

# Package ‘motifTestR’

December 19, 2024

**Title** Perform key tests for binding motifs in sequence data

**Version** 1.3.1

**Description** Taking a set of sequence motifs as PWMs, test a set of sequences for over-representation of these motifs, as well as any positional features within the set of motifs. Enrichment analysis can be undertaken using multiple statistical approaches. The package also contains core functions to prepare data for analysis, and to visualise results.

**License** GPL-3

**Encoding** UTF-8

**URL** <https://github.com/smped/motifTestR>

**BugReports** <https://github.com/smped/motifTestR/issues>

**Depends** Biostrings, GenomicRanges, ggplot2 (>= 3.5.0), R (>= 4.3.0),

**Imports** GenomeInfoDb, graphics, harmonicmeanp, IRanges, matrixStats, methods, parallel, patchwork, rlang, S4Vectors, stats, universalmotif

**Suggests** AnnotationHub, BiocStyle, BSgenome.Hsapiens.UCSC.hg19, extraChIPs, ggdendro, knitr, MotifDb, rmarkdown, rtracklayer, testthat (>= 3.0.0)

**biocViews** MotifAnnotation, ChIPSeq, ChipOnChip, SequenceMatching, Software

**LazyData** false

**RoxygenNote** 7.3.2

**Roxygen** list(markdown = TRUE)

**Config/testthat/edition** 3

**VignetteBuilder** knitr

**git\_url** <https://git.bioconductor.org/packages/motifTestR>

**git\_branch** devel

**git\_last\_commit** 8bd1db5

**git\_last\_commit\_date** 2024-11-01

**Repository** Bioconductor 3.21

**Date/Publication** 2024-12-18

**Author** Stevie Pederson [aut, cre] (ORCID:  
<<https://orcid.org/0000-0001-8197-3303>>)

**Maintainer** Stevie Pederson <[stephen.pederson.au@gmail.com](mailto:stephen.pederson.au@gmail.com)>

## Contents

motifTestR-package . . . . .	2
ar_er_peaks . . . . .	3
ar_er_seq . . . . .	4
clusterMotifs . . . . .	5
countPwmMatches . . . . .	6
ex_pfm . . . . .	7
getClusterMatches . . . . .	8
getPwmMatches . . . . .	10
hg19_mask . . . . .	11
makeRMRanges . . . . .	12
plotMatchPos . . . . .	14
testClusterEnrich . . . . .	16
testClusterPos . . . . .	17
testMotifEnrich . . . . .	19
testMotifPos . . . . .	22
zr75_enh . . . . .	24
<b>Index</b>	<b>25</b>

---

motifTestR-package	<i>motifTestR: Perform Key Analyses on Transcription Factor Binding Motifs</i>
--------------------	--

---

## Description

The package motifTestR has been designed for two primary analyses of TFBMs, testing for positional bias and overall enrichment.

## Details

The package motifTestR provides two primary functions for testing TFBMs within a set of sequences

- `testMotifPos()` for detecting positional bias within a set of test sequences
- `testMotifEnrich()` for testing overall enrichment of a TFBM within a set of test sequences

Motifs are also able to be clustered for analysis as a cluster, or for grouping results. Clusters from external approaches can also be incorporated.

- `testClusterPos()` for detecting positional bias for matches to any motif annotated to a cluster, within a set of test sequences
- `testClusterEnrich()` for testing overall enrichment of any TFBM annotated to a cluster, within a set of test sequences

The main functions rely on lower-level functions such as:

- `countPwmMatches()` simply counts the number of matches within an `XStringSet`
- `getPwmMatches()` returns the position of matches within an `XStringSet`
- `countClusterMatches()` simply counts the number of matches to motifs annotated to a cluster within an `XStringSet`
- `getClusterMatches()` returns the position of matches to motifs annotated to a cluster within an `XStringSet`
- `makeRMRanges()` which produces a set of random, matching ranges based on key characteristics of the set of test sequences/ranges

A simple utility function is provided to enable visualisation of results

- `plotMatchPos()` enables visualisation of the matches within a set of sequences using multiple strategies

#### Author(s)

Stevie Pederson

#### See Also

Useful links:

- <https://github.com/smped/motifTestR>
- Report bugs at <https://github.com/smped/motifTestR/issues>

---

ar\_er\_peaks

*A set of peaks with AR and ER detected*

---

#### Description

A set of ChIP-Seq peaks where AR and ER were both detected

#### Usage

```
data("ar_er_peaks")
```

#### Format

An object of class `GRanges` of length 849.

**Details**

The subset of peaks found on chr1 which contained signal from at least two of AR, ER and H3K27ac, taken from GSE123767. Peaks were resized to a uniform width of 400bp after downloading

Generation of these ranges is documented in `system.file("scripts/ar_er_peaks.R", package = "motifTestR")`

**Source**

<https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE123767>

**Examples**

```
data("ar_er_peaks")
ar_er_peaks
```

---

ar\_er\_seq

*Sequences from peaks with AR and ER detected*

---

**Description**

The genomic sequences obtained from the ar\_er\_peaks

**Usage**

```
data("ar_er_seq")
```

**Format**

An object of class DNASTringSet of length 849.

**Details**

These sequences represent the sequences obtained from BSgenome.Hsapiens.UCSC.hg19 for the peaks supplied as ar\_er\_peaks

Generation of these sequences is documented in `system.file("scripts/ar_er_peaks.R", package = "motifTestR")`

**Examples**

```
data("ar_er_seq")
ar_er_seq
```

---

clusterMotifs	<i>Assign each motif to a cluster</i>
---------------	---------------------------------------

---

### Description

Cluster related motifs for testing as a group

### Usage

```
clusterMotifs(
  motifs,
  type = c("PPM", "ICM"),
  method = c("PCC", "EUCL", "SW", "KL", "ALLR", "BHAT", "HELL", "SEUCL", "MAN",
    "ALLR_LL", "WEUCL", "WPCC"),
  power = 1,
  agglom = "complete",
  thresh = 0.2,
  plot = FALSE,
  labels = FALSE,
  cex = 1,
  linecol = "red",
  ...
)
```

### Arguments

motifs	A list of universal motifs or a list of PWMs
type	Can be ICM or PPM
method	The method to be used for determining similarity/distances
power	Raise correlation matrices to this power before converting to a distance matrix. Only applied if method is either "PCC" or "WPCC"
agglom	Method to be used for agglomeration by <a href="#">hclust</a>
thresh	Tree heights below which motifs are formed into a cluster
plot	Show tree produced by <a href="#">hclust</a> . If requested the value set by thresh will be shown as a horizontal line
labels, cex	Passed to <a href="#">plot.hclust</a>
linecol	Passed to <a href="#">abline</a> as the argument col
...	passed to <a href="#">compare_motifs</a>

### Details

This builds on [compare\\_motifs](#), enabling the assignment of each PWM to a cluster, and subsequent testing of motifs as a cluster, rather than returning individual results.

Internally all matrices are converted to distance matrices and `hclust` is used to form clusters. By default, options such as "EUCL", "MAN" produce distances, whilst similarity matrices are produced when choosing "PCC" and other correlation based metrics. In these cases, the distance matrix is obtained by taking 1 - similarity.

By default PWM labels are hidden (labels = FALSE), however these can be shown using labels = NULL as explained in [plot.hclust](#).

Raising the threshold will lead to fewer, larger clusters whilst leaving this value low will return a more conservative approach, with more smaller clusters. The final decision as the best clustering strategy is highly subjective and left to the user. Manual inspection of motifs within a cluster can be performed using [view\\_motifs](#), as shown in the vignette.

### Value

Named vector with numeric values representing which cluster each motif has been assigned to.

### Examples

```
# Load the example motifs
data("ex_pfm")

# Return a vector with each motif assigned a cluster
# The default uses Pearson's Correlation Coefficient
clusterMotifs(ex_pfm)

# Preview the settings noting that showing labels can clutter the plot
# with large numbers of motifs. The defaults for Euclidean distance
# show the threshold may need raising
clusterMotifs(ex_pfm, plot = TRUE, labels = NULL, method = "EUCL")
```

---

countPwmMatches

*Count the matches to a PWM within an XStringSet*

---

### Description

Count the matches to a PWM within an XStringSet

### Usage

```
countPwmMatches(
  pwm,
  stringset,
  rc = TRUE,
  min_score = "80%",
  mc.cores = 1,
  ...
)
```

**Arguments**

pwm	A Position Weight Matrix
stringset	An XStringSet
rc	logical(1) Also find matches using the reverse complement of pwm
min_score	The minimum score to return a match
mc.cores	Passed to <a href="#">mclapply</a> when analysing a list of PWMs
...	Passed to <a href="#">countPWM</a>

**Details**

Will simply count the matches within an XStringSet and return an integer. All matches are included.

**Value**

An integer vector

**Examples**

```
## Load the example PWM
data("ex_pfm")
esr1 <- ex_pfm$ESR1

## Load the example Peaks
data("ar_er_seq")
countPwmMatches(esr1, ar_er_seq)

## Count all PWMs
countPwmMatches(ex_pfm, ar_er_seq)
```

---

ex\_pfm

*Example Position Frequency Matrices*

---

**Description**

Example Position Frequency Matrices

**Usage**

```
data("ex_pfm")
```

**Format**

An object of class `list` of length 5.

**Details**

This object contains 5 PFMs taken from HOCOMOCOv11-coreA for examples and testing

Generation of this motif list is documented in `system.file("scripts/ex_pfm.R", package = "motifTestR")`

**Examples**

```
data("ex_pfm")
ex_pfm$ESR1
```

---

getClusterMatches	<i>Find matches from a PWM cluster within an XStringSet</i>
-------------------	---

---

**Description**

Find matches from a PWM cluster within a set of sequences

**Usage**

```
getClusterMatches(
  cl,
  stringset,
  rc = TRUE,
  min_score = "80%",
  best_only = FALSE,
  break_ties = c("all", "random", "first", "last", "central"),
  mc.cores = 1,
  ...
)

countClusterMatches(
  cl,
  stringset,
  rc = TRUE,
  min_score = "80%",
  mc.cores = 1,
  ...
)
```

**Arguments**

cl	A list of Position Weight Matrices, universal motifs, with each element representing clusters of related matrices
stringset	An XStringSet



<code>rc</code>	logical(1) Also find matches using the reverse complement of PWMs in the cluster
<code>min_score</code>	The minimum score to return a match
<code>best_only</code>	logical(1) Only return the best match
<code>break_ties</code>	Method for breaking ties when only returning the best match Ignored when all matches are returned (the default)
<code>mc.cores</code>	Passed to <a href="#">mclapply</a>
<code>...</code>	Passed to <a href="#">matchPWM</a>

### Details

This function extends [getPwmMatches](#) by returning a single set of results for set of clustered motifs. This can help remove some of the redundancy in results returned for highly similar PWMs, such as those in the GATA3 family.

Taking a set of sequences as an XStringSet, find all matches above the supplied score (i.e. threshold) for a list of Position Weight Matrices (PWMs), which have been clustered together as highly-related motifs. By default, matches are performed using the PWMs as provided and the reverse complement, however this can easily be disabled by setting `rc = FALSE`.

The function relies heavily on [matchPWM](#) and [Views](#) for speed.

Where overlapping matches are found for the PWMs within a cluster, only a single match is returned. The motif with the highest relative score ( $\text{score} / \text{maxScore}(\text{PWM})$ ) is selected.

When choosing to return the best match (`best_only = TRUE`), only the match with the highest relative score is returned for each sequence. Should there be tied scores, the best match can be chosen as either the first, last, most central, all tied matches, or choosing one at random (the default).

### Value

Output from `getClusterMatches` will be a list of DataFrames with columns: `seq`, `score`, `direction`, `start`, `end`, `from_centre`, `seq_width`, `motif` and `match`

The first three columns describe the sequence with matches, the score of the match and whether the match was found using the forward or reverse PWM. The columns `start`, `end` and `width` describe the where the match was found in the sequence, whilst `from_centre` defines the distance between the centre of the match and the centre of the sequence being queried. The `motif` column denotes which individual motif was found to match in this position, again noting that when matches overlap, only the one with the highest relative score is returned. The final column contains the matching fragment of the sequence as an XStringSet.

Output from `countClusterMatches` will be a simple integer vector the same length as the number of clusters

### Examples

```
# Load example PFMs
data("ex_pfm")
# Cluster using default settings
cl_ids <- clusterMotifs(ex_pfm)
ex_cl <- split(ex_pfm, cl_ids)
```

```

# Add optional names
names(ex_cl) <- vapply(ex_cl, \(x) paste(names(x), collapse = "/"), character(1))

# Load example sequences
data("ar_er_seq")
# Get all matches for each cluster
getClusterMatches(ex_cl, ar_er_seq)
# Or Just count them
countClusterMatches(ex_cl, ar_er_seq)
# Compare this to individual counts
countPwmMatches(ex_pfm, ar_er_seq)

```

---

getPwmMatches	<i>Find all PWM matches within an XStringSet</i>
---------------	--

---

## Description

Find all PWM matches within a set of sequences

## Usage

```

getPwmMatches(
  pwm,
  stringset,
  rc = TRUE,
  min_score = "80%",
  best_only = FALSE,
  break_ties = c("all", "random", "first", "last", "central"),
  mc.cores = 1,
  ...
)

```

## Arguments

pwm	A Position Weight Matrix, list of PWMs or universal motif list
stringset	An XStringSet
rc	logical(1) Also find matches using the reverse complement of pwm
min_score	The minimum score to return a match
best_only	logical(1) Only return the best match
break_ties	Method for breaking ties when only returning the best match Ignored when all matches are returned (the default)
mc.cores	Passed to <a href="#">mclapply</a> if passing multiple PWMs
...	Passed to <a href="#">matchPWM</a>

**Details**

Taking a set of sequences as an XStringSet, find all matches above the supplied score (i.e. threshold) for a single Position Weight Matrix (PWM), generally representing a transcription factor binding motif. By default, matches are performed using the PWM as provided and the reverse complement, however this can easily be disabled by setting `rc = FALSE`.

The function relies heavily on [matchPWM](#) and [Views](#) for speed.

When choosing to return the best match (`best_only = TRUE`), only the match with the highest score is returned for each sequence. Should there be tied scores, the best match can be chosen as either the first, last, most central, all tied matches, or choosing one at random (the default).

**Value**

A DataFrame with columns: `seq`, `score`, `direction`, `start`, `end`, `from_centre`, `seq_width`, and `match`

The first three columns describe the sequence with matches, the score of the match and whether the match was found using the forward or reverse PWM. The columns `start`, `end` and `width` describe the where the match was found in the sequence, whilst `from_centre` defines the distance between the centre of the match and the centre of the sequence being queried. The final column contains the matching fragment of the sequence as an XStringSet.

When passing a list of PWMs, a list of the above DataFrames will be returned.

**Examples**

```
## Load the example PWM
data("ex_pfm")
esr1 <- ex_pfm$ESR1

## Load the example Peaks
data("ar_er_seq")

## Return all matches
getPwmMatches(esr1, ar_er_seq)

## Just the best match
getPwmMatches(esr1, ar_er_seq, best_only = TRUE)

## Apply multiple PWMs as a list
getPwmMatches(ex_pfm, ar_er_seq, best_only = TRUE)
```

---

hg19\_mask

*Regions from hg19 with high N content*


---

**Description**

A GRanges object with regions annotated as telomeres or centromeres

**Usage**

```
data("hg19_mask")
```

**Format**

An object of class GRanges of length 345.

**Details**

The regions defined as centromeres or telomeres in hg19, taken from AnnotationHub objects "AH107360" and "AH107361". These were combined with regions containing Ns from the UCSC 2bit file, and regions with Ns in the BSgenome.Hsapiens.UCSC.hg19 were retained.

Generation of these ranges is documented in `system.file("scripts/hg19_mask.R", package = "motifTestR")`

**Source**

The package AnnotationHub and <https://hgdownload.cse.ucsc.edu/goldenpath/hg19/bigZips/hg19.fa.masked.gz>

**Examples**

```
data("hg19_mask")
hg19_mask
```

---

makeRMRanges

*Form a set of random, matching ranges for bootstrapping or permuting*

---

**Description**

Form a set of ranges from y which (near) exactly match those in x for use as a background set requiring matching

**Usage**

```
makeRMRanges(x, y, ...)

## S4 method for signature 'GRanges,GRanges'
makeRMRanges(
  x,
  y,
  exclude = GRanges(),
  n_iter = 1,
  n_total = NULL,
  replace = TRUE,
  ...,
```

```

    force_ol = TRUE
  )

## S4 method for signature 'GRangesList,GRangesList'
makeRMRanges(
  x,
  y,
  exclude = GRanges(),
  n_iter = 1,
  n_total = NULL,
  replace = TRUE,
  mc.cores = 1,
  ...,
  force_ol = TRUE,
  unlist = TRUE
)

```

### Arguments

x	GRanges/GRangesList with ranges to be matched
y	GRanges/GRangesList with ranges to select random matching ranges from
...	Not used
exclude	GRanges of ranges to omit from testing
n_iter	The number of times to repeat the random selection process
n_total	Setting this value will over-ride anything set using n_iter. Can be vector of any length, corresponding to the length of x, when x is a GRangesList
replace	logical(1) Sample with or without replacement when creating the set of random ranges.
force_ol	logical(1) Enforce an overlap between every site in x and y
mc.cores	Passed to <a href="#">mclapply</a>
unlist	logical(1) Return as a sorted GRanges object, or leave as a GRangesList

### Details

This function uses the width distribution of the 'test' ranges (i.e. x) to randomly sample a set of ranges with matching width from the ranges provided in y. The width distribution will clearly be exact when a set of fixed-width ranges is passed to x, whilst random sampling may yield some variability when matching ranges of variable width.

When both x and y are GRanges objects, they are implicitly assumed to both represent similar ranges, such as those overlapping a promoter or enhancer. When passing two GRangesList objects, both objects are expected to contain ranges annotated as belonging to key features, such that the list elements in y must encompass all elements in x. For example if x contains two elements named 'promoter' and 'intron', y should also contain elements named 'promoter' and 'intron' and these will be sampled as matching ranges for the same element in x. If elements of x and y are not named, they are assumed to be in matching order.

The default behaviour is to assume that randomly-generated ranges are for iteration, and as such, ranges are randomly formed in multiples of the number of 'test' ranges provided in x. The column iteration will be added to the returned ranges. Placing any number into the n\_total argument will instead select a total number of ranges as specified here. In this case, no iteration column will be included in the returned ranges.

Sampling is assumed to be with replacement as this is most suitable for bootstrapping and related procedures, although this can be disabled by setting replace = FALSE

### Value

A GRanges or GRangesList object

### Examples

```
## Load the example peaks
data("ar_er_peaks")
sq <- seqinfo(ar_er_peaks)
## Now sample size-matched ranges for two iterations from chr1
makeRMRanges(ar_er_peaks, GRanges(sq)[1], n_iter = 2)

## Or simply sample 100 ranges if not planning any iterative analyses
makeRMRanges(ar_er_peaks, GRanges(sq)[1], n_total = 100)
```

---

plotMatchPos

*Plot Motif Match Positions*

---

### Description

Plot the distribution of motif matches across sequences

### Usage

```
plotMatchPos(
  matches,
  binwidth = 10,
  abs = FALSE,
  use_totals = FALSE,
  type = c("density", "cdf", "heatmap"),
  geom = c("smooth", "line", "point", "col"),
  cluster = FALSE,
  w = 0.1,
  heat_fill = NULL,
  ...
)
```

## Arguments

matches	Output from <a href="#">getPwmMatches</a>
binwidth	Width of bins to use when plotting
abs	logical(1) Plot absolute distances from centre
use_totals	logical(1). If TRUE, plots will use total counts. The default (FALSE) plots probabilities.
type	Plot match density, the CDF or a binned heatmap
geom	Type of geom to be used for line plots. Ignored for heatmaps
cluster	logical(1) Cluster motifs when drawing a heatmap. If TRUE a dendrogram will be added to the LHS of the plot
w	Relative width of the dendrogram on (0, 1)
heat_fill	scale_fill_continuous object for heatmaps. If not provided, scale_fill_viridis_c() will be added to the heatmap.
...	Passed to individual geom* functions

## Details

Multiple options are provided for showing the distribution of PWM matches within a set of sequences, using either the smoothed probability density, the probability CDF or as a heatmap. Distances can be shown as symmetrical around the centre or using absolute distances from the central position within the sequences.

Heatmaps are only enabled for comparisons across multiple PWMs, with optional clustering enabled. If adding a dendrogram for clustering, the returned plot object will be a patchwork object.

## Value

A ggplot2 object

## Examples

```
## Load the example PWM
data("ex_pfm")
esr1 <- ex_pfm$ESR1

## Load the example sequences from the peaks
data("ar_er_seq")

## Just the best match
bm <- getPwmMatches(esr1, ar_er_seq, best_only = TRUE)
plotMatchPos(bm, se = FALSE)

## Matches can also be shown by distance from centre
plotMatchPos(bm, abs = TRUE)

## Cumulative Probability plots are also implemented
plotMatchPos(bm, type = "cdf", geom = "line", colour = "red") +
  geom_abline(intercept = 0.5, slope = 1/400)
```

---

testClusterEnrich	<i>Test enrichment across a cluster of motifs using a background set of sequences</i>
-------------------	---

---

### Description

Test for enrichment of any motif within a cluster across a set of sequences using a background set to derive a NULL hypothesis

### Usage

```
testClusterEnrich(
  cl,
  stringset,
  bg,
  var = "iteration",
  model = c("quasipoisson", "hypergeometric", "poisson", "iteration"),
  sort_by = c("p", "none"),
  mc.cores = 1,
  ...
)
```

### Arguments

cl	A list of Position Weight Matrices, universal motifs, with each element representing clusters of related matrices
stringset	An XStringSet with equal sequence widths
bg	An XStringSet with the same sequence widths as the test XStringset
var	A column in the mcols element of bg, usually denoting an iteration number
model	The model used for analysis
sort_by	Column to sort results by
mc.cores	Passed to <a href="#">mclapply</a>
...	Passed to <a href="#">getPwmMatches</a> or <a href="#">countPwmMatches</a>

### Details

This extends the analytic methods offered by [testMotifEnrich](#) using PWMs grouped into a set of clusters. As with all cluster-level approaches, hits from multiple PWMs which overlap are counted as a single hit ensuring that duplicated matches are not double-counted, and that only individual positions within the sequences are.

### Value

See [testMotifEnrich](#)



**See Also**

[makeRMRanges\(\)](#), [getClusterMatches\(\)](#), [countClusterMatches\(\)](#), [testMotifEnrich\(\)](#)

**Examples**

```
## Load the example peaks & the sequences
data("ar_er_peaks")
data("ar_er_seq")
sq <- seqinfo(ar_er_peaks)
## Now sample size-matched ranges 10 times larger. In real-world analyses,
## this set should be sampled as at least 1000x larger, ensuring features
## are matched to your requirements. This example masks regions with known N
## content, including centromeres & telomeres
data("hg19_mask")
set.seed(305)
bg_ranges <- makeRMRanges(
  ar_er_peaks, GRanges(sq)[1], exclude = hg19_mask, n_iter = 10
)

## Convert ranges to DNASTringSets
library(BSgenome.Hsapiens.UCSC.hg19)
genome <- BSgenome.Hsapiens.UCSC.hg19
bg_seq <- getSeq(genome, bg_ranges)

## Test for enrichment of clustered motifs
data("ex_pfm")
cl <- list(A = ex_pfm[1], B = ex_pfm[2:3])
testClusterEnrich(cl, ar_er_seq, bg_seq, model = "poisson")
```

---

testClusterPos

*Test positional bias motifs within a cluster*


---

**Description**

Test positional bias for all motifs within a given cluster

**Usage**

```
testClusterPos(
  x,
  stringset,
  binwidth = 10,
  abs = FALSE,
  rc = TRUE,
  min_score = "80%",
  break_ties = "all",
  alt = c("greater", "less", "two.sided"),
```

```

    sort_by = c("p", "none"),
    mc.cores = 1,
    ...
)

```

## Arguments

x	A Position Weight Matrix, universalmotif object or list thereof. Alternatively can be a single DataFrame or list of DataFrames as returned by <a href="#">getClusterMatches</a> with best_only = TRUE
stringset	An XStringSet. Not required if matches are supplied as x
binwidth	Width of bins across the range to group data into
abs	Use absolute positions around zero to find symmetrical enrichment
rc	logical(1) Also find matches using the reverse complement of each PWM
min_score	The minimum score to return a match
break_ties	Choose how to resolve matches with tied scores
alt	Alternative hypothesis for the binomial test
sort_by	Column to sort results by
mc.cores	Passed to <a href="#">mclapply</a>
...	Passed to <a href="#">matchPWM</a>

## Details

This is a reimplementaion of [testMotifPos](#) for sets of motifs which have been clustered for similarity. The positions test the bias of any motifs within the cluster given that overlapping matches are only counted once, and with the match retained being the one with the highest relative score.

It should also be noted that some motif clusters will contain PWMs of varying length. When finding positional bias, the widest motif is taken as the width for all, and any matches from narrower motifs outside of the range allowed by wider motifs are discarded. This reduction in signal will make a small difference in the outer bins, but is not considered to be problematic for the larger analysis.

## Value

A data.frame with columns start, end, centre, width, total\_matches, matches\_in\_region, expected, enrichment, prop\_total, p and consensus\_motif The total matches represent the total number of matches within the set of sequences, whilst the number observed in the final region are also given, along with the proportion of the total this represents. Enrichment is simply the ratio of observed to expected based on the expectation of the null hypothesis

The consensus motif across all matches is returned as a Position Frequency Matrix (PFM) using [consensusMatrix](#).

**Examples**

```
## Load the example PWM
data("ex_pfm")
## Load the example sequences
data("ar_er_seq")

## Cluster the motifs
cl <- list(A = ex_pfm[1], B = ex_pfm[2:3])

## Get the best match and use this data
matches <- getClusterMatches(cl, ar_er_seq, best_only = TRUE)
## Test for enrichment in any position
testClusterPos(matches)

## Or just pass the clustered matrices
## Here we've set abs = TRUE to test absolute distance from the centre
testClusterPos(cl, ar_er_seq, abs = TRUE, binwidth = 10)
```

---

testMotifEnrich	<i>Test motif enrichment using a background set of sequences</i>
-----------------	--

---

**Description**

Test for motif enrichment within a set of sequences using a background set to derive a NULL hypothesis

**Usage**

```
testMotifEnrich(
  pwm,
  stringset,
  bg,
  var = "iteration",
  model = c("quasipoisson", "hypergeometric", "poisson", "iteration"),
  sort_by = c("p", "none"),
  mc.cores = 1,
  ...
)
```

**Arguments**

pwm	A Position Weight Matrix or list of PWMs
stringset	An XStringSet with equal sequence widths
bg	An XStringSet with the same sequence widths as the test XStringset
var	A column in the mcols element of bg, usually denoting an iteration number
model	The model used for analysis

sort_by	Column to sort results by
mc.cores	Passed to <a href="#">mclapply</a>
...	Passed to <a href="#">getPwmMatches</a> or <a href="#">countPwmMatches</a>

## Details

This function offers four alternative models for assessing the enrichment of a motif within a set of sequences, in comparison to a background set of sequences. Selection of the BG sequences plays an important role and, in conjunction with the question being addressed, determines the most appropriate model to use for testing, as described below.

It should also be noted that the larger the BG set of sequences, the larger the computational burden, and results can take far longer to return. For many millions of background sequences, this may run beyond an hour

### Descriptions of Models and Use Cases:

#### *Hypergeometric Tests:*

Hypergeometric tests are best suited to the use case where the test set of sequences represents a subset of a larger set, with a specific feature or behaviour, whilst the BG set may be the remainder of the set without that feature. For example, the test set may represent ChIP-Seq binding sites where signal changes in response to treatment, whilst the BG set represents the sites where no changed signal was observed. Testing is one-sided, for enrichment of motifs within the test set.

Due to these relatively smaller sized datasets, setting model = "hypergeometric", will generally return results quickly

#### *Poisson Tests:*

This approach requires a set of background sequences which should be much larger than the test set of sequences. The parameters for a Poisson model are estimated in a per-sequence manner on the set of BG sequences, and the observed rate of motif-matches within the test set is then tested using [poisson.test](#).

This approach assumes that all matches follow a Poisson distribution, which is often true, but data can also be over-dispersed. Given that this model can also return results relatively quickly, is it primarily suitable for data exploration, such as quickly checking for expected behaviours, but not for final results.

#### *Quasi-Poisson Test:*

The quasipoisson model allows for over-dispersion and will return more conservative results than using the standard Poisson model. Under the method currently implemented here, BG sequences should be divided into blocks (i.e. iterations), identical in size to the test set of sequences. Model parameters are estimated per iteration across the BG set of sequences, with the rate of matches in the test set being compared against these blocks. This ensures more conservative results that if analysing test and bg sequences as collections of individual sequences.

It is expected that the BG set will be matched for the features of interest and chosen using [makeRMRanges](#) with a large number of iterations, e.g. n\_iter = 1000. Due to this parameterisation, quasipoisson approaches can be computationally time-consuming, as this is effectively an iterative approach.

#### *Iteration:*

Setting the model as "iteration" performs a non-parametric analysis, with the exception of returning Z-scores under the Central Limit Theorem. Mean and SD of matches is found for each

iteration, and used to return Z scores, with p-values returned from both a Z-test and from comparing observed values directly to sampled values obtained from the BG sequences. Sampled values are calculated directly and as such, are limited in precision.

As for the QuasiPoisson model, a very large number of iterations is expected to be used, to ensure the CLT holds, again making this a computationally demanding test. Each iteration/block is expected to be identically-sized to the test set, and matched for any features as appropriate using `makeRMRanges()`.

### Value

A data.frame with columns: sequences, matches, expected, enrichment, and p, with additional columns Z, est\_bg\_rate (Poisson), odds\_ratio (Hypergeometric) or Z, sd\_bg, n\_iter and iter\_p (Iterations). The numbers of sequences and matches refer to the test set of sequences, whilst expected is the expected number of matches under the Poisson or iterative null distribution. The ratio of matches to expected is given as enrichment, along with the Z score and p-value. Whilst the Z score is only representative under the Poisson model, it is used to directly estimate p-values under the iterative approach. Under this latter approach, the sd of the matches found in the background sequences is also given, along with the number of iterations and the p-values from permutations testing the one-sided hypothesis hypothesis for enrichment.

It may also be worth noting that if producing background sequences using `makeRMRanges` with `replace = TRUE` and `force_ol = TRUE`, the iterative model corresponds to a bootstrap, given that the test sequences will overlap the background sequences and background ranges are able to be sampled with replacement.

### See Also

`makeRMRanges()`, `getPwmMatches()`, `countPwmMatches()`

### Examples

```
## Load the example peaks & the sequences
data("ar_er_peaks")
data("ar_er_seq")
sq <- seqinfo(ar_er_peaks)
## Now sample size-matched ranges 10 times larger. In real-world analyses,
## this set should be sampled as at least 1000x larger, ensuring features
## are matched to your requirements. This example masks regions with known N
## content, including centromeres & telomeres
data("hg19_mask")
set.seed(305)
bg_ranges <- makeRMRanges(
  ar_er_peaks, GRanges(sq)[1], exclude = hg19_mask, n_iter = 10
)

## Convert ranges to DNASTringSets
library(BSgenome.Hsapiens.UCSC.hg19)
genome <- BSgenome.Hsapiens.UCSC.hg19
bg_seq <- getSeq(genome, bg_ranges)

## Test for enrichment of the ESR1 motif
data("ex_pfm")
```

```

esr1 <- ex_pfm$ESR1
testMotifEnrich(esr1, ar_er_seq, bg_seq, model = "poisson")

## Test all motifs
testMotifEnrich(ex_pfm, ar_er_seq, bg_seq, model = "poisson")

```

---

testMotifPos

*Test for a Uniform Distribution across a set of best matches*


---

## Description

Test for a Uniform Distribution across a set of best matches

## Usage

```

testMotifPos(
  x,
  stringset,
  binwidth = 10,
  abs = FALSE,
  rc = TRUE,
  min_score = "80%",
  break_ties = "all",
  alt = c("greater", "less", "two.sided"),
  sort_by = c("p", "none"),
  mc.cores = 1,
  ...
)

```

## Arguments

x	A Position Weight Matrix, universalmotif object or list thereof. Alternatively can be a single DataFrame or list of DataFrames as returned by <a href="#">getPwmMatches</a> with best_only = TRUE
stringset	An XStringSet. Not required if matches are supplied as x
binwidth	Width of bins across the range to group data into
abs	Use absolute positions around zero to find symmetrical enrichment
rc	logical(1) Also find matches using the reverse complement of each PWM
min_score	The minimum score to return a match
break_ties	Choose how to resolve matches with tied scores
alt	Alternative hypothesis for the binomial test
sort_by	Column to sort results by
mc.cores	Passed to <a href="#">mclapply</a>
...	Passed to <a href="#">matchPWM</a>

## Details

This function tests for an even positional spread of motif matches across a set of sequences, using the assumption (i.e.  $H_0$ ) that if there is no positional bias, matches will be evenly distributed across all positions within a set of sequences. Conversely, if there is positional bias, typically but not necessarily near the centre of a range, this function intends to detect this signal, as a rejection of the null hypothesis.

Input can be provided as the output from [getPwmMatches](#) setting `best_only = TRUE` if these matches have already been identified. If choosing to provide this object to the argument `matches`, nothing is required for the arguments `pwm`, `stringset`, `rc`, `min_score` or `break_ties`. Otherwise, a Position Weight Matrix (PWM) and an XStringSet are required, along with the relevant arguments, with best matches identified within the function.

The set of best matches are then grouped into bins along the range, with the central bin containing zero, and tallied. Setting `abs` to `TRUE` will set all positions from the centre as *absolute values*, returning counts purely as bins with distances from zero, marking this as an inclusive lower bound. Motif alignments are assigned into bins based on the central position of the match, as provided in the column `from_centre` when calling [getPwmMatches](#).

The `binom.test` is performed on each bin using the alternative hypothesis, with the returned p-values across all bins combined using the Harmonic Mean p-value (HMP) (See [p.hmp](#)). All bins with raw p-values below the HMP are identified and the returned values for `start`, `end`, `centre`, `width`, `matches` in region, `expected` and `enrichment` are across this set of bins. The expectation is that where a positional bias is evident, this will be a narrow range containing a non-trivial proportion of the total matches.

It should also be noted that `binom.test()` can return p-values of zero, as beyond machine precision. In these instances, zero p-values are excluded from calculation of the HMP. This will give a very slight conservative bias, and assumes that for these extreme cases, neighbouring bins are highly likely to also return extremely low p-values and no significance will be lost.

## Value

A data.frame with columns `start`, `end`, `centre`, `width`, `total_matches`, `matches_in_region`, `expected`, `enrichment`, `prop_total`, `p` and `consensus_motif`. The total matches represent the total number of matches within the set of sequences, whilst the number observed in the final region are also given, along with the proportion of the total this represents. Enrichment is simply the ratio of observed to expected based on the expectation of the null hypothesis.

The consensus motif across all matches is returned as a Position Frequency Matrix (PFM) using [consensusMatrix](#).

## Examples

```
## Load the example PWM
data("ex_pfm")
esr1 <- ex_pfm$ESR1

## Load the example sequences
data("ar_er_seq")

## Get the best match and use this data
```

```
matches <- getPwmMatches(esr1, ar_er_seq, best_only = TRUE)
## Test for enrichment in any position
testMotifPos(matches)

## Provide a list of PWMs, testing for distance from zero
testMotifPos(ex_pfm, ar_er_seq, abs = TRUE, binwidth = 10)
```

---

zr75\_enh

*Candidate Enhancer Regions from ZR-75-1 Cells*

---

### Description

The chr1 subset of candidate enhancers for ZR-75-1 cells

### Usage

```
data("zr75_enh")
```

### Format

An object of class GRanges of length 5237.

### Details

These enhancers are the chr1 subset of enhancer regions for ZR-75-1 cells as identified by EnhancerAtlas 2.0

```
#' Generation of these ranges is documented in system.file("scripts/zr75_enh.R", package = "motifTestR")
```

### Source

<http://www.enhanceratlas.org/index.php>

### Examples

```
data("zr75_enh")
zr75_enh
```



# Index

- \* **datasets**
  - ar\_er\_peaks, 3
  - ar\_er\_seq, 4
  - ex\_pfm, 7
  - hg19\_mask, 11
  - zr75\_enh, 24
- \* **internal**
  - motifTestR-package, 2
- abline, 5
- ar\_er\_peaks, 3
- ar\_er\_seq, 4
  
- binom.test, 23
  
- clusterMotifs, 5
- compare\_motifs, 5
- consensusMatrix, 18, 23
- countClusterMatches
  - (getClusterMatches), 8
- countClusterMatches(), 3, 17
- countPWM, 7
- countPwmMatches, 6, 16, 20
- countPwmMatches(), 3, 21
  
- ex\_pfm, 7
  
- getClusterMatches, 8, 18
- getClusterMatches(), 3, 17
- getPwmMatches, 9, 10, 15, 16, 20, 22, 23
- getPwmMatches(), 3, 21
  
- hclust, 5, 6
- hg19\_mask, 11
  
- makeRMRanges, 12, 20, 21
- makeRMRanges(), 3, 17, 21
- makeRMRanges, GRanges, GRanges-method
  - (makeRMRanges), 12
- makeRMRanges, GRangesList, GRangesList-method
  - (makeRMRanges), 12
  
- matchPWM, 9–11, 18, 22
- mclapply, 7, 9, 10, 13, 16, 18, 20, 22
- motifTestR (motifTestR-package), 2
- motifTestR-package, 2
  
- p.hmp, 23
- plot.hclust, 5, 6
- plotMatchPos, 14
- plotMatchPos(), 3
- poisson.test, 20
  
- testClusterEnrich, 16
- testClusterEnrich(), 3
- testClusterPos, 17
- testClusterPos(), 3
- testMotifEnrich, 16, 19
- testMotifEnrich(), 2, 17
- testMotifPos, 18, 22
- testMotifPos(), 2
  
- view\_motifs, 6
- Views, 9, 11
  
- zr75\_enh, 24