

Package ‘ArrayTV’

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Title Implementation of wave correction for arrays

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Author Eitan Halper-Stromberg

Maintainer Eitan Halper-Stromberg <Eitan177@gmail.com>

Description Wave correction for genotyping and copy number arrays

License GPL (>= 2)

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Imports foreach, DNACopy, methods, oligoClasses (>= 1.21.3)

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Collate AllGenerics.R calctvScore.R computeGC.R correctionTVscore.R
CorrectM.R gcCorrectMain.R gcFracAllWin.R gcFracOneRange.R
getGCinBestWindowGenome.R methods-gcCorrect.R priorFrac.R
priorFracRestOfGenome.R

biocViews CopyNumberVariation

NeedsCompilation no

R topics documented:

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ArrayTV-package

ArrayTV-implementation of GC correction for arrays

Description

Correct gc biases in array data. The user must provide a range of windows and this function will compute the tv score for each window and correct the arrays based upon the window with the highest tv score (i.e with window showing the most gc bias)

Details

Package: ArrayTV
Type: Package
Version: 1.0
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gcCorrectMain takes a matrix of array signal intensities, each column representing one array, along with vectors corresponding to probe positions and chromosomes. The gcCorrect methods enables array input for objects of class matrix, BeadStudioSet, BafLrrSet, and BafLrrSetList

Author(s)

Eitan Halper-Stromberg Maintainer: Eitan Halper-Stromberg

References

Yuval Benjamini and Terence P. Speed. Summarizing and correcting the GC content bias in high-throughput sequencing. *Nucleic Acids Res*, 40(10):e72, May 2012.

Eitan Halper-Stromberg, Laurence Frelin, Ingo Ruczinski, Robert Scharpf, Chunfa Jie, Benilton Carvalho, Haiping Hao, Kurt Hetrick, Anne Jedlicka, Amanda Dziedzic, Kim Doheny, Alan F. Scott, Steve Baylin, Jonathan Pevsner, Forrest Spencer, and Rafael A. Irizarry. Performance assessment of copy number microarray platforms using a spike-in experiment. *Bioinformatics*, 27(8):1052

Adam B Olshen, E. S. Venkatraman, Robert Lucito, and Michael Wigler. Circular binary segmentation for the analysis of array-based DNA copy number data. *Biostatistics*, 5(4):557

See Also

type ?gcCorrectMain.Rd at the command prompt for an example

computeGC-methods	<i>Compute the GC composition for selected windows</i>
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Description

Compute the GC composition for selected windows around a vector of genomic positions (e.g., locations of markers on an array)

Methods

```
computeGC(x, maxwins, increms, chr, build, ...):
```

x can be numeric, in which case x should be a vector of physical position along the genome. Alternatively, x can be an instance of the `BafLrrSetList` class. When x is numeric, arguments `chr` and `build` are required. Arguments `chr` and `build` are ignored for `BafLrrSetList` objects.

Arguments `maxwins`, `increms`, `build`, and `chr` are as described in the documentation for `gcCorrectMain`

See Also

[BafLrrSetList](#), [gcCorrectMain](#)

Examples

```
if(require(crlmm) && require(ff) && require(doParallel)){
## nodes may be set to the number of cpus available
  nodes<-3
  cl <- makeCluster(nodes)
  registerDoParallel(cl)

data(cnSetExample, package="crlmm")
brList <- BafLrrSetList(cnSetExample)
pos <- unlist(position(brList))
chr <- paste("chr",
  rep(chromosome(brList), elementLengths(brList)),
  sep="")
gc.matrix1 <- ArrayTV:::computeGC(x=pos, maxwins=c(12,10e3),
  increms=c(12,10e3),
  chr=chr,
  build="hg19")
gc.matrix2 <- ArrayTV:::computeGC(brList, c(12, 10e3),
  c(12, 10e3))
  identical(gc.matrix1, gc.matrix2)
  stopCluster(cl)
}
```

Description

Correct gc biases in array data for arrays stored as objects of an acceptable class. The user provides the array data and a range of windows and these methods will format the data and call gcCorrectMain. This function computes the tv score for each window and corrects the arrays based upon the window with the highest tv score (i.e with window showing the most gc bias)

Methods

gcCorrect(object, ...):

Methods have been defined for objects of class `matrix`, `BeadStudioSet`, `BafLrrSet`, and `BafLrrSetList`.

Objects of class `BafLrrSetList` are containers for log R ratios and B allele frequencies (BAFs) stored by chromosome. In particular, each element in this list class contains BAFs and log R ratios as `assayData` and genomic annotation of the markers (`featureData`) for one chromosome.

The `gcCorrect` method for objects of this class extracts the log R ratios for all elements (chromosomes) in the list, and then combines these into a single matrix. If the log R ratios are stored as `ff` objects, the samples will be GC-corrected in chunks determined by the function `ocSamples()`. For example, if `object` contains 100 samples and `ocSamples(10)` was specified prior to calling `gcCorrect`, the wave correction will be performed on 10 samples at a time in order to keep RAM at an acceptable level. When processing large numbers of samples, it can be helpful to evaluate the tv score on a subset of the samples, pick one or two windows for the correction, and precompute the GC content for these windows. See the examples below for the implementation with `BafLrrSetList` objects for details. When the assay data is represented as `ff` objects, the `gcCorrect` method returns the value `NULL`. Note that the assay data stored on disk will have changed as a result of calling this method.

When `object` is a `matrix` of log R ratios, the `gcCorrect` method returns a matrix of wave-adjusted log R ratios.

Additional arguments are passed to `gcCorrectMain` through the `...` operator.

References

Benjamini Y, Speed TP. Summarizing and correcting the GC content bias in high-throughput sequencing. *Nucleic Acids Res* 2012;40:e72

See Also

[BeadStudioSet](#), [BafLrrSetList](#), [BafLrrSet](#) `gcCorrectMain`

Examples

```

    ## Example with 2 iterations of correction
    path <- system.file("extdata", package="ArrayTV")
    load(file.path(path, "array_logratios.rda"))
    nimblegen[, "M"] <- nimblegen[, "M"]/1000
    increms <- c(10,1000,100e3)
    wins <- c(100,10e3,1e6)
    if(require(doParallel)){
      ## nodes may be set to the number of cpus available
      nodes<-3
      cl <- makeCluster(nodes)
      registerDoParallel(cl)
    }

    ## calculate tv scores in many windows and correct M values using the best window
    nimcM1List <- gcCorrectMain(nimblegen[, "M",
    drop=FALSE],
    chr=rep("chr15",
    nrow(nimblegen)),
    starts=nimblegen[, "position"],
    increms=increms,
    maxwins=wins, build='hg18')
    tvScores <- nimcM1List[['tvScore']]
    winsorted <- as.numeric(rownames(tvScores)[order(tvScores, decreasing=TRUE)])
    logwinsorted <- log(as.numeric(winsorted), 10)
    logwinsortdiff <- abs(logwinsorted[1]-logwinsorted)
    ## correct M values a 2nd time using a different window size
    nimcM2List <- gcCorrect(nimcM1List[['correctedVals']],
    chr=rep("chr15",
    nrow(nimblegen)),
    starts=nimblegen[, "position"],
    maxwins=winsorted[logwinsortdiff>=1][1],
    build='hg18')
    ## Refer to the vignette for details

    ## Example using a list container (BafLrrSetList) containing log R
    ## ratios and B allele frequencies

    if(require(crlmm) && require(ff)){
      data(cnSetExample, package="crlmm")
      brList <- BafLrrSetList(cnSetExample)
      is(lrr(brList)[[1]], "ff_matrix")
      rold <- lrr(brList)[[1]][, 1]/100
      ##
      ## To avoid having too much data in RAM it might be
      ## useful to process the samples n at a time
      ##
      ## The number of samples to be processed at a time is
      ## set by the ocSamples function. For example, to
      ## process 20 samples at a time one would do
      ocSamples(20)
    }

```

```

##
## When assay data elements are ff objects, the wave
## corrected values are updated on disk. Currently,
## the data is stored is here:
filename(lrr(brList)[[1]])
##
## To avoid permanently changing the log R ratio
## values for the brList object, we copy the ff files
## to a different path and create a new BafLrrSetList
## object. This is done by the non-exported function
## "duplicateBLList". The path for the new ff objects
## is given by the function ldPath(). Here, we use a
## temp directory
ldPath(tempdir())
brList.copy <- oligoClasses::duplicateBLList(brList, ids=sampleNames(brList))
filename(lrr(brList.copy)[[1]])
##
## wave correct the log R ratios
##
gcCorrect(brList.copy, increms=c(12, 10e3),
          maxwins=c(12, 10e3))
##
##
rupdated <- lrr(brList.copy)[[1]][, 1]/100
## note that rold and rupdated are no longer the same
## since the log R ratios were updated in the
## brList.copy container
plot(rupdated, rold, pch=".")
##
## Remarks on efficiency
##
## If a large number of samples are to be processed,
## the most efficient procedure is to settle on an
## appropriate window size for gc correction using a
## subset of the dataset (e.g., 20 samples). See the
## vignette for details. Here, we assume that two
## windows were already selected using such a
## procedure (here, 12 bp and 10,000 bp) and the goal
## is to efficiently do wave correction using these
## two windows for all the samples in a large study.
## Currently, our recommended approach is to first
## calculate the gc content for these windows:
gc.matrix <- ArrayTV::computeGC(brList.copy, c(12, 10e3),
                               c(12, 10e3))
## The number of columns in gc.matrix will correspond
## to the number of windows
ncol(gc.matrix)
## Having calculated the gc content for these two
## windows, we pass the gc matrix to the method
## gcCorrect. This function will iteratively update
## the log R ratios for the gc content given by the
## columns in gc.matrix (the log R ratios are updated
## for each column in gc.matrix).

```

```

ArrayTV::gcCorrect(brList.copy, increms=c(12, 10e3),
maxwins=c(12, 10e3),
providedGC=gc.matrix)
      stopCluster(cl)
}

```

gcCorrectMain

gcCorrectMain

Description

Correct gc biases in array data. The user must provide a range of windows and this function will compute the tv score for each window and correct the arrays based upon the window with the highest tv score (i.e with window showing the most gc bias)

Usage

```

gcCorrectMain(Ms, chr, starts, samplechr, increms,
              maxwins, jittercorrection=FALSE,
              returnOnlyTV=FALSE, onlyGC=FALSE,providedGC=NULL,build, verbose=FALSE)

```

Arguments

Ms	A matrix holding the array values
chr	a vector indicating the chromosome of each probe (should be the same length as the number of rows in the array matrix)
starts	a vector holding the genomic coordinates of the start positions the probes (should be the same length as the number of rows in the array matrix)
samplechr	a subset (or the entire set) of the chromosomes represented in the arrays. The probes on these chromosomes will be used to compute the tvScore
increms	a vector of integers
maxwins	a vector of integers with each value \geq the corresponding value in increms. Corresponding values of maxwins and increms must have the same quotient. For instance, if increms is <code>c(10,1000)</code> then an acceptable value of maxwins would be <code>(500,5000)</code> because the quotients of the corresponding elements are both 5.
jittercorrection	if TRUE amplify the correction values by a small amount to decrease the auto-correlation of proximal probes
returnOnlyTV	if TRUE only return the tvScores rather than applying corrections
onlyGC	if TRUE return the gc fraction of each window (the result may be subsequently passed into the providedGC argument to be used in correction)
providedGC	a vector of gc values (fractions from zero to 1) to use for computing corrections. This should be the same length as the number of rows in the array, if provided

build	the build matching the array. This is used to load the BSgenome object. If the arrays are annotated based on hg18 this would be 'hg18', if the annotation is based on hg19 this would be 'hg19'.
verbose	provide output throughout computation

Details

see example

Value

The default is to return corrected values for the arrays passed in. This result will have the same dimensions as the first argument (the uncorrected values).if returnOnlyTV is TRUE this will be a matrix of tv score values, one row per window, one column per array. The rownames of the matrix will indicate the size of the window. If onlyGC=TRUE the gc fraction of each window will be returned.

Author(s)

Eitan Halper-Stromberg

References

Optimal window finding was inspired by a similar concept applied to gc correction for next generation sequencing experiments:Benjamini Y, Speed TP. Summarizing and correcting the GC content bias in high-throughput sequencing. Nucleic Acids Res 2012;40:e72

Examples

```
## Example with parallelization, setting the number of processors to 3

path <- system.file("extdata", package="ArrayTV")
load(file.path(path, "array_logratios.rda"))
nimblegen[, "M"] <- nimblegen[, "M"]/1000

increments <- c(10,1000,100e3)
wins <- c(100,10e3,1e6)
  if(require(doParallel)){
    ## nodes may be set to the number of cpus available
    nodes<-3
    cl <- makeCluster(nodes)
    registerDoParallel(cl)
  }

## calculate tv scores in many windows and correct M values
## using the best window
nimcM1List <- gcCorrectMain(nimblegen[, "M", drop=FALSE],
  chr=rep("chr15",nrow(nimblegen)),
  starts=nimblegen[, "position"],
  increments=increments,
```



```
      maxwins=wins,
      build='hg18')
tvScores <- nimcM1List[['tvScore']]
winsorted <- as.numeric(rownames(tvScores)[order(tvScores,decreasing=TRUE)])
logwinsorted <- log(as.numeric(winsorted),10)
logwinsortdiff <- abs(logwinsorted[1]-logwinsorted)
## correct M values a 2nd time using a different window size
nimcM2List <- gcCorrect(nimcM1List[['correctedVals']],
  chr=rep("chr15",nrow(nimblegen)),
  starts=nimblegen[, "position"],
  maxwins=winsorted[logwinsortdiff>=1][1],
  build='hg18')
      stopCluster(cl)
## Refer to the vignette for details
```

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