

# plateCore

April 20, 2011

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`applyControlGates` *Apply control gates to a flowPlate*

---

## Description

Once `setControlGates` has been used to create gates for a `flowPlate` object, gates are applied to test samples using `applyControlGates`. The `applyControlGates` function is separated from `setControlGates` since gates may need be changed outside of `setControlGates`.

## Usage

```
applyControlGates(data, gateType="Negative.Control", ...)
```

## Arguments

<code>data</code>	A <code>flowPlate</code> dataset.
<code>gateType</code>	The type of gate to be applied to the <code>flowPlate</code> . Currently only "Negative.Control" gates are supported.
<code>...</code>	optional arguments

## Value

Returns a `flowPlate` where the `wellAnnotation` now contains additional columns corresponding to total number of events in a well (`Total.Count`), the percentage of cells above background (`Percent.Positive`), and the number of positive cells (`Positive.Count`).

## Author(s)

Errol Strain

## See Also

See Also [setControlGates](#)

**Examples**

```

library(plateCore)
data(plateCore)

## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmcPlate, wellAnnotation, plateName="P1")

# Create a set of negative control gates and then apply them
fp <- setControlGates(fp, gateType="Negative.Control")
fp <- applyControlGates(fp, gateType="Negative.Control")

# Percent Positive and Counts columns are now in the wellAnnotation
head(wellAnnotation(fp))

```

---

 compensate

*Compensate a flowPlate to correct for the effects of spillover.*


---

**Description**

Flow samples are often stained with multiple types of fluorophores. Unfortunately, the emission spectra for these different fluorophores often overlap, and the signals must be corrected before proceeding with the analysis. Compensate adjusts for spillover using the method implemented in the package flowCore. Unlike flowCore, compensate only adjusts for the dyes/fluorophores listed in wellAnnotation.

**Usage**

```

## S4 method for signature 'flowPlate,ANY':
compensate(x, spillover)

```

**Arguments**

x	A flowPlate
spillover	The compensation matrix where the row and column names match the fluorescence channels of the flowPlate.

**Value**

Returns a compensated flowPlate.

**Author(s)**

Errol Strain

**See Also**

See Also [compensation-class](#)

## Examples

```
library(plateCore)
data(plateCore)

# Create the compensation matrix
comp.mat <- spillover(x=compensationSet, unstained=sampleNames(compensationSet)[5],
patt=".*H", fsc="FSC-H", ssc="SSC-H", method="median")

## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmcPlate, wellAnnotation, plateName="P1")

# apply the compensation matrix
fp <- compensate(fp, comp.mat)
```

---

compensationSet	<i>Sample Compensation Data Set</i>
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## Description

Sample Compensation Data Set

## Usage

```
data(plateCore)
```

## Format

The format is an object of class `flowSet` composed of 5 `flowFrames`. The `flowSet` consists of 4 stained and one unstained `flowFrames`. Peripheral Blood Mononucleocytes (PBMCs) were stained with FITC (Fluorescein isothiocyanate), PE (phycoerythrin), PerCp (Peridinin-chlorophyll), and APC (Allophycocyanin).

## Author(s)

Errol Strain

## Source

Sample data set from BD FACS CAP analysis.

## See Also

See Also [compensation-class](#)

**Examples**

```

library(plateCore)
data(plateCore)

# Create the compensation matrix
comp.mat <- spillover(x=compensationSet, unstained=sampleNames(compensationSet)[5],
patt=".*H", fsc="FSC-H", ssc="SSC-H", method="median")

```

---

densityplot

*One-dimensional density plots for flowPlates*


---

**Description**

This function is a modified version of densityplot from the flowViz package that allows for multiple flowFrames per panel. flowViz densityplot plots the density curves in a one per panel style, while the flowPlate densityplot can overlay densities.

**Usage**

```

## S4 method for signature 'formula,flowPlate':
densityplot(x, data, xlab,
  prepanel=prepanel.densityplot.flowPlate,
  panel = panel.densityplot.flowPlate,
  as.table=TRUE,
  filterResult=NULL,
  ...)

```

**Arguments**

x	A formula describing the layout of the plots.
data	A flowPlate.
xlab	Label for the x-axis
prepanel	Lattice-flowViz prepanel function.
panel	Lattice-flowViz panel function.
as.table	Defaults to table layout.
filterResult	filterResult can either take the character string "Negative.Control" and have the negative control wells added to the panels, or if filterResult is a flowFrame then the density curve for the flowFrame will be added to each panel.
...	optional arguments Other arguments are identical to densityPlot from flowViz.

**See Also**

[flowViz::densityplot](#)

## Examples

```
# Load the plateCore package and data
library(plateCore)
data(plateCore)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmPlate, wellAnnotation, plateName="P1")

# Overlay the first 3 flowFrames. If the groups argument was
# omitted, then the flowFrames would be combined into a single
# density curve.
densityplot(~ `FSC-H`, fp[1:3], groups=name, auto.key=TRUE)
```

---

fixAutoFl

*Correct for the effects of cell size (FSC) on autofluorescence*

---

## Description

The `fixAutoFl` function uses the method of Hahne et al. 2006 (Genome Biology) to fit a robust, log-log linear regression to the fluorescence channel of interest versus forward scatter (FSC). The current implementation scales the corrected data so the median fluorescence intensity (MFI) is the same before and after `fixAutoFl`.

## Usage

```
fixAutoFl(fp, fsc="FSC.A", chanCols, unstain, ...)
```

## Arguments

<code>fp</code>	A <code>flowPlate</code>
<code>chanCols</code>	Selected channels to correct for autofluorescence.
<code>unstain</code>	Name(s) of the unstained samples. The function will try to find samples with <code>Sample.Type="Unstained"</code> if no names are given. If there are multiple unstained samples the function will average the slopes.
<code>fsc</code>	Name of the FSC parameter.
<code>...</code>	optional arguments

## Value

Returns a `flowPlate` with autofluorescence due to cell size (FSC) corrected.

## Author(s)

Errol Strain

**Examples**

```

library(plateCore)
data(plateCore)

## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

## Create a flowPlate object from the platePBMC and the wellAnnotation
fp <- flowPlate(pbmcPlate,wellAnnotation,plateName="P1")

## Correct for autofluorescence in FL1.H-FL4.H
fp <- fixAutoFl(fp,fsc="FSC-H",chanCols=c("FL1-H","FL2-H","FL3-H","FL4-H"))

```

---

flowPlate-class      *The flowPlate class.*

---

**Description**

flowPlates are the basic data containers for the plateCore package. A flowPlate is essentially a [flowSet-class](#), plus a `data.frame` describing the layout of the plate and contents of individual wells.

**Slots**

**plateName:** A character string containing the name of the plate.

**plateSet:** A [flowSet-class](#) containing FCS event data. Prior to creating a flowPlate, the FCS files are first read into a [flowSet-class](#) using `read.flowSet`.

**wellAnnotation:** A `data.frame` describing the layout of the plate. Each row describes one channel for a well.

**Methods**

`[, [` Subsetting. `x[i]` where `i` is either a scalar or character corresponding to a sample name, returns a flowPlate object, and `x[[i]]` a [flowFrame](#) object.

*Usage:*

```

flowSet[i]
flowSet[[i]]

```

**Author(s)**

Errol Strain, Florian Hahne, Perry Haaland

**Examples**

```

library(plateCore)
data(plateCore)

##Look at the wellAnnotation
wellAnnotation[1:4,]

```

```
## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700),"SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

## Create a flowPlate object from the platePBMC and the wellAnnotation
fp <- flowPlate(pbmcPlate,wellAnnotation,plateName="P1")

## Subset the flowPlate, creating another flowPlate
fpSmall <- fp["A01"]

## Extract a flowFrame from a flowPlate
ff <-fp[["A01"]]
```

---

flowPlate	<i>Create a flowPlate</i>
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### Description

Constructor for a `flowPlate` object. `sampleNames` for the `flowSet` should match the `Well.Id` column of `wellAnnotation`. `Well.Ids` must be unique to `sampleNames`, which is usually ensured by using the 3 character designations for wells (e.g. "A01","A02",..., "H12").

### Usage

```
flowPlate(data, wellAnnotation, plateName, ...)
```

### Arguments

<code>data</code>	flowSet object to be made into a flowPlate
<code>wellAnnotation</code>	data.frame describing the layout and contents of the flowPlate
<code>plateName</code>	Name of the flowPlate, should be unique within the set flowPlates under consideration
<code>...</code>	optional arguments

### Value

Returns a `flowPlate` object.

### Author(s)

Errol Strain

### Examples

```
library(plateCore)
data(plateCore)

## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700),"SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)
```

```
## Create a flowPlate object from the platePBMC and the wellAnnotation
fp <- flowPlate(pbmcPlate,wellAnnotation,plateName="P1")

## Subset the flowPlate, creating another flowPlate
fpSmall <- fp["A01"]

## Extract a flowFrame from a flowPlate
ff <-fp[["A01"]]

## Retrieve sample names from flowPlate
sampNames <- sampleNames(fp)

## Retrieve the annotatedDataFrame describing the flowPlate
adf <- phenoData(fp)
```

fpbind

*Merge multiple flowPlates into a single virtual flowPlate***Description**

A function to combine multiple flowPlates into a single flowPlate object. The plateName identifiers for the flowPlates must be unique within the set to be bound, otherwise the bind will fail.

**Usage**

```
fpbind(p1, p2, ...)
```

**Arguments**

p1	First flowPlate
p2	Second flowPlate
...	Additional flowPlates

**Value**

Returns a flowPlate

**Author(s)**

Errol Strain

**Examples**

```
library(plateCore)
data(plateCore)

# Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700),"SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

# Create a flowPlate object from the platePBMC and the wellAnnotation
fp1 <- flowPlate(pbmcPlate,wellAnnotation,plateName="P1")
```



```
fp2 <- flowPlate(pbmcPlate,wellAnnotation,plateName="P2")

# Combine the plates.
virtPlate <- fpbind(fp1,fp2)
```

---

`getGroups`*Retrieve Negative control groups from a flowPlate*

---

### Description

Retrieve a list of negative control-based groups from a flowPlate, based on the information in wellAnnotation.

### Usage

```
getGroups(data,type="Negative.Control",chan, ...)
```

### Arguments

<code>data</code>	A flowPlate dataset.
<code>type</code>	Currently only Negative.Control groups are supported.
<code>chan</code>	Fluorescence channel of interest ("FL1-H", "PE-H", etc.)
<code>...</code>	optional arguments

### Value

Returns a list of groups, where each group contains a single negative control well and the associated test well for a particular channel.

### Author(s)

Errol Strain

### Examples

```
library(plateCore)
data(plateCore)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmcPlate,wellAnnotation,plateName="P1")

# Create a set of negative control gates and then apply them
negCon <- getGroups(fp,chan="FL1-H")

negCon[1:2]
```

---

gutterPlot	<i>gutterPlot</i>
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**Description**

A Quality Control plot to check the number of events in each channel that are at either their minimum or maximum value. A large number of these events may indicate a problem with the sample.

**Usage**

```
gutterPlot(fp, chans = c("FSC-H", "SSC-H", "FL1-H", "FL2-H", "FL3-H", "FL4-H"),
```

**Arguments**

fp	A flowPlate.
chans	Channels of interest to show on the gutterPlot.
...	optional arguments

**Value**

Creates a plot where the x-axis is the different wells in a flowPlate, and the y-axis is the fraction of events at the boundary.

**Author(s)**

Jon Gosink and Errol Strain

**Examples**

```
library(plateCore)
data(plateCore)

## Create a flowPlate
fp <- flowPlate(pbmPlate, wellAnnotation, "p1001")

gutterPlot(fp, chans=c("FSC-H", "SSC-H", "FL1-H", "FL2-H", "FL3-H", "FL4-H"))
```

---

mfiPlot	<i>mfiPlot</i>
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---

**Description**

A Quality Control plot that shows the MFI Ratio versus the percentage of positive cells in a flow-Plate. The robust logistic regression is performed using gmlrob from the robustbase package.

**Usage**

```
mfiPlot(fp, thresh=2, Sample.Type="Test", Events="Percentage", ...)
```

**Arguments**

<code>fp</code>	A <code>flowPlate</code> .
<code>thresh</code>	Points more than "thresh" number of standard deviations away from the best fit line will be colored red.
<code>Sample.Type</code>	Type of sample to show on plot. Defaults to "Test"
<code>Events</code>	The robust logistic regression can be performed using either the percentage of events above the negative control gate ("Percentage") or the actual number of events above the gate ("Actual").
<code>...</code>	optional arguments to plot and points.

**Value**

Creates a plot where the x-axis is MFI Ratio and the y-axis is the percentage of cells above the negative control gate.

**Author(s)**

Errol Strain

**Examples**

```
library(plateCore)
data(plateCore)

## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmcPlate, wellAnnotation, plateName="P1")

# Create a set of negative control gates and then apply them
fp <- setControlGates(fp, gateType="Negative.Control")
fp <- applyControlGates(fp, gateType="Negative.Control")

# Compute summary statistics
fp <- summaryStats(fp)

## Create an MFI plot
mfiPlot(fp, thresh=2.5, xlab="MFI Ratio (Test MFI / Isotype MFI)", xlim=c(0.1,250),
        ylab="Percentage of cells above the isotype gate", pch=23)
```

**Description**

This operator is used to construct a `transformFilter` that first applies a `transformList` to the data before applying the filter operation to a `flowPlate`.

**Author(s)**

Errol Strain

**Examples**

```

library(plateCore)
data(plateCore)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmcPlate, wellAnnotation, plateName="P1")

## Create a rectangle filter
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))

xyplot(`FL1-H` ~ `FSC-H` | as.factor(name),
transform("FL1-H"=log10) %on% fp, smooth=FALSE, filter=rectGate, displayFilter=FALSE)

```

---

panel.densityplot.flowPlate

*Lattice-flowViz style panel function for flowPlate densityplot.*


---

**Description**

This function should not be called directly, use [densityplot](#).

**Usage**

```

panel.densityplot.flowPlate(x,
frames, channel, wellAnnotation,
groups=NULL,
subscripts,
col = superpose.symbol$col,
col.points = col,
col.line = col,
filterResult=NULL,
...)

```

**Arguments**

x	character
frames	flowFrames
channel	channel of interest
wellAnnotation	wellAnnotation data.frame
groups	density plot groups parameter
subscripts	densityplot subscripts parameter
col	densityplot col parameter
col.points	densityplot col.points parameter
col.line	densityplot col.line parameter
filterResult	densityplot filterResult parameter
...	optional arguments

**Author(s)**

Errol Strain

**See Also**See Also [densityplot](#)**Examples**

```
# Load the plateCore package and data
library(plateCore)
data(plateCore)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmPlate, wellAnnotation, plateName="P1")

# Overlay the first 3 flowFrames. If the groups argument was
# omitted, then the flowFrames would be combined into a single
# density curve.
densityplot(~ `FSC-H`, fp[1:3], groups=name, auto.key=TRUE)
```

---

panel.xyplot.flowPlate

*Lattice-flowViz style panel function for flowPlate xyplot.*

---

**Description**

This function should not be called directly, use [xyplot](#).

**Usage**

```
panel.xyplot.flowPlate(x,
  frames,
  channel.x, channel.y,
  channel.x.name, channel.y.name,
  filter = NULL,
  filterResults = NULL,
  displayFilter = TRUE,
  pch, smooth,
  wellAnnotation = NULL,
  col = superpose.symbol$col,
  ...)
```

**Arguments**

x	character
frames	flowFrames
channel.x	xyplot channel.x parameter
channel.y	xyplot channel.y parameter

```

channel.x.name      xyplot channel.x.name parameter
channel.y.name      xyplot channel.y.name parameter
filter              xyplot filter parameter
filterResults       xyplot filterResults parameter
displayFilter       xyplot displayFilter parameter
pch                 xyplot pch parameter
smooth              xyplot smooth parameter
wellAnnotation      wellAnnotation data.frame
col                 xyplot col parameter
...                 optional arguments

```

**Author(s)**

Errol Strain

**See Also**

See Also [xyplot](#)

**Examples**

```

library(plateCore)
data(plateCore)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmcPlate,wellAnnotation,plateName="P1")

## Create a rectangle filter
rectGate <- rectangleGate("FSC-H"=c(300,700),"SSC-H"=c(50,400))

xyplot(`SSC-H` ~ `FSC-H` | as.factor(name),
fp[1], smooth=FALSE, filter=rectGate, displayFilter=FALSE)

```

---

pbmcPlate

*pbmcPlate Data Set*

---

**Description**

One 96-well plate from a BD FACS CAP analysis of Peripheral Blood Mononucleocyte (PMBC) cells.

**Usage**

```
data(plateCore)
```

## Format

The format is an object of class `flowSet` composed of 96 `flowFrame`s. Each `flowFrame` corresponds to one well from the plate.

## Details

BD FACS CAP ([http://www.bd.com/technologies/discovery\\_platform/BD\\_FACS\\_CAP.asp](http://www.bd.com/technologies/discovery_platform/BD_FACS_CAP.asp)) is a platform for screening a large number of antibodies (200+) on human samples. Antibodies are arrayed 3-per well on a 96-well plate, along with the appropriate controls. In this experiment, an early version of FACS CAP was used to screen PBMCs from 2 donors for 189 different human cell surface markers. The complete data set is available from the FICCS site shown below. The `pbmcPlate` include with `plateCore` is a lymphocyte enriched subset of one of the replicate plates for donor 1.

## Source

Complete dataset available at [http://www.ficcs.org/software.html#Data\\_Files](http://www.ficcs.org/software.html#Data_Files), the Flow Informatics and Computational Cytometry Society website (FICCS)

## References

Add reference for `plateCore` paper (when/if published).

---

`plateCore-package` *plateCore: A Bioconductor package for high throughput analysis of flow cytometry data*

---

## Description

`plateCore` is a Bioconductor package created to make processing and analysis of large, complex flow datasets in R easier. High throughput flow studies are often run in a 96 or 384-well plate format, with a number of different samples, controls, and antibodies-dye conjugates present on the plate. Analyzing the output from the cytometer requires keeping track of the contents of each well, matching sample wells with control wells, gating each well/channel separately, making the appropriate plots, and summarizing the results. `plateCore` extends the `flowCore` and `flowViz` packages to work on `flowPlate` objects that represent these large flow datasets. For those familiar with `flowCore` and `flowViz`, the gating (filtering), transformation, and other data manipulations for `flowPlates` are very similar to `flowSets`.

## Details

Package: `plateCore`  
Type: Package  
Version: 1.2.1  
Date: 2009-06-29

**Author(s)**

Errol Strain, Florian Hahne, and Perry Haaland Maintainer: Errol Strain <estrain@gmail.com>

**References**

Insert flowCore and flowViz publications.

**Examples**

```
library(plateCore)
data(plateCore)

## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

## Create a flowPlate object from the platePBMC and the wellAnnotation
fp <- flowPlate(pbmcPlate, wellAnnotation, plateName="P1")
```

---

plateSet

*Get the flowSet from a flowPlate object*

---

**Description**

A function to retrieve the flowSet from a flowPlate.

**Usage**

```
plateSet(fp, ...)
```

**Arguments**

fp	A flowPlate
...	optional arguments

**Value**

Returns a flowSet

**Author(s)**

Errol Strain



**Examples**

```

library(plateCore)
data(plateCore)

## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

## Create a flowPlate object from the platePBMC and the wellAnnotation
fp <- flowPlate(pbmcPlate, wellAnnotation, plateName="P1")

## Retrieves the flowSet
fs <- plateSet(fp)

```

---

plotPlate

*plotPlate*


---

**Description**

Make a row vs. column plot of a plate, where the wells are colored according to some value of choice (number of events, median signal intensity, percent positive, etc.).)

**Usage**

```
plotPlate(fp, x = NA, method = "median", main, col, values, width = 1, na.action)
```

**Arguments**

fp	A flowPlate.
x	A character indicating the variable of interest. Valid choices are "events", any single channel name (e.g. FSC-H, SSC-H, FL1-H, etc.), or vector of channel names if the method is mahalanobis.
method	Valid choices are mean, median, sd, mad, mahalanobis, or one of the numeric columns in the wellAnnotation data.frame (e.g. Percent.Positive, Positive.Count, MFI, MFI.Ratio)
main	Main text for the plot
col	Character vector of two colors.
values	Optional list of values, with names corresponding to sampleNames, that will be used for plotting.
width	Width of the well.
na.action	Handling of NA values, either "zero" or "omit".
...	optional arguments

**Value**

Plots the plate to the standard output.

**Author(s)**

Jon Gosink and Errol Strain

## References

The original version of this plot came from the `prada` package.

## Examples

```
library(plateCore)
data(plateCore)

## Create a flowPlate
fp <- flowPlate(pbmcPlate, wellAnnotation, "p1001")

plotPlate(transform("FL1-H"=log10) %on% fp, x="FL1-H", method="mean", col=c("yellow", "darkb
```

---

```
prepanel.densityplot.flowPlate
```

*Lattice-flowViz style panel function for flowPlate densityplot.*

---

## Description

This function should not be called directly, use [densityplot](#).

## Usage

```
prepanel.densityplot.flowPlate(x, frames, channel, ...)
```

## Arguments

<code>x</code>	Character
<code>frames</code>	flowFrames
<code>channel</code>	Character string for channel name.
<code>...</code>	optional arguments

## Author(s)

Errol Strain

## See Also

See Also [densityplot](#)

## Examples

```
# Load the plateCore package and data
library(plateCore)
data(plateCore)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmcPlate, wellAnnotation, plateName="P1")

# Overlay the first 3 flowFrames. If the groups argument was
# omitted, then the flowFrames would be combined into a single
# density curve.
densityplot(~ `FSC-H`, fp[1:3], groups=name, auto.key=TRUE)
```

---

```
prepanel.xyplot.flowPlate
```

*Lattice-flowViz style panel function for flowPlate xyplot.*

---

## Description

This function should not be called directly, use [xyplot](#).

## Usage

```
prepanel.xyplot.flowPlate(x, frames, channel.x, channel.y, ...)
```

## Arguments

x	Character
frames	flowFrames
channel.x	Character string for channel name.
channel.y	Character string for channel name.
...	optional arguments

## Author(s)

Errol Strain

## See Also

See Also [xyplot](#)

## Examples

```
library(plateCore)
data(plateCore)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmPlate, wellAnnotation, plateName="P1")

## Create a rectangle filter
rectGate <- rectangleGate("FSC-H"=c(300, 700), "SSC-H"=c(50, 400))

xyplot(`SSC-H` ~ `FSC-H` | as.factor(name),
fp[1], smooth=FALSE, filter=rectGate, displayFilter=FALSE)
```

---

setControlGates      *Create control gates for a flowPlate*

---

### Description

A function to estimate the threshold between positive and negative cells. This threshold corresponds to a one-dimensional gate, and cells above the gate are considered positive. The default value of numMads=5 generally works well on the linear scale, but will need to be adjusted for transformed data. If each well contains a large number of events for the cell type of interest (>1000), then using the 99.5th quantile usually gives similar values.

### Usage

```
setControlGates(data, gateType, threshType="MAD", numMads=5, isoquantile=.995, .
```

### Arguments

data	A flowPlate
gateType	The type of gate to be set. Currently only "Negative.Control" gates are supported.
threshType	Values can be either "MAD", for median absolute deviation based gating, or "isoQuant" for quantile based gating.
numMads	Number of median absolute deviations above the median to set the initial gate.
isoquantile	Quantile setting for "isoQuant" threshType.
...	optional arguments.

### Value

Returns a flowPlate

### Author(s)

Errol Strain

### Examples

```
library(plateCore)
data(plateCore)

## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmcPlate, wellAnnotation, plateName="P1")

# Create a set of negative control gates and then apply them
fp <- setControlGates(fp, gateType="Negative.Control")

# There should now be a Negative.Control.Gate column in wellAnnotation
head(wellAnnotation(fp))
```

---

Subset	<i>Subset</i>
--------	---------------

---

## Description

Select a subset of events in a flowPlate. If a flowPlate and filter are supplied, then this function calls the Subset function from flowCore. Additionally, the plateCore version of Subset also makes it easy to filter individual flowFrames and keep the flowPlate structure.

## Usage

```
## S4 method for signature 'flowPlate,ANY':  
Subset(x, subset, select=NULL,...)
```

## Arguments

x	A flowPlate
subset	A filter object
select	An optional vector of either sample names or Well.IDs.
...	optional arguments

## Value

Returns a flowPlate

## Author(s)

Errol Strain

## Examples

```
library(plateCore)  
data(plateCore)  
  
# Create a flowPlate from the sample data in plateCore  
fp <- flowPlate(pbmPlate, wellAnnotation, plateName="P1")  
  
## Create a rectangle filter  
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))  
  
## Apply the filter only to sample A01. The other flowFrames  
## are not filtered.  
fp <- Subset(fp, rectGate, 'A01')
```

summaryStats

*Compute summary statistics on a flowPlate***Description**

This function computes the median fluorescence intensity (MFI) and the MFI ratio (ratio of test well MFI to negative control MFI) for each well/channel in a `flowPlate`. The predicted percent positive (`Predict.PP`) and gate score (`Gate.Score`) come from a robust logistic regression of the MFI ratio to either the percentage of positive cells or the actual count of positive cells. `Predict.PP` is the estimated percent positive based on the MFI ratio and `Gate.Score` is the number of standardized residuals the sample data point is away from the best fit line. The `glmrob` function from the `robustbase` package is used for the regression. Results from `summaryStats` are stored in the `wellAnnotation` `data.frame`.

**Usage**

```
summaryStats(data, Events="Percentage", ...)
```

**Arguments**

<code>data</code>	A <code>flowPlate</code>
<code>Events</code>	The robust logistic regression can be performed using either the percentage of events above the negative control gate ("Percentage") or the actual number of events above the gate ("Actual").
<code>...</code>	optional arguments

**Value**

Returns a `flowPlate`

**Author(s)**

Errol Strain

**Examples**

```
library(plateCore)
data(plateCore)

## Get the lymphocytes
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))
pbmcPlate <- Subset(pbmcPlate, rectGate)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmcPlate, wellAnnotation, plateName="P1")

# Create a set of negative control gates and then apply them
fp <- setControlGates(fp, gateType="Negative.Control")
fp <- applyControlGates(fp, gateType="Negative.Control")

# Compute summary statistics
fp <- summaryStats(fp)
```

```
# There should now be MFI and MFI.ratio columns in the wellAnnotation
head(wellAnnotation(fp))
```

---

wellAnnotation	<i>Retrieve a data.frame describing the content of a flowPlate</i>
----------------	--

---

### Description

wellAnnotation returns the tall `data.frame` describing the layout of a `flowPlate`, where each row corresponds to one well-channel.

### Usage

```
wellAnnotation(fp, ...)
```

### Arguments

fp	A flowPlate dataset.
...	optional arguments

### Value

Returns a `data.frame`.

### Author(s)

Errol Strain

### Examples

```
library(plateCore)
data(plateCore)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmPlate, wellAnnotation, plateName="P1")

# Look at the top of wellAnnotation
head(wellAnnotation(fp))
```

xyplot

*Scatter plots (dotplots) for flowPlates.***Description**

A function to create dotplots, and smoothed scatter plots, from flowPlates. This function is a slightly modified version of xyplot from flowViz. The flowPlate xyplot allows users to overlay plots of test samples versus controls, and makes creating informative flowStrips easier. Refer to the documentation for xyplot from flowViz and lattice for more detailed information.

**Usage**

```
## S4 method for signature 'formula,flowPlate':
xyplot(x, data, xlab, ylab,
as.table = TRUE,
prepanel = prepanel.xyplot.flowPlate,
panel = panel.xyplot.flowPlate,
pch = ".", smooth = TRUE,
filter = NULL,
filterResults = NULL,
displayFilter = TRUE,
flowStrip=NULL,
flowStripCex=1,
strip=function(...,style=1) strip.default(...,style=1),
...)
```

**Arguments**

x	A formula describing the layout of the plots. Plots for flowPlates usually condition on either <code>as.factor(name)</code> or <code>as.factor(Well.Id)</code> since only one flowFrame can be shown on each panel (with the exception of Negative.Control overlays).
data	A flowPlate.
xlab	Label for x-axis.
ylab	Label for y-axis.
as.table	Defaults to table layout.
prepanel	Lattice-flowViz prepanel function.
panel	Lattice-flowViz panel function.
pch	Plotting character.
smooth	Plot a smoothed scatterplot by default.
filter	A flowCore filter to apply to each flowFrame.
filterResults	If <code>filterResults="Negative.Control"</code> , the negative control wells corresponding to a test well are overlaid in the test well plots.
displayFilter	Defaults to displaying filter on the plot.



flowStrip	Character vector indicating additional information to be printed on the strip. Values can include any combination of "Well.Id", "MFI", "MFI.Ratio", and "Percent.Positive".
flowStripCex	Font size for the flowStrip.
strip	Lattice strip function.
...	Optional arguments

**Author(s)**

Errol Strain

**See Also**[flowViz::xyplot](#)**Examples**

```
library(plateCore)
data(plateCore)

# Create a flowPlate from the sample data in plateCore
fp <- flowPlate(pbmPlate, wellAnnotation, plateName="P1")

## Create a rectangle filter
rectGate <- rectangleGate("FSC-H"=c(300,700), "SSC-H"=c(50,400))

xyplot(`SSC-H` ~ `FSC-H` | as.factor(name),
fp[1], smooth=FALSE, filter=rectGate, displayFilter=FALSE)
```

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