Package 'flowClean'

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Version 1.2.0		
Title flowClean		
Description A quality control tool for flow cytometry data based on compositional data analysis.		
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Maintainer Kipper Flet	Clean A quality control tool for flow cytometry data based on compositional data analysis. pper Fletez-Brant r Kipper Fletez-Brant <cafletezbrant@gmail.com> R (>= 2.15.0), flowCore it, changepoint, sfsmisc lowViz, grid, gridExtra rtistic-2.0 yes FlowCytometry, QualityControl S documented: ean</cafletezbrant@gmail.com>	
Depends R (>= 2.15.0),	flowCore	
Imports bit, changepoin	t, sfsmisc	
Suggests flowViz, grid,	gridExtra	
License Artistic-2.0		
LazyLoad yes		
biocViews FlowCytome	try, QualityControl	
clean		
clean	clean. For cleaning flow cytometry data.	
Description This function uses co	ompositional data analysis to identify errant collection events.	
Usage		

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Arguments

fF flowFrame object containing experimental data to be cleaned.

vectMarkers A vector of indices representing flow parameters to be examined. These are con-

sidered as columns in the data matrix in which cells are rows and parameters are columns. Generally this vector excludes indices for various 'scatter' parameters

(e.g. 'FSC-A')

filePrefixWithDir

A string containing at least the desired name for the output flow file generated.

Can include directory structure and folder ('/' or '\') characters.

ext The file extension for the output flow file.

binSize A number in [0,1]; represents the fraction of duration of collection per bin.

nCellCutoff An integer; represents the minimum number of cells a population must have to

be included in analysis.

cutoff Method for determining threshold for parameter. Can be "median" (default) or

in [0, 1], which is interpreted as a percentile. Integers > 1 will be interpreted as

the fluorescence value to be used for a threshold.

fcMax Maximum allowable increase relative to presumed 'good' data.

announce If TRUE, will print message to screen if errors detected.

diagnostic If TRUE, will make PNG of populations in time bins, and save with same prefix

as specified in filePrefixWithDir.

Author(s)

Christopher Fletez-Brant, Pratip Chattopadhyay

References

Fletez-Brant C, Spidlen J, Brinkman R, Chattopadhyay P. Quailty Control of flow cytometry data through compositional data analysis. In preparation.

See Also

The package vignette.

Examples

```
data(synPerturbed)
synPerturbed.c <- clean(synPerturbed, vectMarkers=c(5:17),
filePrefixWithDir="sampleName", ext="fcs")</pre>
```

synPerturbed 3

|--|

Description

This is a FCS file in which a subset of one parameter was artificially perturbed so as to have a much higher fluorescent intensity than the remainder of the parameter's observations.

Format

A flowFrame with 17 observables and 76466 cells.

Details

Cells during a specific time period had their fluorescent intensities increased on channel < V705-A>.

Examples

data(synPerturbed)

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