Package 'CytoML'

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Type Package

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2 Contents

'read.gatingML.cytobank.R' 'graphGML_methods.R'
'helperFunctions.R' 'parameter-methods.R' 'transforms.R'
'utils.R' 'writeGatingML.R'
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Contents

addCustomInfo
ce_get_channels
ce_get_compensations
ce_get_markers
ce_get_panels
ce_get_samples
ce_get_transformations
compensate, Gating Set, graph GML-method
constructTree
cytobank_experiment-methods
cytobank_to_gatingset
CytoML-deprecated
diva_get_samples
diva_to_gatingset
diva_workspace-class
extend
fj_ws_get_keywords
fj_ws_get_samples
fj_ws_get_sample_groups
flowjo_workspace-class
gatingset_to_cytobank
gatingset_to_flowjo
getChildren,graphGML,character-method
getCompensationMatrices.graphGML
getGate,graphGML,character-method
getNodes,graphGML-method
getParent,graphGML,character-method
getTransformations.graphGML
graphGML-class
gs_compare_cytobank_counts
matchPath

addCustomInfo	2
addCustommio	3

	open_cytobank_experiment	24
	open_diva_xml	24
	open_flowjo_xml	25
	parse.gateInfo	26
	parseWorkspace	26
	plot,graphGML,missing-method	30
	range.GatingHierarchy	31
	read.gatingML.cytobank	32
	show,graphGML-method	32
Index		34
addCı	ustomInfo add customInfo nodes to each gate node and add BooleanAndGates	

Description

add customInfo nodes to each gate node and add BooleanAndGates

Usage

```
addCustomInfo(root, gs, flowEnv, cytobank.default.scale = TRUE, showHidden)
```

Arguments

root the root node of the XML

gs a GatingSet object

flowEnv the environment that stores the information parsed by 'read.GatingML'.

cytobank.default.scale

logical flag indicating whether to use the default Cytobank asinhtGml2 settings. Currently it should be set to TRUE in order for gates to be displayed properly in Cytobank because cytobank currently does not parse the global scale settings

from GatingML.

showHidden whether to include the hidden population nodes in the output

Value

XML root node

ce_get_channels

Extract channels from cytobank_experiment

Description

Extract channels from cytobank_experiment

Usage

```
ce_get_channels(x, panel_name = NULL)
```

Arguments

x A cytobank_experiment object

panel_name select panel to process

ce_get_compensations Obtain the spillover matrices for the samples in a Cytobank experi-

Description

Obtain the spillover matrices for the samples in a Cytobank experiment

Usage

```
ce_get_compensations(x)
```

Arguments

x A cytobank_experiment object

Value

A named list of spillover matrices

ce_get_markers 5

ce_get_markers

Extract markers from cytobank_experiment

Description

Extract markers from cytobank_experiment

Usage

```
ce_get_markers(x, panel_name = NULL)
```

Arguments

x A cytobank_experiment object

panel_name select panel to process

ce_get_panels

Obtain counts of the number of samples associated with each marker panel in a Cytobank experiment

Description

Obtain counts of the number of samples associated with each marker panel in a Cytobank experiment

Usage

```
ce_get_panels(x)
```

Arguments

x cytobank_experiment object

Value

A tibble of panels with sample counts

 $ce_get_samples$

Obtain a mapping between the samples and marker panels in a Cytobank experiment

Description

Obtain a mapping between the samples and marker panels in a Cytobank experiment

Usage

```
ce_get_samples(x)
```

Arguments

Х

A cytobank_experiment object

Value

A tibble with rows containing sample names and their associated panel names

```
ce\_get\_transformations
```

Obtain the transformations associated with each channel in a Cytobank experiment

Description

Obtain the transformations associated with each channel in a Cytobank experiment

Usage

```
ce_get_transformations(x, panel_name = NULL)
```

Arguments

x A cytobank_experiment object

panel_name select panel to process

Value

A transformerList object containing transformation objects for each transformed channel

```
compensate,GatingSet,graphGML-method
```

compensate a GatingSet based on the compensation information stored in graphGML object

Description

compensate a GatingSet based on the compensation information stored in graphGML object

Usage

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

Arguments

```
x GatingSet spillover graphGML ... unused.
```

Value

compensated GatingSet

constructTree

Reconstruct the population tree from the GateSets

Description

Reconstruct the population tree from the GateSets

Usage

```
constructTree(flowEnv, gateInfo)
```

Arguments

flowEnv the enivornment contains the elements parsed by read.gatingML function gateInfo the data.frame contains the gate name, fcs filename parsed by parse.gateInfo

function

Value

a graphNEL represent the population tree. The gate and population name are stored as nodeData in each node.

```
cytobank_experiment-methods
```

Methods for interacting with cytobank_experiment objects

Description

These methods mirror similar accessor methods for the GatingSet class.

Usage

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

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

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

## S4 method for signature 'cytobank_experiment'
pData(object)
```

Arguments

```
object A cytobank_experiment object
x cytobank_experiment
do.NULL, prefix not used
```

Description

A wrapper that parses the gating ML and FCS files (or cytobank_experiment object) into Gating Set $\,$

Usage

```
## Default S3 method:
cytobank_to_gatingset(x, FCS, trans = NULL, ...)
## S3 method for class 'cytobank_experiment'
cytobank_to_gatingset(x, panel_id = 1, ...)
```

CytoML-deprecated 9

Arguments

x the cytobank_experiment object or the full path of gatingML file

FCS files to be loaded

trans a 'transfomerList' object to override the transformations from gatingML files. it

is typically used by 'cytobank_experiment' parser(i.e. 'cytobank_to_gatingset.cytobank_experiment')

to use the scales info recorded in yaml file.

... other arguments

panel_id select panel to process

Value

a GatingSet

Examples

```
## Not run:
acsfile <- system.file("extdata/cytobank_experiment.acs", package = "CytoML")
ce <- open_cytobank_experiment(acsfile)
xmlfile <- ce$gatingML
fcsFiles <- list.files(ce$fcsdir, full.names = TRUE)
gs <<- cytobank_to_gatingset(xmlfile, fcsFiles)
library(ggcyto)
autoplot(gs[[1]])
## End(Not run)</pre>
```

CytoML-deprecated

Deprecated functions in package CytoML.

Description

```
GatingSet2cytobank -> gatingset_to_cytobank
GatingSet2flowJo -> gatingset_to_flowjo
cytobankExperiment -> open_cytobank_experiment
cytobank2GatingSet -> cytobank_to_gatingset
parseWorkspace -> flowjo_to_gatingset
getKeywords -> fj_ws_get_keywords
getSamples -> fj_ws_get_samples
getSampleGroups -> fj_ws_get_sample_groups
openDiva -> open_diva_xml
parseWorkspace -> diva_to_gatingset
```

10 diva_to_gatingset

diva_get_samples

Get a table of samples from a FACSDiva workspace

Description

Return a data.frame of sample group information from a FACSDiva workspace

Usage

```
diva_get_samples(x)
diva_get_sample_groups(x)
```

Arguments

Х

A diva_workspace

Value

A data. frame with columns tub, name, and specimen

diva_to_gatingset

Parse a FACSDiva Workspace

Description

Function to parse a FACSDiva Workspace, generate a GatingHierarchy or GatingSet object, and associated flowCore gates.

Usage

```
diva_to_gatingset(
  obj,
  name = NULL,
  subset = NULL,
  path = obj@path,
  worksheet = c("normal", "global"),
  swap_cols = list(`FSC-H` = "FSC-W", `SSC-H` = "SSC-W"),
  verbose = FALSE,
  ...
)
```

diva_workspace-class 11

Arguments

obj diva_workspace

name sample group to be parsed, either numeric index or the group name

subset samples to be imported. either numeric index or the sample name. Default is

NULL, which imports all samples.

path the FCS data path

worksheet select worksheet to import. either "normal" or "global"

swap_cols diva seems to swap some data cols during importing fcs to experiments this

argument provide a list to tell the parser which cols to be swapped default is

list('FSC-H' = 'FSC-W', 'SSC-H' = 'SSC-W')

verbose whether print more messages during the parsing

... other arguments

diva_workspace-class An R representation of a BD FACSDiva workspace

Description

Inherited from flowjo_workspace-class

Slots

version: Object of class "character". The version of the XML workspace.

file: Object of class "character". The file name.

.cache: Object of class "environment". An environment for internal use.

path: Object of class "character". The path to the file.

doc: Object of class "XMLInternalDocument". The XML document object.

options: Object of class "integer". The XML parsing options passed to xmlTreeParse.

See Also

GatingSet GatingHierarchy

12 extend

extend

extend the gate to the minimum and maximum limit of both dimensions based on the bounding information.

Description

It is equivalent to the behavior of shifting the off-scale boundary events into the gate boundary that is describled in bounding transformation section of gatingML standard.

Usage

```
extend(
  gate,
  bound,
  data.range = NULL,
  plot = FALSE,
  limits = c("original", "extended")
)
## S3 method for class 'polygonGate'
extend(
  gate,
  bound,
  data.range = NULL,
  plot = FALSE,
  limits = c("original", "extended")
)
## S3 method for class 'rectangleGate'
extend(gate, ...)
## S3 method for class 'ellipsoidGate'
extend(gate, ...)
```

Arguments

gate	a flowCore filter/gate
bound	numeric matrix representing the bouding information parsed from gating ML. Each row corresponds to a channel. rownames should be the channel names. colnames should be $c("min", "max")$
data.range	numeric matrix specifying the data limits of each channel. It is used to set the extended value of vertices and must has the same structure as 'bound'. when it is not supplied, c(Machine\$integer.max,Machine\$integer.max) is used.
plot	whether to plot the extended polygon.
limits	character whether to plot in "extended" or "original" gate limits. Default is "original".
	other arguments

fj_ws_get_keywords 13

Details

The advantage of extending gates instead of shifting data are two folds: 1. Avoid the extra computation each time applying or plotting the gates 2. Avoid changing the data distribution caused by adding the gates

Normally this function is not used directly by user but invoked when parsing GatingML file exported from Cytobank.

Value

a flowCore filter/gate

Examples

```
library(flowCore)
sqrcut <- matrix(c(300,300,600,600,50,300,300,50),ncol=2,nrow=4)
colnames(sqrcut) <- c("FSC-H","SSC-H")
pg <- polygonGate(filterId="nonDebris", sqrcut)
pg
bound <- matrix(c(100,3e3,100,3e3),
    byrow = TRUE, nrow = 2,
    dimnames = list(c("FSC-H", "SSC-H"),
        c("min", "max")))
bound
pg.extened <- extend(pg, bound, plot = TRUE)</pre>
```

fj_ws_get_keywords

Get Keywords

Description

Retrieve keywords associated with a workspace

Usage

```
fj_ws_get_keywords(obj, y, ...)
```

Arguments

obj	A flowjo_workspace
У	ccharacter or numeric specifying the sample name or sample ID
• • •	other arguments sampNloc a character the location where the sample name is specified. See parseWorkspace for more details.

Details

Retrieve a list of keywords from a flowjo_workspace

14 fj_ws_get_samples

Value

A list of keyword - value pairs.

Examples

```
## Not run:
    d<-system.file("extdata",package="flowWorkspaceData")
    wsfile<-list.files(d,pattern="manual.xml",full=TRUE)
    ws <- open_flowjo_xml(wsfile)

fj_ws_get_samples(ws)
    res <- try(fj_ws_get_keywords(ws,"CytoTrol_CytoTrol_1.fcs"), silent = TRUE)
    print(res[[1]])
    fj_ws_get_keywords(ws, 1)

## End(Not run)</pre>
```

fj_ws_get_samples

Get a list of samples from a flowJo workspace

Description

Return a data frame of samples contained in a flowJo workspace

Usage

```
fj_ws_get_samples(x, group_id = NULL)
```

Arguments

```
x A flowjo_workspace
group_id integer specifies the group from which samples are returned
```

Details

The samples with 0 populations are excluded. Returns a data. frame of samples in the flowjo_workspace, including their sampleID, name

Value

A data.frame with columns sampleID, name

Examples

```
## Not run:
    #ws is a flowjo_workspace
    fj_ws_get_samples(ws);
## End(Not run)
```

```
fj_ws_get_sample_groups
```

Get a table of sample groups from a flowJo workspace

Description

Return a data frame of sample group information from a flowJo workspace

Usage

```
fj_ws_get_sample_groups(x)
```

Arguments

Х

A flowjo_workspace object.

Details

Note that the samples with 0 populations are also included (since count populations requires traversing xml for all samples thus can be expensive) Returns a table of samples and groups defined in the flowJo workspace

Value

A data. frame containing the groupName, groupID, and sampleID for each sample in the workspace. Each sample may be associated with multiple groups.

See Also

```
flowjo_workspace-class flowjo_to_gatingset
```

Examples

```
## Not run:
    #ws is a flowjo_workspace
    fj_ws_get_sample_groups(ws);
## End(Not run)
```

```
flowjo_workspace-class
```

An R representation of a flowJo workspace.

Description

Objects can be created by calls of the form new("flowjo_workspace.xml", ...).

Slots

```
doc: Object of class "externalptr".
```

See Also

```
GatingSet GatingHierarchy
```

Examples

```
require(flowWorkspaceData)
d<-system.file("extdata",package="flowWorkspaceData")
wsfile<-list.files(d,pattern="A2004Analysis.xml",full=TRUE)
ws <- open_flowjo_xml(wsfile);
ws
fj_ws_get_samples(ws)</pre>
```

gatingset_to_cytobank Convert a GatingSet to a Cytobank-compatible gatingML

Description

this function retrieves the gates from GatingSet and writes a customed GatingML-2.0 file that can be imported into cytobank.

Usage

```
gatingset_to_cytobank(
   gs,
   outFile,
   showHidden = FALSE,
   cytobank.default.scale = TRUE,
   ...
)
```

gatingset_to_flowjo 17

Arguments

gs a GatingSet object

outFile a file name

showHidden whether to include the hidden population nodes in the output

cytobank.default.scale

logical flag indicating whether to use the default Cytobank asinhtGml2 settings. Currently it should be set to TRUE in order for gates to be displayed properly in Cytobank because cytobank currently does not parse the global scale settings

from GatingML.

rescale.gate default is TRUE. which means the gate is rescaled to the new scale

that is understandable by cytobank. It is recommended not to change this behavior unless user wants to export to a gatingML file used for other purpose other

than being imported into cytobank.

Details

The process can be divided into four steps: 1. Read in gate geometry, compensation and transformation from gatingSet 2. Rescale gate boundaries with flowjo_biexp() so gates can be displayed properly in Cytobank 3. Save gates and hierarchy structure to R environment 4. Write environment out to gatingML using write.GatingML()

Value

nothing

Examples

```
library(flowWorkspace)

dataDir <- system.file("extdata",package="flowWorkspaceData")
gs <- load_gs(list.files(dataDir, pattern = "gs_manual",full = TRUE))

gs_pop_remove(gs, "CD8")

#output to cytobank
outFile <- tempfile(fileext = ".xml")
gatingset_to_cytobank(gs, outFile) #type by default is 'cytobank'</pre>
```

gatingset_to_flowjo

Convert a GatingSet to flowJo workspace

Description

It is a R wrapper for the docker app (https://hub.docker.com/r/rglab/gs-to-flowjo)

18 gatingset_to_flowjo

Usage

```
gatingset_to_flowjo(gs, outFile, showHidden = FALSE, docker_img = NULL, ...)
```

Arguments

a GatingSet object or a folder contains the GatingSet archive (generated by previous save_gs call)

outFile the workspace file path to write

showHidden whether to export hidden gates. Default is FALSE

docker_img the docker image that does the actual work

other arguments passed to save_gs

Details

 $Docker\ images\ for\ gatingset_to_flowjo\ will\ be\ maintained\ at\ https://gallery.ecr.aws/x4k5d9i7/cytoverse/gs-to-wsp$

```
docker pull public.ecr.aws/x4k5d9i7/cytoverse/gs-to-wsp:latest
```

Value

nothing

Examples

```
## Not run:
library(flowWorkspace)

path <- system.file("extdata",package="flowWorkspaceData")
gs_path <- list.files(path, pattern = "gs_manual",full = TRUE)
gs <- load_gs(gs_path)

#output to flowJo
outFile <- tempfile(fileext = ".wsp")
gatingset_to_flowjo(gs, outFile)

#or directly use the archive as the input (to avoid the extra copying inside of the wrapper)
gatingset_to_flowjo(gs_path, outFile)

## End(Not run)</pre>
```

Description

get children nodes

Usage

```
## S4 method for signature 'graphGML, character'
getChildren(obj, y)
```

Arguments

obj graphGML

y character parent node path

Value

```
a graphNEL node
```

Examples

```
## Not run:
g <- read.gatingML.cytobank(xmlfile)
getChildren(g, "GateSet_722326")
getParent(g, "GateSet_722326")
## End(Not run)</pre>
```

 ${\tt getCompensationMatrices.graphGML}$

Extract compensation from graphGML object.

Description

Extract compensation from graphGML object.

Usage

```
## S3 method for class 'graphGML'
getCompensationMatrices(x)
```

Arguments

x graphGML

Value

compensation object or "FCS" when compensation comes from FCS keywords

Description

get gate from the node

Usage

```
## S4 method for signature 'graphGML,character'
getGate(obj, y)
```

Arguments

obj graphGML

y character node path

Value

the gate information associated with the node

Description

```
get nodes from graphGML object
```

Usage

```
## S4 method for signature 'graphGML'
getNodes(x, y, order = c("default", "bfs", "dfs", "tsort"), only.names = TRUE)
```

Arguments

x graphGML

y character node index. When missing, return all the nodes

order character specifying the order of nodes. options are "default", "bfs", "dfs",

"tsort"

only.names logical specifiying whether user wants to get the entire nodeData or just the

name of the population node

Value

It returns the node names and population names by default. Or return the entire nodeData associated with each node.

Examples

```
## Not run:
g <- read.gatingML.cytobank(xmlfile)
getNodes(g)
getNodes(g, only.names = FALSE)
## End(Not run)</pre>
```

```
{\it getParent,graphGML,character-method} \\ {\it get parent nodes}
```

Description

get parent nodes

Usage

```
## S4 method for signature 'graphGML,character'
getParent(obj, y)
```

Arguments

```
obj graphGML
```

y character child node path

Value

```
a graphNEL node
```

```
getTransformations.graphGML
```

Extract transformations from graphGML object.

Description

Extract transformations from graphGML object.

Usage

```
## S3 method for class 'graphGML'
getTransformations(x, ...)
```

Arguments

```
x graphGML ... not used
```

Value

transformerList object

graphGML-class

A graph object returned by 'read.gatingML.cytobank' function.

Description

Each node corresponds to a population(or GateSet) defined in gatingML file. The actual gate object (both global and tailored gates) is associated with each node as nodeData. Compensation and transformations are stored in graphData slot.

Details

The class simply extends the graphNEL class and exists for the purpose of method dispatching.

```
gs_compare_cytobank_counts

compare the counts to cytobank's exported csv so that the parsing result can be verified.
```

Description

compare the counts to cytobank's exported csv so that the parsing result can be verified.

Usage

```
gs_compare_cytobank_counts(
   gs,
   file,
   id.vars = c("FCS Filename", "population"),
   ...
)
```

matchPath 23

Arguments

gs	parsed GatingSet
file	the stats file (contains the populatio counts) exported from cytobank.
id.vars	either "population" or "FCS filename" that tells whether the stats file format is one population per row or FCS file per row.
	arguments passed to data.table::fread function

Value

a data.table (in long format) that contains the counts from openCyto and Cytobank side by side.

Examples

```
acsfile <- system.file("extdata/cytobank_experiment.acs", package = "CytoML")
ce <- open_cytobank_experiment(acsfile)
gs <- cytobank_to_gatingset(ce)
## verify the stats are correct
statsfile <- ce$attachments[1]
dt_merged <- gs_compare_cytobank_counts(gs, statsfile, id.vars = "population", skip = "FCS Filename")
all.equal(dt_merged[, count.x], dt_merged[, count.y], tol = 5e-4)</pre>
```

matchPath	Given the leaf node, try to find out if a collection of nodes can be
	matched to a path in a graph(tree) by the bottom-up searching

Description

Given the leaf node, try to find out if a collection of nodes can be matched to a path in a graph(tree) by the bottom-up searching

Usage

```
matchPath(g, leaf, nodeSet)
```

Arguments

g graphNEL

leaf the name of leaf(terminal) node

nodeSet a set of node names

Value

TRUE if path is found, FALSE if not path is matched.

24 open_diva_xml

```
open_cytobank_experiment
```

Construct a cytobank_experiment object from ACS file

Description

Construct a cytobank_experiment object from ACS file

Usage

```
open_cytobank_experiment(acs, exdir = tempfile())
```

Arguments

acs ACS file exported from Cytobank exdir he directory to extract files to

Value

cytobank_experiment object

open_diva_xml

open Diva xml workspace

Description

open Diva xml workspace

Usage

```
open_diva_xml(file, options = 0, ...)
```

Arguments

file xml file

options argument passed to xmlTreeParse ... arguments passed to xmlTreeParse

Value

```
a diva_workspace object
```

open_flowjo_xml 25

Examples

```
## Not run:
library(flowWorkspace)
library(CytoML)
ws <- open_diva_xml(system.file('extdata/diva/PE_2.xml', package = "flowWorkspaceData"))
ws
diva_get_sample_groups(ws)
gs <- diva_to_gatingset(ws, name = 2, subset = 1)
sampleNames(gs)
gs_get_pop_paths(gs)
plotGate(gs[[1]])
## End(Not run)</pre>
```

open_flowjo_xml

Open/Close a flowJo workspace

Description

Open a flowJo workspace and return a flowjo_workspace object. Close a flowjo_workspace, destroying the internal representation of the XML document, and freeing the associated memory.

Usage

```
open_flowjo_xml(file, options = 0, sample_names_from = "keyword", ...)
```

Arguments

file Full path to the XML flowJo workspace file.

options xml parsing options passed to xmlTreeParse. See http://xmlsoft.org/html/libxml-

parser.html#xmlParserOption for details.

sample_names_from

character specifying where in the XML workspace file to obtain the sample names, either "keyword" for the included \$FIL keyword for each sample, or

"sampleNode" for the name of the sample node

... not used

Details

Open an XML flowJo workspace file and return a flowjo_workspace object. The workspace is represented using a XMLInternalDocument object. Close a flowJoWorkpsace after finishing with it. This is necessary to explicitly clean up the C-based representation of the XML tree. (See the XML package).

Value

```
a flowjo_workspace object.
```

26 parseWorkspace

Examples

```
## Not run:
    file<-"myworkspace.xml"
    ws<-open_flowjo_xml(file);
    ws
## End(Not run)</pre>
```

parse.gateInfo

Parse the cytobank custom_info for each gate

Description

Fcs filename and gate name stored in 'custom_info' element are beyong the scope of the gatingML standard and thus not covered by the default 'read.gatingML'.

Usage

```
parse.gateInfo(file, ...)
```

Arguments

file xml file path

... additional arguments passed to the handlers of 'xmlTreeParse'

Value

a data.frame that contains three columns: id (gateId), name (gate name), fcs (fcs_file_filename).

parseWorkspace

Parse a flowJo Workspace

Description

Function to parse a flowJo Workspace, generate a GatingHierarchy or GatingSet object, and associated flowCore gates. The data are not loaded or acted upon until an explicit call to recompute() is made on the GatingHierarchy objects in the GatingSet.

parseWorkspace 27

Usage

```
parseWorkspace(obj, ...)
## S4 method for signature 'flowjo_workspace'
parseWorkspace(obj, ...)
flowjo_to_gatingset(
 ws,
  name = NULL,
  subset = list(),
  execute = TRUE,
  path = "",
  cytoset = NULL,
  backend_dir = tempdir(),
  backend = get_default_backend(),
  includeGates = TRUE,
  additional.keys = "$TOT",
  additional.sampleID = FALSE,
  keywords = character(),
  keywords.source = "XML",
  keyword.ignore.case = FALSE,
  extend_val = 0,
  extend_to = -4000,
  channel.ignore.case = FALSE,
  leaf.bool = TRUE,
  include_empty_tree = FALSE,
  skip_faulty_gate = FALSE,
  compensation = NULL,
  transform = TRUE,
  fcs_file_extension = ".fcs",
  greedy_match = FALSE,
 mc.cores = 1,
)
```

Arguments

obj flowjo_workspace

... Additional arguments to be passed to FCS parser

ws A flowjo_workspace to be parsed.

name numeric or character. The name or index of the group of samples to be imported. If NULL, the groups are printed to the screen and one can be selected interactively. Usually, multiple groups are defined in the flowJo workspace file.

subset numeric vector specifying the subset of samples in a group to import. Or a character specifying the FCS filenames to be imported. Or an expression to

be passed to 'subset' function to filter samples by 'pData' (Note that the columns

28 parse Workspace

referred by the expression must also be explicitly specified in 'keywords' argu-

ment)

execute TRUE | FALSE a logical specifying if the gates, transformations, and compensa-

tion should be immediately calculated after the flowJo workspace have been

imported. TRUE by default.

path either a character scalar . it is a path to the fcs files that are to be imported.

The code will search recursively, so you can point it to a location above the files.

cytoset a cytoset object that provides the alternative data source other than FCS files.

It is useful sometime to preprocess the raw fcs files (e.g. standardize channels using cytoqc package) and then directly use them for flowJo parsing. when

cytoset is provided, path argument is ignored.

includeGates logical Should gates be imported, or just the data with compensation and trans-

formation?

additional.keys

character vector: The keywords (parsed from FCS header) to be combined(concatenated with "_") with FCS filename to uniquely identify samples. Default is '\$TOT'

(total number of cells) and more keywords can be added to make this GUID.

additional.sampleID

boolean: A boolean specifying whether to include the flowJo sample ID in a GUID to uniquely identify samples. This can be helpful when the filename or other keywords are not enough to differentiate between samples. Default is

FALSE.

keywords character vector specifying the keywords to be extracted as pData of GatingSet

keywords.source

character the place where the keywords are extracted from, can be either

"XML" or "FCS"

keyword.ignore.case

a logical flag indicates whether the keywords matching needs to be case sen-

sitive.

extend_val numeric the threshold that determine wether the gates need to be extended.

default is 0. It is triggered when gate coordinates are below this value.

extend_to numeric the value that gate coordinates are extended to. Default is -4000. Usu-

ally this value will be automatically detected according to the real data range. But when the gates needs to be extended without loading the raw data (i.e.

execute is set to FALSE), then this hard-coded value is used.

channel.ignore.case

a logical flag indicates whether the colnames(channel names) matching needs

to be case sensitive (e.g. compensation, gating..)

leaf.bool a logical whether to compute the leaf boolean gates. Default is TRUE. It helps

to speed up parsing by turning it off when the statistics of these leaf boolean gates are not important for analysis. (e.g. COMPASS package will calculate them by itself.) If needed, they can be calculated by calling recompute method

at later stage.

include_empty_tree

a logical whether to include samples that don't have gates.

parse Workspace 29

skip_faulty_gate

a logical whether to skip the faulty gates so that the parser can still process the

rest of gating tree.

compensation a compensation object, matrix or data.frame or a list of these objects that al-

low the customized compensation () to be used instead of the one specified in flowJo workspace or FCS file. When it is a list, its names is supposed to be matched to sample guids (Default is the fcs filename suffixed by \$TOT. See "additional.keys" arguments for details of guids) When some of the samples don't have the external compensations matched, it will fall back to the flowJo xml or

FCS looking for the compensation matrix.

transform logical to enable/disable transformation of gates and data. Default is TRUE. It

is mainly for debug purpose (when the raw gates need to be parsed.), and only

valid when execute is FALSE.

fcs_file_extension

default is ".fcs"

greedy_match logical: By default, if flowjo_to_gatingset finds multiple FCS files matching

a sample by total event count as well as sampleID and/or keywords specified by additional.keys and additional.sampleID, it will return an error listing the duplicate files. If greedy_match is TRUE, the method will simply take the first file with either filename or \$FIL keyword matching the sample name and having

the correct number of events.

mc.cores numeric the number of threads to pass to the C++ parser to run in parallel

h5_dir the path to write h5 data

Details

A flowjo_workspace is generated with a call to open_flowjo_xml(), passing the name of the xml workspace file. This returns a flowjo_workspace, which can be parsed using the flowjo_to_gatingset() method. The function can be called non-interactively by passing the index or name of the group of samples to be imported via flowjo_to_gatingset(obj,name=x), where x is either the numeric index, or the name. The subset argument allows one to select a set of files from the chosen sample group. The routine will take the intersection of the files in the sample group, the files specified in subset and the files available on disk, and import them.

Value

a GatingSet, which is a wrapper around a list of GatingHierarchy objects, each representing a single sample in the workspace. The GatingHierarchy objects contain graphNEL trees that represent the gating hierarchy of each sample. Each node in the GatingHierarchy has associated data, including the population counts from flowJo, the parent population counts, the flowCore gates generated from the flowJo workspace gate definitions. Data are not yet loaded or acted upon at this stage. To execute the gating of each data file, a call to execute() must be made on each GatingHierarchy object in the GatingSet. This is done automatically by default, and there is no more reason to set this argument to FALSE.

See Also

fj_ws_get_sample_groups,GatingSet

Examples

```
## Not run:
   #f is a xml file name of a flowJo workspace
  ws <- open_flowjo_xml(f)</pre>
  #parse the second group
  gs <- flowjo_to_gatingset(ws, name = 2); #assume that the fcs files are under the same folder as workspace
  gs <- flowjo_to_gatingset(ws, name = 4
                          , path = dataDir
                                                #specify the FCS path
                   , subset = "CytoTrol_CytoTrol_1.fcs") #subset the parsing by FCS filename
  gs <- flowjo_to_gatingset(ws, path = dataDir, name = 4</pre>
                   , keywords = c("PATIENT ID", "SAMPLE ID", "$TOT", "EXPERIMENT NAME") #tell the parser to extract ke
                   , keywords.source = "XML" # keywords are extracted from xml workspace (alternatively can be set to
                   , additional.keys = c("PATIENT ID") #use additional keywords together with FCS filename to uniquel
                   , execute = F) # parse workspace without the actual gating (can save time if just want to get the in
 #subset by pData (extracted from keywords)
 gs <- flowjo_to_gatingset(ws, path = dataDir, name = 4
                            , subset = `TUBE NAME` %in% c("CytoTrol_1", "CytoTrol_2")
                            , keywords = "TUBE NAME")
 #overide the default compensation defined in xml with the customized compenstations
 gs <- flowjo_to_gatingset(ws, name = 2, compensation = comps); #comp is either a compensation object or a list of compensation
 ## End(Not run)
plot,graphGML,missing-method
                          plot the population tree stored in graphGML.
```

Description

The node with dotted order represents the population that has tailored gates (sample-specific gates) defined.

Usage

```
## S4 method for signature 'graphGML,missing'
plot(x, y = "missing", label = c("popName", "gateName"))
```

Arguments

- x a graphNEL generated by constructTree function
- y not used

range.GatingHierarchy

label specifies what to be dispaled as node label. Can be either 'popName' (popu-

lation name parsed from GateSets) or 'gateName' (the name of the actual gate

associated with each node)

Value

nothing

Examples

```
## Not run:
g <- read.gatingML.cytobank(xmlfile)
plot(g)
## End(Not run)</pre>
```

range.GatingHierarchy the parameter range from the flow data associated with GatingHierarchy

Description

the parameter range from the flow data associated with GatingHierarchy

Usage

```
## S3 method for class 'GatingHierarchy'
range(..., na.rm = FALSE, type = c("instrument", "data"), raw.scale = FALSE)
```

Arguments

... GatingHierarchy object

na.rm not used

type character of "instrument" or "data" indicating whether to retrieve the instrument

or the actual data range

raw. scale logical whether convert the range from transformed scale to raw scale

Value

matrix

Examples

```
## Not run:
  range(gh, type = "data")#return data range
  range(gh) #return instrument range
  range(gh, raw.scale = TRUE) #inverse transform the range to the raw scale
## End(Not run)
```

```
read.gatingML.cytobank
```

Parser for gatingML exported by Cytobank

Description

The Default parser (read.gatingML) does not parse the population tree as well as the custom information from cytobank. (e.g. gate name, fcs filename).

Usage

```
read.gatingML.cytobank(file, ...)
```

Arguments

```
file Gating-ML XML file
... additional arguments passed to the handlers of 'xmlTreeParse'
```

Value

a graphGML that represents the population tree. The gate and population name are stored in node-Data of each node. Compensation and transformations are stored in graphData.

Examples

```
## Not run:
g <- read.gatingML.cytobank(xml) #parse the population tree
#plot(g) #visualize it
## End(Not run)</pre>
```

 ${\tt show,graphGML-method} \quad \textit{show method for graphGML}$

Description

show method for graphGML

Usage

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

Arguments

object graphGML

Value

nothing

Index

addCustomInfo,3	<pre>GatingSet2flowJo (gatingset_to_flowjo), 17</pre>
ce_get_channels, 4	gatingset_to_cytobank, 9, 16
ce_get_compensations, 4	gatingset_to_flowjo, 9, 17
ce_get_markers, 5	<pre>getChildren,graphGML,character-method,</pre>
ce_get_panels, 5	19
<pre>ce_get_samples, 6 ce_get_transformations, 6 colnames, cytobank_experiment-method</pre>	getCompensationMatrices.graphGML, 19 getGate,graphGML,character-method, 20 getKeywords (fj_ws_get_keywords), 13 getNodes,graphGML-method, 20 getParent,graphGML,character-method,
<pre>cytobank_to_gatingset, 8, 9 cytobankExperiment</pre>	gs_compare_cytobank_counts, 22
(open_cytobank_experiment), 24 CytoML-deprecated, 9	<pre>markernames,cytobank_experiment-method</pre>
diva_get_sample_groups	8
<pre>(diva_get_samples), 10 diva_get_samples, 10</pre>	matchPath, 23
diva_to_gatingset, 9, 10	open_cytobank_experiment, 9, 24
diva_workspace-class, 11	open_diva_xml, 9, 24
extend, 12	<pre>open_flowjo_xml, 25 openDiva(open_diva_xml), 24</pre>
fj_ws_get_keywords, 9, 13 fj_ws_get_sample_groups, 9, 15, 29	parse.gateInfo,26
fj_ws_get_samples, 9, 14	parseWorkspace, 26
flowjo_to_gatingset, 9, 15	parseWorkspace,diva_workspace-method
<pre>flowjo_to_gatingset(parseWorkspace), 26</pre>	(diva_to_gatingset), 10
flowjo_workspace-class, 11, 16	<pre>parseWorkspace,flowjo_workspace-method</pre>
GatingHierarchy, 11, 16	pData,cytobank_experiment-method
GatingSet, <i>11</i> , <i>16</i> , <i>29</i>	<pre>(cytobank_experiment-methods),</pre>
GatingSet2cytobank	8
(gatingset_to_cytobank), 16	$\verb plot,graphGML,missing-method , 30 $

INDEX 35