

BCRANK

April 19, 2009

BCRANKmatch-class *Class "BCRANKmatch"*

Description

Holds the [bcrank](#) score for one IUPAC consensus sequence. Several objects of this class are collected in a [BCRANKsearch-class](#) object

Objects from the Class

Objects are not intended to be created directly but as a result from running [bcrank](#).

Slots

consensus: consensus sequence in IUPAC coding

bcrankScore: bcrank score for the consensus

Methods

consensus signature(object = "BCRANKmatch"): Returns the consensus sequence.

bcrankScore signature(object = "BCRANKmatch"): Returns the bcrank score.

matchVector signature(object = "BCRANKmatch"): Returns a vector with 0's (no match) and 1's (match) of same length and order as the ranked DNA sequences.

Author(s)

Adam Ameer, <adam.ameur@lcb.uu.se>

See Also

[bcrank](#), [BCRANKsearch-class](#)

BCRANKout

BCRANK results for USF1 ChIP-chip data

Description

Results from running `bcrank` on USF1 whole genome ChIP-chip data for the human liver cell line HepG2.

Usage

```
data(BCRANKout)
```

Source

Data from whole genome ChIP-chip experiments on human liver cell line HepG2. (Rada-Iglesias, A., et al. 2007)

References

Rada-Iglesias, A., et al. (2007) Whole-genome maps of USF1 and USF2 binding and histone H3 acetylation reveal new aspects of promoter structure and candidate genes for common human disorders. *Genome Research*, Accepted

See Also

[bcrank](#)

BCRANKresult-class *Class "BCRANKresult"*

Description

Holds the results from running `bcrank`. Contains a number of `bcrank` object, one for each restart of the `bcrank` search.

Slots

fname: The name of the fasta file used for running `bcrank`.

toplist: A list of `BCRANKsearch-class` objects, ranked by their scores.

Methods

fname signature(object = "BCRANKmatch"): Returns the fasta file name.

toplist signature(object = "BCRANKmatch", i=NULL): If `i` is NULL, returns a data frame containing consensus and score for the results for each restart of the `bcrank` search. Otherwise, the `i`'th `BCRANKsearch-class` object in the `toplist` is returned.

Author(s)

Adam Ameur, adam.ameur@lcb.uu.se

See Also

[bcrank](#), [BCRANKsearch-class](#),

BCRANKsearch-class *Class "BCRANKsearch"*

Description

Holds the whole search path from a single [bcrank](#) run. Each individual search step is stored in a [BCRANKmatch-class](#) object. Several objects of this class are collected in a [BCRANKresult-class](#) object

Objects from the Class

Objects are not intended to be created directly but as a result from running [bcrank](#).

Slots

searchPath: a collection of [BCRANKmatch-class](#) objects, containing all [bcrank](#) search steps from a start guess to a locally optimal solution.

finalPWM: position weight matrix containing all DNA sites for the highest scoring consensus in this [bcrank](#) run.

Methods

searchPath signature(object = "BCRANKsearch", i=NULL): If *i* is NULL, returns a data frame containing consensus and score for the whole search path. Otherwise, the *i*'th [BCRANKmatch-class](#) object in the search path is returned.

pwm signature(object = "BCRANKsearch", normalize=TRUE): Returns the position weight matrix (pwm) for the highest scoring consensus in this [bcrank](#) run. Matrix positions are between 0 and 1 when *normalize* is TRUE. When FALSE, the number of matching sequences is reported.

plot signature(x = "BCRANKsearch", y = "missing"): A plot method for the search-Path.

Author(s)

Adam Ameer, <adam.ameur@lcb.uu.se>

See Also

[bcrank](#), [BCRANKmatch-class](#), [BCRANKresult-class](#)

bcrank

*BCRANK: predicting binding site consensus from ranked DNA sequences***Description**

This function implements an algorithm for detection of short DNA sequences that are overrepresented in some part of the list. Starting from some initial consensus DNA sequence coded in IUPAC symbols, the method uses a heuristic search to improve the consensus until a local optimum is found. Individual predicted binding sites can be reported by the function `matchingSites`.

Usage

```
bcrank(fafile, startguesses=c(), restarts=10, length=10,
       reorderings=500, silent=FALSE, plot.progress=FALSE,
       do.search=TRUE, use.P1=FALSE, use.P2=TRUE, strip.desc=TRUE)
```

Arguments

<code>fafile</code>	a ranked fasta file containing DNA sequences.
<code>startguesses</code>	a character vector with consensus sequences in IUPAC coding to be used as starting sequences in the search. If empty, random start guesses will be generated.
<code>restarts</code>	number restarts of the algorithm when using random start guesses.
<code>length</code>	length of random start guess.
<code>reorderings</code>	number of random reorderings of the DNA sequences performed when calculating score.
<code>silent</code>	reports progress status if FALSE.
<code>plot.progress</code>	if TRUE, the progress is displayed in a plot.
<code>do.search</code>	if FALSE, no search is performed. In that case the start guesses are assigned with scores and reported as results.
<code>use.P1</code>	Use penalty for bases other than A,C,G,T.
<code>use.P2</code>	Use penalty for motifs matching repetitive sequences.
<code>strip.desc</code>	An argument used by the function <code>HYPERLINK(readFASTA())(readFASTA())</code> . It specifies whether or not the ">" marking at the beginning of description lines should be removed.

Value

The method returns an object of class `BCRANKresult-class`.

Author(s)

Adam Ameer, <adam.ameur@lcb.uu.se>

References

Ameer, A., Rada-Iglesias, A., Komorowski, J., Wadelius, C. Novel algorithm and ChIP-analysis identifies candidate functional SNPs. Submitted

See Also

[matchingSites](#), [BCRANKresult-class](#)

Examples

```
## Load example fasta file
fastaFile <- system.file("Exfiles/USF1_small.fa", package = "BCRANK")
## Run BCRANK
## Not run: BCRANKout <- bcrank(fastaFile, restarts=20)

## Show BCRANK results
toptable(BCRANKout)
## The top scoring result
topMotif <- toptable(BCRANKout,1)
## Plot BCRANK search path
plot(topMotif)
## Position Weight Matrix
pwm(topMotif, normalize=FALSE)
```

matchingSites

Report IUPAC consensus occurrences in a fasta file

Description

This function reports all occurrences of a consensus sequence in a fasta file. It can be used to extract transcription factor binding sites predicted by BCRANK or other motif search methods.

Usage

```
matchingSites(fafile, motifSequence, revComp=TRUE, strip.desc=TRUE)
```

Arguments

fafile	a ranked fasta file containing DNA sequences.
motifSequence	a character vector in IUPAC coding representing a DNA sequence.
revComp	set to TRUE if the reverse complement also be matched.
strip.desc	An argument used by the function <code>HYPERLINK(readFASTA())(readFASTA())</code> . It specifies whether or not the ">" marking at the beginning of description lines should be removed.

Value

Returns a data frame with positions, strand and DNA sequence for the matching sites.

Author(s)

Adam Ameer, <adam.ameur@lcb.uu.se>

References

Ameer, A., Rada-Iglesias, A., Komorowski, J., Wadelius, C. Novel algorithm and ChIP-analysis identifies candidate functional SNPs. Submitted

See Also[bcrank](#)

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