encoDnaseI

April 2, 2010

ALICOR   *aligned and interpolated correlation for local maximum traces for two scatterplots*

Description

aligned and interpolated correlation for local maximum traces from two scatterplots

Usage

```
alicor(x1, y1, x2, y2, bin = 50000)
ALICOR(ssr, dns = rawCD4, bin = 50000)
```

Arguments

x1   domain points for first scatterplot

y1   range points for first scatterplot

x2   domain points for second scatterplot

y2   range points for second scatterplot

ssr   an instance of class snpScreenResult

dns   an instance of class hg18track

bin   bin size, units are base pairs

Details

We define a scatterplot to be a pair of vectors (x, y). We are interested in measuring the distance between two scatterplots, focusing on the locations of local peaks and valleys. Two scatterplots are close if their peaks and valleys in y are nearby in x.

We have no restrictions on commonalities between the scatterplots, but this only makes sense if there is reasonable overlap between their x ranges.

The algorithm, implemented in low-level function alicor, is as follows. Use parameter bin to define a grid in x for each scatterplot, and compute the maximum y value in each x-grid interval. Compute a common domain for the two scatterplots based solely on x1. Linearly interpolate the maximal series for each scatterplot on the common domain. Compute the correlation coefficient for the resulting interpolated series.

The higher-level function ALICOR adapts this to snpScreenResult plot_mlp display data and to hg18 annotation track data.

1

Value

scalar correlation coefficient

Note

Many variations on this algorithm are possible, but the code is not very flexible at this time.

Author(s)

VJ Carey <stvjc@channing.harvard.edu>

Examples

```
data(sOSR2)
data(c19g)
juxtaPlot(c19g, sOSR2)
ALICOR(sOSR2,  c19g)
```

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rawCd4DnaseI	*A data frame with information on the UCSC browser track related to DNaseI hypersensitivity*

Description

A data frame with information on the UCSC browser track related to DNaseI hypersensitivity; the rawCD4 object is an eSet extension representing the same information; rawHelaDnaseI is like rawCD4 but results on Hela cells.

Usage

```
data(rawCd4DnaseI)
data(rawHelaDnaseI)
data(rawCD4)
```

Details

Obtained from a MySQL representation of the data distributed at the Genome Browser FTP site

Value

a data.frame

Author(s)

Vince Carey <stvjc@channing.harvard.edu>

References

hgdownload.cse.ucsc.edu ... it appears that they do not offer the MYD/MYI representations, just the txt.gz and sql files now. So if you obtain the encodeNhgriDnaseHsChipRawCd4.txt and .sql files at goldenPath/currentGenomes/Homo_Sapiens/encode/database, you can reconstruct the underlying data for this data.frame (hg18, Nov 2007).

Examples

```
data(rawCd4DnaseI)
dim(rawCd4DnaseI)
rawCd4DnaseI[1:5,]
library(lattice)
xyplot(dataValue~chromStart|chrom, data=rawCd4DnaseI, subset=chrom %in%
  c("chr1", "chr10", "chr19", "chr20"), scales=list(x=list(relation="free")))
```

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hg18track-class   *Class "hg18track" container for hg18 annotation found in genome browser track files; class "chrnum" extends numeric for an indexing application on hg18track objects.*

Description

container for hg18 annotation found in genome browser track files

Objects from the Class

Objects can be created by calls of the form `new("hg18track", assayData, featureData, experimentData, annotation, dataVals, ...)`. These are single-sample eSet instances.

Note that `demoTrk19` is a restriction of the `rawCD4` structure to the interval of chromosome 19 that was assayed in the ENCODE project for DnaseI hypersensitivity.

Slots

`assayData`: Object of class `"AssayData"` ~~

`phenoData`: Object of class `"AnnotatedDataFrame"` ~~

`featureData`: Object of class `"AnnotatedDataFrame"` ~~

`experimentData`: Object of class `"MIAME"` ~~

`annotation`: Object of class `"character"` ~~

`.__classVersion__`: Object of class `"Versions"` ~~

Extends

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

Methods

[`signature(x = "hg18track")`: select using numeric, logical, or chrnum indices.

chrnum `signature(object = "hg18track")`: extract numeric tokens for chromosome number at which data values are obtained; note that chrnum is also used as name of a class.

dataVals `signature(object = "hg18track")`: actual data values

getTrkXY `signature(object = "hg18track", type = "character")`: obtain a list with components x, y indicating location and data value respectively; location is within chromosome; default type is 'midpoint' of locations given as intervals

getTrkXY `signature(object = "hg18track", type = "missing")`: take default midpoint x values corresponding to data values

rangeLocs `signature(object = "hg18track")`: if measures from only one chromosome are present , this returns low and high values of chromStart and chromEnd respectively, otherwise error.

clipTrk `signature(obj = "hg18track", low="numeric", hi="numeric", attr="ANY")`: create a restriction of the track using an interval specification. by default the `chromStart` featureData component is used for coordinates to clip; if `attr` is non-missing, the featureData component named by `attr` will be used.

initialize `signature(.Object = "hg18track")`: create a new instance

Author(s)

VJ Carey <stvjc@channing.harvard.edu>

Examples

```
showClass("hg18track")
data(rawCD4)
rawCD4
rawCD4.chr1 = rawCD4[ chrnum(1), ]
rangeLocs(rawCD4.chr1)
plot(getTrkXY(rawCD4.chr1), ylab="data value", xlab="interval midpt on chr 1" )
c52 = clipTrk(rawCD4[ chrnum(5), ], 1.30e8, 1.33e8 )
plot(getTrkXY(c52))
```

---

juxtaPlot	*two-panel plot with track info and snp screen t-values*

Description

two-panel plot with track info and snp screen t-values

Usage

```
juxtaPlot(trk, ssr)
```

Arguments

trk	instance of [hg18track](#)
ssr	instance of **GGtools** `snpScreenResult`

Details

`xyplot` of lattice package is used.

Value

`xyplot` output; use `print` in Sweave.

Author(s)

VJ Carey <stvjc@channing.harvard.edu>

Examples

```
data(sOSR2)
data(c19g)
juxtaPlot(c19g, sOSR2)
```

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