

# Package ‘flowWorkspace’

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**Type** Package

**Title** Infrastructure for representing and interacting with gated and ungated cytometry data sets.

**Version** 4.4.0

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**Description** This package is designed to facilitate comparison of automated gating methods against manual gating done in flowJo. This package allows you to import basic flowJo workspaces into BioConductor and replicate the gating from flowJo using the flowCore functionality. Gating hierarchies, groups of samples, compensation, and transformation are performed so that the output matches the flowJo analysis.

**License** file LICENSE

**License\_restricts\_use** yes

**LazyLoad** yes

**Imports** Biobase, BiocGenerics, cytolib (>= 2.3.9), lattice, latticeExtra, XML, ggplot2, graph, graphics, grDevices, methods, stats, stats4, utils, RBGL, tools, Rgraphviz, data.table, dplyr, Rcpp, scales, matrixStats, RcppParallel, RProtoBufLib, digest, aws.s3, aws.signature, flowCore(>= 2.1.1), ncdfFlow(>= 2.25.4), DelayedArray, S4Vectors

**Collate** 'cytoframe.R' 'cytoset.R' 'AllClasses.R' 'getStats.R'  
'GatingHierarchy\_Methods.R' 'GatingSet\_Methods.R'  
'GatingSetList\_Methods.R' 'RcppExports.R'  
'filterObject\_Methods.R' 'add\_Methods.R' 'copyNode.R'  
'cytoctx.R' 'deprecated.R' 'flow\_trans.R' 'getDescendants.R'  
'getSingleCellExpression.R' 'identifier.R' 'load\_fcs.R'  
'load\_gs.R' 'merge\_GatingSet.R' 'merge\_gslist.R' 'moveNode.R'  
'parse\_transformer.R' 'setGate\_Methods.R' 'updateIndices.R'  
'utils.R' 'zzz.R'

**Suggests** testthat, flowWorkspaceData (>= 2.23.2), knitr, ggcryo, parallel, CytoML, openCyto

**LinkingTo** Rcpp, BH(>= 1.62.0-1), RProtoBufLib(>= 1.99.4), cytolib (>= 2.3.7), Rhdf5lib, RcppArmadillo, RcppParallel(>= 4.4.2-1)

**VignetteBuilder** knitr

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---

flowWorkspace-package *Import and replicate flowJo workspaces and gating schemes using flowCore.*

---

## Description

Import flowJo workspaces into R. Generate the flowJo gating hierarchy and gates using flowCore functionality. Transform and compensate data in accordance with flowJo settings. Plot gates, gating hierarchies, population statistics, and compare flowJo vs flowCore population summaries.

## Details

Package:	flowWorkspace
Type:	Package
Version:	0.5.40
Date:	2011-03-04
License:	Artistic 2.0
LazyLoad:	yes
Depends:	R (>= 2.16.0), Rcpp (>= 0.9.9)

## Author(s)

Greg Finak, Mike Jiang

## References

<http://www.rglab.org/>

---

asinhtGml2\_trans      *Inverse hyperbolic sine transformation.*

---

## Description

Used to construct inverse hyperbolic sine transform object.

## Usage

```
asinhtGml2_trans(..., n = 6, equal.space = FALSE)
```

**Arguments**

...	parameters passed to asinh_Gml2
n	desired number of breaks (the actual number will be different depending on the data range)
equal.space	whether breaks at equal-spaced intervals

**Value**

asinhGml2 transformation object

**Examples**

```
trans.obj <- asinhGml2_trans(equal.space = TRUE)
data <- 1:1e3
brks.func <- trans.obj[["breaks"]]
brks <- brks.func(data)
brks # fasinh space displayed at raw data scale

#transform it to verify it is equal-spaced at transformed scale
trans.func <- trans.obj[["transform"]]
brks.trans <- trans.func(brks)
brks.trans
```

---

asinh\_Gml2

*inverse hyperbolic sine transform function generator (GatingML 2.0 version)*

---

**Description**

hyperbolic sine/inverse hyperbolic sine transform function constructor. It is simply a special form of flowjo\_fasinh with length set to 1 and different default values for parameters t, m, a.

**Usage**

```
asinh_Gml2(T = 262144, M = 4.5, A = 0, inverse = FALSE)
```

**Arguments**

T	numeric the maximum value of input data
M	numeric the full width of the transformed display in asymptotic decades
A	numeric Additional negative range to be included in the display in asymptotic decades
inverse	whether to return the inverse function

**Value**

fasinh/fsinh transform function

## Examples

```
trans <- asinh_Gml2()
data.raw <- c(1,1e2,1e3)
data.trans <- trans(data.raw)
data.trans

inverse.trans <- asinh_Gml2(inverse = TRUE)
inverse.trans(data.trans)
```

---

booleanFilter-class	<i>A class describing logical operation (&amp; or !) of the reference populations</i>
---------------------	---------------------------------------------------------------------------------------

---

## Description

booleanFilter class inherits class [expressionFilter](#) and exists for the purpose of methods dispatching.

## Usage

```
booleanFilter(expr, ..., filterId = "defaultBooleanFilter")

char2booleanFilter(expr, ..., filterId = "defaultBooleanFilter")
```

## Arguments

expr	expression
...	further arguments to the expression
filterId	character identifier

## See Also

[add GatingHierarchy](#)

## Examples

```
# "4+/TNFa+" and "4+/IL2+" are two existing gates
#note: no spaces between node names and & , ! operators
booleanFilter(`4+/TNFa+&!4+/IL2+`)

#programmatically
n1 <- "4+/TNFa+"
n2 <- "4+/IL2+"
exprs <- paste0(n1, "&!", n2)
call <- substitute(booleanFilter(v), list(v = as.symbol(exprs)))
eval(call)
```

---

cf_append_cols	<i>Append data columns to a flowFrame</i>
----------------	-------------------------------------------

---

### Description

Append data columns to a flowFrame

### Usage

```
cf_append_cols(cf, cols, ctx = .cytoctx_global)
```

### Arguments

cf	A cytoframe.
cols	A numeric matrix containing the new data columns to be added. Must has column names to be used as new channel names.

### Details

It is used to add extra data columns to the existing flowFrame. It handles keywords and parameters properly to ensure the new flowFrame can be written as a valid FCS through the function `write.FCS`

.

### Examples

```
library(flowCore)
data(GvHD)
tmp <- GvHD[[1]]
cf <- flowFrame_to_cytoframe(tmp)
kf <- kmeansFilter("FSC-H"=c("Pop1", "Pop2", "Pop3"), filterId="myKmFilter")
fres <- filter(cf, kf)
cols <- as.integer(fres@subSet)
cols <- matrix(cols, dimnames = list(NULL, "km"))
cf <- cf_append_cols(cf, cols)
```

---

cf_backend_type	<i>return the cytoframe backend storage format</i>
-----------------	----------------------------------------------------

---

### Description

return the cytoframe backend storage format

**Usage**

```
cf_backend_type(cf)
```

**Arguments**

cf	cytoframe
----	-----------

**Value**

one of "mem", "h5", "tile"

---

*cf\_get\_uri*

*Return the file path of the underlying h5 file*

---

**Description**

Return the file path of the underlying h5 file

**Usage**

```
cf_get_uri(cf)  
cf_get_h5_file_path(cf)
```

**Arguments**

cf	cytoframe object
----	------------------

**Details**

For the in-memory version of cytoframe, it returns an empty string. This can be used to check whether it is on-disk format.

**See Also**

Other cytoframe/cytoset IO functions: [cf\\_write\\_disk\(\)](#), [cf\\_write\\_h5\(\)](#), [cf\\_write\\_tile\(\)](#), [cs\\_get\\_uri\(\)](#), [load\\_cytoframe\\_from\\_fcs\(\)](#), [load\\_cytoframe\(\)](#), [load\\_cytoset\\_from\\_fcs\(\)](#)

---

cf_is_subsetted	<i>check whether a cytoframe/cytoset is a subsetted(by column or by row) view</i>
-----------------	-----------------------------------------------------------------------------------

---

**Description**

check whether a cytoframe/cytoset is a subsetted(by column or by row) view

**Usage**

```
cf_is_subsetted(x)
cs_is_subsetted(x)
```

**Arguments**

x                    a cytoset or cytoframe

---

cf_write_disk	<i>Save the cytoframe to disk</i>
---------------	-----------------------------------

---

**Description**

Save the cytoframe to disk

**Usage**

```
cf_write_disk(
  cf,
  filename,
  backend = get_default_backend(),
  ctx = .cytoctx_global
)
```

**Arguments**

cf	cytoframe object
filename	the full path of the output file
backend	either "h5" or "tile"
ctx	cytoctx object, see [cytoctx] for details

**See Also**

Other cytoframe/cytoset IO functions: [cf\\_get\\_uri\(\)](#), [cf\\_write\\_h5\(\)](#), [cf\\_write\\_tile\(\)](#), [cs\\_get\\_uri\(\)](#), [load\\_cytoframe\\_from\\_fcs\(\)](#), [load\\_cytoframe\(\)](#), [load\\_cytoset\\_from\\_fcs\(\)](#)

---

cf_write_h5	<i>Save the cytoframe as h5 format</i>
-------------	----------------------------------------

---

## Description

Save the cytoframe as h5 format

## Usage

```
cf_write_h5(cf, filename, ctx = .cytoctx_global)
```

## Arguments

cf	cytoframe object
filename	the full path of the output h5 file
ctx	cytoctx object, see [cytoctx] for details

## See Also

Other cytoframe/cytoset IO functions: [cf\\_get\\_uri\(\)](#), [cf\\_write\\_disk\(\)](#), [cf\\_write\\_tile\(\)](#), [cs\\_get\\_uri\(\)](#), [load\\_cytoframe\\_from\\_fcs\(\)](#), [load\\_cytoframe\(\)](#), [load\\_cytoset\\_from\\_fcs\(\)](#)

---

cf_write_tile	<i>Save the cytoframe as h5 format</i>
---------------	----------------------------------------

---

## Description

Save the cytoframe as h5 format

## Usage

```
cf_write_tile(cf, filename, ctx = .cytoctx_global)
```

## Arguments

cf	cytoframe object
filename	the full path of the output file
ctx	cytoctx object, see [cytoctx] for details

## See Also

Other cytoframe/cytoset IO functions: [cf\\_get\\_uri\(\)](#), [cf\\_write\\_disk\(\)](#), [cf\\_write\\_h5\(\)](#), [cs\\_get\\_uri\(\)](#), [load\\_cytoframe\\_from\\_fcs\(\)](#), [load\\_cytoframe\(\)](#), [load\\_cytoset\\_from\\_fcs\(\)](#)

---

cleanup	<i>Remove on-disk files associatated with flowWorkspace data classes</i>
---------	--------------------------------------------------------------------------

---

### Description

These methods immediately delete the on-disk storage associated with [cytoframe](#), [cytoset](#), [GatingHierarchy](#), or [GatingSet](#) objects

### Usage

```
cf_cleanup(cf, ctx = .cytoctx_global)
```

### Arguments

cf	a cytoframe, cytoset, GatingHierarchy, or GatingSet object
ctx	cytoctx object, see [cytoctx] for details

### Details

this will override tempdir() in determining the top directory under which files can safely be removed.

---

cleanup_temp	<i>Remove temporary files associatated with flowWorkspace data classes</i>
--------------	----------------------------------------------------------------------------

---

### Description

These methods immediately delete the on-disk h5 storage associated with [cytoframe](#), [cytoset](#), [GatingHierarchy](#), or [GatingSet](#) objects, but only if it is under the directory pointed to by tempdir() or alternatively specified by the temp\_dir option. The temp\_dir option should be used with caution as it acts as a guard against accidental removal of non-temporary storage.

### Usage

```
cf_cleanup_temp(x, temp_dir = NULL)

cs_cleanup_temp(x, temp_dir = NULL)

gh_cleanup_temp(x, temp_dir = NULL)

gs_cleanup_temp(x, temp_dir = NULL)
```

### Arguments

x	a cytoframe, cytoset, GatingHierarchy, or GatingSet object
temp_dir	an optional argument designating another path as temporary storage. If specified this will override tempdir() in determining the top directory under which files can safely be removed.

## Details

Use of these functions will generally be unnecessary for most users, but they are provided for workflows that involve repeated creation of such data structures within the same R session to avoid overwhelming temporary storage.

---

compensate

*compensate the flow data associated with the GatingSet*

---

## Description

The compensation is saved in the GatingSet and can be retrieved by [gh\\_get\\_compensations](#).

## Usage

```
## S4 method for signature 'GatingSet,ANY'  
compensate(x, spillover)
```

## Arguments

x	GatingSet, GatingSetList, cytoframe, or cytoset
spillover	compensation object or spillover matrix or a list of compensation objects

## Value

a GatingSet, GatingSetList, cytoframe, or cytoset object with the underling flow data compensated.

## Examples

```
## Not run:  
  
cfile <- system.file("extdata","compdata","compmatrix", package="flowCore")  
comp.mat <- read.table(cfile, header=TRUE, skip=2, check.names = FALSE)  
## create a compensation object  
comp <- compensation(comp.mat, compensationId="comp1")  
#add it to GatingSet  
gs <- compensate(gs, comp)  
  
## End(Not run)
```

## convert

---

*Methods for conversion between flowCore and flowWorkspace data classes*

---

**Description**

These methods perform conversions between flowWorkspace classes ([cytoframe/cytoset](#)) and flowCore classes ([flowFrame/flowSet](#)) as well as between single-sample and aggregated classes (e.g. between cytoset and a list of cytoframes)

**Usage**

```
cytoframe_to_flowFrame(cf)

flowFrame_to_cytoframe(fr, ...)

cytoset_to_flowSet(cs)

flowSet_to_cytoset(
  fs,
  path = tempfile(),
  backend = get_default_backend(),
  tmp = tempfile()
)

cytoset_to_list(cs)
```

**Arguments**

cf	cytoframe object
fr	flowframe
...	arguments passed to 'load_cytoframe_from_fcs' call
cs	cytoset
fs	flowSet or ncdfFlowSet
path	the h5 path for cytoset
tmp	the temp folder when the temporary files are written to during conversion by default, it is system temp path. And it can be changed to the customized location when there is not enough space at system path.

**Details**

The first set of methods consist of a pair of methods to coerce a cytoframe to or from a flowFrame and another pair to coerce a cytoset to or from a flowSet.

The conversion between the two sets of data container classes mostly entails a conversion of the back-end representation of the data. cytoframe and cytoset objects contain flowFrame and

flowSet objects respectively, so coercion of a cytoframe to flowFrame entails moving the data from the 'C'-level data structure to the corresponding exprs, description, and parameters slots. Coercion of a flowFrame to a cytoframe entails creation of the 'C'-level data structure from the flowFrame slots. The names of each of the methods are pretty self-explanatory.

The second set of methods perform disaggregation of data objects that represent multiple samples in to lists of data objects that represent a single sample. The opposite direction is handled by the constructors for the aggregate data classes.

## Methods

**cytoframe\_to\_flowFrame(object = "cytoframe")** Returns a flowFrame object coerced from a cytoframe object.

**flowFrame\_to\_cytoframe(object = "flowFrame")** Returns a cytoframe object coerced from a flowFrame object.

**cytoset\_to\_flowSet(object = "cytoset")** Returns a flowSet object coerced from a cytoset object.

**flowSet\_to\_cytoset(object = "flowSet")** Returns a cytoset object coerced from a flowSet object.

**flowSet\_to\_list(object = "flowSet")** Returns a list of cytoframe objects with names provided by the sampleNames of the original cytoset

**flowSet(object = "list")** Constructs a cytoset object from a list of cytoframe objects. See documentation for [cytoset](#)

**cytoset\_to\_list(object = "cytoset")** Returns a list of cytoframe objects with names provided by the sampleNames of the original cytoset

**cytoset(object = "list")** Constructs a cytoset object from a list of cytoframe objects. See documentation for [flowSet](#)

## See Also

[merge\\_list\\_to\\_gs](#)

## Examples

```
library(flowCore)
data("GvHD")
fs <- GvHD[1]
cs <- flowSet_to_cytoset(fs)
cf <- cs[[1, returnType="cytoframe"]]
ff <- cytoframe_to_flowFrame(cf)
```

---

convert_backend	<i>convert h5 based gs archive to tiledb</i>
-----------------	----------------------------------------------

---

### Description

convert h5 based gs archive to tiledb

### Usage

```
convert_backend(gs_dir, output_dir)
```

### Arguments

gs_dir	existing gs archive path
output_dir	the new gs path

---

convert_legacy_gs	<i>convert the legacy GatingSet archive (mixed with R and C++ files) to the new format (C++ only)</i>
-------------------	-------------------------------------------------------------------------------------------------------

---

### Description

Older versions of flowWorkspace represented [GatingSet-class](#) objects using a combination of R and C++ files, while newer versions have moved the representation entirely to the C++ level for the sake of efficiency. In order to use GatingSet or GatingSetList archives created in older versions, they will need to be converted to the new format.

### Usage

```
convert_legacy_gs(from, to, ...)
convert_legacy_gslist(from, to, ...)
```

### Arguments

from	the old archive path
to	the new archive path
...	tmp the path where the temporary files will be written to during the conversion. By default it is system temp folder and sometime it is helpful to be able to customize it to other location when system temp folder is full or not sufficient when converting big data sets.

### Details

Note that it is likely some of the keyword values (mainly offsets e.g. BEGINDATA) may change slightly after the conversion due to the process of rewriting data to FCS files through write.FCS.

**Examples**

```
## Not run:  
convert_legacy_gs(old_gs_path, new_gs_path)  
  
## End(Not run)
```

---

cs\_add\_cytoframe      *Add a cytoframe to a cytoset*

---

**Description**

Add a cytoframe to a cytoset

**Usage**

```
cs_add_cytoframe(cs, sn, cf)
```

**Arguments**

cs	cytoset
sn	sample name to be added
cf	cytoframe to be added

---

cs\_get\_uri      *Return the path of the underlying data files*

---

**Description**

Return the path of the underlying data files

**Usage**

```
cs_get_uri(x)  
  
cs_get_h5_file_path(x)  
  
gs_get_uri(x)
```

**See Also**

Other cytoframe/cytoset IO functions: [cf\\_get\\_uri\(\)](#), [cf\\_write\\_disk\(\)](#), [cf\\_write\\_h5\(\)](#), [cf\\_write\\_tile\(\)](#), [load\\_cytoframe\\_from\\_fcs\(\)](#), [load\\_cytoframe\(\)](#), [load\\_cytoset\\_from\\_fcs\(\)](#)

---

cs_set_cytoframe	<i>update a cytoframe in a cytoset</i>
------------------	----------------------------------------

---

### Description

update a cytoframe in a cytoset

### Usage

```
cs_set_cytoframe(cs, sn, cf)
```

### Arguments

cs	cytoset
sn	sample name
cf	cytoframe

---

cytoctx	<i>Create cyto context that carries tiledb context parameters</i>
---------	-------------------------------------------------------------------

---

### Description

Create cyto context that carries tiledb context parameters

Convert cytoctx to a list

### Usage

```
cytoctx(cred = NULL, num_threads = 1L)

print.cytoctx(x, ...)

ctx_to_list(x)
```

### Arguments

cred	credentials for s3 access. It is a list containing elements of "AWS_ACCESS_KEY_ID", "AWS_SECRET_ACCESS_KEY", "AWS_REGION" when NULL, read the default credential file from disk (e.g., <code>~/.aws/credentials</code> )
num_threads	tiledb multithread parameters
x	cytoctx

### Value

a cytoctx object that is a wrapper around the external pointer, thus it is copy-by-reference object

---

**cytoframe**

*cytoframe: A reference class for efficiently managing the data representation of a flowFrame*

---

## Description

This class serves the same purpose as the [flowFrame](#) class from the [flowCore](#) package: to store quantitative data on cell populations from a single FCS run. The primary difference is in the underlying representation of the data. While [flowFrame](#) objects store the underlying data matrix in the `exprs` slot as an R object, [cytoframe](#) objects store the matrix (as well as the data from the other slots) in a C data structure that is accessed through an external pointer. This allows for greater optimization of data operations including I/O, parsing, transformation, and gating.

## Details

From the user's standpoint, interacting with a [cytoframe](#) is very similar to interacting with a [flowframe](#), with one important difference. While operations such as subsetting or copying a [flowFrame](#) using the standard R assignment operator (`<-`) will perform a deep copy of the data in its slots, the same operations on a [cytoframe](#) will produce a view to the same underlying data as the original object. This means that changes made to the [cytoframe](#) resulting from subsetting or copying will affect the original [cytoframe](#). If a deep copy of the underlying data is desired, the `realize_view` method will accomplish this.

Because the [cytoframe](#) class inherits from [flowFrame](#), the [flowFrame](#) slots are present but not utilized. Thus, attempting to access them directly will yield empty data structures. However, the `exprs`, `parameters`, or `description` methods work in a manner similar to a [flowFrame](#) by accessing the same information from the underlying data structure.

## Methods

Many of the methods here have their own documentation pages or are more extensively explained in the documentation for [flowFrame](#), so those documentation pages may be consulted as well for more details.

[ Subsetting. Returns an object of class [cytoframe](#). The syntax for subsetting is similar to that of [data.frames](#). In addition to the usual index vectors (integer and logical by position, character by parameter names), [cytoframes](#) can be subset via [filterResult](#) and [filter](#) objects.

*Usage:*

```
cytoframe[i,j]
```

```
cytoframe[filter,]
```

```
cytoframe[filterResult,]
```

Note that the value of argument `drop` is ignored when subsetting [cytoframes](#).

**\$** Subsetting by channel name. This is similar to subsetting of columns of [data.frames](#), i.e., `frame$FSC.H` is equivalent to `frame[, "FSC.H"]`. Note that column names may have to be quoted if they are not valid R symbols (e.g. `frame$"FSC-H"` or `frame$`FSC-H``).

**exprs, exprs<-** `exprs` returns an object of class `matrix` containing the measured intensities. Rows correspond to cells, columns to the different measurement channels. The `colnames` attribute of the matrix should hold the names or identifiers for the channels. The `rownames` attribute would usually not be set.

`exprs<-` replaces the raw data intensities. The replacement value must be a numeric matrix with `colnames` matching the parameter definitions. Implicit subsetting is allowed (i.e. less columns in the replacement value compared to the original `cytoframe`), but all columns must be defined in the original `cytoframe`.

*Usage:*

```
exprs(cytoframe)
exprs(cytoframe) <-value
```

**head, tail** Show first/last elements of the raw data matrix

*Usage:*

```
head(cytoframe)
tail(cytoframe)
```

**keyword, keyword<-** Extract all entries or a single entry from the annotations by keyword or replace the entire list of key/value pairs with a new named list. See [keyword](#) for details.

*Usage:*

```
keyword(cytoframe)
keyword(cytoframe, character)
keyword(cytoframe) <-list(value)
```

**parameters, parameters<-** Extract parameters and return an object of class [AnnotatedDataFrame](#) containing information about each column of the `cytoframe`, or replace such an object.

This information will generally be filled in by `load_cytoframe_from_fcs` or similar functions using data from the FCS keywords describing the parameters. To access the actual pa-

parameter annotation, use `pData(parameters(cytoframe))`.

Replacement is only valid with `AnnotatedDataFrames` containing all `varLabels` name, desc, range, `minRange` and `maxRange`, and matching entries in the name column to the colnames of the `exprs` matrix. See `parameters` for more details.

*Usage:*

```
parameters(cytoframe)  
parameters(cytoframe) <-value
```

**show** Display details about the `cytoframe` object.

**summary** Return descriptive statistical summary (min, max, mean and quantile) for each channel

*Usage:*

```
summary(cytoframe)
```

**plot** Basic plots for `cytoframe` objects. If the object has only a single parameter this produces a `histogram`. For exactly two parameters we plot a bivariate density map (see `smoothScatter`) and for more than two parameters we produce a simple `splom` plot. To select specific parameters from a `flowFrame` for plotting, either subset the object or specify the parameters as a character vector in the second argument to `plot`. The `smooth` parameters lets you toggle between density-type `smoothScatter` plots and regular scatterplots. For far more sophisticated plotting of flow cytometry data, see the `ggcryo` package.

*Usage:*

```
plot(cytoframe, ...)  
plot(cytoframe, character, ...)  
plot(cytoframe, smooth=FALSE, ...)
```

**ncol, nrow, dim** Extract the dimensions of the data matrix.

*Usage:*

```
ncol(cytoframe)  
nrow(cytoframe)
```

```
dim(cytoframe)
```

**featureNames, colnames, colnames<-** colnames and featureNames are synonyms. They extract parameter names (i.e., the colnames of the data matrix). For colnames there is also a replacement method. This will update the name column in the parameters slot as well.

*Usage:*

```
featureNames(cytoframe)
colnames(cytoframe)
colnames(cytoframe) <-value
```

**markernames, markernames<-** Access or replace the marker names associated with the channels of the cytoframe. For replacement, value should be a named list or character vector where the names correspond to the channel names and the values correspond to the marker names.

*Usage:*

```
markernames(object)
markernames(object) <-value
```

**names** Extract pretty formatted names of the parameters including parameter descriptions.

*Usage:*

```
names(cytoframe)
```

**identifier** Extract GUID of a cytoframe. Returns the file name if no GUID is available. See [identifier](#) for details.

*Usage:*

```
identifier(cytoframe)
```

**range** Get instrument or actual data range of the cytoframe. Note that instrument dynamic range is not necessarily the same as the range of the actual data values, but the theoretical range of values the measurement instrument was able to capture. The values of the dynamic range will be transformed when using the transformation methods for cytoframe objects.

*Parameters:*

`x`: cytoframe object.

`type`: Range type. either "instrument" or "data". Default is "instrument"

*Usage:*

```
range(x, type = "data")
```

**each\_row, each\_col** Apply functions over rows or columns of the data matrix. These are convenience methods. See [each\\_col](#) for details.

*Usage:*

```
each_row(cytoframe, function, ...)
```

```
each_col(cytoframe, function, ...)
```

**transform** Apply a transformation function on a cytoframe object. This uses R's [transform](#) function by treating the cytoframe like a regular `data.frame`. `flowCore` provides an additional inline mechanism for transformations (see [%on%](#)) which is strictly more limited than the out-of-line transformation described here.

*Usage:*

```
transform(cytoframe, translist, ...)
```

**filter** Apply a [filter](#) object on a cytoframe object. This returns an object of class [filterResult](#), which could then be used for subsetting of the data or to calculate summary statistics. See [filter](#) for details.

*Usage:*

```
filter(cytoframe, filter)
```

**split** Split cytoframe object according to a [filter](#), a [filterResult](#) or a factor. For most types of filters, an optional `flowSet=TRUE` parameter will create a [flowSet](#) rather than a simple list. See [split](#) for details.

*Usage:*

```
split(cytoframe, filter, flowSet=FALSE, ...)
```

```
split(cytoframe,filterResult,flowSet=FALSE,...)
split(cytoframe,factor,flowSet=FALSE,...)
```

**Subset** Subset a cytoframe according to a filter or a logical vector. The same can be done using the standard subsetting operator with a filter, filterResult, or a logical vector as first argument.

*Usage:*

```
Subset(cytoframe,filter)
Subset(cytoframe,logical)
```

**cbind2 Not yet implemented.**

Expand a cytoframe by the data in a numeric matrix of the same length. The matrix must have column names different from those of the cytoframe. The additional method for numerics only raises a useful error message.

*Usage:*

```
cbind2(cytoframe, matrix)
cbind2(cytoframe, numeric)
```

**compensate** Apply a compensation matrix (or a [compensation](#) object) on a cytoframe object. This returns a compensated cytoframe.

*Usage:*

```
compensate(cytoframe, matrix)
compensate(cytoframe, data.frame)
compensate(cytoframe, compensation)
```

**decompensate Not yet implemented.**

Reverse the application of a compensation matrix (or a [compensation](#) object) on a cytoframe object. This returns a decompensated cytoframe.

*Usage:*

```
decompensate(cytoframe, matrix)
```

```
decompensate(cytoframe, data.frame)
```

**spillover** Extract spillover matrix from description slot if present. It is equivalent to keyword(x,c("spillover","SPILL"))  
Thus will simply return a list of keyword values for "spillover" and "SPILL".

*Usage:*

```
spillover(cytoframe)
```

**realize\_view** Returns a new `cytoframe` with its own copy of the underlying data (a deep copy).  
The optional `filepath` argument accepts a string to specify a full filename for storing the new  
copy of the data in h5 format.

*Usage:*

```
realize_view(cytoframe,filepath)
```

## See Also

[flowSet](#), [read.FCS](#)

---

`cytoframe-labels`

*Methods to change channel and marker names for cytoframe and cytoset objects*

---

## Description

The methods allow direct alteration of channel names or marker names of `cytoframe` and `cytoset` objects. These objects are accessed by reference and changed in place, so there is no need to assign the return value of these methods.

## Usage

```
cf_swap_colnames(x, col1, col2)  
  
cf_rename_channel(x, old, new)  
  
cf_rename_marker(x, old, new)  
  
cs_swap_colnames(x, col1, col2)
```

## Arguments

x	a cytoframe
col1	first channel name to swap
col2	second channel name to swap
old	old channel or marker name to be changed
new	new channel or marker name after change

---

cytoset	<i>cytoset: a reference class for efficiently managing the data representation of a flowSet</i>
---------	-------------------------------------------------------------------------------------------------

---

## Description

This class is a container for a set of [cytoframe](#) objects, analogous to a [flowSet](#).

## Details

Similar to the distinction between the [cytoframe](#) and [flowFrame](#) classes, the primary difference between the [cytoset](#) and [flowSet](#) classes is in the underlying representation of the data. Because [cytoset](#) is a reference class, copying or subsetting a [cytoset](#) object will return a [cytoset](#) pointing to the same underlying data. A deep copy of the data can be obtained via the `realize_view` method.

There is one notable exception to the typical behavior of most methods returning a [cytoframe](#). The standard extraction operator (`[[[]]]`) will by default perform a deep copy of the subset being extracted and return a [flowFrame](#). This is for the sake of compatibility with existing user scripts.

## Creating Objects

Objects can be created using `cytoset()` and then adding samples by providing a [cytoframe](#) and sample name to `cs_add_cytoframe`:

```
cs <- cytoset()
cs_add_cytoframe(cs, "Sample Name", cytoframe)
```

The safest and easiest way to create [cytosets](#) directly from FCS files is via the [load\\_cytoset\\_from\\_fcs](#) function, and there are alternative ways to specify the files to read. See the separate documentation for details.

## Methods

**[, [[** Subsetting.  $x[i]$  where  $i$  is a scalar, returns a `cytoset` object, and  $x[[i]]$  a `flowFrame` object. In this respect the semantics are similar to the behavior of the subsetting operators for lists.  $x[i, j]$  returns a `cytoset` for which the parameters of each `cytoframe` have been subset according to  $j$ ,  $x[[i, j]]$  returns the subset of a single `flowFrame` for all parameters in  $j$ .

The reason for the default behavior of the extraction operator `[[[]]]` returning a `flowFrame` rather than `cytoframe` is for backwards compatibility with existing user scripts. This behavior can be overridden to instead return a `cytoframe` with the additional `returnType` argument.

*Usage:*

```
cytoset[i]
cytoset[i, j]
cytoset[[i]]
cytoset[[i, returnType = "cytoframe"]]
```

**get\_cytoframe\_from\_cs** Extract a `cytoframe` from a `cytoset` by supplying either a sample name or index and optionally supplying a subset of columns.

The `cytoframe` to be extracted ( $i$  argument) can be specified using its sample name (character) or index in the `cytoset` (int/numeric). Columns ( $j$  argument) can be specified using channel name (character), index (int/numeric), or logical vector. If this argument is missing, all columns will be selected.

*Usage:*

```
(Assuming cs is a cytoset and cf is the extracted cytoframe) cf <-get_cytoframe_from_cs(cs, i, j)
cf <-get_cytoframe_from_cs(cs, i)
```

**\$** Subsetting by frame name. This will return a single `cytoframe` object. Note that names may have to be quoted if they are not valid R symbols (e.g. `cytoset$"sample 1"`).

**colnames, colnames<-** Extract or replace the character object with the (common) column names of all the data matrices in the `cytoframes`.

*Usage:*

```
colnames(cytoset)
colnames(cytoset) <-value
```

**identifier, identifier<-** Extract or replace the name item from the environment.

*Usage:*

```
identifier(cytoset)

identifier(cytoset) <-value
```

**phenoData, phenoData<-** Extract or replace the [AnnotatedDataFrame](#) containing the phenotypic data for the whole data set. Each row corresponds to one of the [cytoframes](#). The `sampleNames` of `phenoData` (see below) must match the names of the `cytoframes` in the `frames` environment.

*Usage:*

```
phenoData(cytoset)

phenoData(cytoset) <-value
```

**pData, pData<-** Extract or replace the data frame (or columns thereof) containing actual phenotypic information from the `phenoData` of the underlying data.

*Usage:*

```
pData(cytoset)

pData(cytoset)$someColumn <-value
```

**varLabels, varLabels<- Not yet implemented.**

Extract and set `varLabels` in the [AnnotatedDataFrame](#) of the `phenoData` of the underlying data.

*Usage:*

```
varLabels(cytoset)

varLabels(cytoset) <-value
```

**sampleNames** Extract and replace sample names from the `phenoData`. Sample names correspond to frame identifiers, and replacing them will also replace the GUID for each `cytoframe`. Note that each sample name needs to be unique.

*Usage:*

```
sampleNames(cytoset)
```

```
sampleNames(cytoset) <-value
```

**keyword** Extract or replace keywords specified in a character vector or a list from the `description` slot of each frame. See [keyword](#) for details.

*Usage:*

```
keyword(cytoset, list(keywords))  
keyword(cytoset, keywords)  
keyword(cytoset) <-list(foo="bar")
```

**length** The number of [cytoframe](#) objects in the set.

*Usage:*

```
length(cytoset)
```

**show** display object summary.

**summary** Return descriptive statistical summary (min, max, mean and quantile) for each channel of each [cytoframe](#).

*Usage:*

```
summary(cytoset)
```

**fsApply** Apply a function on all frames in a cytoset object. Similar to [sapply](#), but with additional parameters. See [fsApply](#) for details.

*Usage:*

```
fsApply(cytoset, function, ...)  
fsApply(cytoset, function, use.exprs=TRUE, ...)
```

**compensate** Apply a compensation matrix on all frames in a cytoset object. See [compensate](#) for details.

*Usage:*

```
compensate(cytoset, matrix)
```

**transform** Apply a transformation function on all frames of a cytoset object. See [transform](#) for details.

*Usage:*

```
transform(cytoset, ...)
```

**filter** Apply a filter on a cytoset object. There are methods for [filter](#) objects, and lists of filter objects. The latter has to be a named list, where names of the list items are matching the `sampleNames` of the cytoset. See [filter](#) for details.

*Usage:*

```
filter(cytoset, filter)
```

```
filter(cytoset, list(filters))
```

**split** Split all cytoframe objects according to a [filter](#), [filterResult](#) or a list of such objects, where the length of the list has to be the same as the length of the cytoset. This returns a list of [cytoframes](#) or an object of class `cytoset` if the `flowSet` argument is set to `TRUE`. Alternatively, a cytoset can be split into separate subsets according to a factor (or any vector that can be coerced into a factor), similar to the behaviour of [split](#) for lists. This will return a list of cytosets. See [split](#) for details.

*Usage:*

```
split(cytoset, filter)
```

```
split(cytoset, filterResult)
```

```
split(cytoset, list(filters))
```

```
split(cytoset, factor)
```

**Subset** Returns a cytoset of [cytoframes](#) that have been subset according to a [filter](#) or [filterResult](#), or according to a list of such items of equal length as the cytoset. See [Subset](#) for details.

*Usage:*

```
Subset(cytoset, filter)
```

```
Subset(cytoset,filterResult)  
Subset(cytoset,list(filters))
```

**rbind2 Not yet implemented.**

Combine two cytoset objects, or one cytoset and one [cytoframe](#) object.

*Usage:*

```
rbind2(cytoset,cytoset)  
rbind2(cytoset,cytoframe)
```

**spillover** Compute spillover matrix from a compensation set. See [spillover](#) for details.

**realize\_view** Returns a new cytoset with its own copy of the underlying data (a deep copy). The optional `filepath` argument accepts a string to specify a full directory name for storing the new copies of the data from the FCS files in h5 format.

*Usage:*

```
realize_view(cytoset,filepath)
```

**cs\_add\_cytoframe** Adds a [cytoframe](#) to the cytoset with sample name given by a string.

*Usage:*

```
cs_add_cytoframe(cytoset,"SampleName",cytoframe)
```

---

```
delete_gs
```

*delete the archive of GatingSet*

---

**Description**

delete the archive of GatingSet

**Usage**

```
delete_gs(path, ctx = .cytoctx_global)
```

**Arguments**

path	either a local path or s3 path (e.g. "s3://bucketname/gs_path")
------	-----------------------------------------------------------------

---

estimateLogicle	<i>Compute logicle transformation from the flowData associated with a GatingHierarchy</i>
-----------------	-------------------------------------------------------------------------------------------

---

## Description

See details in [estimateLogicle](#)

## Usage

```
## S3 method for class 'GatingHierarchy'
estimateLogicle(x, channels, ...)
```

## Arguments

x	a GatingHierarchy
channels	channels or markers for which the logicle transformation is to be estimated.
...	other arguments

## Value

transformerList object

## Examples

```
## Not run:
# gs is a GatingSet
trans.list <- estimateLogicle(gs[[1]], c("CD3", "CD4", "CD8"))
# trans.list is a transformerList that can be directly applied to GatingSet
gs <- transform(gs, trans.list)

## End(Not run)
```

---

extract_cluster_pop_name_from_node	
------------------------------------	--

---

	<i>Extract the population name from the node path It strips the parent path and cluster method name.</i>
--	----------------------------------------------------------------------------------------------------------

---

## Description

Extract the population name from the node path It strips the parent path and cluster method name.

## Usage

```
extract_cluster_pop_name_from_node(node, cluster_method_name)
```

**Arguments**

node	population node path
cluster_method_name	the name of the clustering method

**Examples**

```
extract_cluster_pop_name_from_node("cd3/flowClust_pop1", "flowClust")
#returns "pop1"
```

---

filter_to_list	<i>convert flowCore filter to a list</i> It convert the flowCore gate to a list whose structure can be understood by underlying c++ data structure.
----------------	-----------------------------------------------------------------------------------------------------------------------------------------------------

---

**Description**

convert flowCore filter to a list

It convert the flowCore gate to a list whose structure can be understood by underlying c++ data structure.

**Usage**

```
filter_to_list(x)
```

**Arguments**

x	filter a flowCore gate. Currently supported gates are: "rectangleGate", "polygonGate", "ellipsoidGate" and "booleanFilter"
---	----------------------------------------------------------------------------------------------------------------------------

**Value**

a list

---

flowjo.biexp	<i>construct the flowJo-type biexponentioal transformation function</i>
--------------	-------------------------------------------------------------------------

---

**Description**

Normally it was parsed from flowJo xml workspace. This function provides the alternate way to construct the flowJo version of logicle transformation function within R.

**Usage**

```
flowjo.biexp(
  channelRange = 4096,
  maxValue = 262144,
  pos = 4.5,
  neg = 0,
  widthBasis = -10,
  inverse = FALSE
)
```

**Arguments**

channelRange	numeric	the maximum value of transformed data
maxValue	numeric	the maximum value of input data
pos	numeric	the full width of the transformed display in asymptotic decades
neg	numeric	Additional negative range to be included in the display in asymptotic decades
widthBasis	numeric	unkown.
inverse	logical	whether to return the inverse transformation function.

**Examples**

```
trans <- flowjo.biexp()
data.raw <- c(-1, 1e3, 1e5)
data.trans <- trans(data.raw)
round(data.trans)
inv <- flowjo.biexp(inverse = TRUE)
round(inv(data.trans))
```

flowjo.biexp\_trans *flowJo biexponential transformation.*

**Description**

Used for constructing biexponential transformation object.

**Usage**

```
flowjo.biexp_trans(..., n = 6, equal.space = FALSE)
flowJo.biexp_trans(...)
```

**Arguments**

...	parameters passed to <code>flowJoTrans</code>
n	desired number of breaks (the actual number will be different depending on the data range)
equal.space	whether breaks at equal-spaced intervals

**Value**

biexponential transformation object

**Examples**

```
library(flowCore)
data(GvHD)
fr <- GvHD[[1]]
data.raw <- exprs(fr)[, "FL1-H"]
trans.obj <- flowjo_biexp_trans(equal.space = TRUE)
brks.func <- trans.obj[["breaks"]]
brks <- brks.func(data.raw)
brks # biexp space displayed at raw data scale

#transform it to verify it is equal-spaced at transformed scale
trans.func <- trans.obj[["transform"]]

print(trans.func(brks))
```

**flowjo\_fasinh** *inverse hyperbolic sine transform function*

**Description**

hyperbolic sine/inverse hyperbolic sine (flowJo-version) transform function constructor

**Usage**

```
flowjo_fasinh(m = 4, t = 12000, a = 0.7, length = 256)
flowjo_fsinh(m = 4, t = 12000, a = 0.7, length = 256)
```

**Arguments**

<b>m</b>	numeric the full width of the transformed display in asymptotic decades
<b>t</b>	numeric the maximum value of input data
<b>a</b>	numeric Additional negative range to be included in the display in asymptotic decades
<b>length</b>	numeric the maximum value of transformed data

**Value**

fasinh/fsinh transform function

### Examples

```
trans <- flowjo_fasinh()
data.raw <- c(1,1e2,1e3)
data.trans <- trans(data.raw)
data.trans

inverse.trans <- flowjo_fsinh()
inverse.trans(data.trans)
```

---

flowjo\_fasinh\_trans    *flowJo inverse hyperbolic sine transformation.*

---

### Description

Used to construct the inverse hyperbolic sine transform object.

### Usage

```
flowjo_fasinh_trans(..., n = 6, equal.space = FALSE)

flowJo_fasinh_trans(...)
```

### Arguments

...	parameters passed to flowjo_fasinh
n	desired number of breaks (the actual number will be different depending on the data range)
equal.space	whether breaks at equal-spaced intervals

### Value

fasinh transformation object

### Examples

```
trans.obj <- flowjo_fasinh_trans(equal.space = TRUE)
data <- 1:1e3
brks.func <- trans.obj[["breaks"]]
brks <- brks.func(data)
brks # fasinh space displayed at raw data scale

#transform it to verify it is equal-spaced at transformed scale
trans.func <- trans.obj[["transform"]]
round(trans.func(brks))
```

---

flowjo_log_trans	<i>flog transform function</i>
------------------	--------------------------------

---

### Description

flog transform function constructor. It is different from flowCore version of [logGml2](#) in the way that it reset negative input so that no NAN will be returned.

### Usage

```
flowjo_log_trans(
  decade = 4.5,
  offset = 1,
  scale = 1,
  n = 6,
  equal.space = FALSE
)
```

### Arguments

decade	total number of decades (i.e. log(max)-log(min))
offset	offset to the original input(i.e. min value)
scale	the linear scale factor
n	desired number of breaks (the actual number will be different depending on the data range)
equal.space	whether breaks at equal-spaced intervals

### Value

flog(or its inverse) transform function

### Examples

```
trans <- flowjo_log_trans()
data.raw <- c(1,1e2,1e3)
data.trans <- trans[["transform"]](data.raw)
data.trans

inverse.trans <- trans[["inverse"]]
inverse.trans(data.trans)

#negative input
data.raw <- c(-10,1e2,1e3)
data.trans <- trans[["transform"]](data.raw)
data.trans
inverse.trans(data.trans)#we lose the original value at lower end since flog can't restore negative value
```

```
#different
trans <- flowjo_log_trans(decade = 3, offset = 30)
data.trans <- trans[["transform"]](data.raw)
data.trans
inverse.trans <- trans[["inverse"]]
inverse.trans(data.trans)
```

---

flowWorkspace-deprecated

*Deprecated functions in package **flowWorkspace**.*

---

### Description

```
getStats -> gs(/gh)_pop_get_stats
getProp -> gh_pop_get_proportion
getTotal -> gh_pop_get_count
getPopStats -> gs(/gh)_pop_get_stats
getNodes -> gs_get_pop_paths
getParent -> gs_pop_get_parent
getChildren -> gs_pop_get_children
getGate -> gs(/gh)_get_gate
getIndices -> gh_pop_get_indices
isGated -> gh_pop_is_gated
isNegated -> gh_pop_is_negated
isHidden -> gh_pop_is_hidden
getData -> gs(/gh)_get_data
getTransformations -> gh_get_transformations
getCompensationMatrices -> gh_get_compensations
plotGate -> autoplot
setNode -> gs(/gh)_set_node_name/gs(/gh)_set_node_visible
isNcdf -> gs_is_h5
flowData -> gs_cyto_data
flowData<- -> gs_cyto_data<-
getLogLevel -> get_log_level
setLogLevel -> set_log_level
rbind2 -> gslist_to_gs
filterObject -> filter_to_list
add -> gs_pop_add
```

```

Rm -> gs_pop_remove
copyNode -> gh_copy_gate
openWorkspace -> open_flowjo_xml
flowJo.flog -> flowjo_log_trans
flowJoTrans -> flowjo.biexp
flowJo.biexp_trans -> flowjo.biexp_trans
flowJo.fasinh -> flowjo.fasinh
flowJo.fsinh -> flowjo.fsinh
flowJo.fasinh_trans -> flowjo.fasinh_trans
getDescendants -> gh_pop_get_descendants
getSingleCellExpression -> gs_get_singlecell_expression
groupByTree -> gs_split_by_tree
groupByChannels -> gs_split_by_channels
checkRedundantNodes -> gs_check_redundant_nodes
dropRedundantNodes -> gs_remove_redundant_nodes
dropRedundantChannels -> gs_drop_redundant_channels
updateChannels -> gs_update_channels
moveNode -> gh_pop_move
setGate -> gs(/gh)_pop_set_gate
updateIndices -> gh_pop_set_indices
getMergedStats -> gs_pop_get_count_with_meta
set.count.xml -> gh_pop_set_xml_count

```

---

flow\_breaks

*Generate the breaks that makes sense for flow data visualization*

---

## Description

It is mainly used as helper function to construct breaks function used by 'trans\_new'.

## Usage

```
flow_breaks(x, n = 6, equal.space = FALSE, trans.fun, inverse.fun)
```

## Arguments

x	the raw data values
n	desired number of breaks (the actual number will be different depending on the data range)
equal.space	whether breaks at equal-spaced intervals
trans.fun	the transform function (only needed when equal.space is TRUE)
inverse.fun	the inverse function (only needed when equal.space is TRUE)

**Value**

either  $10^n$  intervals or equal-spaced(after transformed) intervals in raw scale.

**Examples**

```
library(flowCore)
data(GvHD)
fr <- GvHD[[1]]
data.raw <- exprs(fr)[, "FL1-H"]
flow_breaks(data.raw)

trans <- logicleTransform()
inv <- inverseLogicleTransform(trans = trans)
myBrks <- flow_breaks(data.raw, equal.space = TRUE, trans = trans, inv = inv)
round(myBrks)
#to verify it is equally spaced at transformed scale
print(trans(myBrks))
```

**flow\_trans**

*helper function to generate a trans objects Used by other specific trans constructor*

**Description**

helper function to generate a trans objects Used by other specific trans constructor

**Usage**

```
flow_trans(name, trans.fun, inverse.fun, equal.space = FALSE, n = 6)
```

**Arguments**

name	transformation name
trans.fun	the transform function (only needed when equal.space is TRUE)
inverse.fun	the inverse function (only needed when equal.space is TRUE)
equal.space	whether breaks at equal-spaced intervals
n	desired number of breaks (the actual number will be different depending on the data range)

---

**GatingHierarchy-class Class GatingHierarchy**

---

## Description

GatingHierarchy is a class for representing the gating hierarchy, which can be either imported from a flowJo workspace or constructed in R.

## Details

There is a one-to-one correspondence between GatingHierarchy objects and FCS files in the flowJo workspace. Each sample (FCS file) is associated with its own GatingHierarchy. It is also more space efficient by storing gating results as logical/bit vector instead of copying the raw data.

Given a GatingHierarchy, one can extract the data associated with any subpopulation, extract gates, plot gates, and extract population proportions. This facilitates the comparison of manual gating methods with automated gating algorithms.

## See Also

[GatingSet](#)

## Examples

```
## Not run:
require(flowWorkspaceData)
d<-system.file("extdata",package="flowWorkspaceData")
wsfile<-list.files(d,pattern="A2004Analysis.xml",full=TRUE)
library(CytoML)
ws <- open_flowjo_xml(wsfile);
G<-try(flowjo_to_gatingset(ws,path=d,name=1));
gh <- G[[1]]
gh_pop_compare_stats(gh);
gh_plot_pop_count_cv(gh)
nodes <- gs_get_pop_paths(gh)
thisNode <- nodes[4]
require(ggcyto)
autoplot(gh,thisNode);
gh_pop_get_gate(gh,thisNode);
gh_pop_get_data(gh,thisNode)

## End(Not run)
```

---

GatingSet-class	<i>Class "GatingSet"</i>
-----------------	--------------------------

---

## Description

GatingSet holds a set of GatingHierarchy objects, representing a set of samples and the gating scheme associated with each.

## Details

Objects stores a collection of GatingHierarchies and represent a group in a flowJo workspace. A GatingSet can have two “states”. After a call to `flowjo_to_gatingset(...,execute=FALSE)` , the workspace is imported but the data is not. Setting `execute` to TRUE is needed in order to load, transform, compensate, and gate the associated data. Whether or not a GatingHierarchy has been applied to data is encoded in the `flag` slot. Some methods will warn the user, or may not function correctly if the GatingHierarchy has not been executed. This mechanism is in place, largely for the purpose of speed when working with larger workspaces. It allows the use to load a workspace and subset desired samples before proceeding to load the data.

## Slots

**pointer:** Object of class "externalptr". points to the gating hierarchy stored in C data structure.  
**transformation:** Object of class "list". a list of transformation objects used by GatingSet.

## See Also

[GatingHierarchy](#)

## Examples

```
## Not run:
require(flowWorkspaceData)
d<-system.file("extdata",package="flowWorkspaceData")
wsfile<-list.files(d,pattern="A2004Analysis.xml",full=TRUE)
library(CytoML)
ws <- open_flowjo_xml(wsfile);
G<-try(flowjo_to_gatingset(ws,execute=TRUE,path=d,name=1));
gs_plot_pop_count_cv(G);

## End(Not run)
```

---

GatingSet-methods      *constructors for GatingSet*

---

### Description

construct a gatingset with empty trees (just root node)

### Usage

```
## S4 method for signature 'cytoset,ANY'
GatingSet(x)
```

### Arguments

x	a flowSet, ncdfFlowSet, or cytoset
...	arguments passed to flowSet_to_cytoset() when x is a flowSet

### Examples

```
## Not run:
#fdata could be a flowSet, ncdfFlowSet, or GatingSet
gs <- GatingSet(fdata)

## End(Not run)
```

---

GatingSetList-class      *Class "GatingSetList"*

---

### Description

A list of of GatingSet objects. This class exists for method dispatching.

use GatingSetList constructor to create a GatingSetList from a list of GatingSet

### Usage

```
GatingSetList(x, samples = NULL)
```

### Arguments

x	a list of GatingSet
samples	character vector specifying the order of samples. if not specified, the samples are ordered as the underlying stored order.

## Details

Objects store a collection of GatingSets, which usually has the same gating trees and markers. Most GatingSets methods can be applied to GatingSetList.

## See Also

[GatingSet](#) [GatingHierarchy](#)

## Examples

```
## Not run:
#load several GatingSets from disk
gs_list<-lapply(list.files("../gs_toMerge",full=T) ,function(this_folder){
  load_gs(this_folder)
})

#gs_list is a list
gs_groups <- merge(gs_list)
#returns a list of GatingSetList objects
gslist2 <- gs_groups[[2]]
#gslist2 is a GatingSetList that contains multiple GatingSets and they share the same gating and data structure
gslist2
class(gslist2)
sampleNames(gslist2)

#reference a GatingSet by numeric index
gslist2[[1]]
#reference a GatingSet by character index
gslist2[["30104.fcs"]]

#loop through all GatingSets within GatingSetList
lapply(gslist2,sampleNames)

#subset a GatingSetList by [
sampleNames(gslist2[c(4,1)])
sampleNames(gslist2[c(1,4)])
gslist2[c("30104.fcs")]

#get flow data from it
gs_pop_get_data(gslist2)
#get gated flow data from a particular population
gs_pop_get_data(gslist2, "3+")

#get the gates associated with one population
gs_pop_get_gate(gslist2,"3+")
gs_pop_get_gate(gslist2,5)

#get the pheno data
pData(gslist2[3:1])
#modify the pheno data
pd <- pData(gslist2)
pd$id <- 1:nrow(pd)
```

```

pData(gslist2) <- pd
pData(gslist2[3:2])

#plot the gate
autoplot(gslist2[1:2],5)

#remove cerntain gates by loop through GatingSets
gs_get_pop_paths(gslist2[[1]])
lapply(gslist2,function(gs)gs_pop_remove("Excl",gs = gs))

#extract the stats
gs_pop_get_count_fast(gslist2)
#extract statistics by using getQAStats defined in QUALIFIER package
res<-getQAStats(gslist2[c(4,2)],isMFI=F,isSpike=F,nslaves=1)

#archive the GatingSetList
save_gslist(gslist2, path ="~/rglab/workspace/flowIncubator/output/gslist",overwrite=T)
gslist2 <- load_gslist(path ="~/rglab/workspace/flowIncubator/output/gslist")

#convert GatingSetList into one GatingSet by merge_list_to_gs
gs_merged2 <- merge_list_to_gs(gslist2)
gs_merged2

## End(Not run)

## Not run:
samleNames(gsA) # return A1, A2
samleNames(gsB) # return B1, B2
gs.list <- list(gsA, gsB)
gslist<- GatingSetList(gs.list)
sampleNames(gslist) #return A1,A2,B1,B2

#set different order when create the GatingSetList
gslist<- GatingSetList(gs.list, samples = c("A1", "B1", "A2", "B2"))
sampleNames(gslist) #return A1,B1,A2,B2

## End(Not run)

```

get\_default\_backend    *get/set the default backend format of cytoframe*

### Description

get/set the default backend format of cytoframe

### Usage

```

get_default_backend()

set_default_backend(backend = c("h5", "mem", "tile"))

```

**Arguments**

backend	one of c("h5", "mem", "tile")
---------	-------------------------------

---

get_log_level	<i>get/set the log level</i>
---------------	------------------------------

---

**Description**

It is helpful sometime to get more detailed print out for the purpose of trouble shooting

**Usage**

```
get_log_level()
set_log_level(level = "none")
```

**Arguments**

level	a character that represents the log level , can be value of c("none", "GatingSet", "GatingHierarchy", "Population", "gate") default is "none" , which does not print any information from C parser.
-------	-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

**Value**

a character that represents the internal log level

**Examples**

```
get_log_level()
set_log_level("Population")
get_log_level()
```

---

gh_apply_to_cs	<i>Construct a GatingSet using a template</i>
----------------	-----------------------------------------------

---

**Description**

This uses a [GatingHierarchy](#) as a template to apply to other loaded samples in the form of a [cytoset](#), resulting in a [GatingSet](#). The transformations and gates from the template are applied to all samples. The compensation applied to each of the samples can be controlled via the `compensation_source` argument.

**Usage**

```
gh_apply_to_cs(x, cs, swap_cols = FALSE, compensation_source = "sample", ...)
```

**Arguments**

x GatingHierarchy  
 cs a cytoset  
 swap\_cols for internal usage  
 compensation\_source  
     One of the following options:  
     • "sample" – each cytoframe will be compensated with the spillover matrix included in its own FCS  
     • "template" – all cytoframes will be compensated with the spillover matrix of the template GatingHierarchy  
     • "none" – no compensation will be applied  
 ... not currently used

**Value**

a GatingSet

gh\_apply\_to\_new\_fcs *Construct a GatingSet using a template and FCS files*

**Description**

This uses a [GatingHierarchy](#) as a template to apply to other loaded samples in the form of a list of FCS files, resulting in a [GatingSet](#). The transformations and gates from the template are applied to all samples.

**Usage**

```
gh_apply_to_new_fcs(  
  x,  
  files,  
  swap_cols = FALSE,  
  backend = get_default_backend(),  
  compensation_source = "sample",  
  ...  
)
```

**Arguments**

x GatingHierarchy  
 swap\_cols for internal usage  
 backend the backend storage mode to use for [load\\_cytoset\\_from\\_fcs](#)  
 compensation\_source  
     One of the following options:

- "sample" – each cytoframe will be compensated with the spillover matrix included in its own FCS
- "template" – all cytoframes will be compensated with the spillover matrix of the template GatingHierarchy
- "none" – no compensation will be applied

... other arguments passed to `load_cytoset_from_fcs`

## Details

This method is still included to support legacy scripts but will be deprecated for the more modular workflow of loading a `cytoset` via `load_cytoset_from_fcs` followed by `gh_apply_to_cs`.

---

`gh_copy_gate`

*Copy a node along with all of its descendant nodes to the given ancestor*

---

## Description

Copy a node along with all of its descendant nodes to the given ancestor

## Usage

`gh_copy_gate(gh, node, to)`

## Arguments

<code>gh</code>	GatingHierarchy
<code>node</code>	the node to be copied
<code>to</code>	the new parent node under which the node will be copied

## Examples

```
library(flowWorkspace)
dataDir <- system.file("extdata", package="flowWorkspaceData")
suppressMessages(gs <- load_gs(list.files(dataDir, pattern = "gs_manual", full = TRUE)))
gh <- gs[[1]]
old.parent <- gs_pop_get_parent(gh, "CD4")
new.parent <- "singlets"
gh_copy_gate(gh, "CD4", new.parent)
gs_get_pop_paths(gh)
```

---

gh\_get\_cluster\_labels *Retrieve the cluster labels from the cluster nodes*

---

### Description

Clustering results are stored as individual gated nodes. This helper function collect all the gating indices from the same clustering run (identified by 'parent' node and 'cluster\_method\_name' and merge them as a single factor.

### Usage

```
gh_get_cluster_labels(gh, parent, cluster_method_name)
```

### Arguments

gh	GatingHierarchy
parent	the parent population/node name or path
cluster_method_name	the name of the clustering method

---

gh\_get\_compensations *Retrieve the compensation matrices from a GatingHierarchy or GatingSet*

---

### Description

Retrieve the compensation matrices from a GatingHierarchy or GatingSet.

### Usage

```
gh_get_compensations(x)  
gs_get_compensations(x)
```

### Arguments

x	A GatingHierarchy or GatingSet object.
---	----------------------------------------

### Details

Return all the compensation matrices in a GatingHierarchy or GatingSet

### Value

A list of `matrix` representing the spillover matrix in GatingHierarchy or GatingSet

## Examples

```
## Not run:
# Assume gh is a GatingHierarchy and gs is a GatingSet
gh_get_compensations(gh)
gs_get_compensations(gs)

## End(Not run)
```

### gh\_get\_transformations

*Return a list of transformations or a transformation in a GatingHierarchy*

## Description

Return a list of all the transformations or a transformation in a GatingHierarchy

## Usage

```
gh_get_transformations(
  x,
  channel = NULL,
  inverse = FALSE,
  only.function = TRUE,
  ...
)
```

## Arguments

x	A GatingHierarchy object
channel	character channel name
inverse	logical whether to return the inverse transformation function. Valid when only.function is TRUE
only.function	logical whether to return the function or the entire transformer object(see scales package) that contains transform and inverse and breaks function.
...	other arguments equal.spaced logical passed to the breaks function to determine whether to break at $10^n$ or equally spaced intervals

## Details

Returns a list of the transformations or a transformation in the flowJo workspace. The list is of length L, where L is the number of distinct transformations applied to samples in the flowjo\_workspace. Each element of L is itself a list of length M, where M is the number of parameters that were transformed for a sample or group of samples in a flowjo\_workspace. For example, if a sample has 10 parameters, and 5 are transformed during analysis, using two different sets of transformations, then L will be of length 2, and each element of L will be of length 5. The elements of L represent channel- or parameter-specific transformation functions that map from raw intensity values to channel-space used by flowJo.

**Value**

lists of functions(or transform objects when only.function is FALSE), with each element of the list representing a transformation applied to a specific channel/parameter of a sample.

**Examples**

```
## Not run:
#Assume gh is a GatingHierarchy
gh_get_transformations(gh); # return a list transformation functions
gh_get_transformations(gh, inverse = TRUE); # return a list inverse transformation functions
gh_get_transformations(gh, channel = "FL1-H") # only return the transformation associated with given channel
gh_get_transformations(gh, channel = "FL1-H", only.function = FALSE) # return the entire transform object

## End(Not run)
```

**gh\_plot\_pop\_count\_cv** *Plot the coefficient of variation between xml and openCyto population statistics for each population in a gating hierarchy.*

**Description**

This function plots the coefficient of variation calculated between the xml population statistics and the openCyto population statistics for each population in a gating hierarchy extracted from a xml Workspace.

**Usage**

```
gh_plot_pop_count_cv(x, path = "auto", ...)
gs_plot_pop_count_cv(x, scales = list(x = list(rot = 90)), path = "auto", ...)
```

**Arguments**

x	A GatingHierarchy from or a GatingSet.
path	character see <a href="#">gs_get_pop_paths</a>
...	Additional arguments to the barplot methods.
scales	list see <a href="#">barchart</a>

**Details**

The CVs are plotted as barplots across panels on a grid of size m by n.

**Value**

Nothing is returned.

**See Also**[gs\\_pop\\_get\\_count\\_fast](#)**Examples**

```
## Not run:
#G is a GatingHierarchy
gs_plot_pop_count_cv(G, 4, 4);

## End(Not run)
```

**gh\_pop\_compare\_stats** *Compare the stats(count/freq) between the version parsed from xml and the one recalculated/gated from R*

**Description**

Compare the stats(count/freq) between the version parsed from xml and the one recalculated/gated from R

**Usage**

```
gh_pop_compare_stats(x, path = "auto", ...)
```

**Arguments**

x	GatingHierarchy
path	see <a href="#">gs_get_pop_paths</a>
...	not used

**gh\_pop\_get\_cluster\_name**  
*check if a node is clustering node*

**Description**

check if a node is clustering node

**Usage**

```
gh_pop_get_cluster_name(gh, node)
```

**Arguments**

gh	GatingHierarchy
node	the population/node name or path

**Value**

the name of the clustering method. If it is not cluster node, returns NULL

---

gh_pop_get_data	<i>get gated flow data from a GatingHierarchy/GatingSet/GatingSetList</i>
-----------------	---------------------------------------------------------------------------

---

**Description**

get gated flow data from a GatingHierarchy/GatingSet/GatingSetList

**Usage**

```
gh_pop_get_data(obj, y = "root", inverse.transform = FALSE, ...)
```

**Arguments**

obj	A GatingHierarchy, GatingSet or GatingSetList object.
y	character the node name or full(/partial) gating path. If not specified, will return the complete flowFrame/flowSet at the root node.
inverse.transform	logical flag indicating whether to inverse transform the data
...	arguments passed to ncdfFlow::[]

**Details**

Returns a flowFrame/flowSet containing the events in the gate defined at node *y*. Subset membership can be obtained using `gh_pop_get_indices`. Population statistics can be obtained using `getPop` and `gh_pop_compare_stats`. When calling `gh_pop_get_data` on a GatingSet, the trees representing the GatingHierarchy for each sample in the GatingSet are presumed to have the same structure. To update the data, use `gs_cyto_data` method.

**Value**

A flowFrame object if *obj* is a GatingHierarchy. A flowSet or ncdfFlowSet if a GatingSet. A ncdfFlowList if a GatingSetList.

**See Also**

[gs\\_cyto\\_data](#) [gh\\_pop\\_get\\_indices](#) [gh\\_pop\\_compare\\_stats](#)

## Examples

```
## Not run:
#G is a GatingSet
geData(G,3) #get a flowSet constructed from the third node / population in the tree.
geData(G,"cd4")

#gh is a GatingHierarchy
gh_pop_get_data(gh)

## End(Not run)
```

---

gh\_pop\_get\_descendants

*get all the descendant nodes for the given ancestor*

---

## Description

get all the descendant nodes for the given ancestor

## Usage

```
gh_pop_get_descendants(gh, node, showHidden = TRUE, ...)
```

## Arguments

gh	GatingHierarchy
node	the node path
showHidden	whether show hidden nodes
...	passed to getNode call

## Examples

```
library(flowWorkspace)
dataDir <- system.file("extdata", package="flowWorkspaceData")
suppressMessages(gs <- load_gs(list.files(dataDir, pattern = "gs_manual", full = TRUE)))
gh_pop_get_descendants(gs[[1]], "CD4")
gh_pop_get_descendants(gs[[1]], "CD8", path = "auto")
```

---

gh\_pop\_get\_full\_path convert the partial gating path to the full path

---

**Description**

convert the partial gating path to the full path

**Usage**

```
gh_pop_get_full_path(gh, path)
```

**Arguments**

gh	GatingHierarchy object
path	the partial gating path

**Value**

the full gating path

---

gh\_pop\_get\_indices *Get the membership indices for each event with respect to a particular gate in a GatingHierarchy*

---

**Description**

Returns a logical vector that describes whether each event in a sample is included or excluded by this gate.

**Usage**

```
gh_pop_get_indices(obj, y)
```

**Arguments**

obj	A GatingHierarchy representing a sample.
y	A character giving the name or full(/partial) gating path of the population / node of interest.

**Details**

Returns a logical vector that describes whether each event in the data file is included in the given gate of this GatingHierarchy. The indices are for all events in the file, and do not reflect the population counts relative to the parent but relative to the root. To get population frequencies relative to the parent one cross-tabulate the indices of y with the indices of its parent.

**Value**

A logical vector of length equal to the number of events in the FCS file that determines whether each event is or is not included in the current gate.

**Note**

Generally you should not need to use *gh\_pop\_get\_indices* but the more convenient methods *gh\_pop\_get\_proportion* and *gh\_pop\_compare\_stats* which return population frequencies relative to the parent node. The indices returned reference all events in the file and are not directly suitable for computing population statistics, unless subsets are taken with respect to the parent populations.

**See Also**

[gh\\_pop\\_compare\\_stats](#)

**Examples**

```
## Not run:
#G is a gating hierarchy
#Return the indices for population 5 (topological sort)
gh_pop_get_indices(G,gs_get_pop_paths(G,tsort=TRUE)[5]);

## End(Not run)
```

**gh\_pop\_get\_indices\_mat**

*Return the single-cell matrix of 1/0 dichotomized expression*

**Description**

Return the single-cell matrix of 1/0 dichotomized expression

**Usage**

`gh_pop_get_indices_mat(gh, y)`

**Arguments**

<code>gh</code>	GatingHierarchy object
<code>y</code>	character vector containing the node names

---

gh\_pop\_get\_proportion *Get count or proportion from populations*

---

## Description

Get count or proportion from populations

## Usage

```
gh_pop_get_proportion(x, y, xml = FALSE)
```

```
gh_pop_get_count(x, y, xml = FALSE)
```

## Arguments

x	GatingHierarchy
y	character node name or path
xml	whether to extract xml stats or openCyto stats

---

gh\_pop\_move *move a node along with all of its descendant nodes to the given ancestor*

---

## Description

move a node along with all of its descendant nodes to the given ancestor

## Usage

```
gh_pop_move(gh, node, to, recompute = TRUE)
```

## Arguments

gh	GatingHierarchy
node	the node to be moved
to	the new parent node under which the node will be moved to
recompute	whether to recompute the gates after the node is moved. Default is TRUE.

## Examples

```
library(flowWorkspace)
dataDir <- system.file("extdata", package="flowWorkspaceData")
suppressMessages(gs <- load_gs(list.files(dataDir, pattern = "gs_manual", full = TRUE)))
gh <- gs[[1]]
old.parent <- gs_pop_get_parent(gh, "CD4")
new.parent <- "singlets"
gh_pop_move(gh, "CD4", new.parent)
gs_pop_get_parent(gh, "CD4")
```

---

gh\_pop\_set\_indices      *directly update event indices without changing gates*

---

## Description

It is useful when we want to alter the population at events level yet without removing or adding the existing gates.

## Usage

```
gh_pop_set_indices(obj, y, z)
```

## Arguments

obj	GatingHierarchy object
y	character node name or path
z	logical vector as local event indices relative to node y

## Examples

```
library(flowWorkspace)
dataDir <- system.file("extdata", package="flowWorkspaceData")
suppressMessages(gs <- load_gs(list.files(dataDir, pattern = "gs_manual", full = TRUE)))
gh <- gs[[1]]
#get pop counts
pop.stats <- gh_pop_get_stats(gh, nodes = c("CD3+", "CD4", "CD8"))
pop.stats

# subsample 30% cell events at CD3+ node
total <- gh_pop_get_count(gh, "root")
gInd <- seq_len(total) #create integer index for cd3
gInd <- sample.int(total, size = total * 0.3) #randomly select 30%
#convert it to logic index
gInd.logical <- rep(FALSE, total)
gInd.logical[gInd] <- TRUE
#replace the original index stored at GatingHierarchy
gh_pop_set_indices(gh, "CD3+", gInd.logical)
#check the updated pop counts
```

```
gh_pop_get_stats(gs[[1]], nodes = c("CD3+", "CD4", "CD8")) #note that CD4, CD8 are not updated
#update all the descendants of CD3+
nodes <- gh_pop_get_descendants(gh, "CD3+")
for (node in nodes) suppressMessages(recompute(gh, node))
gh_pop_get_stats(gs[[1]], nodes = c("CD3+", "CD4", "CD8")) #now all are update to date
```

`gh_pop_set_xml_count` *save the event counts parsed from xml into c++ tree structure*

## Description

It is for internal use by the diva parser

## Usage

```
gh_pop_set_xml_count(gh, node, count)
```

## Arguments

gh	GatingHierarchy
node	the unique gating path that uniquely identifies a population node
count	integer number that is events count for the respective gating node directly parsed from xml file

## Examples

```
## Not run:
gh_pop_set_xml_count(gh, "CD3", 10000)

## End(Not run)
```

`gslist_to_gs` *Merge a GatingSetList into a single GatingSet*

## Description

Merge a GatingSetList into a single GatingSet

## Usage

```
gslist_to_gs(x, ...)
```

## Arguments

x	GatingSetList
...	other arguments passed to gslist_to_gs method for ncdfFlowList

---

gs\_check\_redundant\_nodes

*try to determine the redundant terminal(or leaf) nodes that can be removed*

---

## Description

These leaf nodes make the gating trees to be different from one another and can be removed by the subsequent convenient call [gs\\_remove\\_redundant\\_nodes](#).

## Usage

```
gs_check_redundant_nodes(x, path = "auto", ...)
```

## Arguments

x	GatingSet or list of groups(each group is a list of 'GatingSet'). When it is a list, it is usually the outcome from <a href="#">gs_split_by_tree</a> .
path	argumented passed to <a href="#">gs_get_pop_paths</a> . The default value is "auto".
...	other arguments passed to <a href="#">gs_get_pop_paths</a> .

## Value

a list of the character vectors inicating the nodes that are considered to be redundant for each group of GatingSets.

## Examples

```
## Not run:
gslist <- list(gs1, gs2, gs3, gs4, gs5)
gs_groups <- gs_split_by_tree(gslist)
toRm <- gs_check_redundant_nodes(gs_groups)

## End(Not run)
```

---

gs\_cyto\_data

*Fetch or replace the flowData object associated with a GatingSet .*

---

## Description

Accessor method that gets or replaces the [cytoset/flowSet/ncdfFlowSet](#) object in a GatingSet or GatingHierarchy

**Usage**

```
gs_cyto_data(x, ...)

## S4 method for signature 'GatingSet'
gs_cyto_data(x, inverse.transform = FALSE)

gs_cyto_data(x) <- value
```

**Arguments**

x	A GatingSet
...	other arguments
inverse.transform	logical flag indicating whether to inverse transform the data
value	The replacement flowSet or ncdfFlowSet object

**Details**

Accessor method that sets or replaces the ncdfFlowSet object in the GatingSet or GatingHierarchy.

**Value**

the object with the new flowSet in place.

---

gs\_get\_compensation\_internal  
*extract compensation object from GatingSet*

---

**Description**

extract compensation object from GatingSet

**Usage**

```
gs_get_compensation_internal(gs, sampleName)
```

**Arguments**

gs	GatingSet
sampleName	sample name

---

gs_get_leaf_nodes	<i>get all the leaf nodes</i>
-------------------	-------------------------------

---

### Description

get all the leaf nodes

### Usage

```
gs_get_leaf_nodes(x, ancestor = "root", ...)
gh_get_leaf_nodes(x, ancestor = "root", ...)
```

### Arguments

x	GatingHierarchy/GatingSet object
ancestor	ancestor node where the leaf nodes descend from. Default is 'root'.
...	arguments passed to 'gs_get_pop_paths' method

### Value

the leaf nodes

---

gs_get_pop_paths	<i>Get the names of all nodes from a gating hierarchy.</i>
------------------	------------------------------------------------------------

---

### Description

gs\_get\_pop\_paths returns a character vector of names of the nodes (populations) in the GatingSet.

### Usage

```
gs_get_pop_paths(
  x,
  y = NULL,
  order = "regular",
  path = "full",
  showHidden = FALSE,
  ...
)

gh_get_pop_paths(
  x,
  y = NULL,
  order = "regular",
```

```

  path = "full",
  showHidden = FALSE,
  ...
)

```

### Arguments

x	A GatingSet Assuming the gating hierarchy are identical within the GatingSet, the Gating tree of the first sample is used to query the node information.
y	A character not used.
order	order=c("regular","tsort","bfs") returns the nodes in regular, topological or breadth-first sort order. "regular" is default.
path	A character or numeric scalar. when numeric, it specifies the fixed length of gating path (length 1 displays terminal name). When character, it can be either 'full' (full path, which is default) or 'auto' (display the shortest unique gating path from the bottom of gating tree).
showHidden	logical whether to include the hidden nodes
...	Additional arguments.

### Details

integer indices of nodes are based on regular order, so whenever need to map from character node name to integer node ID, make sure to use default order which is regular.

### Value

gs\_get\_pop\_paths returns a character vector of node/population names, ordered appropriately.

### Examples

```

## Not run:
# G is a gating hierarchy
gs_get_pop_paths(G, path = 1)#return node names (without prefix)
gs_get_pop_paths(G, path = "full")#return the full path
gs_get_pop_paths(G, path = 2)#return the path as length of two
gs_get_pop_paths(G, path = "auto")#automatically determine the length of path
gs_pop_set_name(G, "L", "lymph")

## End(Not run)

```

---

### gs\_get\_singlecell\_expression

*Return the cell events data that express in any of the single populations defined in y*

---

## Description

Returns a list of matrix containing the events that expressed in any one of the populations defined in y

## Usage

```
gs_get_singlecell_expression(
  x,
  nodes,
  other.markers = NULL,
  swap = FALSE,
  threshold = TRUE,
  marginal = TRUE,
  mc.cores = getOption("mc.cores", 1L),
  inverse.transform = FALSE,
  ...
)

gs_get_singlecell_expression_by_gate(...)
```

## Arguments

x	A GatingSet or GatingSetList object .
nodes	character vector specifying different cell populations
other.markers	character vector specifying the extra markers/channels to be returned besides the ones derived from "nodes" and "map" argument. It is only valid when threshold is set to FALSE.
swap	logical indicates whether channels and markers of flow data are swapped.
threshold	logical indicates whether to threshold the flow data by setting intensity value to zero when it is below the gate threshold.
marginal	logical indicates whether the gate is treated as 1d marginal gate. Default is TRUE, which means markers are determined either by node name or by 'map' argument explained below. When FALSE, the markers are determined by the gate dimensions. and node name and 'map' argument are ignored.
mc.cores	passed to mclapply. Default is 1, which means the process runs in serial mode. When it is larger than 1, parallel mode is enabled.
inverse.transform	logical flag indicating whether to inverse transform the data
...	other arguments map a named list providing the mapping between node names (as specified in the gating hierarchy of the gating set) and channel names (as specified in either the desc or name columns of the parameters of the associated flowFrames in the GatingSet). see examples.
	ignore.case whether to ignore case when match the marker names. Default is FALSE.

**Value**

A list of numerci matrices

**Author(s)**

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**See Also**

[gh\\_pop\\_get\\_indices](#) [gs\\_pop\\_get\\_count\\_fast](#)

**Examples**

```
## Not run:
#G is a GatingSet
nodes <- c("4+/TNFa+", "4+/IL2+")
res <- gs_get_singlecell_expression(gs, nodes)
res[[1]]

# if it fails to match the given nodes to the markers, then try to provide the mapping between node and marker explicitly
res <- gs_get_singlecell_expression(gs, nodes, map = list("4+/TNFa+" = "TNFa", "4+/IL2+" = "IL2"))

# It can also operate on the 2d gates by setting marginal to FALSE
# The markers are no longer deduced from node names or supplied by map
# Instead, it retrieves the markers that are associated with the gates
nodes <- c("4+/TNFa+IFNg+", "4+/IL2+IL3+")
res <- gs_get_singlecell_expression(gs, nodes, marginal = FALSE)
#or simply call convenient wrapper
gs_get_singlecell_expression_by_gate(gs, nodes)

## End(Not run)
```

**gs\_is\_persistent** *determine whether the flow data associated with a GatingSet is persistent(on-disk) or in-memory*

**Description**

determine whether the flow data associated with a GatingSet is persistent(on-disk) or in-memory

**Usage**

```
gs_is_persistent(x)

gs_is_h5(x)

isNcdf(x)
```

**Arguments**

x GatingSet object

**Value**

logical

---

**gs\_plot\_diff\_tree** *visualize the tree structure differnece among the GatingSets*

---

**Description**

visualize the tree structure differnece among the GatingSets

**Usage**

`gs_plot_diff_tree(x, path = "auto", ...)`

**Arguments**

x list of groups(each group is a list of 'GatingSet'). it is usually the outcome from [gs\\_split\\_by\\_tree](#).  
 path passed to getNodes  
 ... passed to getNodes

**Examples**

```
## Not run:
gslist <- list(gs1, gs2, gs3, gs4, gs5)
gs_groups <- gs_split_by_tree(gslist)
gs_plot_diff_tree(gs_groups)

## End(Not run)
```

---

**gs\_pop\_add** *Create a GatingSet and add/remove the flowCore gate(or population) to/from a GatingHierarchy/GatingSet.*

---

**Description**

GatingSet method creates a gatingset from a flowSet with the ungated data as the root node. add method add the flowCore gate to a GatingHierarchy/GatingSet. `gs_pop_set_gate` method update the gate of one population node in GatingHierarchy/GatingSet. `Rm` method Remove the population node from a GatingHierarchy/GatingSet. They are equivalent to the `workFlow`,`add` and `Rm` methods in `flowCore` package. `recompute` method does the actual gating after the gate is added,i.e. calculating the event indices according to the gate definition.

**Usage**

```
gs_pop_add(gs, gate, validityCheck = TRUE, ...)
gs_pop_remove(gs, node, ...)
```

**Arguments**

gs	A GatingSet
gate	A flowCore::filter or a list of flowCore::filters or logical vectors to be added to the GatingSet. when logical vectors, they represent the indices of events to be included in the populations. It can be global that represents the index to the original full events or local index that is relative to the parent population cell events. See examples for more details.
validityCheck	logical whether to check the consistency of tree structure across samples. default is TRUE. Can be turned off when speed is prefered to the robustness.
...	some other arguments to specify how the gates are added to the gating tree. <ul style="list-style-type: none"> <li>names a character vector of length four,which specifies the population names resulted by adding a quadGate.The order of the names is clock-wise starting from the top left quadrant population.</li> <li>parent a character scalar to specify the parent node name where the new gate to be added to, by default it is NULL,which indicates the root node</li> <li>name a character scalar to specify the node name of population that is generated by the gate to be added.</li> <li>recompute a logical flag</li> <li>negated: a logical scalar to specify whether the gate is negated,which means the the population outside of the gate will be kept as the result population. It is FALSE by default.</li> </ul>
node	A character identifies the population node in a GatingHierarchy or GatingSet to remove

**Value**

GatingSet method returns a GatingSet object with just root node. add method returns a population node ID (or four population node IDs when adding a quadGate) that uniquely identify the population node within a GatingHierarchy.

**See Also**

[GatingSet-class](#)

**Examples**

```
## Not run:
library(flowCore)
data(GvHD)
#select raw flow data
fs<-GvHD[1:3]
```

```

#transform the raw data
  tf <- transformList(colnames(fs[[1]])[3:6], asinh, transformationId="asinh")
  fs_trans<-transform(fs,tf)

#add transformed data to a gatingset
  gs <- GatingSet(fs_trans)
  gs
  gs_get_pop_paths(gs[[1]]) #only contains root node

#add one gate
  rg <- rectangleGate("FSC-H"=c(200,400), "SSC-H"=c(250, 400),
    filterId="rectangle")

  nodeID<-gs_pop_add(gs, rg)#it is added to root node by default if parent is not specified
  nodeID
  gs_get_pop_paths(gs[[1]]) #the second population is named after filterId of the gate

#add a quadGate
  qg <- quadGate("FL1-H"=2, "FL2-H"=4)
  nodeIDs<-gs_pop_add(gs,qg,parent="rectangle")
  nodeIDs #quadGate produces four population nodes
  gs_get_pop_paths(gs[[1]]) #population names are named after dimensions of gate if not specified

#add a boolean Gate
  bg<-booleanFilter(`CD15 FITC-CD45 PE+|CD15 FITC+CD45 PE-`)
  bg
  nodeID2<-gs_pop_add(gs, bg, parent="rectangle")
  nodeID2
  gs_get_pop_paths(gs[[1]])
#do the actual gating
  recompute(gs)

#plot one gate for one sample
  autoplot(gs[[1]],"rectangle")
  autoplot(gs[[1]],nodeID) #may be smoothed automatically if there are not enough events after gating

#plot gates across samples using lattice plot
  autoplot(gs,nodeID)
#plot all gates for one sample
  autoplot(gs[[1]])#boolean gate is skipped by default
  autoplot(gs[[1]],bool=TRUE)

#plot the gating hierarchy
  plot(gs[[1]])
#remove one node causing the removal of all the descendants
  gs_pop_remove('rectangle', gs = gs)
  gs_get_pop_paths(gs[[1]])

#add logical vectors as gate
  lg <- sapply(sampleNames(gs), function(sn){
    gh <- gs[[sn]]
    dat <- exprs(gh_pop_get_data(gh, "cd3+"))#get events data matrix for this sample at cd3+
  })

```

```

  vec <- dat[, "FSC-A"] > 1e4 & data[, "SSC-A"] > 1e5
  vec
  })
  gs_pop_add(gs, lg, name = "new_bool", parent = "cd3+")

## End(Not run)

```

---

**gs\_pop\_get\_count\_fast** *Return a table of population statistics for all populations in a GatingHierarchy/GatingSet or the population proportions or the total number of events of a node (population) in a GatingHierarchy*

---

## Description

`gs_pop_get_count_fast` is more useful than `getPop`. Returns a table of population statistics for all populations in a `GatingHierarchy`/`GatingSet`. Includes the `xml` counts, `openCyto` counts and frequencies.

## Usage

```

gs_pop_get_count_fast(
  x,
  statistic = c("count", "freq"),
  xml = FALSE,
  subpopulations = NULL,
  format = c("long", "wide"),
  path = "full",
  ...
)
gs_pop_get_count_with_meta(x, ...)

```

## Arguments

<code>x</code>	a <code>GatingSet</code> or <code>GatingSetList</code>
<code>statistic</code>	character specifies the type of population statistics to extract.(only valid when format is "wide"). Either "freq" or "count" is currently supported.
<code>xml</code>	logical indicating whether the statistics come from <code>xml</code> (if parsed from <code>xml</code> workspace) or from <code>openCyto</code> .
<code>subpopulations</code>	character vector to specify a subset of populations to return. (only valid when format is "long")
<code>format</code>	character value of <code>c("wide", "long")</code> specifying whether to organize the output in long or wide format
<code>path</code>	character see <a href="#">gs_get_pop_paths</a>
<code>...</code>	additional arguments passed to <code>gs_pop_get_count_fast</code>

## Details

`gs_pop_get_count_fast` returns a table population statistics for all populations in the gating hierarchy. The output is useful for verifying that the import was successful, if the xml and openCyto derived counts don't differ much (i.e. if they have a small coefficient of variation.) for a GatingSet, returns a matrix of proportions for all populations and all samples

## Value

`gs_pop_get_count_fast` returns a `data.frame` with columns for the population name, xml derived counts, openCyto derived counts, and the population proportions (relative to their parent population).

a `data.table` of merged population statistics with sample metadata.

## See Also

[gs\\_get\\_pop\\_paths](#)

## Examples

```
## Not run:
#gh is a GatingHierarchy
gs_pop_get_count_fast(gh);
gh_pop_get_stats(gh,gs_get_pop_paths(gh,tsort=T)[5])

#gs is a GatingSet
gs_pop_get_count_fast(gs)
#optionally output in long format as a data.table
gs_pop_get_count_fast(gs, format = "long", path = "auto")
#only get stats for a subset of populations
gs_pop_get_count_fast(gs, format = "long", subpopulations = gs_get_pop_paths(gs)[4:6])

## End(Not run)
## Not run:
#G is a GatingSetList
stats = gs_pop_get_count_with_meta(G)

## End(Not run)
```

---

<code>gs_pop_get_gate</code>	<i>Return the flowCore gate definition associated with a node in a GatingHierarchy/GatingSet.</i>
------------------------------	---------------------------------------------------------------------------------------------------

---

## Description

Return the flowCore gate definition object associated with a node in a GatingHierarchy or GatingSet object.

**Usage**

```
gh_pop_get_gate(obj, y)
gs_pop_get_gate(obj, y)
```

**Arguments**

obj	A GatingHierrachy or GatingSet
y	A character the name or full(/partial) gating path of the node of interest.

**Value**

A gate object from flowCore. Usually a polygonGate, but may be a rectangleGate. Boolean gates are represented by a "BooleanGate" S3 class. This is a list boolean gate definition that references populations in the GatingHierarchy and how they are to be combined logically. If obj is a GatingSet, assuming the trees associated with each GatingHierarchy are identical, then this method will return a list of gates, one for each sample in the GatingSet corresponding to the same population indexed by y.

**See Also**

[gh\\_pop\\_get\\_data](#) [gs\\_get\\_pop\\_paths](#)

**Examples**

```
## Not run: #gh is a GatingHierarchy
gh_pop_get_gate(gh, "CD3") #return the gate for the fifth node in the tree, but fetch it by name.
#G is a GatingSet
gs_pop_get_gate(G, "CD3") #return a list of gates for the fifth node in each tree

## End(Not run)
```

gs\_pop\_get\_gs                    *subset gs by population node*

**Description**

Basically it returns a new GatingSet with only the subtree of the given population node

**Usage**

```
gs_pop_get_gs(gs, pop)
```

**Arguments**

gs	GatingSet
pop	the population node that will become the new root node

**Value**

a new GatingSet that share the underlying events data

---

gs_pop_get_parent	<i>Return the name of the parent population or a list of child populations of the current population in the GatingHierarchy</i>
-------------------	---------------------------------------------------------------------------------------------------------------------------------

---

**Description**

Returns the name of the parent population or a character/numeric vector of all the children of the current population in the given GatingHierarchy

**Usage**

```
gs_pop_get_parent(obj, y, ...)
gh_pop_get_parent(obj, y, ...)
gs_pop_get_children(obj, y, showHidden = TRUE, ...)
gh_pop_get_children(obj, y, showHidden = TRUE, ...)
```

**Arguments**

obj	A GatingHierarchy
y	a character/numeric the name or full(/partial) gating path or node indices of the node / population.
...	other arguments passed to <a href="#">gs_get_pop_paths</a> methods
showHidden	logical whether to include the hidden children nodes.

**Value**

`gs_pop_get_parent` returns a character vector, the name of the parent population. `gs_pop_get_children` returns a character or numeric vector of the node names or node indices of the child nodes of the current node. An empty vector if the node has no children.

**See Also**

[gs\\_get\\_pop\\_paths](#)

## Examples

```
## Not run:
# G is a GatingHierarchy
# return the name of the parent of the fifth node in the hierarchy.
gs_pop_get_parent(G,gs_get_pop_paths(G[[1]])[5])
n<-gs_get_pop_paths(G,tsort=T)[4]
#Get the names of the child nodes of the 4th node in this gating hierarchy.
gs_pop_get_children(G,n)
#Get the ids of the child nodes
gs_pop_get_children(G,4)

## End(Not run)
```

---

gs_pop_get_stats	<i>Extract stats from populations(or nodes)</i>
------------------	-------------------------------------------------

---

## Description

Extract stats from populations(or nodes)

## Usage

```
gs_pop_get_stats(x, ...)

gh_pop_get_stats(
  x,
  nodes = NULL,
  type = "count",
  xml = FALSE,
  inverse.transform = FALSE,
  stats.fun.arg = list(),
  ...
)
```

## Arguments

x	a GatingSet or GatingHierarchy
...	arguments passed to <a href="#">gs_get_pop_paths</a> method.
nodes	the character vector specifies the populations of interest. default is all available nodes
type	the character vector specifies the type of pop stats or a function used to compute population stats. when character, it is expected to be either "count" or "percent". Default is "count" (total number of events in the populations). when a function, it takes a flowFrame object through 'fr' argument and return the stats as a named vector.
xml	whether to extract xml stats or openCyto stats

```

inverse.transform
logical flag . Whether inverse transform the data before computing the stats.

stats.fun.arg a list of arguments passed to ‘type‘ when ‘type‘ is a function.

```

### Value

a data.table that contains stats values (if MFI, for each marker per column) along with ‘pop’ column and ‘sample’ column (when used on a ‘GatingSet’)

### Examples

```

## Not run:
dataDir <- system.file("extdata", package="flowWorkspaceData")
suppressMessages(gs <- load_gs(list.files(dataDir, pattern = "gs_manual", full = TRUE)))

# get stats all nodes
dt <- gs_pop_get_stats(gs) #default is "count"

nodes <- c("CD4", "CD8")
gs_pop_get_stats(gs, nodes, "percent")

# pass a build-in function
gs_pop_get_stats(gs, nodes, type = pop.MFI)

# compute the stats based on the raw data scale
gs_pop_get_stats(gs, nodes, type = pop.MFI, inverse.transform = TRUE)

# supply user-defined stats fun
pop.quantiles <- function(fr){
  chnls <- colnames(fr)
  res <- matrixStats::colQuantiles(exprs(fr), probs = 0.75)
  names(res) <- chnls
  res
}
gs_pop_get_stats(gs, nodes, type = pop.quantiles)

## End(Not run)

```

---

### gs\_pop\_get\_stats\_tfilter

*Extract stats from populations(or nodes) within a restricted time window*

---

### Description

Extract stats from populations(or nodes) within a restricted time window

**Usage**

```
gs_pop_get_stats_tffilter(x, ...)

gh_pop_get_stats_tffilter(
  x,
  nodes = NULL,
  type = c("Count", "Frequency"),
  inverse.transform = FALSE,
  stats.fun.arg = list(),
  tfilter = NULL,
  path = c("full", "auto"),
  ...
)
```

**Arguments**

x	GatingSet or GatingHierarchy
nodes	the character vector specifies the populations of interest. default is all available nodes
type	the character vector specifies the type of pop stats or a function used to compute population stats. When it is a character, it is expected to be either "Count" or "Frequency". Default is "Count" (total number of events in the populations). When it is a function, it takes a flowFrame object through the 'fr' argument and returns the stats as a named vector.
inverse.transform	logical flag . Whether to inverse transform the data before computing the stats.
stats.fun.arg	a list of arguments passed to 'type' when 'type' is a function.
tfilter	Either a list (tmin, tmax) specifying the minimum and maximum of a the time window filter or a GatingHierarchy, whose minimum and maximum time will be used to determine the window. For both x and the reference GatingHierarchy in tfilter, the only channels that will match this filter are "Time" or "time" and the filter will be applied to each event such that only events with time value t where tmin <= t <= tmax will be evaluated.
path, ...	arguments passed to 'gh_get_pop_paths()'

---

gs_pop_set_gate	<i>update the gate</i>
-----------------	------------------------

---

**Description**

update the population node with a flowCore-compatible gate object

**Usage**

```
gh_pop_set_gate(obj, y, value, negated = FALSE, ...)
gs_pop_set_gate(obj, y, value, ...)
```

**Arguments**

obj	GatingHierarchy or GatingSet
y	character node name or path
value	filter or filterList or list of filter objects
negated	logical see <a href="#">add</a>
...	other arguments

**Details**

Usually [recompute](#) is followed by this call since updating a gate doesn't re-calculating the cell events within the gate automatically. see [filterObject](#) for the gate types that are currently supported.

**Examples**

```
## Not run:
rg1 <- rectangleGate("FSC-H"=c(200,400), "SSC-H"=c(250, 400), filterId="rectangle")
rg2 <- rectangleGate("FSC-H"=c(200,400), "SSC-H"=c(250, 400), filterId="rectangle")
flist <- list(rg1,rg2)
names(flist) <- sampleNames(gs[1:2])
gs_pop_set_gate(gs[1:2], "lymph", flist)
recompute(gs[1:2], "lymph")

## End(Not run)
```

gs\_pop\_set\_name

*Update the name of one node in a gating hierarchy/GatingSet.*

**Description**

gh\_pop\_set\_name/gs\_pop\_set\_name update the name of one node in a gating hierarchy/GatingSet.

**Usage**

```
gh_pop_set_name(x, y, value)
gs_pop_set_name(x, y, value)
```

**Arguments**

x	GatingHierarchy
y	pop name/path
value	A character the name of the node

### Examples

```
## Not run:
# G is a GatingHierarchy
gs_get_pop_paths(G[[1]])#return node names
gh_pop_set_name(G,"L","lymph")

## End(Not run)
```

---

gs\_pop\_set\_visibility  *hide/unhide a node*

---

### Description

hide/unhide a node

### Usage

```
gh_pop_set_visibility(x, y, value)

gs_pop_set_visibility(x, y, value)
```

### Arguments

x	GatingHierarchy object
y	character node name or path
value	TRUE/FALSE to indicate whether to hide a node

### Examples

```
## Not run:
gh_pop_set_visibility(gh, 4, FALSE) # hide a node
gh_pop_set_visibility(gh, 4, TRUE) # unhide a node

## End(Not run)
```

---

gs\_remove\_redundant\_channels

*Remove the channels from flow data that are not used by gates*

---

### Description

Removing these redundant channels can help standardize the channels across different GatingSet objects and make them mergable.

**Usage**

```
gs_remove_redundant_channels(gs, ...)
```

**Arguments**

gs	a GatingSet
...	other arguments passed to gs_get_pop_paths method

**Value**

a new GatingSet object that has redundant channels removed. Please note that this new object shares the same reference (or external pointers) with the original GatingSets.

**Examples**

```
## Not run:
gs_new <- gs_remove_redundant_channels(gs)

## End(Not run)
```

---

gs\_remove\_redundant\_nodes

*Remove the terminal leaf nodes that make the gating trees to be different from one another.*

---

**Description**

It is usually called after [gs\\_split\\_by\\_tree](#) and [gs\\_check\\_redundant\\_nodes](#). The operation is done in place through external pointers which means all the original GatingSets are modified.

**Usage**

```
gs_remove_redundant_nodes(x, toRemove)
```

**Arguments**

x	GatingSet or list of groups(each group is a list of 'GatingSet'). When it is a list, it is usually the outcome from <a href="#">gs_split_by_tree</a> .
toRemove	list of the node sets to be removed. its length must equals to the length of 'x'. When x is a list, toRemove is usually the outcome from <a href="#">gs_check_redundant_nodes</a> .

## Examples

```
## Not run:
gslist <- list(gs1, gs2, gs3, gs4, gs5)
gs_groups <- gs_split_by_tree(gslist)
toRm <- gs_check_redundant_nodes(gs_groups)
gs_remove_redundant_nodes(gs_groups, toRm)

#Now they can be merged into a single GatingSetList.
#Note that the original gs objects are all modified in place.
GatingSetList(gslist)

## End(Not run)
```

---

gs\_split\_by\_channels *split GatingSets into groups based on their flow channels*

---

## Description

Sometime it is gates are defined on the different dimensions across different GatingSets, (e.g. ‘FSC-W’ or ‘SSC-H’ may be used for Y axis for cytokines) These difference in dimensions may not be critical since they are usually just used for visualization(instead of thresholding events) But this prevents the gs from merging because they may not be collected across batches Thus we have to separate them if we want to visualize the gates.

## Usage

```
gs_split_by_channels(x)
```

## Arguments

x	a list of GatingSets
---	----------------------

## Examples

```
## Not run:
gslist <- list(gs1, gs2, gs3, gs4, gs5)
gs_groups <- gs_split_by_channels(gslist)

## End(Not run)
```

---

gs_split_by_tree	<i>split GatingSets into groups based on their gating schemes Be careful that the splitted resluts still points to the original data set!!</i>
------------------	------------------------------------------------------------------------------------------------------------------------------------------------

---

## Description

It allows isomorphism in Gating tree and ignore difference in hidden nodes i.e. tree is considered to be the same as long as `gs_get_pop_paths(gh, path = "auto", showHidden = F)` returns the same set

## Usage

```
gs_split_by_tree(x)
```

## Arguments

x	a list of GatingSets or one GatingSet
---	---------------------------------------

## Value

when x is a GatingSet, this function returns a list of sub-GatingSets When x is a list of GatingSets, it returns a list of list, each list itself is a list of GatingSets, which share the same gating tree.

## Examples

```
## Not run:
gslist <- list(gs1, gs2, gs3, gs4, gs5)
gs_groups <- gs_split_by_tree(gslist)

## End(Not run)
```

---

gs_update_channels	<i>Update the channel information of a GatingSet (c++ part)</i>
--------------------	-----------------------------------------------------------------

---

## Description

It updates the channels stored in gates, compensations and transformations based on given mapping between the old and new channel names.

## Usage

```
gs_update_channels(gs, map, all = TRUE)
```

**Arguments**

gs	a GatingSet object
map	data.frame contains the mapping from old (case insensitive) to new channel names Note: Make sure to remove the '<' or '>' characters from 'old' name because the API tries to only look at the raw channel name so that the gates with both prefixed and non-prefixed names could be updated.
all	logical whether to update the flow data as well

**Value**

when 'all' is set to TRUE, it returns a new GatingSet but it still shares the same underlying c++ tree structure with the original GatingSet otherwise it returns nothing (less overhead.)

**Examples**

```
## Not run:
##this will update both "Qdot 655-A" and "<Qdot 655-A>"
gs <- gs_update_channels(gs, map = data.frame(old = c("Qdot 655-A")
                                              , new = c("QDot 655-A")
                                              )
)
## End(Not run)
```

identifier-methods      *Retrieve/replace the GUID of a GatingSet or GatingSetList*

**Description**

Retrieve or replace the GUID (globally unique identifier) for a [GatingSet](#) or [GatingSetList](#)

**Usage**

```
identifier(object)

## S4 replacement method for signature 'GatingSet,ANY'
identifier(object) <- value

## S4 replacement method for signature 'GatingSetList,character'
identifier(object) <- value
```

**Arguments**

object	a GatingSet or GatingSetList
value	string

---

is_tiledb_support	<i>check whether cytolib is build with tiledb support</i>
-------------------	-----------------------------------------------------------

---

**Description**

check whether cytolib is build with tiledb support

**Usage**

```
is_tiledb_support()
```

**Value**

TRUE or FALSE

---

keyword	<i>Retrieve a specific keyword for a specific sample in a GatingHierarchy or or set of samples in a GatingSet or GatingSetList</i>
---------	------------------------------------------------------------------------------------------------------------------------------------

---

**Description**

Retrieve a specific keyword for a specific sample in a GatingHierarchy or or set of samples in a GatingSet or GatingSetList

**Usage**

```
## S4 method for signature 'GatingHierarchy,character'
keyword(object, keyword)

## S4 method for signature 'GatingHierarchy,missing'
keyword(object, keyword = "missing", ...)
```

**Arguments**

object	GatingHierarchy or GatingSet or GatingSetList
keyword	character specifying keyword name. When missing, extract all keywords.
...	other arguments passed to <a href="#">keyword-methods</a>

**Details**

See keyword in Package ‘flowCore’

**See Also**

[keyword-methods](#)

## Examples

```
## Not run:
# get all the keywords from all samples
keyword(G)
# get all the keywords from one sample
keyword(G[[1]])
# filter the instrument setting
keyword(G[[1]], compact = TRUE)
# get single keyword from all samples
keyword(G, "FILENAME")
# get single keyword from one sample
keyword(G[[1]], "FILENAME")

## End(Not run)
```

---

keyword-mutators

*Methods to alter keywords in cytoframe, cytoset, GatingHierarchy, or GatingSet objects*

---

## Description

These methods allow for direct insertion, deletion, or renaming of keywords in [cytoframe](#), [cytoset](#), [GatingHierarchy](#), or [GatingSet](#) objects.

## Usage

```
cf_keyword_insert(cf, keys, values)

cf_keyword_delete(cf, keys)

cf_keyword_rename(cf, old_keys, new_keys)

cf_keyword_set(cf, keys, values)

cs_keyword_insert(cs, keys, values)

cs_keyword_delete(cs, keys)

cs_keyword_rename(cs, old_keys, new_keys)

cs_keyword_set(cs, keys, values)

gh_keyword_insert(gh, keys, values)

gh_keyword_delete(gh, keys)

gh_keyword_rename(gh, old_keys, new_keys)
```

```
gh_keyword_set(gh, keys, values)

gs_keyword_insert(gs, keys, values)

gs_keyword_delete(gs, keys)

gs_keyword_rename(gs, old_keys, new_keys)

gs_keyword_set(gs, keys, values)
```

## Arguments

cf	a <a href="#">cytoframe</a>
keys	the keyword names to insert/delete/replace – single value or vector
values	the values to associate with the supplied keywords – single value or vector of sample length as keys
old_keys	the old keyword name (for renaming)
new_keys	the new keyword name (for renaming)
gh	a <a href="#">GatingHierarchy</a>
gs	a <a href="#">GatingSet</a>

## Details

Each of the methods taking two character vectors (keys/values or old\_keys/new\_keys) will also accept a single named vector for flexibility in usage.

For the functions that take a vector of keys and a vector of values (the `keyword_insert` and `keyword_set` functions), the names of this vector should be the keys to which the values of the vector will be assigned.

For the `keyword_rename` functions, the names of this vector should be the existing keyword names (`old_keys`) while the values should be the replacement keyword names (`new_keys`).

See examples for details

## Examples

```
library(flowCore)
data(GvHD)
cs <- flowSet_to_cytoset(GvHD[1:2])

keys <- c("CYTNUM", "CREATOR")

# Values before changes
keyword(cs, keys)

# Set two keyword values using separate key and values vectors
values <- c("E3598", "CELLQuest 3.4")
cs_keyword_set(cs, keys, values)
```

```

# Values after changes
keyword(cs, keys)

# Change the values again using a single named vector
values <- c("E3599", "CELLQuest 3.5")
names(values) <- keys
cs_keyword_set(cs, values)

# Values after changes
keyword(cs, keys)

```

---

<b>lapply-methods</b>	<i>apply FUN to each sample (i.e. GatingHierarchy or cytoframe) in a GatingSet or cytoset</i>
-----------------------	-----------------------------------------------------------------------------------------------

---

## Description

sample names are used for names of the returned list

## Usage

```
lapply(X, FUN, ...)
```

## Arguments

X	GatingSet or cytoset
FUN	function to be applied to each sample in 'GatingSet' or 'cytoset'
...	other arguments to be passed to 'FUN'

---

<b>length</b>	<i>Methods to get the length of a GatingSet</i>
---------------	-------------------------------------------------

---

## Description

Return the length of a GatingSet or GatingSetList object (number of samples).

## Usage

```

## S4 method for signature 'GatingSet'
length(x)

## S4 method for signature 'GatingSet'
show(object)

```

**Arguments**

x	GatingSet
object	object

---

load_cytoframe	<i>Load the cytoframe from disk</i>
----------------	-------------------------------------

---

**Description**

Load the cytoframe from disk

**Usage**

```
load_cytoframe(uri, on_disk = TRUE, readonly = on_disk, ctx = .cytoctx_global)
```

**Arguments**

uri	path to the cytoframe file
on_disk	logical flag indicating whether to keep the data on disk and load it on demand. Default is TRUE.
readonly	logical flag indicating whether to open h5 data as readonly. Default is TRUE. And it is valid when on_disk is set to true.
ctx	cytoctx object, see [cytoctx] for details

**See Also**

Other cytoframe/cytoset IO functions: [cf\\_get\\_uri\(\)](#), [cf\\_write\\_disk\(\)](#), [cf\\_write\\_h5\(\)](#), [cf\\_write\\_tile\(\)](#), [cs\\_get\\_uri\(\)](#), [load\\_cytoframe\\_from\\_fcs\(\)](#), [load\\_cytoset\\_from\\_fcs\(\)](#)

---

load_cytoframe_from_fcs	<i>Read a single FCS file in to a cytoframe</i>
-------------------------	-------------------------------------------------

---

**Description**

Similar to [read.FCS](#), this takes a filename for a single FCS file and returns a cytoframe.

## Usage

```
load_cytoframe_from_fcs(
  filename,
  transformation = "linearize",
  which.lines = NULL,
  alter.names = FALSE,
  column.pattern = NULL,
  invert.pattern = FALSE,
  decades = 0,
  is_h5 = NULL,
  backend = get_default_backend(),
  uri = NULL,
  h5_filename = NULL,
  min.limit = NULL,
  truncate_max_range = TRUE,
  dataset = NULL,
  emptyValue = TRUE,
  num_threads = 1,
  ignore.text.offset = FALSE,
  text.only = FALSE
)
```

## Arguments

<code>filename</code>	The filename of the single FCS file to be read
<code>transformation</code>	A character string that defines the type of transformation. Valid values are <code>linearize</code> (default), <code>linearize-with-PnG-scaling</code> , or <code>scale</code> . The <code>linearize</code> transformation applies the appropriate power transform to the data. The <code>linearize-with-PnG-scaling</code> transformation applies the appropriate power transform for parameters stored on log scale, and also a linear scaling transformation based on the "gain" (FCS \$PnG keywords) for parameters stored on a linear scale. The <code>scale</code> transformation scales all columns to $[0, 10^{decades}]$ , defaulting to <code>decades = 0</code> as in the FCS4 specification. A logical can also be used: <code>TRUE</code> is equal to <code>linearize</code> and <code>FALSE</code> (or <code>NULL</code> ) corresponds to no transformation. Also, when the transformation keyword of the FCS header is set to "custom" or "applied", no transformation will be used.
<code>which.lines</code>	Numeric vector to specify the indices of the lines to be read. If it is <code>NULL</code> , all the records are read. If it is of length 1, a random sample of the size indicated by <code>which.lines</code> is read in.
<code>alter.names</code>	Logical indicating whether or not we should rename the columns to valid R names using <code>make.names</code> . The default is <code>FALSE</code> .
<code>column.pattern</code>	An optional regular expression defining parameters we should keep when loading the file. The default is <code>NULL</code> .
<code>invert.pattern</code>	Logical. By default, <code>FALSE</code> . If <code>TRUE</code> , inverts the regular expression specified in <code>column.pattern</code> . This is useful for indicating the channel names that we do not want to read. If <code>column.pattern</code> is set to <code>NULL</code> , this argument is ignored.
<code>decades</code>	When scaling is activated, the number of decades to use for the output.

is_h5	Logical indicating whether the data should be stored in h5 format
h5_filename	String specifying a name for the h5 file if is_h5 is TRUE
min.limit	The minimum value in the data range that is allowed. Some instruments produce extreme artifactual values. The positive data range for each parameter is completely defined by the measurement range of the instrument and all larger values are set to this threshold. The lower data boundary is not that well defined, since compensation might shift some values below the original measurement range of the instrument. This can be set to an arbitrary number or to NULL (the default value), in which case the original values are kept.
truncate_max_range	Logical. Default is TRUE. can be optionally turned off to avoid truncating the extreme positive value to the instrument measurement range, i.e. '\$PnR'.
dataset	The FCS file specification allows for multiple data segments in a single file. Since the output of load_cytoframe_from_cytoset is a single cytoframe we can't automatically read in all available sets. This parameter allows the user to choose one of the subsets for import. Its value should be an integer in the range of available data sets. This argument is ignored if there is only a single data segment in the FCS file.
emptyValue	Logical indicating whether or not to allow empty values for keywords in TEXT segment. It affects how double delimiters are treated. If TRUE, double delimiters are parsed as a pair of start and end single delimiters for an empty value. Otherwise, double delimiters are parsed as one part of the string of the keyword value. The default is TRUE.
num_threads	Integer allowing for parallelization of the parsing operation by specifying a number of threads
ignore.text.offset	Logical indicating whether to ignore the keyword values in TEXT segment when they don't agree with the HEADER. Default is FALSE, which throws the error when such a discrepancy is found. Users can turn it on to ignore the TEXT segment when they are sure of the accuracy of the HEADER segment so that the file still can be read.
text.only	whether to only parse text section of FCS (default is FALSE), it is sometime useful to skip loading data section for the faster loading meta data from FCS <a href="#">read.AnnotatedDataFrame</a> , see details

## Details

The function `load_cytoframe_from_fcs` works with the output of the FACS machine software from a number of vendors (FCS 2.0, FCS 3.0 and List Mode Data LMD). However, the FCS 3.0 standard includes some options that are not yet implemented in this function. If you need extensions, please let us know. The output of the function is an object of class `cytoframe`.

For specifications of FCS 3.0 see <http://www.isac-net.org> and the file `../doc/fcs3.html` in the doc directory of the package.

The `which.lines` arguments allow you to read a subset of the record as you might not want to read the thousands of events recorded in the FCS file. It is mainly used when there is not enough memory to read one single FCS (which probably will not happen). It will probably take more time than reading the entire FCS (due to the multiple disk IO).

**Value**

An object of class [cytoframe](#) that contains the data, the parameters monitored, and the keywords and values saved in the header of the FCS file.

**See Also**

Other cytoframe/cytoSET IO functions: [cf\\_get\\_uri\(\)](#), [cf\\_write\\_disk\(\)](#), [cf\\_write\\_h5\(\)](#), [cf\\_write\\_tile\(\)](#), [cs\\_get\\_uri\(\)](#), [load\\_cytoframe\(\)](#), [load\\_cytoSET\\_from\\_fcs\(\)](#)

---

`load_cytoSET_from_fcs` *Read one or several FCS files in to a cytoSET*

---

**Description**

Similar to [read.flowSet](#), this takes a list of FCS filenames and returns a cytoSET.

**Usage**

```
load_cytoSET_from_fcs(  
  files = NULL,  
  path = ".",  
  pattern = NULL,  
  phenoData = NULL,  
  descriptions,  
  name.keyword,  
  transformation = "linearize",  
  which.lines = NULL,  
  alter.names = FALSE,  
  column.pattern = NULL,  
  invert.pattern = FALSE,  
  decades = 0,  
  is_h5 = NULL,  
  h5_dir = NULL,  
  backend = get_default_backend(),  
  backend_dir = tempdir(),  
  min.limit = NULL,  
  truncate_max_range = TRUE,  
  dataset = NULL,  
  emptyValue = TRUE,  
  num_threads = 1,  
  ignore.text.offset = FALSE,  
  sep = "\t",  
  as.is = TRUE,  
  name,  
  file_col_name = NULL,  
  ...  
)
```

## Arguments

<code>files</code>	Optional character vector with filenames.
<code>path</code>	Directory where to look for the files.
<code>pattern</code>	This argument is passed on to <code>dir</code> , see details.
<code>phenoData</code>	An object of class <code>AnnotatedDataFrame</code> , character or a list of values to be extracted from the <code>cytoframe</code> object, see details.
<code>descriptions</code>	Character vector to annotate the object of class <code>cytoSET</code> .
<code>name.keyword</code>	An optional character vector that specifies which FCS keyword to use as the sample names. If this is not set, the GUID of the FCS file is used for sampleNames, and if that is not present (or not unique), then the file names are used.
<code>transformation</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>which.lines</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>alter.names</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>column.pattern</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>invert.pattern</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>decades</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>is_h5</code>	logical indicating whether the data should be stored in h5 format
<code>h5_dir</code>	String specifying a name for the h5 directory for the h5 files if <code>is_h5</code> is TRUE
<code>min.limit</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>truncate_max_range</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>dataset</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>emptyValue</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>num_threads</code>	Integer allowing for parallelization of the parsing operation by specifying a number of threads
<code>ignore.text.offset</code>	see <code>load_cytoframe_from_fcs</code> for details.
<code>sep</code>	Separator character that gets passed on to <code>read.AnnotatedDataFrame</code> .
<code>as.is</code>	logical that gets passed on to <code>read.AnnotatedDataFrame</code> . This controls the automatic coercion of characters to factors in the <code>phenoData</code> .
<code>name</code>	An optional character scalar used as name of the object.
<code>file_col_name</code>	optionally specify the column name that stores the fcs filename when <code>phenoData</code> is supplied <code>read.AnnotatedDataFrame</code> , see details.
<code>...</code>	Further arguments that get passed on to

## Details

There are four different ways to specify the file from which data is to be imported:

First, if the argument `phenoData` is present and is of class `AnnotatedDataFrame`, then the file names are obtained from its sample names (i.e. row names of the underlying data.frame). Also column name will be generated based on sample names if it is not there. This column is mainly

used by visualization methods in flowViz. Alternatively, the argument `phenoData` can be of class `character`, in which case this function tries to read a `AnnotatedDataFrame` object from the file with that name by calling `read.AnnotatedDataFrame(file.path(path, phenoData), ...{})`.

In some cases the file names are not a reasonable selection criterion and the user might want to import files based on some keywords within the file. One or several keyword value pairs can be given as the `phenoData` argument in form of a named list.

Third, if the argument `phenoData` is not present and the argument `files` is not `NULL`, then `files` is expected to be a character vector with the file names.

Fourth, if neither the argument `phenoData` is present nor `files` is not `NULL`, then the file names are obtained by calling `dir(path, pattern)`.

### Value

An object of class `cytoset`.

### See Also

Other cytoframe/cytoset IO functions: `cf_get_uri()`, `cf_write_disk()`, `cf_write_h5()`, `cf_write_tile()`, `cs_get_uri()`, `load_cytoframe_from_fcs()`, `load_cytoframe()`

---

load\_meta

*Flush/load meta data (keywords, pData, channels/markers) to/from disk (only valid for on-disk cytoset/cytoframe)*

---

### Description

Flush/load meta data (keywords, `pData`, channels/markers) to/from disk (only valid for on-disk cytoset/cytoframe)

### Usage

```
cf_flush_meta(cf)  
  
cf_load_meta(cf)  
  
cs_flush_meta(cs)  
  
cs_load_meta(cs)
```

### Arguments

<code>cf</code>	cytoframe object
<code>cs</code>	cytoset object

---

lock	<i>Lock/Unlock the cytoset/cytoframe by turning on/off its read-only flag</i>
------	-------------------------------------------------------------------------------

---

**Description**

Lock/Unlock the cytoset/cytoframe by turning on/off its read-only flag

**Usage**

cf\_lock(cf)

cf\_unlock(cf)

cs\_lock(cs)

cs\_unlock(cs)

**Arguments**

cf                    cytoframe object

cs                    cytoset object

---

logicleGml2_trans	<i>GatingML2 version of logicle transformation.</i>
-------------------	-----------------------------------------------------

---

**Description**

The only difference from [logicle\\_trans](#) is it is scaled to c(0,1) range.

**Usage**

```
logicleGml2_trans(
  T = 262144,
  M = 4.5,
  W = 0.5,
  A = 0,
  n = 6,
  equal.space = FALSE
)
```

**Arguments**

T, M, W, A        see [logicleGml2](#)

n                    desired number of breaks (the actual number will be different depending on the data range)

equal.space        whether breaks at equal-spaced intervals

## Value

a `logicleGml2` transformation object

## Examples

```

trans.obj <- logicleGml2_trans(equal.space = TRUE)
data <- 1:1e3
brks.func <- trans.obj[["breaks"]]
brks <- brks.func(data)
brks # logicle space displayed at raw data scale
#transform it to verify the equal-spaced breaks at transformed scale
print(trans.obj[["transform"]](brks))

```

`logicle_trans` *logicle transformation.*

## Description

Used for construct logicle transform object.

## Usage

```
logicle_trans(..., n = 6, equal.space = FALSE)
```

## Arguments

...	arguments passed to <code>logicTransform</code> .
<code>n</code>	desired number of breaks (the actual number will be different depending on the data range)
<code>equal.space</code>	whether breaks at equal-spaced intervals

## Value

a logicle transformation object

## Examples

```

trans.obj <- logicle_trans(equal.space = TRUE)
data <- 1:1e3
brks.func <- trans.obj[["breaks"]]
brks <- brks.func(data)
brks # logicle space displayed at raw data scale
#transform it to verify the equal-spaced breaks at transformed scale
print(trans.obj[["transform"]](brks))

```

---

`logtGml2_trans`*Gating-ML 2.0 Log transformation.*

---

## Description

Used to construct GML 2.0 flog transformer object.

## Usage

```
logtGml2_trans(t = 262144, m = 4.5, n = 6, equal.space = FALSE)
```

## Arguments

<code>t</code>	top scale value
<code>m</code>	number of decades
<code>n</code>	desired number of breaks (the actual number will be different depending on the data range)
<code>equal.space</code>	whether breaks at equal-spaced intervals

## Details

GML 2.0 standard log transform function constructor. The definition is as in the GML 2.0 standard section 6.2 "parametrized logarithmic transformation – flog" This deviates from standard only in the following way. Before applying the logarithmic transformation, non-positive values are assigned the smallest positive value from the input rather than having undefined values (NA) under the transformation.

## Value

`logtGml2` transformation object

## Examples

```
trans.obj <- logtGml2_trans(t = 1e3, m = 1, equal.space = TRUE)
data <- 1:1e3
brks.func <- trans.obj[["breaks"]]
brks <- brks.func(data)
brks # fasinh space displayed at raw data scale

#transform it to verify it is equal-spaced at transformed scale
trans.func <- trans.obj[["transform"]]
brks.trans <- trans.func(brks)
brks.trans
```

---

markernames	<i>Get/set the column(channel) or marker names</i>
-------------	----------------------------------------------------

---

## Description

It simply calls the methods for the underlying flow data (flowSet/ncdfFlowSet/ncdfFlowList).

## Usage

```
## S4 method for signature 'GatingHierarchy'
markernames(object)

## S4 replacement method for signature 'GatingHierarchy'
markernames(object) <- value

## S4 method for signature 'GatingHierarchy'
colnames(x, do.NULL = "missing", prefix = "missing")

## S4 replacement method for signature 'GatingHierarchy'
colnames(x) <- value
```

## Arguments

value	named character vector for markernames<-; regular character vector for colnames<-
.	
x, object	GatingHierarchy/GatingSet/GatingSetList
do.NULL, prefix	not used.

## Examples

```
## Not run:

markers.new <- c("CD4", "CD8")
chnls <- c("<B710-A>", "<R780-A>")
names(markers.new) <- chnls
markernames(gs) <- markers.new

chnls <- colnames(gs)
chnls.new <- chnls
chnls.new[c(1,4)] <- c("fsc", "ssc")
colnames(gs) <- chnls.new

## End(Not run)
```

---

<code>merge_list_to_gs</code>	<i>Merge a list of GatingSets into a single GatingSet</i>
-------------------------------	-----------------------------------------------------------

---

### Description

It also checks the consistency of the cyto data and gates.

### Usage

`merge_list_to_gs(x, ...)`

### Arguments

<code>x</code>	a list of GatingSets
<code>...</code>	other arguments (not used)

---

<code>ncFlowSet</code>	<i>Fetch the flowData object associated with a GatingSet .</i>
------------------------	----------------------------------------------------------------

---

### Description

Deprecated by `flowData` method

Deprecated by `flowData` method

---

<code>nodeflags</code>	<i>The flags of gate nodes</i>
------------------------	--------------------------------

---

### Description

`gh_pop_is_gated` checks if a node is already gated. `gh_pop_is_negated` checks if a node is negated. `gh_pop_is_hidden` checks if a node is hidden.

### Usage

```
gh_pop_is_gated(obj, y)
gh_pop_is_negated(obj, y)
gh_pop_is_hidden(obj, y)
gh_pop_is_bool_gate(obj, y)
```

### Arguments

<code>obj</code>	GatingHierarchy
<code>y</code>	node/gating path

---

<code>openWorkspace</code>	<i>It is now moved along with entire flowJo parser to CytoML package</i>
----------------------------	--------------------------------------------------------------------------

---

**Description**

It is now moved along with entire flowJo parser to CytoML package

**Usage**

```
openWorkspace(file, ...)
```

**Arguments**

<code>file</code>	xml file
<code>...</code>	other arguments

---

<code>pData-methods</code>	<i>read/set pData of flow data associated with GatingHierarchy, GatingSet, or GatingSetList</i>
----------------------------	-------------------------------------------------------------------------------------------------

---

**Description**

Accessor method that gets or replaces the pData of the flowset/ncdfFlowSet object in a GatingHierarchy, GatingSet, or GatingSetList

**Usage**

```
pData(object)  
pData(object) <- value
```

**Arguments**

<code>object</code>	GatingSet or GatingSetList
<code>value</code>	<code>data.frame</code> The replacement of pData for flowSet or ncdfFlowSet object

**Value**

a `data.frame`

---

plot-methods

*plot a gating tree*

---

## Description

Plot a tree/graph representing the GatingHierarchy

## Usage

```
plot(x, y, ...)
```

## Arguments

x GatingHierarchy or GatingSet. If GatingSet, the first sample will be used to extract gating tree.

y missing or character specifies.

... other arguments:

- boolean: TRUE | FALSE logical specifying whether to plot boolean gate nodes. Defaults to FALSE.
- showHidden: TRUE | FALSE logical whether to show hidden nodes
- layout: See [layoutGraph](#) in package Rgraphviz
- width: See [layoutGraph](#) in package Rgraphviz
- height: See [layoutGraph](#) in package Rgraphviz
- fontsize: See [layoutGraph](#) in package Rgraphviz
- labelfontsize: See [layoutGraph](#) in package Rgraphviz
- fixedsize: See [layoutGraph](#) in package Rgraphviz

## Examples

```
## Not run:
#ggs is a GatingSet
plot(gs) # the same as plot(gs[[1]])
#plot a subtree rooted from 'CD4'
plot(gs, "CD4")

## End(Not run)
```

---

plotGate-methods-defunct

*Plot gates and associated cell population contained in a GatingHierarchy or GatingSet*

---

**Description**

**Important:** The plotGate methods are now defunct and gates should instead be plotted using the [autoplots](#) method from the ggcryo package. The plotGate documentation has been left here to ease the transition.

When applied to a GatingHierarchy, `arrange` is set as TRUE, then all the gates associated with it are plotted as different panel on the same page. If `arrange` is FALSE, then it plots one gate at a time. By default, `merge` is set as TRUE, plot multiple gates on the same plot when they share common parent population and axis. When applied to a GatingSet, if `lattice` is TRUE, it plots one gate (multiple samples) per page, otherwise, one sample (with multiple gates) per page.

**Usage**

```
plotGate(x, y, ...)
```

**Arguments**

<code>x</code>	<a href="#">GatingSet</a> or <a href="#">GatingHierarchy</a> object
<code>y</code>	character the node name or full(/partial) gating path or numeric representing the node index in the GatingHierarchy. or missing which will plot all gates and one gate per page. It is useful for generating plots in a multi-page pdf. Nodes can be accessed with <a href="#">gs_get_pop_paths</a> .
<code>...</code>	<ul style="list-style-type: none"> <li>• <code>bool</code> logical specifying whether to plot boolean gates.</li> <li>• <code>arrange.main</code> character The title of the main page of the plot. Default is the sample name. Only valid when <code>x</code> is GatingHierarchy</li> <li>• <code>arrange</code> logical indicating whether to arrange different populations/nodes on the same page via <code>arrangeGrob</code> call.</li> <li>• <code>merge</code> logical indicating whether to draw multiple gates on the same plot if these gates share the same parent population and same x,y dimensions/parameters;</li> <li>• <code>projections</code> list of character vectors used to customize x,y axis. By default, the x,y axis are determined by the respective gate parameters. The elements of the list are named by the population name or path (see <code>y</code>). Each element is a pair of named character specifying the channel name(or marker name) for x, y axis. Short form of channel or marker names (e.g. "APC" or "CD3") can be used as long as they can be uniquely matched to the dimensions of flow data. For example, <code>projections = list("lymph" = c(x = "SSC-A", y = "FSC-A"), "CD3" = c(x = "CD3", y = "SSC-A"))</code></li> <li>• <code>par.settings</code> list of graphical parameters passed to <a href="#">lattice</a>;</li> </ul>

- gpar list of grid parameters passed to `grid.layout`;
- lattice logical deprecated;
- formula formula a formula passed to `xyplot` function of `flowViz`, by default it is NULL, which means the formula is generated according to the x,y parameters associated with gate.
- cond character the conditioning variable to be passed to lattice plot.
- overlayNode names. These populations are plotted on top of the existing gates(defined by y argument) as the overlaid dots.
- overlay.symbolA named (lattice graphic parameter) list that defines the symbol color and size for each overlaid population. If not given, we automatically assign the colors.
- keyLattice legend paraemter for overlay symbols.
- default.y character specifying y channel for `xyplot` when plotting a 1d gate. Default is "SSC-A" and session-wise setting can be stored by `'flowWorkspace.par.set("plotGate", list(default.y = "FSC-A"))'`
- type character either "xyplot" or "densityplot". Default is "xyplot" and session-wise setting can be stored by `'flowWorkspace.par.set("plotGate", list(type = "xyplot"))'`
- fitGate used to disable behavior of plotting the gate region in 1d densityplot. Default is FALSE and session-wise setting can be stored by `'flowWorkspace.par.set("plotGate", list(fitGate = FALSE))'`
- strip ligcal specifies whether to show pop name in strip box,only valid when x is `GatingHierarchy`
- strip.texteither "parent" (the parent population name) or "gate "(the gate name).
- raw.scale logical whether to show the axis in raw(untransformed) scale. Default is TRUE and can be stored as session-wise setting by `'flowWorkspace.par.set("plotGate", list(raw.scale = TRUE))'`
- xlim, ylim character can be either "instrument" or "data" which determines the x, y axis scale either by instrument measurement range or the actual data range. or numeric which specifies customized range. They can be stored as session-wise setting by `'flowWorkspace.par.set("plotGate", list(xlim = "instrument"))'`
- ...
  - path A character or numeric scalar passed to `gs_get_pop_paths` method (used to control how the gating/node path is displayed)
  - ... The other additional arguments to be passed to `xyplot`.

### Value

a `trellis` object if `arrange` is FALSE,

### References

<http://www.rglab.org/>

**Examples**

```
## Not run:  
#G is a GatingHierarchy  
plotGate(G,gs_get_pop_paths(G)[5]);#plot the gate for the fifth node  
  
## End(Not run)
```

---

**pop\_add***Add populations to a GatingHierarchy*

---

**Description**

Add populations to a GatingHierarchy

**Usage**

```
pop_add(gate, gh, ...)  
  
## S3 method for class 'filter'  
pop_add(gate, gh, ...)  
  
## S3 method for class 'filters'  
pop_add(gate, gh, names = NULL, ...)  
  
## S3 method for class 'quadGate'  
pop_add(gate, gh, names = NULL, ...)  
  
## S3 method for class 'logical'  
pop_add(gate, gh, parent, name, recompute, cluster_method_name = NULL, ...)  
  
## S3 method for class 'factor'  
pop_add(gate, gh, name = NULL, ...)  
  
## S3 method for class 'logicalFilterResult'  
pop_add(gate, gh, ...)  
  
## S3 method for class 'multipleFilterResult'  
pop_add(gate, gh, name = NULL, ...)  
  
gh_pop_remove(gh, node, ...)
```

**Arguments**

gate	a gate object that extends flowCore::filter or flowCore::filters
gh	GatingHierarchy
...	other arguments

names	a character vector of length four, which specifies the population names resulted by adding a quadGate. The order of the names is clock-wise starting from the top left quadrant population.
parent	a character scalar to specify the parent node name where the new gate to be added to, by default it is NULL, which indicates the root node
name	the population name
recompute	whether to recompute the gates
cluster_method_name	when adding the logical vectors as the gates, the name of the cluster method can be used to tag the populations as the extra meta information associated with the gates.
node	population name/path

---

**prettyAxis***Determine tick mark locations and labels for a given channel axis*

---

**Description**

Determine tick mark locations and labels for a given channel axis

**Usage**

```
prettyAxis(gh, channel)
```

**Arguments**

gh	GatingHiarchy
channel	character channel name

**Value**

when there is transformation function associated with the given channel, it returns a list of that contains positions and labels to draw on the axis other wise returns NULL

**Examples**

```
## Not run:
prettyAxis(gh, "<B710-A>")

## End(Not run)
```

---

**recompute***Compute the cell events by the gates stored within the gating tree.*

---

## Description

Compute each cell event to see if it falls into the gate stored within the gating tree and store the result as cell count.

## Usage

```
recompute(  
  x,  
  y = "root",  
  alwaysLoadData = FALSE,  
  verbose = FALSE,  
  leaf.bool = TRUE  
)  
  
## S3 method for class 'GatingSet'  
recompute(  
  x,  
  y = "root",  
  alwaysLoadData = FALSE,  
  verbose = FALSE,  
  leaf.bool = TRUE  
)  
  
## S3 method for class 'GatingSetList'  
recompute(x, ...)
```

## Arguments

x	GatingSet or GatingSetList
y	character node name or node path. Default "root". Optional.
alwaysLoadData	logical. Specifies whether to load the flow raw data for gating boolean gates. Default 'FALSE'. Optional. Sometime it is more efficient to skip loading the raw data if all the reference nodes and parent are already gated. 'FALSE' will check the parent node and reference to determine whether to load the data. This check may not be sufficient since the further upstream ancestor nodes may not be gated yet. In that case, we allow the gating to fail and prompt user to recompute those nodes explicitly. When TRUE, then it forces data to be loaded to guarantee the gating process to be uninterrupted at the cost of unnecessary data IO.
verbose	default is FALSE
leaf.bool	whether to compute the leaf boolean gate, default is TRUE
...	arguments

## Details

It is usually used immediately after `add` or `gs_pop_set_gate` calls.

---

rotate\_gate

*Simplified geometric rotation of gates associated with nodes*

---

## Description

Rotate a gate associated with a node of a `GatingHierarchy` or `GatingSet`. This method is a wrapper for `rotate_gate` that enables updating of the gate associated with a node of a `GatingHierarchy` or `GatingSet`.

`rotate_gate` calls `gs_pop_set_gate` to modify the provided `GatingHierarchy` or `GatingSet` directly so there is no need to re-assign its output. The arguments will be essentially identical to the `flowCore` method, except for the specification of the target gate. Rather than being called on an object of type `flowCore:filter`, here it is called on a `GatingHierarchy` or `GatingSet` object with an additional character argument for specifying the node whose gate should be transformed. The rest of the details below are taken from the `flowCore` documentation.

## Usage

```
## S3 method for class 'GatingHierarchy'
rotate_gate(obj, y, deg = NULL, rot_center = NULL, ...)
```

## Arguments

obj	A <code>GatingHierarchy</code> or <code>GatingSet</code> object
y	A character specifying the node whose gate should be modified
deg	An angle in degrees by which the gate should be rotated in the counter-clockwise direction
rot_center	A separate 2-dimensional center of rotation for the gate, if desired. By default, this will be the center for <code>ellipsoidGate</code> objects or the centroid for <code>polygonGate</code> objects. The <code>rot_center</code> argument is currently only supported for <code>polygonGate</code> objects.
...	not used

## Details

This method allows for geometric rotation of filter types defined by simple geometric gates (`ellipsoidGate`, and `polygonGate`). The method is not defined for `rectangleGate` or `quadGate` objects, due to their definition as having 1-dimensional boundaries.

The angle provided in the `deg` argument should be in degrees rather than radians. By default, the rotation will be performed around the center of an `ellipsoidGate` or the centroid of the area encompassed by a `polygonGate`. The `rot_center` argument allows for specification of a different center of rotation for `polygonGate` objects (it is not yet implemented for `ellipsoidGate` objects) but it is usually simpler to perform a rotation and a translation individually than to manually specify the composition as a rotation around a shifted center.

**See Also**`transform_gate flowCore::rotate\_gate`**Examples**

```
## Not run:  
#' # Rotates the original gate 15 degrees counter-clockwise  
rotate_gate(gs, node, deg = 15)  
# Rotates the original gate 270 degrees counter-clockwise  
rotate_gate(gs, node, 270)  
  
## End(Not run)
```

---

`sampleNames`*Get/update sample names in a GatingSet*

---

**Description**

Return a sample names contained in a GatingSet

**Usage**

```
sampleNames(object)  
  
sampleNames(object) <- value
```

**Arguments**

object	a GatingSet
value	character new sample names

**Details**

The sample names comes from pdata of fs.

**Value**

A character vector of sample names

**Examples**

```
## Not run:  
#G is a GatingSet  
sampleNames(G)  
  
## End(Not run)
```

---

save_cytoset	<i>save/load a cytoset to/from disk. load_cytoset() can load a cytoset from either the archive previously saved by save_cytoset() call or from a folder that contains a collection of individual cytoframe files (either in h5 format or tiledb format)</i>
--------------	-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

---

## Description

Save/load a cytoset to/from the disk.

## Usage

```
save_cytoset(cs, path, ...)
load_cytoset(path, verbose = FALSE, ...)
```

## Arguments

cs	A cytoset
path	A character scalar giving the path to save/load the cytoset to/from.
...	other arguments passed to save_gs/load_gs
verbose	whether to print details. Default is FALSE.

## Value

load\_cytoset returns a cytoset object

## Examples

```
## Not run:
#cs is a cytoset
save_cytoset(cs, outdir)
cs <- load_cytoset(outdir)

#or from cytoframe on-disk files
# e.g. h5_dir contains the cytoframes in h5 format
cs <- load_cytoset(h5_dir)

## End(Not run)
```

---

save_gs	<i>save/load a GatingSet/GatingSetList to/from disk.</i>
---------	----------------------------------------------------------

---

## Description

Save/load a GatingSet/GatingSetList which is the gated flow data including gates and populations to/from the disk. The GatingSet object The internal C data structure (gating tree),ncdfFlowSet object(if applicable)

Retrieve sample names by scanning h5 files from a GatingSet folder

## Usage

```
save_gs(
  gs,
  path,
  cdf = NULL,
  backend_opt = c("copy", "move", "skip", "symlink", "link"),
  ctx = .cytoctx_global,
  ...
)

load_gs(
  path,
  h5_readonly = NULL,
  backend_READONLY = TRUE,
  select = character(),
  verbose = FALSE,
  ctx = .cytoctx_global
)

## S4 method for signature 'character'
sampleNames(object)

save_gslist(gslist, path, ...)

load_gslist(path)
```

## Arguments

gs	A GatingSet
path	A character scalar giving the path to save/load the GatingSet to/from.
backend_opt	a character scalar. The valid options are :"copy","move","skip","symlink" specifying what to do with the backend data file. Sometimes it is more efficient to move or create a symlink of the existing backend file to the archived folder. It is useful to "skip" archiving backend file if raw data has not been changed.
ctx	cytoctx object, see [cytoctx] for details

...	other arguments: not used.
h5_READONLY	whether to open h5 data as read-only. Default is TRUE
select	an integer or character vector to select a subset of samples to load
verbose	logical flag to optionally print the versions of the libraries that were used to archive the GatingSet for troubleshooting purpose.
object	a GatingSet folder
gslist	A GatingSetList

## See Also

[GatingSet-class](#), [GatingSetList-class](#)

## Examples

```
## Not run:
#G is a GatingSet
save_gs(G, path="tempFolder")
G1<-load_gs(path="tempFolder")

#G is a GatingSet

save_gslist(gslist1, path="tempFolder")
gslist2<-load_gslist(path="tempFolder")

## End(Not run)
## Not run:
sampleNames(gsd)

## End(Not run)
```

---

scale\_gate

*Simplified geometric scaling of gates associated with nodes*

---

## Description

Scale a gate associated with a node of a GatingHierarchy or [GatingSet](#). This method is a wrapper for [scale\\_gate](#) that enables updating of the gate associated with a node of a GatingHierarchy or GatingSet.

scale\_gate calls [gs\\_pop\\_set\\_gate](#) to modify the provided GatingHierarchy or GatingSet directly so there is no need to re-assign its output. The arguments will be essentially identical to the flowCore method, except for the specification of the target gate. Rather than being called on an object of type [filter](#), here it is called on a GatingHierarchy or GatingSet object with an additional character argument for specifying the node whose gate should be transformed. The rest of the details below are taken from the flowCore documentation.

## Usage

```
## S3 method for class 'GatingHierarchy'  
scale_gate(obj, y, scale = NULL, ...)
```

## Arguments

obj	A GatingHierarchy or GatingSet object
y	A character specifying the node whose gate should be modified
scale	Either a numeric scalar (for uniform scaling in all dimensions) or numeric vector specifying the factor by which each dimension of the gate should be expanded (absolute value > 1) or contracted (absolute value < 1). Negative values will result in a reflection in that dimension.
...	not used

## Details

This method allows uniform or non-uniform geometric scaling of filter types defined by simple geometric gates ([quadGate](#), [rectangleGate](#), [ellipsoidGate](#), and [polygonGate](#)) Note that these methods are for manually altering the geometric definition of a gate. To easily transform the definition of a gate with an accompanying scale transformation applied to its underlying data, see [?ggcyt:::rescale\\_gate](#).

The scale argument passed to `scale_gate` should be either a scalar or a vector of the same length as the number of dimensions of the gate. If it is scalar, all dimensions will be multiplicatively scaled uniformly by the scalar factor provided. If it is a vector, each dimension will be scaled by its corresponding entry in the vector.

The scaling behavior of `scale_gate` depends on the type of gate passed to it. For `rectangleGate` and `quadGate` objects, this amounts to simply scaling the values of the 1-dimensional boundaries. For `polygonGate` objects, the values of `scale` will be used to determine scale factors in the direction of each of the 2 dimensions of the gate (`scale_gate` is not yet defined for higher-dimensional `polytopeGate` objects). **Important:** For `ellipsoidGate` objects, `scale` determines scale factors for the major and minor axes of the ellipse, *in that order*. Scaling by a negative factor will result in a reflection in the corresponding dimension.

## See Also

`transform_gate` [flowCore:::scale\\_gate](#)

## Examples

```
## Not run:  
# Scales both dimensions by a factor of 5  
scale_gate(gs, node, 5)  
  
# Shrinks the gate in the first dimension by factor of 1/2  
# and expands it in the other dimension by factor of 3  
scale_gate(gs, node, c(0.5,3))  
  
## End(Not run)
```

shift\_gate

*Simplified geometric translation of gates associated with nodes***Description**

Shift the location of a gate associated with a node of a GatingHierarchy or GatingSet. This method is a wrapper for `shift_gate` that enables updating of the gate associated with a node of a GatingHierarchy or GatingSet.

`shift_gate` calls `gs_pop_set_gate` to modify the provided GatingHierarchy or GatingSet directly so there is no need to re-assign its output. The arguments will be essentially identical to the `flowCore` method, except for the specification of the target gate. Rather than being called on an object of type `flowCore::filter`, here it is called on a GatingHierarchy or GatingSet object with an additional character argument for specifying the node whose gate should be transformed. The rest of the details below are taken from the `flowCore` documentation.

**Usage**

```
## S3 method for class 'GatingHierarchy'
shift_gate(obj, y, dx = NULL, dy = NULL, center = NULL, ...)
```

**Arguments**

obj	A GatingHierarchy or GatingSet object
y	A character specifying the node whose gate should be modified
dx	Either a numeric scalar or numeric vector. If it is scalar, this is just the desired shift of the gate in its first dimension. If it is a vector, it specifies both dx and dy as (dx,dy). This provides an alternate syntax for shifting gates, as well as allowing shifts of ellipsoidGate objects in more than 2 dimensions.
dy	A numeric scalar specifying the desired shift of the gate in its second dimension.
center	A numeric vector specifying where the center or centroid should be moved (rather than specifying dx and/or dy)
...	not used

**Details**

This method allows for geometric translation of filter types defined by simple geometric gates (`rectangleGate`, `quadGate`, `ellipsoidGate`, or `polygonGate`). The method provides two approaches to specify a translation. For `rectangleGate` objects, this will shift the `min` and `max` bounds by the same amount in each specified dimension. For `quadGate` objects, this will simply shift the dividing boundary in each dimension. For `ellipsoidGate` objects, this will shift the center (and therefore all points of the ellipse). For `polygonGate` objects, this will simply shift all of the points defining the polygon.

The method allows two different approaches to shifting a gate. Through the `dx` and/or `dy` arguments, a direct shift in each dimension can be provided. Alternatively, through the `center` argument, the gate can be directly moved to a new location in relation to the old center of the gate. For `quadGate`

objects, this center is the intersection of the two dividing boundaries (so the value of the boundary slot). For rectangleGate objects, this is the center of the rectangle defined by the intersections of the centers of each interval. For ellipsoidGate objects, it is the center of the ellipsoid, given by the mean slot. For polygonGate objects, the centroid of the old polygon will be calculated and shifted to the new location provided by center and all other points on the polygon will be shifted by relation to the centroid.

## See Also

transform\_gate [flowCore::shift\\_gate](#)

## Examples

```
## Not run:
# Moves the entire gate +500 in its first dimension and 0 in its second dimension
shift_gate(gs, node, dx = 500)

#Moves the entire gate +250 in its first dimension and +700 in its second dimension
shift_gate(gs, node, dx = 500, dy = 700)

# Same as previous
shift_gate(gs, node, c(500,700))

# Move the gate based on shifting its center to (700, 1000)
shift_gate(gs, node, center = c(700, 1000))

## End(Not run)
```

---

standardize-GatingSet *The tools to standardize the tree structures and channel names.*

---

## Description

```
gs_split_by_tree(x)
gs_split_by_channels(x)
gs_check_redundant_nodes(x)
gs_remove_redundant_nodes(x, toRemove)
gs_remove_redundant_channels(gs)
gs_update_channels(gs, map, all = TRUE)
gh_pop_move(gh, node, to)
gs_pop_set_visibility(x, y, FALSE)
```

## Details

In order to merge multiple GatingSets into single [GatingSetList](#), the gating trees and channel names must be consistent. These functions help removing the discrepancies and standardize the GatingSets so that they are mergable.

[gs\\_split\\_by\\_tree](#) splits the GatingSets into groups based on the gating tree structures.

[gs\\_split\\_by\\_channels](#) split GatingSets into groups based on their flow channels.

[gs\\_check\\_redundant\\_nodes](#) returns the terminal(or leaf) nodes that makes the gating trees to be different among GatingSets and thus can be considered to remove as redundant nodes.

[gs\\_remove\\_redundant\\_nodes](#) removes the terminal(or leaf) nodes that are detected as redundant by [gs\\_check\\_redundant\\_nodes](#).

[gs\\_remove\\_redundant\\_channels](#) remove the redundant channels that are not used by any gate defined in the GatingSet.

[gs\\_update\\_channels](#) modifies the channel names in place. (Usually used to standardize the channels among GatingSets due to the letter case discrepancies or typo).

[gh\\_pop\\_move](#) inserts a dummy gate to the GatingSet. Is is useful trick to deal with the extra non-leaf node in some GatingSets that can not be simply removed by [gs\\_remove\\_redundant\\_nodes](#)

[gs\\_pop\\_set\\_visibility](#) hide a node/gate in a GatingSet. It is useful to deal with the non-leaf node that causes the tree structure discrepancy.

## Description

`pop.MFI` computes and returns the median fluorescence intensity for each marker. They are typically used as the arguments passed to `gh_pop_get_stats` method to perform the sample-wise population stats calculations.

## Usage

```
pop.MFI(fr)
```

## Arguments

<code>fr</code>	a flowFrame represents a gated population
-----------------	-------------------------------------------

## Value

a named numeric vector

---

subset	<i>subset the GatingSet/GatingSetList based on 'pData'</i>
--------	------------------------------------------------------------

---

**Description**

subset the GatingSet/GatingSetList based on 'pData'

**Usage**

```
## S3 method for class 'GatingSet'  
subset(x, subset, ...)
```

**Arguments**

x	GatingSet or GatingSetList
subset	logical expression(within the context of pData) indicating samples to keep. see <a href="#">subset</a>
...	other arguments. (not used)

**Value**

a codeGatingSet or GatingSetList object

---

swap_data_cols	<i>Swap the colnames Perform some validity checks before returning the updated colnames</i>
----------------	---------------------------------------------------------------------------------------------

---

**Description**

Swap the colnames Perform some validity checks before returning the updated colnames

**Usage**

```
swap_data_cols(cols, swap_cols)
```

**Arguments**

cols	the original colname vector
swap_cols	a named list specifying the pairs to be swapped

**Value**

the new colname vector that has some colnames swapped

## Examples

```
library(flowCore)
data(GvHD)
fr <- GvHD[[1]]
colnames(fr)
new <- swap_data_cols(colnames(fr), list(`FSC-H` = "SSC-H", `FL2-H` = "FL2-A"))
colnames(fr) <- new
```

---

transform

*transform the flow data associated with the GatingSet*

---

## Description

The transformation functions are saved in the GatingSet and can be retrieved by [gh\\_get\\_transformations](#). Currently only flowJo-type biexponential transformation(either returned by [gh\\_get\\_transformations](#) or constructed by [flowJoTrans](#)) is supported.

## Usage

```
## S4 method for signature 'GatingSet'
transform(`_data`, translist, ...)
```

## Arguments

_data	GatingSet or GatingSetList
translist	expect a transformList object or a list of transformList objects(with names matched to sample names)
...	other arguments passed to 'transform' method for 'ncdfFlowSet'.(e.g. 'ncdfFile')

## Value

a GatingSet or GatingSetList object with the underling flow data transformed.

## Examples

```
## Not run:
library(flowCore)
data(GvHD)
fs <- GvHD[1:2]
gs <- GatingSet(fs)

#construct biexponential transformation function
biexpTrans <- flowjo_biexp_trans(channelRange=4096, maxValue=262144, pos=4.5, neg=0, widthBasis=-10)

#make a transformList object
chnls <- c("FL1-H", "FL2-H")
transList <- transformerList(chnls, biexpTrans)
```

```
#add it to GatingSet
gs_trans <- transform(gs, transList)

## End(Not run)
```

---

transformerList	<i>Constructor for transformerList object</i>
-----------------	-----------------------------------------------

---

## Description

Similar to transformList function, it constructs a list of transformer objects generated by trans\_new method from scales so that the inverse and breaks functions are also included.

## Usage

```
transformerList(from, trans)
```

## Arguments

from	channel names
trans	a trans object or a list of trans objects constructed by trans_new method.

## Examples

```
library(flowCore)
library(scales)
#create tranformer object from scratch
trans <- logicleTransform(w = 0.5, t = 262144, m = 4.5, a = 0)
inv <- inverseLogicleTransform(trans = trans)
trans.obj <- flow_trans("logicle", trans, inv, n = 5, equal.space = FALSE)

#or simply use convenient constructor
#trans.obj <- logicle_trans(n = 5, equal.space = FALSE, w = 0.5, t = 262144, m = 4.5, a = 0)

transformerList(c("FL1-H", "FL2-H"), trans.obj)

#use different transformer for each channel
trans.obj2 <- asinhtGml2_trans()
transformerList(c("FL1-H", "FL2-H"), list(trans.obj, trans.obj2))
```

## transform\_gate

*Simplified geometric transformations of gates associated with nodes***Description**

Perform geometric transformations of a gate associated with a node of a [GatingHierarchy](#) or [GatingSet](#). This method is a wrapper for [transform\\_gate](#) that enables updating of the gate associated with a node of a GatingHierarchy or GatingSet.

`transform_gate` calls [gs\\_pop\\_set\\_gate](#) to modify the provided GatingHierarchy or GatingSet directly so there is no need to re-assign its output. The arguments will be essentially identical to the `flowCore` method, except for the specification of the target gate. Rather than being called on an object of type `flowCore::filter`, here it is called on a GatingHierarchy or GatingSet object with an additional character argument for specifying the node whose gate should be transformed. The rest of the details below are taken from the `flowCore` documentation.

**Usage**

```
## S3 method for class 'GatingHierarchy'
transform_gate(
  obj,
  y,
  scale = NULL,
  deg = NULL,
  rot_center = NULL,
  dx = NULL,
  dy = NULL,
  center = NULL,
  ...
)
```

**Arguments**

<code>obj</code>	A <code>GatingHierarchy</code> or <code>GatingSet</code> object
<code>y</code>	A character specifying the node whose gate should be modified
<code>scale</code>	Either a numeric scalar (for uniform scaling in all dimensions) or numeric vector specifying the factor by which each dimension of the gate should be expanded (absolute value $> 1$ ) or contracted (absolute value $< 1$ ). Negative values will result in a reflection in that dimension.
<code>deg</code>	For <code>rectangleGate</code> and <code>quadGate</code> objects, this amounts to simply scaling the values of the 1-dimensional boundaries. For <code>polygonGate</code> objects, the values of <code>scale</code> will be used to determine scale factors in the direction of each of the 2 dimensions of the gate ( <code>scale_gate</code> is not yet defined for higher-dimensional <code>polytopeGate</code> objects). <b>Important:</b> For <code>ellipsoidGate</code> objects, <code>scale</code> determines scale factors for the major and minor axes of the ellipse, in that order.
<code>center</code>	An angle in degrees by which the gate should be rotated in the counter-clockwise direction.

rot_center	A separate 2-dimensional center of rotation for the gate, if desired. By default, this will be the center for ellipsoidGate objects or the centroid for polygonGate objects. The rot_center argument is currently only supported for polygonGate objects. It is also usually simpler to perform a rotation and a translation individually than to manually specify the composition as a rotation around a shifted center.
dx	Either a numeric scalar or numeric vector. If it is scalar, this is just the desired shift of the gate in its first dimension. If it is a vector, it specifies both dx and dy as (dx, dy). This provides an alternate syntax for shifting gates, as well as allowing shifts of ellipsoidGate objects in more than 2 dimensions.
dy	A numeric scalar specifying the desired shift of the gate in its second dimension.
center	A numeric vector specifying where the center or centroid should be moved (rather than specifying dx and/or dy)
...	Assignments made to the slots of the particular Gate-type filter object in the form "<slot_name> = <value>"

## Details

This method allows changes to the four filter types defined by simple geometric gates ([quadGate](#), [rectangleGate](#), [ellipsoidGate](#), and [polygonGate](#)) using equally simple geometric transformations (shifting/translation, scaling/dilation, and rotation). The method also allows for directly resetting the slots of each Gate-type object. Note that these methods are for manually altering the geometric definition of a gate. To easily transform the definition of a gate with an accompanying scale transformation applied to its underlying data, see [?ggcyto::rescale\\_gate](#).

First, `transform_gate` will apply any direct alterations to the slots of the supplied Gate-type filter object. For example, if "mean = c(1, 3)" is present in the argument list when `transform_gate` is called on a `ellipsoidGate` object, the first change applied will be to shift the `mean` slot to (1, 3). The method will carry over the dimension names from the gate, so there is no need to provide column or row names with arguments such as `mean` or `cov` for `ellipsoidGate` or `boundaries` for `polygonGate`.

`transform_gate` then passes the geometric arguments (dx, dy, deg, rot\_center, scale, and center) to the methods which perform each respective type of transformation: [shift\\_gate](#), [scale\\_gate](#), or [rotate\\_gate](#). The order of operations is to first scale, then rotate, then shift. The default behavior of each operation follows that of its corresponding method but for the most part these are what the user would expect. A few quick notes:

- `rotate_gate` is not defined for `rectangleGate` or `quadGate` objects, due to their definition as having 1-dimensional boundaries.
- The default center for both rotation and scaling of a `polygonGate` is the centroid of the polygon. This results in the sort of scaling most users expect, with a uniform scale factor not distorting the shape of the original polygon.

## See Also

[flowCore::transform\\_gate](#)

## Examples

```
## Not run:
# Scale the original gate non-uniformly, rotate it 15 degrees, and shift it
transform_gate(gs, node, scale = c(2,3), deg = 15, dx = 500, dy = -700)

# Scale the original gate (in this case an ellipsoidGate) after moving its center to (1500, 2000)
transform_gate(gs, node, scale = c(2,3), mean = c(1500, 2000))

## End(Not run)
```

[,GatingSet,ANY,ANY,ANY-method

*Bracket operators on GatingSet and GatingSetList objects*

## Description

[ subsets a GatingSet or GatingSetList using the familiar bracket notation  
 [[ extracts a GatingHierarchy object from a GatingSet.

## Usage

```
## S4 method for signature 'GatingSet,ANY,ANY,ANY'
x[i, j, ... , drop = TRUE]

## S4 method for signature 'GatingSet,numeric'
x[[i, j, ...]]
```

## Arguments

x	a GatingSet or GatingSetList
i	numeric or logical or character used as sample indices
j, ..., drop	unused

## Value

The [ operator returns an object of the same type as x corresponding to the subset of indices in i, while the [[ operator returns a single GatingHierarchy

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