Package ‘ACME’

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Title Algorithms for Calculating Microarray Enrichment (ACME)
Author Sean Davis <sdavis2@mail.nih.gov>
Maintainer Sean Davis <sdavis2@mail.nih.gov>
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Imports graphics, stats
Description ACME (Algorithms for Calculating Microarray Enrichment) is a set of tools for analysing tiling array ChIP/chip, DNase hypersensitivity, or other experiments that result in regions of the genome showing "enrichment". It does not rely on a specific array technology (although the array should be a "tiling" array), is very general (can be applied in experiments resulting in regions of enrichment), and is very insensitive to array noise or normalization methods. It is also very fast and can be applied on whole-genome tiling array experiments quite easily with enough memory.
License GPL (>= 2)
URL http://watson.nci.nih.gov/~sdavis
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R topics documented:

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ACMECalcSet-class

Description

A subclass of ACMESet that can also store the parameters and results of an ACME calculation.

Objects from the Class

Objects can be created by calls of the form new("ACMECalcSet", assayData, phenoData, featureData, experimentData, annotation, cutpoints, threshold, exprs, vals, ...). In addition to the constraints defined by ACMESet, this class can also hold the results (in the assay-DataElement vals) and the threshold and cutpoints from an ACME do.aGFF.calc run.

Slots

cutpoints: Object of class "numeric" The values of the cutpoints used in an analysis by do.aGFF.calc, one per sample.
threshold: Object of class "numeric" The threshold used in an analysis.
assayData: Object of class "AssayData". See ExpressionSet for details.
phenoData: Object of class "AnnotatedDataFrame" See ExpressionSet for details.
featureData: Object of class "AnnotatedDataFrame" See ExpressionSet for details.
experimentData: Object of class "MIAME" See ExpressionSet for details.
annotation: Object of class "character" See ExpressionSet for details.
__classVersion__: Object of class "Versions" See ExpressionSet for details.

Extends


Methods

cutpoints signature(x = "ACMECalcSet"): A simple getter for the cutpoints.
plot signature(x = "ACMECalcSet"): A convenience plotting method that also takes sample and chrom
show signature(object = "ACMECalcSet"): A show method
threshold signature(x = "ACMECalcSet"): A simple getter for the threshold
vals signature(x = "ACMECalcSet"): an accessor for the p-values from a run of do.aGFF.calc. Returns a matrix with samples in columns and probes in rows.

Author(s)

Sean Davis <sdavis2@mail.nih.gov>

See Also

ACMESet
ACMESet-class

Examples

```r
class("ACMECalcSet")
data(example.agff)
b <- do.aGFF.calc(example.agff, thresh=0.95, window=1000)
b
head(vals(b))
threshold(b)
cutpoints(b)
```

ACMESet-class  Class "ACMESet"

Description
An extension of ExpressionSet to deal with ACME data including chromosome locations

Objects from the Class

Objects can be created by calls of the form `new("ACMESet", assayData, phenoData, featureData, experimentData, annotation, exprs, ...)`. The exprs assayDataElement stores the data. The featureData slot stores the chromosome location. In practice, the data.frame underlying the featureData MUST contain three columns named chromosome, start, and end; this is enforced by the class validity method.

Slots

- `assayData`: Object of class "AssayData". See ExpressionSet for details.
- `phenoData`: Object of class "AnnotatedDataFrame" See ExpressionSet for details.
- `featureData`: Object of class "AnnotatedDataFrame" See ExpressionSet for details.
- `experimentData`: Object of class "MIAME" See ExpressionSet for details.
- `annotation`: Object of class "character" See ExpressionSet for details.
- `.__classVersion__`: Object of class "Versions" See ExpressionSet for details.

Extends

Class "ExpressionSet", directly. Class "eSet", by class "ExpressionSet", distance 2. Class "VersionedBiobase", by class "ExpressionSet", distance 3. Class "Versioned", by class "ExpressionSet", distance 4.

Methods

- `chromosome` signature(object = "ACMESet"): Accessor for the chromosome. Returns a vector of chromosomes.
- `end` signature(x = "ACMESet"): Accessor for the end location for a probe. If that is not known, this could be set to the same value as the start location.
- `plot` signature(x = "ACMESet"): A convenience plotting method that takes a sample name and chrom as well.
- `start` signature(x = "ACMESet"): Accessor for the start location for a probe.
aGFF-class

Author(s)
Sean Davis <sdavis2@mail.nih.gov>

See Also
ExpressionSet, ACMECalcSet

Examples

```
showClass("ACMESet")
data(example.agff)
example.agff
head(chromosome(example.agff))
head(start(example.agff))
head(end(example.agff))
```

---

**aGFF-class**  
Class for storing GFF-like data

**Description**

The GFF format is quite versatile while remaining simple. This class simply stores the annotation associated with a set of GFF files from the same regions of the genome along with some information about the samples from which the data came and the data (from the "score" column of the GFF file) themselves.

**Objects from the Class**

Objects can be created by calls of the form `new("aGFF", ...)`. Also, the `read.resultsGFF()` function returns aGFF objects.

**Slots**

- **annotation**: Object of class "data.frame" with two columns absolutely necessary, "Chromosome" and "Location". Other columns can be included.
- **data**: Object of class "matrix" of the same number of rows as the annotation slot and the same number of columns as the number of rows in the samples slot, containing data for later analysis
- **samples**: Object of class "data.frame" for describing the samples, one row per sample

**Methods**

- **plot** signature(x = "aGFF"): to plot a region along the genome.
- **print** signature(x = "aGFF"): simple method to display summary of aGFF object
- **show** signature(object = "aGFF"): simple method to display summary of aGFF object

**Author(s)**

Sean Davis

**See Also**

read.resultsGFF and aGFFCalc-class
aGFFCalc-class

Examples

# Load an example
data(example.agff)
example.agff

aGFFCalc-class  Class "aGFFCalc"

Description

Store results of ACME calculations

Objects from the Class

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

Slots

call: Object of class "call", contains the exact call to do.aGFF.calc, for historical purposes
threshold: Object of class "numeric", the threshold used in the calculation
cutpoints: Object of class "numeric", the data value above which probes were considered positive
vals: Object of class "matrix", equivalent in size to the original data matrix, containing the calculated p-values from the ACME algorithm
annotation: Object of class "data.frame", currently a copy of the original annotation, possibly reordered in chromosome order
data: Object of class "matrix", the original data, possibly reordered
samples: Object of class "data.frame", sample metadata

Extends

Class "aGFF", directly.

Methods

plot signature(x = "aGFFCalc", ask=FALSE): plot the results of an ACME calculation
print signature(x = "aGFFCalc"): brief overview of the object
show signature(object = "aGFFCalc"): brief overview of the object

Author(s)

Sean Davis <sdavis2@mail.nih.gov>

See Also

do.aGFF.calc, aGFF-class

Examples

data(example.agff)
example.agffcalc <- do.aGFF.calc(example.agff,window=1000,thresh=0.9)
exa...
do.aGFF.calc

Perform ACME calculation

Description

This function performs the moving window chi-square calculation. It is written in C, so is quite fast.

Usage

```
do.aGFF.calc(x, window, thresh)
```

Arguments

- `x` An aGFF class object
- `window` An integer value, representing the number of basepairs to include in the windowed chi-square calculation
- `thresh` The quantile of the data distribution for each sample that will be used to classify a probe as positive

Details

A window size on the order of 2-3 times the average size of fragments from sonication, digestion, etc. and containing at least 8-10 probes is the recommended size. Larger size windows are probably more sensitive, but obviously reduce the accuracy with which boundaries of signal can be called.

A threshold of between 0.9 and 0.99 seems empirically to be adequate. If one plots the histogram of data values and there is an obvious better choice (such as a bimodal distribution, with one peak representing enrichment), a more data-driven approach may yield better results.

Value

An object of class aGFFCalc

Author(s)

Sean Davis <sdavis2@mail.nih.gov>

Examples

```
data(example.agff)
example.agffcalc <- do.aGFF.calc(example.agff, window=1000, thresh=0.9)
example.agffcalc
```
Example AGFF

An example ACME data structure of class ACMESet

Description
An ACMESet data structure from two Nimblegen arrays, custom tiled to include multiple HOX genes.

Usage
data(example.agff)

Format
The format is: chr "example.agff"

Source
From Scacheri et al., PloS Genet, 2006. Pubmed ID 16604156

Examples
data(example.agff)
example.agff

---

**findClosestGene**

*Find closest refseq gene*

Description
This function is used to find the nearest refseq transcript(s) to a point in the genome specified. Note that it is limited to the refseq transcripts listed at genome.ucsc.edu, where this function goes for information.

Usage
findClosestGene(chrom, pos, genome = "hg17", position = "txStart")

Arguments

<table>
<thead>
<tr>
<th>Argument</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>chrom</td>
<td>Usually specified like 'chr1', 'chr2', etc.</td>
</tr>
<tr>
<td>pos</td>
<td>A position in base pairs in the genome</td>
</tr>
<tr>
<td>genome</td>
<td>Something like 'hg16', 'hg17', 'mm6', etc.</td>
</tr>
<tr>
<td>position</td>
<td>The location to measure distance from: one of 'txStart', 'txEnd', 'cdsStart', 'cdsEnd'</td>
</tr>
</tbody>
</table>

Details
The first time the function is run, it checks to see if the refflat table for the given genome is present in the package environment. If not, it downloads it to the /tmp directory and gunzips it (using getRefflat). It is then stored so that in future calls, there is no re-download required.
findRegions

Value

A data frame with the gene name, refseq id(s), txStart, txEnd, cdsStart, cdsEnd, exon count, and
distance. Note that distance is measured as pos-position, so negative values mean that the point in
the gene is to the left of the point specified in the function call (with the p-tel on the left).

Note

The function may return more than one transcript, as several transcripts may have the same start site

Author(s)

Sean Davis <sdavis2@mail.nih.gov>

Examples

findClosestGene('chr1',100000000,'hg17')

findRegions

Find all regions in data above p-value threshold

Description

After the ACME calculation, each probe is associated with a p-value of enrichment. However, one
often wants the contiguous regions associated with runs of p-values above a given p-value threshold.

Usage

findRegions(x, thresh = 1e-04)

Arguments

x An ACMESetCalc object
thresh The p-value threshold

Details

Runs of p-values above the p-value threshold will be reported as one "region". These can be used
for downstream analyses, export to browsers, submitted for transcription factor binding enrichment
analyses, etc.

Value

A data frame with these columns:

<table>
<thead>
<tr>
<th>Column</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Length</td>
<td>The length of the region in probes</td>
</tr>
<tr>
<td>TF</td>
<td>Either TRUE or FALSE; TRUE regions represent regions of enrichment while FALSE regions are the regions between the TRUE regions</td>
</tr>
<tr>
<td>StartInd</td>
<td>The starting Index of the region</td>
</tr>
<tr>
<td>EndInd</td>
<td>The ending Index of the region</td>
</tr>
<tr>
<td>Sample</td>
<td>The sample containing the region</td>
</tr>
</tbody>
</table>
Chromosome  The Chromosome of the region
Start  The starting basepair of the region
End  The ending basepair of the region
Median  The median p-value in the region
Mean  The mean p-value in the region

Author(s)

Sean Davis <sdavis2@mail.nih.gov>

See Also

do.aGFF.calc, findClosestGene

Examples

data(example.agff)
example.agffcalc <- do.aGFF.calc(example.agff, window=1000, thresh=0.9)
foundregions <- findRegions(example.agffcalc, thresh=0.001)
foundregions[1:6,]

datasets 

generics  

Generics defined within ACME 

Description

See methods descriptions for details.

Usage

vals(x, ...)
chromosome(object, ...)
end(x, ...)
start(x, ...)
plot(x, y, ...)
cutpoints(x, ...)
threshold(x, ...)

Arguments

x  An ACMESet or ACMECalcSet object (for cutpoints and threshold)
object  An ACMESet or ACMECalcSet object (for cutpoints and threshold)
y  Treated as missing for plotting these types of objects
...  Passed into method

Details

These are all getters for ACMESet and ACMECalcSet objects.
getRefflat

Value
See methods descriptions for details

Author(s)
Sean Davis <sdavis2@mail.nih.gov>

See Also
ACMESet, ACMECalcSet

Examples
```r
data(example.agff)
head(chromosome(example.agff))
head(end(example.agff))
head(start(example.agff))
```

---

**getRefflat**  
*Get the refflat table from ucsc for the given genome*

Description
Fetches the refflat table from ucsc, stores in temp dir and then gunzips it and reads it in.

Usage
```r
getRefflat(genome = "hg17")
```

Arguments
- **genome**  
The genome code from ucsc, like 'hg16', 'mm6', etc.

Value
A data frame mirroring the UCSC table structure.

Author(s)
Sean Davis <sdavis2@mail.nih.gov>

References
http://genome.ucsc.edu

See Also
findClosestGene

Examples
```r
rf <- getRefflat("hg17")
```
read.resultsGFF  

Read Nimblegen GFF files

Description
A GFF format file is a quite flexible format for storing genomic data. Nimblegen uses these format files as one format for making chip-chip data available. This function reads these files, one per experiment and creates a resulting aGFF-class object.

Usage
read.resultsGFF(fnames, path = ".", samples = NULL, notes = NULL, skip = 0, sep = "\t", quote = "\"")

Arguments
- fnames: A vector of filenames
- path: The path to the filenames
- samples: A data.frame containing sample information, one row per sample, in the same order as the files in fnames
- notes: A character vector for notes--not currently stored
- skip: Number of lines to skip if the file contains a header
- sep: The field separator--should be a tab character for gff files, but can be set if necessary.
- quote: The text quote character--again not used for gff file, typically

Details
The output is an ACMESet object.

Value
A single ACMESet object.

Author(s)
Sean Davis <sdavis2@mail.nih.gov>

References
http://www.sanger.ac.uk/Software/formats/GFF/

See Also
ACMESet

Examples
datdir <- system.file('extdata', package='ACME')
fnames <- dir(datdir)
example.agff <- read.resultsGFF(fnames, path=datdir)
write.bedGraph  Write bedGraph format tracks for UCSC genome browser

Description

Generate bedGraph format files for the UCSC genome browser. This function will write the bedGraph files associated with a aGFFcalc object. There will be either one or two files (default two) representing the raw data and the calculated data (which is output as -log10(val) for visualization purposes for EACH sample).

Usage

write.bedGraph(x, raw = TRUE, vals = TRUE, directory = ".")

Arguments

x  An ACMESet or ACMECalcSet object
raw  Boolean. Create a file for the raw data?
vals  Boolean. Create a file for the calculated p-values?
directory  Give a directory for storing the files

Author(s)

Sean Davis

Examples

data(example.agff)
write.bedGraph(example.agff)

write.sgr  Write Affy IGB .sgr format files

Description

The affy Integrated Genome Browser (IGB) is a powerful, fast browser for genomic data. The file format is simple (three columns: chromosome, location, and score) to generate. This function will write the sgr files associated with a aGFFcalc object. There will be either one or two files (default two) representing the raw data and the calculated data (which is output as -log10(val) for visualization purposes).

Usage

write.sgr(x, raw = TRUE, vals = TRUE, directory = ".")
write.sgr

Arguments

- **x**: An ACMESet or ACMECalcSet object
- **raw**: Boolean. Create a file for the raw data?
- **vals**: Boolean. Create a file for the calculated p-values?
- **directory**: Give a directory for storing the files

Author(s)

Sean Davis

Examples

```r
data(example.agff)
write.sgr(example.agff)
```
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