This function aims to import acquired cell events into a `Celldata` object.

Input files can be tab-separated or FCS files. Different transformations can be applied such as logicle, arcsinh or logarithmic. Importantly, a downsampling of cell events can be performed using uniformly-based or density-based random selections. Cell marker having technical or biological biases can be excluded during the import.

## Usage

```
import(
  files,
  filetype = "fcs",
  transform = c("logicle", "arcsinh", "logarithmic", "none"),
  d.method = c("none", "uniform", "density"),
  parameters.method = list(exclude.pctile = 0.01, target.pctile = 0.05, target.number =
    NULL, target.percent = 0.1),
  exclude.markers = NULL,
  seed = 42
)
```

## Arguments

<code>files</code>	a character vector specifying the path of the tab-separated or FCS files to load
<code>filetype</code>	a character vector specifying the format of the loaded files. By default, FCS is used
<code>transform</code>	a character value containing the type of the transformation to apply. Possible values are: 'logicle', 'arcsinh', 'logarithmic' or 'none'
<code>d.method</code>	a character value containing the type of the downsampling to apply. Possible values are: 'none', 'uniform' or 'density'
<code>parameters.method</code>	a list value containing the parameters to use for downsampling
<code>exclude.markers</code>	a character vector providing the marker names to be excluded during the import
<code>seed</code>	a numeric value providing the random seed to use during stochastic operations

## Value

a S4 object of class 'Celldata'



---

performUpsampling	<i>Performs the upsampling of downsampled events</i>
-------------------	--

---

### Description

This function aims to perform upsample downsampled events based on an existing Celldata object and existing cell events stored in tab-separated or FCS files

Importantly, the identification of cell clusters must have been performed prior to this operation

### Usage

```
performUpsampling(
  Celldata,
  files,
  transform = c("logicle", "arcsinh", "logarithmic", "none")
)
```

### Arguments

Celldata	a Celldata object
files	a character vector providing the path of the tab-separated or FCS files
transform	a character value containing the type of the transformation to apply. Possible values are: 'logicle', 'arcsinh', 'logarithmic' or 'none'

### Value

a S4 object of class 'Celldata'

---

plot	<i>Plots graphics for all Celldata objects</i>
------	--

---

### Description

Generates a graphical representation for a Celldata object The displayed representation depends on the current analysis status of the Celldata object

- If the manifold has not been calculated, then the number of cells per sample will be displayed
- If the manifold has been calculated but not the clustering, then the manifold representation will be displayed
- If the manifold and clustering have been calculated, then a heatmap of marker expressions will be displayed

### Usage

```
## S4 method for signature 'Celldata,ANY'
plot(x)
```

### Arguments

x	a Celldata object
---	-------------------

**Value**

a ggplot2 object

---

plotBoxplot	<i>Plots cell cluster abundances using a boxplot representation</i>
-------------	---

---

**Description**

This function aims to visualize and compare the cell cluster abundances for each biological condition using boxplot and jitter representations.

The abundance of a specific cell cluster or a set of cell clusters can be displayed. The representation can be restricted to a specific set of samples. Moreover boxplot can be constructed based on sample meta information. Statistic can be computed for all comparisons.

**Usage**

```
plotBoxplot(
  Celldata,
  clusters,
  samples = NULL,
  observation = c("individual", "condition", "timepoint"),
  test.statistics = c("wilcox.test", "t.test"),
  paired = FALSE,
  hide.ns = TRUE
)
```

**Arguments**

<code>Celldata</code>	a Celldata object
<code>clusters</code>	a character vector containing the identifiers of the clusters to use
<code>samples</code>	a character vector containing the names of biological samples to use. By default, all samples are used
<code>observation</code>	a character value containing the parameters to use
<code>test.statistics</code>	a character value providing the type of statistical test to use. Possible values are: 'wilcox.test' or 't.test'
<code>paired</code>	a boolean value indicating if a paired or unpaired comparison should be applied
<code>hide.ns</code>	a boolean value indicating xxx

**Value**

a ggplot2 object

---

plotCellCounts	<i>Plots the numbers of cells for each sample</i>
----------------	---

---

## Description

This function aims to visualize the number of cells associated to each sample.

This representation displays the samples in the X-axis and the number of associated cells in the Y-axis. Several statistics can be computed and shown.

## Usage

```
plotCellCounts(  
  Celldata,  
  stats = c("min", "median", "mean", "q75", "max"),  
  samples = NULL,  
  sort = TRUE  
)
```

## Arguments

Celldata	a Celldata object
stats	a character vector providing the statistics to display. Possible values are: 'min', 'median', 'mean', 'q75', 'max'
samples	a character vector containing the names of biological samples to use. By default, all samples are used
sort	a boolean value indicating if clusters must be sorted by the number associated sample

## Details

The following statistic can be computed:

- 'min' corresponds to the lowest number of cells within a data set
- 'median' corresponds to the number of cells separates the lower half from the upper half within data set
- 'mean' corresponds to the number of cells quantity shared within data set
- 'q75' corresponds to the number of cells separates the quantiles 75
- 'max' corresponds to the largest number of cells within a data set

## Value

a ggplot2 object

---

plotClustersCounts	<i>Plots the numbers of cells for each clusters</i>
--------------------	---

---

**Description**

This function aims to visualize the number of cells associated to each clusters.

This representation displays the clusters in the X-axis and the total number of associated cells in the Y-axis.

**Usage**

```
plotClustersCounts(
  Celldata,
  clusters = NULL,
  sort = TRUE,
  legend.max.samples = 10
)
```

**Arguments**

Celldata	a Celldata object
clusters	a character vector containing the identifiers of the clusters to use. By default, all clusters are used
sort	a boolean value indicating if clusters must be sorted by the number associated cluster
legend.max.samples	xxx

**Value**

a ggplot2 object

---

plotCombineHM	<i>Plots a combined expression and statistic heatmaps</i>
---------------	---

---

**Description**

This function aims to combine the expression and statistic heatmaps.

**Usage**

```
plotCombineHM(HM1, HM2)
```

**Arguments**

HM1	a ggplot object containing the expression heatmap
HM2	a ggplot object containing the statistic heatmap

**Value**

a ggplot2 object

---

plotCoordinates	<i>Plots of phenotype of cell clusters using parallels coordinates</i>
-----------------	--

---

**Description**

This function aims to visualize xxx

**Usage**

```
plotCoordinates(
  Celldata,
  condition.samples = c("condition", "timepoint"),
  samples = NULL,
  clusters
)
```

**Arguments**

Celldata	a Celldata object
condition.samples	a character vector containing the variable to be studied for the samples. Possible values are: 'condition' or 'timepoint'
samples	a character vector containing the names of biological samples to use. By default, all samples are used
clusters	a character vector containing the identifiers of the clusters to use

**Value**

xx

---

plotDistogram	<i>Plots of a distogram of marker co-expression</i>
---------------	---

---

**Description**

This function aims to visualize the pairwise co-expression between all markers using a distogram representation. Each tile corresponds to the co-expression between two markers and is gradient-colored based on the Pearson correlation

**Usage**

```
plotDistogram(Celldata, clusters = NULL)
```

**Arguments**

Celldata	a Celldata object
clusters	a character vector containing the identifier of the cluster to use

**Details**

The Pearson correlation is computed based on the marker expressions

**Value**

a ggplot2 object

---

plotHmAbundances	<i>Plots an heatmap of cell cluster abundances</i>
------------------	--

---

**Description**

This function aims to visualize the abundances of cell clusters using an heatmap representation

In such heatmap each column corresponds a cell cluster and he row corresponds the different samples The heatmap can be restricted to specific cell clusters and samples. The levels of abundance of each sample in each cluster is represented using a color gradient scale Abundance values can be centered and reduced.

**Usage**

```
plotHmAbundances(
  Celldata,
  clusters = NULL,
  samples = NULL,
  saturation = 2.5,
  rescale = FALSE
)
```

**Arguments**

Celldata	a Celldata object
clusters	a character vector containing the identifiers of the clusters to use. By default, all clusters are used
samples	a character vector containing the names of biological samples to use. By default, all samples are used
saturation	a numeric value providing the saturation threshold of cell cluster abundances
rescale	a boolean specifying if cell cluster abundances must be centered and reduced

**Value**

a ggplot2 object

---

plotHmExpressions	<i>Plots an heatmap of cell marker expressions</i>
-------------------	--

---

## Description

This function aims to visualize the cell marker expressions for selected markers and clusters

The mean of median marker expressions is computed for each cluster, and marker expressions displayed using a categorical heatmap (5 categories are defined by default) The range expression of each cell marker is discretized into several categories between bounds of marker expressions To hierarchical clustering, shown using dendrogramm, can be computed on both marker and cluster levels

## Usage

```
plotHmExpressions(  
  Celldata,  
  markers = NULL,  
  clusters = NULL,  
  method.hclust = c("ward.D", "ward.D2", "single", "complete", "average", "mcquitty",  
    "median", "centroid"),  
  nb.cat = 5,  
  seed = 42  
)
```

## Arguments

Celldata	a Celldata object
markers	a character vector providing the marker names to use. By default, all markers are used
clusters	a character vector containing the identifiers of the clusters to use. By default, all clusters are used
method.hclust	a character value providing the agglomeration method to be use. Possible values are: 'ward.D', 'ward.D2', 'single', 'complete', 'average', 'mcquitty', 'median' or 'centroid' (please refer to the function 'hclust' of the 'stats' package)
nb.cat	a numeric specifying the number of categories to use
seed	a numeric value providing the random seed to use during stochastic operations

## Value

a ggplot2 object

---

plotHmStatistics	<i>Plots an heatmap of a statistical analysis results</i>
------------------	---

---

### Description

This function aims to visualize the results of differential cell clusters analysis

This representation displays statistical information for each cell cluster for a given comparison of samples. Different statistics can be visualized, such as the p-value, the log2(fold-change), and effect size.

### Usage

```
plotHmStatistics(
  Celldata,
  clusters = NULL,
  statistics = c("pvalue", "lfc", "effsize"),
  saturation = 3
)
```

### Arguments

<code>Celldata</code>	a <code>Celldata</code> object
<code>clusters</code>	a character vector containing the identifiers of the clusters to use. By default, all clusters are used
<code>statistics</code>	a character value providing the name of the statistic to display. Possible values are: 'pvalue' for p-value, 'lfc' for log2 fold change or 'eff' for effect size
<code>saturation</code>	a numeric value providing the saturation value for statistics to display

### Value

a `ggplot2` object

---

plotLDA	<i>Plots a LDA representation based cell cluster abundances</i>
---------	---

---

### Description

This function aims to represent a Linear Discriminant Analysis representation based on cell cluster abundances.

### Usage

```
plotLDA(
  Celldata,
  levels = c("predictions", "coefficients"),
  ref.condition,
  condition,
  clusters = NULL
)
```



**Arguments**

Celldata	a Celldata object
levels	a character value containing the variable to be displayed. Possible values are: 'clusters' or 'samples'
ref.condition	a character value providing the name of reference condition
condition	a character value providing the name of the condition to be compared
clusters	a character vector containing the identifiers of the clusters to use. By default, all clusters are used

**Value**

xx

---

plotManifold	<i>Plots a representation of a computed manifold</i>
--------------	--

---

**Description**

This function aims to visualize a computed manifold representation for given analysis.

This representation can be used on a Celldata object for which a manifold analysis has been performed.

If a cell clustering has been performed, then the clusters are delineated using concave hulls. Additionally, the manifold can be colored based on the local cell density or marker expressions. It is possible to centred and reduce the values of expressions.

**Usage**

```
plotManifold(
  Celldata,
  markers = "density",
  samples = NULL,
  observation = c("individual", "condition", "timepoint"),
  scale = FALSE,
  quant.low = 0.05,
  quant.high = 0.95
)
```

**Arguments**

Celldata	a Celldata object
markers	a character value providing the name of the marker to use for the colouring. By default, cells are colored based on their local density
samples	a character vector containing the names of biological samples to use
observation	a character value containing the parameters to use
scale	a boolean value specifying if expression value must be rescaled
quant.low	a numeric value providing the number of first quantile
quant.high	a numeric value providing the number of last quantile

**Value**

a ggplot2 object

---

plotMDS	<i>Plots a MDS representation based on cell cluster abundances</i>
---------	--

---

**Description**

This function aims to visualize the similarities between samples or clusters based on their abundances, using a Multidimensional Scaling representation. Each dot represents a sample or a cluster and the distances between the dots are proportional to the Euclidean distance between these objects. The representation can be restricted to specific cell clusters and samples. In addition, it is possible to choose the levels displayed, clusters or samples.

**Usage**

```
plotMDS(
  Celldata,
  matrix = c("abundance", "expression"),
  levels = c("clusters", "samples"),
  condition.samples = c("condition", "timepoint"),
  clusters = NULL,
  samples = NULL,
  plot.text = TRUE
)
```

**Arguments**

<code>Celldata</code>	a Celldata object
<code>matrix</code>	a character vector containing the matrix to be studied. Possible values are: 'abundance' or 'expression'
<code>levels</code>	a character value containing the variable to be displayed. Possible values are: 'clusters' or 'samples'
<code>condition.samples</code>	a character vector containing the variable to be studied for the samples. Possible values are: 'condition' or 'timepoint'
<code>clusters</code>	a character vector containing the identifiers of the clusters to use. By default, all clusters are used
<code>samples</code>	a character vector containing the names of biological samples to use. By default, all samples are used
<code>plot.text</code>	a boolean value specifying if adds text directly at the plot

**Value**

a ggplot2 object

**Description**

This function aims to represent a Principal Component Analysis representation based on cell cluster abundances. In such representation, clusters or samples are positioned based on computed principal components. The representation can be displayed based on specific principal components. The representation can be restricted to specific cell clusters and samples. In addition, it is possible to choose the levels displayed, clusters or samples.

**Usage**

```
plotPCA(
  Celldata,
  levels = c("both", "clusters", "samples"),
  clusters = NULL,
  samples = NULL,
  components = c(1, 2),
  condition.samples = c("condition", "timepoint"),
  cor.radius.th = 0.6,
  plot.text = TRUE
)
```

**Arguments**

<code>Celldata</code>	a <code>Celldata</code> object
<code>levels</code>	a character value containing the variable to be displayed. Possible values are: 'both', 'clusters' or 'samples'
<code>clusters</code>	a character vector containing the identifier of the cluster to use. By default, all clusters are used
<code>samples</code>	a character vector containing the names of biological samples to use. By default, all samples are used
<code>components</code>	a numeric vector providing the components to display
<code>condition.samples</code>	a character vector containing the variable to be studied for the samples. Possible values are: 'condition' or 'timepoint'
<code>cor.radius.th</code>	a numeric value specifying xxx
<code>plot.text</code>	a boolean value specifying if adds text directly at the plot

**Value**

a `ggplot2` object

---

plotPhenoClusters	<i>Plots of phenotype of identified cell clusters</i>
-------------------	---

---

### Description

This function aims to visualize xxx

### Usage

```
plotPhenoClusters(
  Celldata,
  clusters,
  quant.low = 0.05,
  quant.high = 0.95,
  dip.th = 0.01
)
```

### Arguments

Celldata	a Celldata object
clusters	a character vector containing the identifier of the cluster to use
quant.low	a numeric value providing the number of first quantile
quant.high	a numeric value providing the number of last quantile
dip.th	a numeric value specifying xxx

### Value

a ggplot2 object

---

plotScatter	<i>Plots of a scatter plot of marker co-expression</i>
-------------	--

---

### Description

This function aims to visualize co-expression between two markers using a scatter representation

### Usage

```
plotScatter(Celldata, marker1, marker2, samples = NULL, clusters = NULL)
```

### Arguments

Celldata	a Celldata object
marker1	a character value specifying the first marker to be visualised
marker2	a character value specifying the second marker to be visualised
samples	a character vector containing the names of biological samples to use. By default, all samples are used
clusters	a character vector containing the identifiers of the clusters to use. By default, all clusters are used

**Value**

a ggplot2 object

---

plotVolcano	<i>Plots of a volcano plot of statistical analysis</i>
-------------	--

---

**Description**

This function aims to visualize the results of a differentially abundant analysis using a Volcano plot. In such representation, each dot corresponds to a cell cluster and dots are positioned in two dimensional space where the X-axis represents the  $\log_2(\text{fold-change})$  and the Y-axis represents the  $-\log_{10}$  of the p-value. A horizontal line is displayed accordingly to the p-value threshold and two vertical lines are displayed accordingly to the fold-change threshold.

**Usage**

```
plotVolcano(Celldata, comparison, th.pv = 1.3, th.fc = 1.5, plot.text = TRUE)
```

**Arguments**

Celldata	a Celldata object
comparison	a character value containing the comparison to study
th.pv	a numeric value containing the p-value threshold to use
th.fc	a numeric value containing the fold-change threshold to use
plot.text	a boolean value specifying if adds text directly at the plot

**Value**

a ggplot2 object

---

print	<i>Prints information for a given Celldata object</i>
-------	---

---

**Description**

Prints a preview for a Celldata object

**Usage**

```
## S4 method for signature 'Celldata'
print(x)
```

**Arguments**

x	a Celldata object
---	-------------------

**Value**

none

---

QCMarkerNames	<i>Verifies the consistency of the marker names within cell event files</i>
---------------	---

---

### Description

This function aims to check the consistency of marker names across multiple tab-separated or FCS files.

Additionally, the number of cells associated to each sample is displayed.

### Usage

```
QCMarkerNames(files)
```

### Arguments

files	a character vector specifying the path of the tab-separated or FCS files to check
-------	---

### Value

a data.frame containing the marker names and the associated number of cells for each sample (rownames = samples and colnames = markers)

---

QCMarkerRanges	<i>Verifies the consistency of marker expressions integrity within cell event files</i>
----------------	---

---

### Description

This function aims to check the consistency of marker expressions ranges across multiple tab-separated or FCS files.

The marker expressions ranges are calculated based on the user-defined quantiles.

### Usage

```
QCMarkerRanges(files, probs = c(0.05, 0.95))
```

### Arguments

files	a character vector specifying the path of the FCS files to verified
probs	a numerical vector providing the quantiles used to define marker expressions ranges

### Value

a list containing two data.frame for the lower and upper marker expression ranges (rownames = samples and colnames = markers)

---

QC <small>Small</small> Clusters	<i>Computes the percentage of cell clusters with low number of cells</i>
----------------------------------	--

---

**Description**

This function aims to compute and show cell clusters having a number of associated cells lower than a specific threshold

**Usage**

```
QCSmallClusters(Celldata, th.size = 50, plot.device = TRUE)
```

**Arguments**

Celldata	a Celldata object
th.size	a numeric value providing the minimum number of cells needed for a cluster to be considered a small cluster
plot.device	a boolean value specifying a results representation must be displayed

**Value**

a numerical value corresponding to the percentage of cell cluster with low number of cells

---

QCUniformClusters	<i>Computes the percentage of clusters with uniform phenotypes</i>
-------------------	--

---

**Description**

This function aims to identify and show cell clusters having a uniform phenotype

A uniform cluster corresponds to a cluster that have a unimodal expression and a low spread of expression for all its markers

**Usage**

```
QCUniformClusters(  
  Celldata,  
  uniform.test = c("both", "uniform", "IQR"),  
  th.pvalue = 0.05,  
  th.IQR = 2,  
  plot.device = TRUE  
)
```

**Arguments**

<code>Celldata</code>	a <code>Celldata</code> object
<code>uniform.test</code>	a character providing the name of test assessment to perform. Possible value are : 'both', 'uniform', 'IQR'
<code>th.pvalue</code>	a numeric value providing the p-value threshold of the Hartigan's dip test (unimodal if $pvalue > th.pvalue$ )
<code>th.IQR</code>	a numeric value providing the IQR (interquartile range) threshold to assume a distribution as uniform
<code>plot.device</code>	a boolean value specifying if result representation must be displayed

**Details**

- 'uniform' corresponds to the verification of the unimodal distribution of markers with a Hartigans test

- 'IQR' corresponds to the verification of the distribution of markers so that they are not below the IQR threshold (interquartile range)

- 'both' corresponds to the combination of the two parameters : uniform and IQR

**Value**

a numerical value corresponding to the percentage of cell cluster with unimodal expression and a low spread

---

<code>renameMarkers</code>	<i>Renames markers within a <code>Celldata</code> object</i>
----------------------------	--

---

**Description**

This function aims to rename cell markers stored within a `Celldata` object.

This function is interesting to remove the names of the fluorochromes or metals recorded during the acquisition process.

**Usage**

```
renameMarkers(Celldata, marker.names)
```

**Arguments**

<code>Celldata</code>	a <code>Celldata</code> object
<code>marker.names</code>	a character vector providing the new marker names to use

**Value**

a S4 object of class 'Celldata'



---

show	<i>Prints information for a Celldata objects</i>
------	--

---

**Description**

Shows a preview for a Celldata object

**Usage**

```
## S4 method for signature 'Celldata'  
show(object)
```

**Arguments**

object            a Celldata object

**Value**

none

# Index

`assignMetadata`, [3](#)

`Celldata` (`Celldata`-class), [4](#)  
`Celldata`-class, [4](#)  
`computeStatistics`, [4](#)  
`CreateMetaclusters`, [5](#)

`export`, [5](#)  
`export`, `Celldata`-method (`export`), [5](#)

`generateManifold`, [6](#)

`identifyClusters`, [7](#)  
`import`, [8](#)

`performUpsampling`, [9](#)  
`plot`, [9](#)  
`plot`, `Celldata`, ANY-method (`plot`), [9](#)  
`plotBoxplot`, [10](#)  
`plotCellCounts`, [11](#)  
`plotClustersCounts`, [12](#)  
`plotCombineHM`, [12](#)  
`plotCoordinates`, [13](#)  
`plotDistogram`, [13](#)  
`plotHmAbundances`, [14](#)  
`plotHmExpressions`, [15](#)  
`plotHmStatistics`, [16](#)  
`plotLDA`, [16](#)  
`plotManifold`, [17](#)  
`plotMDS`, [18](#)  
`plotPCA`, [19](#)  
`plotPhenoClusters`, [20](#)  
`plotScatter`, [20](#)  
`plotVolcano`, [21](#)  
`print`, [21](#)  
`print`, `Celldata`-method (`print`), [21](#)

`QCMarkerNames`, [22](#)  
`QCMarkerRanges`, [22](#)  
`QCSmallClusters`, [23](#)  
`QCUniformClusters`, [23](#)

`renameMarkers`, [24](#)

`show`, [25](#)  
`show`, `Celldata`-method (`show`), [25](#)