

# Package ‘flowDensity’

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**Type** Package

**Title** Sequential Flow Cytometry Data Gating

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**Description** This package provides tools for automated sequential gating analogous to the manual gating strategy based on the density of the data.

**Imports** flowCore, graphics, car, sp, rgeos, gplots, RFOC, GEOmap, flowWorkspace (>= 3.21.5), methods, grDevices

**Depends** R (>= 2.10.0), methods

**SystemRequirements** GEOS (>= 3.2.0); for building from source: GEOS from <http://trac.osgeo.org/geos/>; GEOS OSX frameworks built by William Kyngesburye at <http://www.kyngchaos.com/> may be used for source installs on OSX.

**License** Artistic-2.0

**biocViews** Bioinformatics, FlowCytometry, CellBiology, Clustering, Cancer, FlowCytData, StemCells, DensityGating

**LazyLoad** yes

**NeedsCompilation** no

## R topics documented:

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CellPopulation-class    *Class "CellPopulation"*

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### Description

This class represents the output of 'flowDensity(.)' function from flowDensity package.

### Objects from the Class

Objects can be created by calls of the form `new("CellPopulation", ...)`.

### Slots

`flow.frame`: Object of class "flowFrame" representing the flow cytometry data of the cell population

`proportion`: Object of class "numeric" representing proportion of the cell population with respect to its parent cell population

`cell.count`: Object of class "numeric" representing cell count of the cell population

`channels`: Object of class "character" representing channel names corresponding to the 2 dimensions where the cell population is extracted

`position`: Object of class "logical" representing position of the cell population in the 2-dimensional space

`gates`: Object of class "numeric" representing thresholds on each channel used to gate the cell population

`filter`: Object of class "matrix" representing boundary of the cell population using a convex polygon

`index`: Object of class "numeric" representing indices of the data points in the cell population with respect to its parent cell population

### Methods

**flowDensity** signature(obj = "CellPopulation", channels = "ANY", position = "logical", singlet.gate ...  
 ...

**flowDensity** signature(obj = "CellPopulation", channels = "missing", position = "missing", singlet.g ...  
 ...

**getflowFrame** signature(obj = "CellPopulation"): ...

**plot** signature(x = "flowFrame", y = "CellPopulation"): ...

### Author(s)

Jafar Taghiyar <email: <jtaghiyar@bccrc.ca>

### Examples

```
showClass("CellPopulation")
```

---

deGate *1D density gating method*

---

## Description

Find the best threshold for a single channel in flow cytometry data based on its density distribution.

## Usage

```
deGate(obj,channel, n.sd = 1.5, use.percentile = FALSE, percentile = 0.95,use.upper=FALSE, upper :
all.cuts = FALSE, tinypeak.removal=1/25,node=NA, adjust.dens = 1,count.lim=20,magnitude=.3, ...)
```

## Arguments

obj	obj: a 'FlowFrame' object, 'CellPopulation' or 'GatingHierarchy'
channel	a channel's name or its corresponding index in the 'flow.frame'.
n.sd	an integer coefficient for the standard deviation to determine the threshold based on the standard deviation if 'sd.threshold' is TRUE.
use.percentile	if TRUE, forces to return the 'percentile'th threshold.
percentile	a value in [0,1] that is used as the percentile. The default value is 0.95.
use.upper	Logical. If TRUE, forces to return the inflection point based on the first (last) peak if upper=F (upper=T). Default value is set to 'FALSE'
upper	if TRUE, finds the change in the slope at the tail of the density curve, if FALSE, finds it at the head. Default value is set to 'NA'.
verbose	Logical. If TRUE, Prints a message if only one peak is found, or when inflection point is used to set the gates.
twin.factor	a value in [0,1] that is used to exclude twinpeaks
bimodal	Logical. If TRUE, it returns a cutoff that splits population closer to 50-50, when there are more than two peaks.
after.peak	Logical. If TRUE, it returns a cutoff that is after the maximum peaks, when there are more than two peaks.
alpha	a value in [0,1) specifying the significance of change in the slope being detected. This is by default 0.1, and typically need not be changed.
sd.threshold	if TRUE, uses 'n.sd' times standard deviation as the threshold. Default value is set to 'FALSE'.
all.cuts	if TRUE, returns all the identified cutoff points, i.e. potential thresholds for that channel. Default value is set to 'FALSE'.
tinypeak.removal	A number in [0,1] to exclude/include tiny peaks in density distribution.
node	A character defining the parent population when extracting data from GatingHierarchy.
adjust.dens	The smoothness of density in [0,Inf] to be used in density(.). The default value is 1 and should not be changed unless necessary
count.lim	minimum limit for events count in order to calculate the threshold. Default is 20, returning NA as threshold.
magnitude	A value between 0 and 1, for tracking a slope and reporting changes that are smaller than magnitude*peak_height
...	Extra arguments to be passed to smoothSpline function.

**Value**

an integer value (vector) of cutoff(s), i.e. threshold(s), on the specified channel

**Author(s)**

Mehrnoush Malek <<mmalekes@bccrc.ca>>

**See Also**

[getflowFrame](#) [notSubFrame](#) [flowDensity](#)

**Examples**

```
data_dir <- system.file("extdata", package = "flowDensity")
load(list.files(pattern = 'sampleFCS_1', data_dir, full = TRUE))
#Find the threshold for CD20
cd19.gate <- deGate(f,channel="PerCP-Cy5-5-A")
# Gate out the CD20- populations using the notSubFrame
plotDens(f,c("APC-H7-A","PerCP-Cy5-5-A"))
abline(h=cd19.gate,lty=3,col=2)
```

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flowDensity

*Automated Sequential Gating Tool for Flow Cytometry*

---

**Description**

flowDensity is an automated clustering algorithm which aims to emulate the current practice of manual sequential gating. It is designed to identify the predefined cell subsets based on the density distribution of the parent cell population by analyzing the peaks of the density curve.

When the density distribution has only one peak:

- the first argument that would be used is the percentile, the default value is 95th.
- If set to 'NA' then upper will be used when set to 'FALSE/TRUE'.
- If both upper and percentiles are set to 'NA' sd.threshold is used when set to 'TRUE'.
- If either all these arguments are 'NA', or one of them fails, the algorithm find the best cutoff based on the inflection points and by comparing the position of the peak and the mean of the population.

**Usage**

```
flowDensity(obj, channels, position,node, ...)
```

**Arguments**

obj	a 'CellPopulation', 'flowFrame' or 'GatingHierarchy' object.
channels	a vector of two channel names or their corresponding indices.
position	a vector of two logical values specifying the position of the cell subset of interest on the 2D plot.

node A character defining the parent population when extracting data from GatingHierarchy.

...

This can be used to pass one of the following arguments:

- 'use.percentile' if TRUE, returns the 'percentile'th threshold.
- 'percentile' a value in [0,1] that is used as the percentile if 'use.percentile' is TRUE.
- 'upper' if 'TRUE', it finds the change in the slope after the peak with index 'peak.ind'.
- 'use.upper' if 'TRUE', forces to return the inflection point based on the first (last) peak if upper=F (upper=T)
- 'twin.factor' a value in [0,1] that is used to exclude twinpeaks.
- 'bimodal' If TRUE, it returns a cutoff that splits population closer to 50-50, when there are more than two peaks.
- 'after.peak' If TRUE, it returns a cutoff that is after the maximum peaks, when there are more than two peaks.
- 'sd.threshold' if TRUE, it uses 'n.sd' times standard deviation for gating.
- 'n.sd' an integer that is multiplied to the standard deviation to determine the place of threshold if 'sd.threshold' is 'TRUE'.
- 'tynypeak.removal' a vector of length 2, for sensitivity of peak finding for each channel. See deGate() for more information.
- 'filter' If provided it uses the given filter to gate the population.
- 'use.control' if TRUE, it finds the threshold using a matched control population and uses it for gating.
- 'control' a 'flowFrame' or 'CellPopulation' object used for calculating the gating threshold when 'use.control' is set to TRUE. If a control population is used, the other arguments ('upper', 'percentile', etc.) are applied to the control data when finding the threshold (i.e. not to 'obj').
- 'alpha' a value in [0,1] specifying the significance of change in the slope which would be detected. This is by default 0.1, and typically need not be changed.
- 'ellip.gate' if TRUE, it fits an ellipse on the data as a gate, otherwise the rectangle gating results are returned
- 'scale' a value in [0,1] that scales the size of ellipse to fit if 'ellip.gate' is TRUE

### Value

A CellPopulation object

### Author(s)

Mehrnoush Malek <<mmalekes@bccrc.ca>> Jafar Taghiyar <<jtaghiyar@bccrc.ca>>

### See Also

[deGate](#) [getFlowFrame](#) [notSubFrame](#)

**Examples**

```
data_dir <- system.file("extdata", package = "flowDensity")
load(list.files(pattern = 'sampleFCS_1', data_dir, full = TRUE))
lymph <- flowDensity(obj=f, channels=c('FSC-A', 'SSC-A'),
                    position=c(TRUE, FALSE), upper= c(NA, TRUE), debris.gate=c(TRUE, FALSE))
slotNames(lymph)
```

---

flowDensity-methods      *Methods for Function flowDensity in Package **flowDensity***

---

**Description**

Methods for function flowDensity in package **flowDensity**

**Usage**

```
flowDensity(obj, channels, position,node, ...)
## S4 method for signature 'CellPopulation, ANY, logical, missing'
flowDensity(obj, channels, position, ...)
## S4 method for signature 'flowFrame,ANY, logical, missing)'
flowDensity(obj, channels, position, ...)
## S4 method for signature 'GatingHierarchy, ANY, logical, ANY'
flowDensity(obj, channels, position,node, ...)
```

**Arguments**

obj	GatingHierarchy or <a href="#">CellPopulation</a> object
channels	a vector of two channel names or their corresponding indices.
position	a vector of two logical values specifying the position of the cell subset of interest on the 2D plot.
node	A character defining the parent population when extracting data from GatingHierarchy.
...	Check flowDensity description.

**Value**

a CellPopulation object.

---

getflowFrame                    *'CellPopulation' class accessor.*

---

### Description

an accessor for 'CellPopulation' class to get its 'FlowFrame' object. This will remove all the NA values in the frame.

### Usage

```
getflowFrame(obj)
```

### Arguments

obj                    a 'CellPopulation' object.

### Value

a 'FlowFrame' object.

### Author(s)

Jafar Taghiyar <<jtaghiyar@bccrc.ca>>

### Examples

```
data_dir <- system.file("extdata", package = "flowDensity")
load(list.files(pattern = 'sampleFCS_1', data_dir, full = TRUE))
lymph <- flowDensity(obj=f, channels=c('FSC-A', 'SSC-A'),
                    position=c(TRUE, FALSE), upper= c(NA, TRUE), debris.gate=c(TRUE, FALSE))
f.lymph <- getflowFrame(lymph)
```

---

getPeaks                    *Finding Peaks*

---

### Description

Find all peaks in density along with their indices

### Usage

```
getPeaks(frame, channel, tinypeak.removal=1/25, ...)
```

### Arguments

frame                    a 'FlowFrame' object.  
channel                    a channel's name or its corresponding index in the 'flow.frame'.  
tinypeak.removal                    A number in [0,1] to exclude/include tiny peaks in density distribution.  
...                    Arguments passed to smoothSpline function, spar is 0.4.

**Value**

a list, including peaks and their corresponding indices

**Author(s)**

Mehrnoush Malek <<mmalekes@bccrc.ca>>

**See Also**

[deGate](#) [notSubFrame](#) [flowDensity](#)

**Examples**

```
data_dir <- system.file("extdata", package = "flowDensity")
load(list.files(pattern = 'sampleFCS_1', data_dir, full = TRUE))
#Find the threshold for CD20
peaks <- getPeaks(f,channel="PerCP-Cy5-5-A",tinypeak.removal=1/30)
peaks
```

---

nmRemove

*Preprocessing helper function for flow cytometry data*

---

**Description**

Remove the margin events on the axes. Usually, these events are considered as debris or artifacts. This is specifically useful for 'FSC' and 'SSC' channels in a 'FlowFrame' object. However, any channel can be input as an argument.

**Usage**

```
nmRemove( flow.frame, channels, neg=FALSE, verbose=FALSE,return.ind=FALSE)
```

**Arguments**

flow.frame	a 'FlowFrame' object.
channels	a vector of channel names or their corresponding indices.
neg	if TRUE, negative events are also removed
verbose	if TRUE, it prints the margin event in each channel
return.ind	if TRUE, it return indices of margin events for each channel.

**Value**

a 'FlowFrame' object, or a 'list' of indices identifying margin events for each channel.

**Author(s)**

Jafar Taghiyar <<jtaghiyar@bccrc.ca>> Mehrnoush Malek <<mmalekes@bccrc.ca>>



## Examples

```
data_dir <- system.file("extdata", package = "flowDensity")
load(list.files(pattern = 'sampleFCS_2', data_dir, full = TRUE))
#Removing margin events of FSC-A and SSC-A channels
no.margin <- nmRemove(f2, c("FSC-A", "SSC-A"), verbose=TRUE)
plotDens(f2, c("FSC-A", "SSC-A"))
# Scatter plot of FSC-A vs. SSC-A after removing margins
plotDens(no.margin, c("FSC-A", "SSC-A"))
```

---

notSubFrame

*Removing a subset of a FlowFrame object*


---

## Description

Remove a subset of a FlowFrame object specified by gates from the flowDensity method. It comes in handy when one needs the complement of a cell population in the input flow cytometry data.

## Usage

```
notSubFrame(obj, channels, position = NA, gates, filter)
```

## Arguments

obj	a 'FlowFrame' or 'cellPopulation' object.
channels	a vector of two channel names or their corresponding indices in the 'flow.frame'.
position	a vector of two logical values specifying the position of the cell subset of interest on the 2D plot.
gates	the gates slot in the CellPopulation object which is output by flowDensity function. It can also be a vector of two integer values each of which specifies a threshold for the corresponding channel in 'channels' argument.
filter	boundary of the subset to be removed. This value is stored in the 'filter' slot of a 'CellPopulation' object.

## Value

a CellPopulation object.

## Author(s)

Mehrnoush Malek <<mmalekes@bccrc.ca>>

## Examples

```
data_dir <- system.file("extdata", package = "flowDensity")
load(list.files(pattern = 'sampleFCS_1', data_dir, full = TRUE))
#Find the threshold for CD20
cd20.gate <- deGate(f, channel="APC-H7-A")
# Gate out the CD20- populations using the notSubFrame
CD20.pos <- notSubFrame(f, channels=c("APC-H7-A", "PerCP-Cy5-5-A"), position=c(FALSE, NA), gates=c(cd20.gate, NA))
#Plot the CD20+ cells on same channels
plotDens(CD20.pos@flow.frame, c("APC-H7-A", "PerCP-Cy5-5-A"))
```

---

`plotDens`*Plot flow cytometry data with density-based colors*

---

**Description**

Generate a scatter dot plot with colors based on the distribution of the density of the provided channels.

**Usage**

```
plotDens(obj, channels,node=NA , col, main, xlab, ylab, pch=".", ...)
```

**Arguments**

<code>obj</code>	a 'FlowFrame','GatingHierarchy' or 'cellPopulation' object.
<code>channels</code>	a vector of two channel names or their corresponding indices in the 'flow.frame'.
<code>node</code>	Only used if <code>gatingHierarchy</code> is provided as an input.It subtractes the <code>flowFrame</code> from the node of <code>gatingHierarchy</code> . see <code>getNode()</code> in <code>flowWorkspace</code> .
<code>col</code>	A specification for the default plotting color: see '?par'.
<code>main</code>	an overall title for the plot: see '?plot'
<code>xlab</code>	a title for the x axis: see '?plot'
<code>ylab</code>	a title for the y axis: see '?plot'
<code>pch</code>	Either an integer specifying a symbol or a single character to be used as the default in plotting points: see '?par'.
<code>...</code>	can be used to provide desired arguments for the <code>plot()</code> function used to plot the output results.

**Value**

a scatter dot plot with density-based colors.

**Author(s)**

Mehrnoush Malek <<mmalekes@bccrc.ca>> Jafar Taghiyar <<jtaghiyar@bccrc.ca>>

**Examples**

```
data_dir <- system.file("extdata", package = "flowDensity")
load(list.files(pattern = 'sampleFCS_1', data_dir, full = TRUE))
#Plot CD3 vs. CD19 to see the distribution of cell populations and their density
plotDens(f,c("V450-A","PerCP-Cy5-5-A"))
```

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