Package 'peakCallingPipeline'

May 12, 2022

Title Call peaks for variety of ATAC-Seq data	
Version 0.1	
URL https://github.com/	
Description ADD	
License GPL-3	
Depends ArchR, parallel, data.table, assertthat	
Suggests knitr, rmarkdown, testthat	
Encoding UTF-8	
LazyData true	
VignetteBuilder knitr	
Roxygen list(markdown = TRUE)	
RoxygenNote 7.1.2	
R topics documented:	
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peakCaller Calls peaks currently utilizing ArchR	

Description

Sets global variables about specific genome build and number of threads to utilze on the machine.

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Usage

```
peakCaller(
  archr.proj,
  archr.genome,
  groupBy,
  archr.threads = 4,
  arrow.file.dir = NULL,
  cell.annotation.file = NULL,
  archr.clustering = FALSE,
  varFeatures = 15000,
  resolution = c(0.2, 1, 2),
  tileSize = 25,
  normMethod = "ReadsInTSS",
  maxCells = NULL,
  archr.visualize = TRUE,
  output.folder = NULL
)
```

Arguments

archr.proj An already setup ArchR project or a name (string) to build a new ArchR project.

If passing an already defined archr.proj then arrow.file.dir and cell.annotation.file

will not be used.

archr.genome Valid genome name for ArchR: mm10, hg38

groupBy Metadata field to group cells by

archr. threads Number of threads to utilize, warning increases memory usage as well.

arrow.file.dir Directory where all the Arrow files are stored

cell.annotation.file

File that contains sample_id and additional cell annotations to append to the

ArchRProject

archr.clustering

Boolean (T/F) determining if clustering/LSI should be run.

varFeatures Number of variable features to use for cluster

resolution Vector of clustering resolutions for multi-round clusters

tileSize Width of tiles that will section the genome normMethod Normalization method for fragment pileups maxCells Number of cells to use for each group.

archr.visualize

Should marker peak visualizations be produced

output.folder If supplied we will save the marker peak table and ArchRProject to this location.

Value

ArchRProject, markerPeaks

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