

Package ‘spliceSites’

October 12, 2016

Type Package

Title A bioconductor package for exploration of alignment gap positions from RNA-seq data

Version 1.20.0

Date 2012-10-28

Author Wolfgang Kaisers

Maintainer Wolfgang Kaisers <kaisers@med.uni-duesseldorf.de>

Description Performs splice centered analysis on RNA-seq data.

License GPL-2

biocViews RNAseq, GeneExpression, DifferentialExpression, Proteomics

Depends methods, rbamtools (>= 2.14.3), refGenome (>= 1.6.0), Biobase, Biostrings (>= 2.28.0)

Imports BiocGenerics, doBy, seqLogo, IRanges

Collate allClasses.r allGenerics.r c-methods.r dim-methods.r head-methods.r show-methods.r spliceSites.r

NeedsCompilation yes

R topics documented:

| | |
|----------------------------------|----|
| spliceSites-package | 2 |
| aaGapSites-class | 4 |
| addGeneAligns | 5 |
| addHbond | 6 |
| addMaxEnt | 7 |
| alt_X_ranks | 8 |
| annGapSites-class | 9 |
| annotate-ExpressionSet | 11 |
| annotation | 12 |
| as.data.frame-methods | 13 |
| c-methods | 13 |
| caRanges-class | 13 |
| cdRanges-class | 15 |

| | |
|--------------------------------|----|
| countByGeneName | 16 |
| cRanges-class | 17 |
| dim-methods | 19 |
| dnaGapSites-class | 19 |
| dnaRanges | 20 |
| extractByGeneName | 21 |
| extractRange | 22 |
| gapSites | 23 |
| gapSites-class | 25 |
| getGapSites | 27 |
| hbond-class | 28 |
| head-methods | 29 |
| initialize-methods | 29 |
| keyProfiler-class | 30 |
| lrCodons | 31 |
| maxEnt-class | 32 |
| merge-methods | 33 |
| plotGeneAlignDepth | 33 |
| rangeByGeneName | 35 |
| readCuffGeneFpkm | 36 |
| readExpSet | 37 |
| readMergedBamGaps | 38 |
| readTabledBamGaps | 39 |
| seqlogo | 40 |
| silic_try | 41 |
| sortTable-methods | 41 |
| SpliceCountSet-class | 42 |
| trim | 42 |
| truncateSeq | 43 |
| truncate_seq | 44 |
| trypsinCleave | 45 |
| uniqueJuncAnn | 46 |
| write.files | 47 |
| xCodons | 48 |
| xJunc | 49 |
| xJuncStrand | 50 |
| [-methods | 52 |

Index**53**

spliceSites-package *Calculate information on splice-sites from gapped alignments in RNA-seq data.*

Description

The package defines 'cRanges' the (centered ranges) class which represents a genomic range that contains a highlighted position (center): This will usually be the boundary between an exon and an intron. The second defined type is the class 'gapSites' which represents two exonic regions divided by a gap (usually an intron). There are subclasses which additionally contain DNA or AA sequences.

Details

Package: spliceSites
Type: Package
Version: 1.0
Date: 2012-10-28
License: GPL-2
Depends: methods,rbamtools,refGenome,Biobase,BiocGenerics,Biostrings,seqLogo

Author(s)

Wolfgang Kaisers Maintainer: Wolfgang Kaisers <kaisers@med.uni-duesseldorf.de>

References

Yeo G, Burge CB Maximum entropy modeling of short sequence motifs with applications to RNA splicing signals. J Comput Biol 2004; 11(2-3):377-94 http://genes.mit.edu/burgelab/maxent/Xmaxentscan_scoreseq.html

See Also

[rbamtools](#) [refGenome](#)

Examples

```
bam <- system.file("extdata", "rna_fem.bam", package="spliceSites")
reader <- bamReader(bam, idx=TRUE)
ga <- alignGapList(reader)
bamClose(reader)
dnafile <- system.file("extdata", "dna_small.RData", package="spliceSites")
load(dnafile)
ucf <- system.file("extdata", "uc_small_junc.RData", package="spliceSites")
ucj <- loadGenome(ucf)
annotation(ga) <- annotate(ga, ucj)
ga
```

aaGapSites-class *Class "aaGapSites"*

Description

Contains gapAligns data and a AAStringSet.

Objects from the Class

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

Slots

`seq`: "AAStringSet": Contains amino acid sequences.

`nAligns`: "numeric": Contains total number of aligns.

`nAlignGaps`: "numeric": Contains total number of align gaps.

`dt`: "data.frame": Contains data for all gap sites.

Extends

Class "[gapSites](#)", directly.

Methods

head signature(`x = "aaGapSites"`): Returns the first lines of object.

show signature(`object = "aaGapSites"`): Returns the last lines of object.

truncateSeq signature(`x="caRanges"`, `rme=TRUE`, `trunc=42L`): Truncates contained sequence when character (given by ASCII code in `trunc`). The default (42L) encodes for character '*' which indicates stop-codon.

trypsinCleave signature(`x = "caRanges"`, `minLen = 5`): Performs in silico trypsinization of contained sequence. The sequence fragment which contains the (position depicted) exon-intron boundary is returned. Datasets for which the truncated sequence is shorter than `minLen` are excluded.

write.files signature(`x = "caRanges"`): Exports contained data into "csv" file.

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read gap-sites from BAM-file
bam <- system.file("extdata", "rna_fem.bam", package="spliceSites")
reader <- bamReader(bam, idx=TRUE)
ga <- alignGapList(reader)
bamClose(reader)

# B) Load reference dna
dnafile <- system.file("extdata", "dna_small.RData", package="spliceSites")
load(dnafile)

# C) Calculate cross junctional ranges
lrj <- lrJunc(ga, lfeatlen=6, rfeatlen=6, strand='+')
lr1 <- lrCodons(lrj, frame=1, strand='+')
lr2 <- lrCodons(lrj, frame=2, strand='+')
lr3 <- lrCodons(lrj, frame=3, strand='+')
lr <- c(lr1, lr2, lr3)

# D) Add DNA-sequence
lrd <- dnaGapSites(lr, dna_small)

# E) Translate DNA to amino acid
lra <- translate(lrd)
```

| | |
|---------------|--|
| addGeneAligns | <i>Reads a bamRange object for a given bamReader, refGenome and gene name.</i> |
|---------------|--|

Description

Locates gene in genome via refGenome and reads a bamRange from the determined region.

Usage

```
addGeneAligns(x)
```

Arguments

x gapSites. The result contains a copy of the passed object.

Details

The function adds a gene_aligns column to the contained data.frame.

Value

gapSites

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read gapSites
bam <- system.file("extdata", "rna_fem.bam", package="spliceSites")
reader <- bamReader(bam, idx=TRUE)
ga <- alignGapList(reader)
bamClose(reader)

# B) Annotate
ucf <- system.file("extdata", "uc_small_junc.RData", package="spliceSites")
ucj <- loadGenome(ucf)
annotation(ga) <- annotate(ga, ucj)

# C) align part
gal <- addGeneAligns(ga)
gal
```

addHbond

Class "hbond": Provides data and functions for calculation of HBond scores for 5' splice-sites.

Description

The addHbond methods add HBond scores to gapSites and cdRanges objects. HBond scores provide a measure for the capability of a 5' splice-site to form H-bonds with the U1 snRNA. The function requires at least 3 exon nucleotides and 8 intron nucleotides. The first two intron nucleotides are expected to be 'GT' (for other values the returned score will be 0). The routine equally accepts upper and lower case characters.

Usage

```
addHbond(x, dna)
```

Arguments

| | |
|-----|---|
| x | gapSites. The object to which HBond scores are added. |
| dna | DNAStrngSet. Reference sequence identifier. |

Details

In cdRanges objects, the function adds a hbond column. In gapSites objects, the function adds a lhbond (left side) and a rhbond (right side) column. The lhbond values always assume '+'-strand (because HBond works on the 5' side). The rhbond values always assume '-'-strand. Therefore, there will be discrepancies in the output of write.annDNA.tables because the leftseq and rightseq sequences are reverse-complemented according to the strand column: The xhbond may be > 0 without GT at position 4 (but with AC at position 7).

Author(s)

Wolfgang Kaisers

Referenceshttp://www.uni-duesseldorf.de/rna/html/hbond_score.php**Examples**

```
# A) Read gapSites
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)

# B) Load DNA
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)

# C) HBond
gab<-addHbond(ga,dna_small)

# D) cdRanges
lj<-lJunc(ga,featlen=3,gaplen=8,strand='+')
ljd<-dnaRanges(lj,dna_small)
ljdh<-addHbond(ljd)
```

addMaxEnt

addMaxEnt: Extract subset of data contained in given range given object.

Description

addMaxEnt adds new columns to object data which contain MaxEnt-Score derived values. mxe_ps5 contains score5 values for left align-gap (exon-intron) boundary (i.e. assumed to reside on '+'-strand. mxe_ps3 contains score3 (maxent) values for right align-gap (intron-exon) boundary (i.e. assumed to reside on '+'-strand).

mxe_ms5 contains score5 values for right align-gap (exon-intron) boundary on reverseComplement transformed sequence (i.e. assumed to reside on '-'-strand).

mxe_ms3 contains score3 values for left align-gap (intron-exon) boundary on reverseComplement transformed sequence (i.e. assumed to reside on '-'-strand).

From these values, s3strand, s5strand and meStrand are derived: s3strand is '+' when mxe_ps5 >= mxe_ms5 and '-' otherwise; s3strand is '+' when mxe_ps3 >= mxe_ms3 and '-' otherwise.

meStrand equals s5strand when s5strand=s3strand and '*' otherwise.

The function setMeStrand copies existing meStrand values into strand column (and throws an error when meStrand does not exist).

Usage

```
addMaxEnt(x,dna,maxent,digits=1)
```

Arguments

| | |
|--------|---|
| x | gapSites. |
| dna | DNAStrngSet. Reference sequence identifier. |
| maxent | maxEnt. Contains score table which are internally used by score3 and score5 methods. |
| digits | Numeric. Default value: 1. Internally calculated maxent scores are rounded to given number of decimal places. |

Value

gapSites

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read gapSites
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)

# B) Load DNA
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)

# C) maxEnt
mes<-load.maxEnt()
gae<-addMaxEnt(ga,dna_small,mes)
getMeStrand(gae)
sae<-setMeStrand(gae)
```

alt_X_ranks

alt_left_ranks and alt_right_ranks functions: Identification of alternative splicing events from gapped alignments.

Description

alt_X_ranks covers the functions alt_left_ranks and alt_right_ranks. Both functions identify alternative splice-sites. alt_left_ranks finds sites which share the same rstart value (on the same seqid). alt_right_ranks finds sites which share the same lend value (on the same seqid). alt_ranks combines the results of both functions together with seqid, lend and rstart values in one table.

Usage

```
alt_left_ranks(x)
```

Arguments

x gapSites. Object for which alternative ranks are calculated

Details

The function `alt_left_ranks` groups align-gaps (splice-sites) which share identical `rstart` position and have different `lend` position. Each Group is assigned a unique `alt_id` (integer value beginning from 1). The first column in the returned `data.frame` is an `id`-column which facilitates table merging with the source table. The result has the same number of rows as the source and the `id`-column.

Value

`data.frame`. The table contains the columns `nr_alt`, `alt_id`, `id`, `diff_ranks` and `gap_diff`.

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read gapSites
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)

# B) alt_ranks
alr<-alt_left_ranks(ga)
ar<-alt_ranks(ga)
```

annGapSites-class *Class "annGapSites"*

Description

Contains data from align gaps together with annotation data (and optional data about alternative splice positions). Objects of this class are returned from the annotation member function for class `gapSites`.

Details

`plot_diff` plots tabulated distance between inner gap-site border and annotated exon-intron boundaries.

Objects from the Class

Objects can be created by calls of the form `annotation` on `gapSites` objects.

Slots

`nAligns`: Object of class "numeric" Total number of aligns.

`nAlignGaps`: Object of class "numeric" Total number of gapped aligns.

`dt`: "data.frame". Contains gap-positions, annotation data and optional alternative position data.

`annotation`: "data.frame". Contains annotation data.

`profile`: "data.frame". Contains descriptive data for source probes (BAM-files).

Extends

Class "`gapSites`", directly.

Methods

`as.data.frame` signature(`x = "annGapSites"`): Returns the contained data.

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read gapSites from BAM
bam <- system.file("extdata", "rna_fem.bam", package="spliceSites")
reader <- bamReader(bam, idx=TRUE)
ga <- alignGapList(reader)
bamClose(reader)

# B) Load annotation data
ucf <- system.file("extdata", "uc_small_junc.RData", package="spliceSites")
ucj <- loadGenome(ucf)

# C) Add Annotation
annotation(ga) <- annotate(ga, ucj)

# D) Retrieve annotation
aga <- annotation(ga)
aga

# D) plot_diff
aga <- annotation(ga)
plot_diff(aga)
```

`annotate-ExpressionSet`

Adds annotation data to existing ExpressionSet (created by readExpSet)

Description

Reads featureData from incoming Expression set which should contain range data on embedding exons for gap-sites. The annotate function then overlaps the ranges with given annotation data. The result of overlapping is written into a AnnotatedDataFrame.

Arguments

| | |
|--------|---------------|
| object | ExpressionSet |
| genome | refGenome |

Value

AnnotatedDataFrame

Author(s)

Wolfgang Kaisers

Examples

```
# A) Names of BAM-files
bam <- character(2)
bam[1] <- system.file("extdata", "rna_fem.bam", package="spliceSites")
bam[2] <- system.file("extdata", "rna_mal.bam", package="spliceSites")

# B) Experiment Profile
prof <- data.frame(gender=c("f", "m"))
meta <- data.frame(labelDescription=names(prof), row.names=names(prof))
pd<-new("AnnotatedDataFrame", data=prof, varMetadata=meta)

# C) Read ExpressionSet
es <- readExpSet(bam, phenoData=pd)

# D) Annotate ExpressionSet
ucf <- system.file("extdata", "uc_small.RData", package="spliceSites")
uc <- loadGenome(ucf)
juc <- getSplICETable(uc)
ann <- annotate(es, juc)
```

annotation

Annotation functions for gapSites objects

Description

The `annotate` function takes a `gapSites` and a `refGenome` object and returns a list which additionally contains a 'class' attribute 'annotationResult'. The object is intended as input for the annotation member function of class `gapSites`. The annotation member functions act as writing and reading accessor for annotation data inside `gapSites` objects.

Usage

```
annotate(object, junc)
```

Arguments

`object` [gapSites]. Align-gap data for which annotations are provided via overlap.
`junc` [refJunctions]. Object which provides annotated splice site positions.

Details

The annotation reading accessor takes a `gapSites` object and returns a `annAlignGaps` object. The annotation writing accessor takes a `gapSites` and a `annotationResult` object and copies the contained table into the annotation slot of the `gapSites` object.

Value

`annAlignGaps`

Author(s)

Wolfgang Kaisers

Examples

```
# A) Create gapSites object
bam <- system.file("extdata", "rna_fem.bam", package="spliceSites")
reader <- bamReader(bam[1], idx=TRUE)
ga <- alignGapList(reader)
bamClose(reader)

# B) Read refGenome object
ucf <- system.file("extdata", "uc_small_junc.RData", package="spliceSites")
ucj <- loadGenome(ucf)

# C) Add annotation data
annotation(ga) <- annotate(ga, ucj)
```

as.data.frame-methods as.data.frame *Returning content of data.frame.*

Description

Methods for function as.data.frame

Methods

signature(x = "gapSites") Method for 'gapSites'.

signature(x = "annGapSites") Method for 'annGapSites'.

c-methods *Coercing functions c.*

Description

Coerce objects by binding contained data.

Methods

signature(x = "cRanges") Method for 'cRanges'.

signature(x = "gapSites") Method for 'gapSites'.

caRanges-class *Class "caRanges"*

Description

"caRanges" Objects that contain a centered genomic range and amino acid sequences.

Objects from the Class

Objects are usually created from objects of class "cdRanges" by the "translate" function.

Slots

dt: Object of class "data.frame". Contains the columns "seqid", "start", "end", "strand", "position", "id", "frame"

seq: Object of class "AAStringSet". Contains amino-acid-sequence of ranges described in dt.

Extends

Class "[cRanges](#)", directly.

Methods

- c** signature(x = "caRanges"): Generic combining for caRanges objects.
- getSequence** signature(x="caRanges"): Returns contained sequence (DNAStrngSet).
- head** signature(x = "aaGapAligns"): Returns the first lines of object.
- show** signature(object = "aaGapAligns"): Returns the last lines of object.
- truncateSeq** signature(x="caRanges", rme=TRUE, trunc=42L): Truncates contained sequence when character (given by ASCII code in trunc). The default (42L) encodes for character '*' which indicates stop-codon.
- trypsinCleave** signature(x = "caRanges", minLen = 5): Performs in silico trypsinization of contained sequence. The sequence fragment which contains the (position depicted) exon-intron boundary is returned. Datasets for which the truncated sequence is shorter than minLen are excluded.
- write.files** signature(x = "caRanges"): Exports contained data into "csv" file.

Author(s)

Wolfgang Kaisers

See Also

cRanges

Examples

```
# A) Read gapSites object
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
ga
# B) Create cRanges object
lj<-lJunc(ga,featlen=21,gaplen=21,strand='+')
ljc<-lCodons(lj,frame=1,keepStrand=TRUE)
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
# C) Add DNA sequence
cdr<-dnaRanges(ljc,dna_small)
# D) Translate into AA sequence
ar<-translate(cdr)
# E) Truncate and cleave...
tra<-truncateSeq(ar)
tyc<-trypsinCleave(tra)
```

| | |
|----------------|------------------|
| cdRanges-class | Class "cdRanges" |
|----------------|------------------|

Description

"cdRanges" Objects that contain centered Ranges (exon-intron junctions) and dna-sequences.

Objects from the Class

Objects are usually created from "cRanges" with the function "dnaRanges".

Slots

dt: Object of class "data.frame". Contains the columns "seqid", "start", "end", "strand", "position", "id", "frame".

seq: Object of class "DNAStrngSet". Contains the dna-sequence of ranges described in dt.

Extends

Class "cRanges", directly.

Methods

c signature(x = "cdRanges"): Generic combining for cdRanges objects.

getSequence signature(x="cdRanges"): Returns contained sequence (DNAStrngSet).

head signature(x = "cdRanges"): Prints first items from object.

initialize signature(.Object = "cdRanges"): Create an instance of class using new.

seqlogo signature(x = "cdRanges"): Show a seqlogo of contained sequences

translate signature(x = "cdRanges"): Translates dna-sequence into amino-acid-sequence. Returns an object of class "caRanges".

Author(s)

Wolfgang Kaisers

See Also

cRanges

Examples

```
# A) Read gapSites object
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
ga
# B) Create cRanges object
```

```

ljc<-lJunc(ga, featlen=21, gaplen=21, strand='+')
ljc<-lCodons(ljc, frame=1, keepStrand=TRUE)
dnafile<-system.file("extdata", "dna_small.RData", package="spliceSites")
load(dnafile)
# C) Add DNA sequence
cdr<-dnaRanges(ljc, dna_small)
# D) seqLogo ...
seqlogo(cdr)

```

| | |
|-----------------|--|
| countByGeneName | <i>Reads align number for selected gene from multiple BAM-files.</i> |
|-----------------|--|

Description

Opens multiple BAM-files and reads aligns for selected gene for each file. The function counts the tag-selected value which either is a BAM-cigar operation (like "N" or "M") or the total number of aligns.

Usage

```
countByGeneName(object, infiles, idxInfiles=paste(infiles, ".bai", sep=""), gene, tag="N")
```

Arguments

| | |
|------------|--|
| object | Object of class "refGenome" |
| infiles | Vector of BAM-files |
| idxInfiles | (Optional) Vector of BAM-index files. |
| gene | Gene name |
| tag | Character. Passed to (rbamtools) 'bamCountAll' function. Default value is "N". Other accepted values include "nAligns", "M", "I", "D". |

Details

countByGeneName first uses the extractByGeneName and getGenePositions from 'refGenome' in order to calculate coordinates from the given gene name. Then for each given BAM-file name, the functions calls the bamCount function and returns a vector with a count value for each given file. Internally countByGeneName also checks for existing BAM-index file and tries to create index files which do not exist.

Value

Numeric vector. Length equals number of BAM-input files.

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read filenames
ucf<-system.file("extdata","uc_small.RData",package="spliceSites")
uc<-loadGenome(ucf)
bam<-character(2)
bam[1]<-system.file("extdata","rna_fem.bam",package="spliceSites")
bam[2]<-system.file("extdata","rna_mal.bam",package="spliceSites")
# B) count
countByGeneName(uc,bam,gene="WASH7P",tag="N")
countByGeneName(uc,bam,gene="WASH7P",tag="nAligns")
```

cRanges-class

Class "cRanges": Centered ranges.

Description

"cRanges" Objects that contain centered genomic ranges. The center position marks a prominent position inside the range, generally an exon-intron junction. Position values represent the 0-based position of last exon nucleotide.

Objects from the Class

Objects can be created by calls of the form `new("cRanges", seqnames, start, end, width, strand, position, id)`.

Slots

dt: Object of class "data.frame". The data.frame contains the columns id, seqnames, start, end, width, strand and position. Each row contains data for one centered range.

Methods

as.data.frame signature(x = "cRanges"): Returns a copy of the contained data inside a data.frame object.

c signature(x = "cRanges"): Generic combining for cRanges objects.

count signature(x = "cRanges"): Returns the number of contained ranges (number of rows).

dim signature(x = "cRanges"): Returns the dim of the contained data.frame.

dnaRanges signature(x = "cRanges", dnaset="DNAStrngSet", useStrand="logical", removeUnknownStrand=logical): Takes a cRanges object and a DNAStrngSet (a reference sequence) and adds the appropriate DNA sequence to the genomic ranges. Returns a cdRanges object.

end signature(x = "cRanges"): Returns end column of data.

head signature(x = "cRanges", n="numeric", digits="numeric"): Returns first n (default: n=6) lines of contained data.frame.

id signature(x = "cRanges"): Returns id column from contained data.frame.

initialize signature(.Object = "cRanges"): Generic class initialisation method.

lCodons signature(x = "cRanges", frame="numeric", keepStrand="logical"): Returns cRanges object which represents ranges truncated to codon size. When 'keepStrand' is set to FALSE, strand is set to '+'. The intention is that appended DNA sequences which then can be translated into amino acids.

rCodons signature(x = "cRanges", frame="numeric", keepStrand="logical"): Returns cRanges object which represents ranges truncated to codon size. When 'keepStrand' is set to FALSE, strand is set to '+'. The intention is that appended DNA sequences which then can be translated into amino acids.

seqid signature(x = "cRanges"): Returns vector with seqid's.

show signature(object = "cRanges"): Generic print function.

sortTable signature(x="cRanges"): Sort contained tables by seqid, lend and rstart.

start signature(x = "cRanges"): Returns start column from contained data.frame.

strand signature(x = "cRanges"): Returns strand column from contained data.frame.

width signature(x = "cRanges"): Returns width of contained ranges (=end-start+1).

Author(s)

Wolfgang Kaisers

See Also

gapRanges

Examples

```
# A) Create cRanges object from scratch
sq<-factor(c(1,1,2,2,3,3),labels=c("chr1","chr2","chr3"))
st<-c(100,200,100,300,100,400)
en<-c(120,210,110,310,110,410)
pos<-c(2,3,4,5,6,7)
cr<-new("cRanges",seqid=sq,start=st,end=en,position=pos)
cr
seqid(cr)
start(cr)
end(cr)
width(cr)
strand(cr)
id(cr)
lCodons(cr,frame=1,keepStrand=TRUE)
lCodons(cr,frame=1,keepStrand=FALSE)
lCodons(cr,frame=2,keepStrand=TRUE)
rCodons(cr,frame=1,keepStrand=FALSE)
# + + + + + + + + + + + + + + + + + + + + + + + + + + + + #
# B) Intended way to create a cRanges object from BAM data
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
ga
```

```

lj<-lJunc(ga, featlen=3, gaplen=6, strand='+')
lj
# C) ...
table(strand(lj))

```

dim-methods

dim: *Return dimensions of contained data.frame.*

Description

Methods for function dim

Methods

signature(x = "cRanges") Method for 'cRanges'.
signature(x = "gapSites") Method for 'gapSites'.

dnaGapSites-class

Class "dnaGapSites"

Description

dnaGapSites contains all data which is stored in objects of class "gapSites" plus additional DNA sequences in the "seq" slot.

Objects from the Class

Objects are usually created from gapSites via dnaGapSites.

Slots

seq: "DNAStrngSet". Contains DNA sequence.
nAligns: code"numeric". Contains total number of aligns.
nAlignGaps: "numeric". Contains total number of align gaps.
dt: code"data.frame". Contains data on gap-sites.

Extends

Class "[gapSites](#)", directly.

Methods

head signature(x = "dnaGapSites"): Returns head of dt.
seqlogo signature(x = "dnaGapSites"): Prints seq-logo of stored dna-sequence.
show signature(object = "dnaGapSites"): Prints head of dt.
translate signature(x = "dnaGapSites"): Returns an object of class aalignGaps by translating seq into amino acids.

Author(s)

Wolfgang Kaisers

See Also

gapSites

Examples

```

# A) Read gapSites
bam<-system.file("extdata", "rna_fem.bam", package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
# B) Load DNA sequence
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
# C 1) Add DNA
dga<-dnaGapSites(ga,dna_small)
dga
# C 2) Calculate codon positions
lrj<-lrJunc(ga,lfeatlen=6,rfeatlen=6,strand='+')
lrc<-lrCodons(lrj,frame=1,strand='+')
# D) Add DNA sequence and translate
lrd<-dnaGapSites(lrc,dna_small)
lra<-translate(lrd)
lra

```

| | |
|-----------|--|
| dnaRanges | <i>Reads a bamRange object for a given bamReader, refGenome and gene name.</i> |
|-----------|--|

Description

Locates gene in genome via refGenome and reads a bamRange from the determined region.

Usage

```
dnaRanges(x,dnaset,useStrand=TRUE,removeUnknownStrand=TRUE,verbose=TRUE,...)
```

Arguments

| | |
|---------------------|--|
| x | cRanges. Range-data will be copied from this object. |
| dnaset | DNAStrngSet. Contains the reference sequence from which the DNA-sequence is extracted. |
| useStrand | logical. When TRUE, sequences for which strand='-' are reverse-complemented. |
| removeUnknownStrand | logical. When TRUE, sequences for which strand='-' are removed. |
| verbose | logical. Determines amount of console output during routine runtime. |
| ... | Optional additional arguments (currently unused). |

Value

cdRanges

Author(s)

Wolfgang Kaisers

Examples

```

bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
lj<-lJunc(ga,featlen=6,gaplen=6,strand='+')
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
ljd<-dnaRanges(lj,dna_small)
seqlogo(ljd)

```

| | |
|-------------------|--|
| extractByGeneName | <i>extractByGeneName: Extract subset for sites which lie in range(s) defined by given gene list.</i> |
|-------------------|--|

Description

The function takes a 'cRanges' object (or derived) and searches inside of given 'refGenome' object for gene names. From identified gene-name matches genomic target regions can be defined for which in turn the contained sites are extracted.

Usage

```
extractByGeneName(object, geneNames, src, ...)
```

Arguments

| | |
|-----------|---|
| object | gapSites or cRanges (or derived). Object inside which the data is searched for. |
| geneNames | Character. Vector of gene names for which data is to be extracted. |
| src | refGenome. Contains gene annotation (for conversion of gene-name to genomic coordinates). |
| ... | (currently unused) |

Details

The function internally calls 'extractByGeneName' on 'refGenome'. This function also prints out non matching gene names. On the result, the function calls 'getGenePositions' from which the genomic regions can be extracted. For each gene, data is extracted via 'extractRange' and the resulting objects are then concatenated.

Value

Same type as object

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read gapSites from BAM
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
# B) Load DNASTringSet
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
# C) load refGenome
ucf<-system.file("extdata","uc_small.RData",package="spliceSites")
uc<-loadGenome(ucf)
# D) For cRanges
lj<-lJunc(ga,featlen=6,gaplen=3,strand='+')
ljw<-extractByGeneName(lj, geneNames="WASH7P", src=uc)
# E) For cdRanges
ljc<-lCodons(lj,frame=2)
ljcd<-dnaRanges(ljc,dna_small)
ljcdw<-extractByGeneName(ljcd, geneNames="WASH7P", src=uc)
# F) For caRanges
ljca<-translate(ljcd)
ljcaw<-extractByGeneName(ljca, geneNames="WASH7P", src=uc)
# G) For gapSites
lrj<-lRJunc(ga,lfeatlen=6,rfeatlen=6,strand='+')
lrjw<-extractByGeneName(lrj, geneNames="WASH7P", src=uc)
```

extractRange

extractRange: Extract subset from object where records lie in given range.

Description

Searches in object for data which lie inside the given range and returns an object of same type containing extracted data.

Usage

```
extractRange(object, seqid, start, end)
```

Arguments

| | |
|--------|---|
| object | gapSites or cRanges (or derived). Object inside which the data is searched for. |
| seqid | character. Reference sequence identifier. |
| start | numeric. Start position of given range. |
| end | numeric. End position of given range. |

Value

Same type as object

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read gapSites
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
# B) Load refGenome
ucf<-system.file("extdata","uc_small.RData",package="spliceSites")
uc<-loadGenome(ucf)
# C) For gapSites
extractRange(ga,seqid="chr1",start=14000,end=30000)
# D) For cRanges
lj<-lJunc(ga,featlen=3,gaplen=6,strand='+')
extractRange(lj,seqid="chr1",start=14000,end=30000)
```

gapSites

Creating 'gapSites' and 'dnaGapSites' objects.

Description

gapSites creates objects of class gapSites from scratch. dnaGapSites creates objects of class dnaGapSites from gapSites objects.

Usage

```
gapSites(seqid=factor(),lstart=integer(),lend=integer(),
         rstart=integer(),rend=integer(),gaplen,strand,
         nr_aligns=1,nAligns=sum(nr_aligns),
         nAlignGaps=sum(nr_aligns),nProbes=1)
```

Arguments

| | |
|------------|--|
| seqid | Character. Identifies reference sequence. |
| lstart | Coordinates for start of left range. |
| lend | Coordinates for end of left range. Usually exon-intron boundary. |
| rstart | Coordinates for start of right range. Usually exon-intron boundary. |
| rend | Coordinates for end of right range. |
| gaplen | Length of enclosed gap. Should equal rstart-lend-1. |
| strand | + or - or * (for unknown). Default: '*'. |
| nr_aligns | Number of gapped aligns which have the same exon-intron boundaries (lend and rstart) |
| nAligns | Total number of aligns for probeset. |
| nAlignGaps | Total number of gapped aligns for probeset. |
| nProbes | Numeric. Number of probes in which this gapped position is present. |

Details

The intended way to create a gapSites object is to use the alignGapList function which in turn calls the (rBamTools) bamGapList function. When a BAM file almost exclusively contains gapped aligns which sometimes are multiply gapped, possibly the 'nAlignGaps' value is greater than the 'nAligns'. When reading BAM files which contain the complete data of an alignment, usually the 'nAlignGaps' value is about 1/3 of the 'nAligns' value.

Value

An object of class 'gapSites'.

Author(s)

Wolfgang Kaisers

Examples

```
# A) Construct source data from scratch
seqid<-c("chr1","chr1","chr2","chr2","chr2")
lstart<-c(900, 1900,900 ,900, 1900)
lend <-c(1000,2000,1000,1000,2000)
rstart<-c(1100,2100,1100,1200,2100)
rend <-c(1200,2200,1200,1300,2200)
nr_aligns<-c(10,20,30,40,10)

# B) Construct gapSites object
ga<-gapSites(seqid,lstart,lend,rstart,rend,nr_aligns=nr_aligns)
ga

# C) Use gapSites accessors
seqid(ga)
lend(ga)
```



```

rstart(ga)
strand(ga)
gptm(ga)
rpmg(ga)
nAligns(ga)
nAlignGaps(ga)

# D) Create
bam<-system.file("extdata", "rna_fem.bam", package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
ga
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
dga<-dnaGapSites(ga,dna_small)

```

| | |
|----------------|---|
| gapSites-class | <i>Class "gapSites": Container for tabulated alignment gap positions on RNA-seq data.</i> |
|----------------|---|

Description

Contains tabulated data on alignment gaps on RNA - seq data. "getalignGaps(reader,seqid)" reads gapped alignments for the specified seqid from a BAM file (via CRAN rbamtools) into an object of class "gapSites".

Objects from the Class

Objects can be created by calls of the form alignGapList(reader).

Slots

nAligns: Object of class "numeric" Total number of aligns in alignment.

nAlignGaps: Object of class "numeric" Total number of gapped aligns in alignment.

dt: Object of class "data.frame" Table containing basic data for object.

annotation: Object of class "dataFrameOrNULL" Optional data.frame containing annotation data.

profile: dataFrameOrNULL Optional. Contains probe information (Name of BAM-file, group affiliation, number of sites).

Methods

as.data.frame signature(x = "gapSites"): Returns copy of contained data.frame.

c signature(x = "gapSites"): Specialisation of generic combine function.

dim signature(x = "gapSites"): Specialisation of generic dim function.

dnaGapSites signature(x = "gapSites", dnaset="DNAStringSet"): Create dnaGapSites object by adding DNA sequences.

- getAnnStrand** signature(x): Return strand vector based on annotation content.
- getProfile** signature(x): Return profile table (data.frame) which contains BAM-file names, group affiliation and number of Sites.
- gptm** signature(x = "gapSites"): Reading accessor for gptm values.
- head** signature(x = "gapSites"): Specialisation of generic head function.
- lRCodons** signature(x = "gapSites"): Returns gapSites object where lstart and rend positions are truncated toward the next smaller full codon position (used for preparation of translation to amino acid sequence)
- lJunc** signature(x = "gapSites", featlen="numeric", gaplen="numeric", keepStrand="logical" (F
featlen: Number nucleotides of feature (=exon). gaplen: Number of nucleotides of gap (=intron). keepStrand: Values for strand are copied from argument, otherwise all positions are marked as "+". unique: Multiple identical positions (arising from alternative splice sites on the right side) are collapsed to one line (number of sites is counted in "mult"). Position: 0-based position of last exon nucleotide in DNA sequence.
- lRJunc** signature(x = "gapSites", lfeatlen="numeric", rfeatlen="numeric", "strand"): Returns gapSites object where positions are shifted so that given feature length's are present for lstart and rend positions (used as preparatory steps for obtaining sensible seq - logo's on exonic junction regions).
- addGeneAligns** signature(object="gapSites"): Adds number of alignments per gene as new column to alignment gap position table. Annotation tables must be present. Otherwise an error occurs.
- merge** signature(x = "gapSites", y = "ANY"): Specialisation of generic merge (data.frame) function.
- nAligns** signature(object = "gapSites"): Reading accessor for nAligns value.
- nAlignGaps** signature(object = "gapSites"): Reading accessor for nAlignGaps value.
- rpmg** signature(x = "gapSites"): Reading accessor for rpmg values.
- show** signature(object = "gapSites"): Specialisation of generic show function.
- sortTable** signature(x="gapSites"): Sorts all contained tables by seqid, lend and rstart.
- write.annDNA.tables** signature(x="gapSites", dnaset="DNAStrngSet", filename="charac
Writes csv file with gap-positions, annotations and dna-sequence.

Author(s)

Wolfgang Kaisers

See Also

dnaGapSites

Examples

```
bam<-character(2)
bam[1]<-system.file("extdata", "rna_fem.bam", package="spliceSites")
bam[2]<-system.file("extdata", "rna_mal.bam", package="spliceSites")
reader<-bamReader(bam[1], idx=TRUE)
```

```
agl<-alignGapList(reader)
agl
bamClose(reader)

mbs<-readMergedBamGaps(bam)
mbs
getProfile(mbs)
```

getGapSites

Read gapSites

Description

getGapSites and alignGapList read gap-site data from single BAM-files (given as bamReader) and return a gapSites object. getGapSites reads data for one seqid (given as 1-based numeric value). alignGapList reads the whole BAM-file. The functions test for opened reader and initialized index.

Usage

```
getGapSites(reader,seqid,startid=1)
```

Arguments

| | |
|---------|--|
| reader | bamReader (rbamtools). An opened instance of bamReader with initialized index. |
| seqid | Numeric. 1-based index of reference sequence for which gap-sites are to be read. |
| startid | Numeric. Default: 1. Determines start value for id column from which the values are ascending enumerated. startid greater than 1 allow to produce unique values over multiple BAM-files. |

Details

getGapSites internally calls rbamtools::gapList. alignGapList internally calls rbamtools::bamGapList. 'nProbes' values are set to 1.

Value

gapSites

Author(s)

Wolfgang Kaisers

Examples

```

bam<-system.file("extdata", "rna_fem.bam", package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
gal<-getGapSites(reader,1,startid=10)
gal
gal<-alignGapList(reader)
gal

```

hbond-class

Class "hbond"

Description

Provides methods and data for calculation of HBond 5' splice-site scores. HBond scores provide a measure for the capability of a 5' splice-site to form H-bonds with the U1 snRNA. The function requires at least 3 exon nucleotides and 8 intron nucleotides. The hbond function takes a vector DNA sequences and a vector of position (pos) values. The position values represent the 1-based position of the last exon nucleotide. Therefore all position values must be ≥ 3 and the sequence length must be $\geq \text{pos} + 8$.

Details

The first two intron nucleotides must be 'GT' otherwise returned value is 0. All other sequence characters must be in "ATCG" (capitalization does not matter). When any other character (such as N) is found, the function also returns 0.

Creation of hbond objects

Objects can be created by `load.hbond()`.

Slots

ev: Object of class "environment" Contains external score data.

basedir: Object of class "character" Directory from which external data is restored.

Methods

basedir signature(x = "hbond"): Returns basedir value.

basedir<- signature(x = "hbond",value="character"): Sets basedir value.

hbond signature(x = "hbond",seq="character",pos="integer"): Calculates score5 value for seq at given position.

Author(s)

Wolfgang Kaisers

References

http://www.uni-duesseldorf.de/rna/html/hbond_score.php

Examples

```
hb<-load.hbond()
seq<-c("CAGGTGAGTTC","ATGCTGGAGAA","AGGGTGCGGGC","AAGGTAACGTC","AAGGTGAGTTC")
hbond(hb,seq,3)
```

| | |
|--------------|---|
| head-methods | head <i>Return first lines of contained data.frame.</i> |
|--------------|---|

Description

Methods for function head.

Methods

signature(x = "cRanges") Method for 'cRanges'.

signature(x = "aaGapSites") Method for 'aaGapSites'.

signature(x = "cdRanges") Method for 'cdRanges'.

signature(x = "cRanges") Method for 'cRanges'.

signature(x = "dnaGapSites") Method for 'dnaGapSites'.

signature(x = "gapSites") Method for 'gapSites'.

| | |
|--------------------|---|
| initialize-methods | initialize <i>Initializing objects.</i> |
|--------------------|---|

Description

Methods for function initialize

Methods

signature(.Object = "cdRanges") Method for 'cdRanges'.

signature(.Object = "cRanges") Method for 'cRanges'.

signature(.Object = "keyProfiler") Method for 'keyProfiler'.

signature(.Object = "SpliceCountSet") Method for 'SpliceCountSet'.

keyProfiler-class *Class "keyProfiler"*

Description

Internal class that counts occurrence of profile factors (e.g. gender male and female) successively for added key-tables. The columns of the key-tables define the groups (e.g. genomic positions: seqid, start, end) for each all profile factors are counted.

Objects from the Class

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

Slots

ev: Environment: contains the main data of each object. The environment contains the data.frames 'dtb' (key-tabled profiles) and 'prof' (profiles: a table that contains the profile definition for each added key-table) as well as 'groupExpr', an unevaluated Expression which does the data.frame-grouping after addition of a new key-table.

unique: Logical: When true, there can be maximal one table added for each indexed profile

counted: Logical: Stores the information which profile already has been counted. Is only used when 'unique' is 'TRUE'.

useValues: Logical: When TRUE, the object tables the values given together with each key-table, otherwise the profiles are simply counted.

Methods

addKeyTable signature(x = "keyProfiler", keyTable="data.frame", index="numeric", values="numeric"): Adds keyed data to key-table and counts values according to profile (which is defined by index via profile table).

getKeyTable signature(x = "keyProfiler"): Returns key-table.

appendKeyTable signature(x = "keyProfiler", keytable="data.frame", prefix="character", valFactor="numeric"): Appends internal key-table to keytable-argument. A prefix can be added to column-names. A given valFactor is multiplied with the counted values. A given rateFactor causes counted values to be converted into rates (i.e. divided by column-sums and multiplied with rateFactor value. Values are rounded when a digits argument is provided.)

Author(s)

Wolfgang Kaisers

Examples

```
# Loads profile, position data (key) and aggregated values (ku) data.frames
load(system.file("extdata", "key.RData", package="spliceSites"))
# Group positions
kpc<-new("keyProfiler",keyTable=key1[,c("seqid","lend","rstart")],prof=prof)
addKeyTable(kpc,keyTable=key2[,c("seqid","lend","rstart")],
            index=2,values=key2$nAligns)
addKeyTable(kpc,keyTable=key3[,c("seqid","lend","rstart")],
            index=4,values=key3$nAligns)
cp<-appendKeyTable(kpc,ku,prefix="c.")
```

Ircodons

*Ircodon methods***Description**

The Ircodon function works on gapSites objects. gapSites manage data on align-gaps which represent data on RNA splice sites. On the contained ranges the function can have two effects: an upstream frame-shift of 0 to 2 positions and a downstream trim to full codons (i.e. (end-start+1)%3==0). The strand argument controls direction of effects: '+' strand mode means left frame-shift and right truncation. '-' strand mode means right frame-shift and left truncation.

Usage

```
Ircodons(x, frame=1L, strand="+")
```

Arguments

| | |
|--------|---|
| x | gapSites. Object in which codon positions are calculated |
| frame | Numeric. Default is 1. Accepted values are 1,2 or 3. The value causes a frame-shift of size (frame-1). |
| strand | Character or numeric. Default is '+' which is equivalent to 0. Any other value will be interpreted as '-' which is equivalent to 1. |

Details

The function causes an upstream frameshift and a downstream truncation. gapSites objects contain data on gap aligns which represent a related pair of exon-intron boundaries. The returned object is of the same class as the input. Supplemented DNA sequence gapSites objects will omit introns and will represent the 'spliced' DNA around the splice site. Ircodon function is intended to shift coordinates, so that the resulting DNA-sequence can readily be translated in a putative amino-acid sequence which contains the splice-site.

Author(s)

Wolfgang Kaisers

Examples

```
# A) Create gapSites object
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)

# B) lr-Junctions for '+' -strand
lrj<-lrJunc(ga,lfeatlen=6,rfeatlen=6,strand='+')
lr1<-lrCodons(lrj,frame=1)
lr2<-lrCodons(lrj,frame=2)
lr3<-lrCodons(lrj,frame=3)
lr<-c(lr1,lr2,lr3)

# C) lr-Junctions for '-' -strand
lrj<-lrJunc(ga,lfeatlen=6,rfeatlen=6,strand='-')
lr1<-lrCodons(lrj,frame=1)
lr2<-lrCodons(lrj,frame=2)
lr3<-lrCodons(lrj,frame=3)
lr<-c(lr1,lr2,lr3)
```

maxEnt-class

Class "maxEnt"

Description

Provides methods for calculation of Splice-site scores. Both functions (score5 and score3) are intended to work on the '+' strand. score5 scores the 5' side (i.e. the splice donor, left) and the score3 scores the 3' side (i.e. the splice acceptor, right).

Creation of maxEnt objects

Objects can be created by `load.maxEnt()`.

Slots

ev: Object of class "environment" Contains external score data.

basedir: Object of class "character" Directory from which external data is restored.

Methods

basedir signature(x = "maxEnt"): Returns basedir value.

basedir<- signature(x = "maxEnt", value="character"): Sets basedir value.

score5 signature(x = "maxEnt", seq="character", pos="integer"): Calculates score5 value for seq at given position.

scoreSeq5 signature(x="maxEnt", seq="character", frame="integer"): Calculates score5 values for a single sequence and a series of positions (frame).

score3 signature(x = "maxEnt", seq="character", pos="integer", which="character"): Calculates score3 value for seq at given position. Accepted values for which are: "ent", "wmm" and "emm".

scoreSeq3 signature(x = "maxEnt", seq="character", frame="integer", which="character"): Calculates score3 values for a single sequence and a series of positions (frame). Accepted values for which are: "ent", "wmm" and "emm".

Author(s)

Wolfgang Kaisers

Examples

```
mes<-load.maxEnt()
score5(mes, "CCGGTAAGAA", 4) # 9.844127
score3(mes, "CTCTACTACTATCTATCTAGATC", pos=20) # 6.706947

# scoreSeq functions
sq5<-scoreSeq5(mes, seq="ACGGTAAGTCAGGTAAGT")
sq3<-scoreSeq3(mes, seq="TTTTTTTTCTCACTTTTAGAGACTTCATTCTTCTCAAATAGGTT")
```

merge-methods

merge *Merging two objects into one.*

Description

Methods for function merge

Methods

signature(x = "gapSites", y = "ANY") Method for 'gapSites'.

plotGeneAlignDepth

plotGeneAlignDepth: Plots of read alignment depth for genetic regions

Description

The function takes a bamReader and a refGenome object together with a gene name and an optional transcript name and plots the read alignment depth for the region of the gene in the opened BAM file. When transcript data is present, the exonic ranges are added as rectangles on a chromosomal line.

Usage

```
plotGeneAlignDepth(reader, genome,
                   gene=NULL, transcript=NULL,
                   log="y", cex.main=2,
                   col="grey50", fill="grey90", grid=TRUE,
                   box.col="grey20", box.border="grey80")
```

Arguments

| | |
|------------|---|
| reader | bamReader (rbamtools). Must be opened and have initialized index. |
| genome | refGenome. Object which contains genomic annotation data. |
| gene | character. Name of one single gene. |
| transcript | character (optional). Name of one single transcript. |
| log | character. Name of one single gene. |
| cex.main | numeric. Determines size of main title. |
| col | color. A color for align depth line. |
| fill | color. A color for the interior of align depth area. |
| grid | logical. When TRUE, a grid is drawn. |
| box.col | color. A color for the interior of exon rectangles. |
| box.border | color. A color for the border of exon rectangles. |

Details

The function checks for opened bamReader and initialized index. When transcript name is given, the function will plot the positions of the transcript beneath the alignment depth.

Author(s)

Wolfgang Kaisers

Examples

```
# Open bamReader
bam <- system.file("extdata", "rna_fem.bam", package="spliceSites")
reader <- bamReader(bam, idx=TRUE)
# Load annotation data
ucf <- system.file("extdata", "uc_small.RData", package="spliceSites")
uc <- loadGenome(ucf)
plotGeneAlignDepth(reader, uc, gene="WASH7P", transcript="uc001aac.4")
```

| | |
|-----------------|--|
| rangeByGeneName | <i>Reads a bamRange object for a given bamReader, refGenome and gene name.</i> |
|-----------------|--|

Description

Locates gene in genome via refGenome and reads a bamRange from the determined region.

Usage

```
rangeByGeneName(reader, genome, gene, complex=TRUE)
```

Arguments

| | |
|---------|--|
| reader | Object of class (rbamtools) bamReader. The reader must pass isOpen and index.initialized test. |
| genome | Object of class (refgenome) refGenome. |
| gene | Single gene name (character) |
| complex | Logical. Passed to 'bamRange' function. When TRUE, only aligns with nCigar>1 are counted. |

Value

bamRange

Author(s)

Wolfgang Kaisers

Examples

```
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ucf<-system.file("extdata","uc_small.RData",package="spliceSites")
uc<-loadGenome(ucf)
range<-rangeByGeneName(reader,uc,"WASH7P")
size(range)
```

readCuffGeneFpkm *Reads FPKM values into ExpressionSet.*

Description

Opens fpkm_tracking files and collects FPKM values into ExpressionSet. The function is intended to work with genes.fpkm_tracking files. In order to get unique gene identifier, the contained values are grouped and for each gene the maximum FPKM values is selected. There should only be a few hundred multiple occurring genes and the maximum value should give a (slight) underestimation of the real value.

Usage

```
readCuffGeneFpkm(cuff, phenoData, summ="max")
```

Arguments

| | |
|-----------|--|
| cuff | character: Vector of cufflinks files |
| phenoData | AnnotatedDataFrame: Requirement for construction of an ExpressionSet |
| summ | character: Must be either 'max' or 'sum'. A handful of tracking id's occur multiple times due to multiple transcripts which partially are non-overlapping. The summ (summarize) Argument determines the way the multiplerts are handled. |

Value

ExpressionSet

Author(s)

Wolfgang Kaisers

Examples

```
n<-10
cuff <- system.file("extdata", "cuff_files",
  paste(1:n, "genes", "fpkm_tracking", sep="."), package="spliceSites")

## Create Pheno - data
gr <- system.file("extdata", "cuff_files", "groups.csv", package="spliceSites")
groups <- read.table(gr, sep="\t", header=TRUE)
meta <- data.frame(labelDescription=c("gender", "age-group", "location"),
  row.names=c("gen", "agg", "loc"))
phenoData <- new("AnnotatedDataFrame", data=groups, varMetadata=meta)

## Read ExpressionSet
exset <- readCuffGeneFpkm(cuff, phenoData)
```

| | |
|------------|--|
| readExpSet | <i>Reads align number or gptm or rpmg value from all given BAM-files and all identified align gaps into ExpressionSet.</i> |
|------------|--|

Description

Opens multiple BAM-files and reads aligns for selected gene for each file. Number of alignes is counted.

Usage

```
readExpSet(bam, idx, val="nAligns", phenoData, expData)
```

Arguments

| | |
|-----------|--|
| bam | Vector of BAM-files |
| idx | Vector of index files (optional) |
| val | "gptm", "rpmg" or "nAligns". Value type which is written to ExpressionSet matrix (nAligns = read count). |
| phenoData | AnnotatedDataFrame. Each BAM-file must correspond to one identifier. |
| expData | MIAME. Optional. Experiment data which can be added to ExpressionSet |

Value

ExpressionSet

Author(s)

Wolfgang Kaisers

Examples

```
# A) Names of BAM-files
bam <- character(2)
bam[1] <- system.file("extdata", "rna_fem.bam", package="spliceSites")
bam[2] <- system.file("extdata", "rna_mal.bam", package="spliceSites")

# B) Experiment Profile
prof <- data.frame(gender=c("f", "m"))
meta <- data.frame(labelDescription=names(prof), row.names=names(prof))
pd <- new("AnnotatedDataFrame", data=prof, varMetadata=meta)

# C) Read ExpressionSet
es <- readExpSet(bam, phenoData=pd)

# D) Annotate ExpressionSet
ucf <- system.file("extdata", "uc_small_junc.RData", package="spliceSites")
juc <- loadGenome(ucf)
```

```
ann <- annotate(es, juc)
phenoData(es) <- ann
```

readMergedBamGaps *Reads an object of type gapSites using a vector of BAM file names.*

Description

The function takes a vector of BAM-file names and corresponding BAM-index file names. For each given filename, the BAM-file will be opened. The functions uses the bamGapList function (rbamtools) to obtain a data.frame from an bamReader. Values for 'gptm' and 'rpmg' are added. Both are rounded to the number of given digits. The function tests for open connection to BAM-file and for initialized index.

Usage

```
readMergedBamGaps(infiles,idxInfiles=paste(infiles, ".bai", sep=""), digits=3)
```

Arguments

| | |
|------------|---|
| infiles | character. Name of BAM-files to be opened. |
| idxInfiles | character. Name of corresponding BAM-index files. Default: paste(infiles, ".bai", sep="") |
| digits | numeric. gptm and rpmg values will be rounded to the number of decimal places given. |

Value

gapSites

Author(s)

Wolfgang Kaisers

Examples

```
bam<-character(2)
bam[1]<-system.file("extdata", "rna_fem.bam", package="spliceSites")
bam[2]<-system.file("extdata", "rna_mal.bam", package="spliceSites")
mbg<-readMergedBamGaps(bam)
```

readTabledBamGaps *readTabledBamGaps function*

Description

readTabledBamGaps

Usage

```
readTabledBamGaps(infiles, idxInfiles=paste(infiles, ".bai", sep=""), prof, rpmg=TRUE)
```

Arguments

| | |
|------------|--|
| infiles | character. Names of BAM-files. |
| idxInfiles | character. Names of BAM-index files. When given index file is not found, the function attempts to create a BAM-index file with the depicted name. |
| prof | data.frame. Contains group affiliations for each BAM-file. Each column describes an entity by which values are grouped. The row-number in prof must be equal to the number of given BAM-files. The order of BAM infiles and prof defines the group classification for each BAM file. All prof columns must be factors. |
| rpmg | logical. When TRUE, there will be group specific rpmg align-rates be added to the result table |

Details

The function reads gap-align data from all given BAM-files. For each factor level, the number of probes and aligns are counted. When gptm=TRUE also the gptm values are written for each group. The result table contains for each prof factor level 2 (or 3) extra columns.

Value

gapSites

Author(s)

Wolfgang Kaisers

Examples

```
bam<-character(2)
bam[1]<-system.file("extdata", "rna_fem.bam", package="spliceSites")
bam[2]<-system.file("extdata", "rna_mal.bam", package="spliceSites")
prof<-data.frame(gender=c("f", "m"))
rtbg<-readTabledBamGaps(bam, prof=prof, rpmg=TRUE)
rtbg
getProfile(rtbg)
```

| | |
|---------|--|
| seqlogo | <i>seqlogo: Plotting sequence logo for 'cdRanges' and 'dnaGapSites' objects.</i> |
|---------|--|

Description

The function produces a sequence logo plot based on the contained sequences.

Usage

```
seqlogo(x, strand="+", useStrand=TRUE, ...)
```

Arguments

| | |
|-----------|---|
| x | cdRanges or dnaGapSites Object. |
| strand | Character. Determines the subset for which the seqlogo is plotted. This option is only used when useStrand is given as 'TRUE'. |
| useStrand | Logical. Determines whether the given strand information is used. For useStrand=FALSE the plot is made up from all contained sequences. |
| ... | (Currently unused) |

Details

The function fails with an error message when the dataset does not contain any records with the given strand (except useStrand=FALSE).

Value

None

Author(s)

Wolfgang Kaisers

Examples

```
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
lj<-lJunc(ga,featlen=6,gaplen=6,strand='+')
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
ljd<-dnaRanges(lj,dna_small)
seqlogo(ljd)
```

| | |
|------------|----------------------------|
| silic_tryp | <i>silic_tryp function</i> |
|------------|----------------------------|

Description

silic_tryp performs silicon trypsination and returns the fragments to which the position coordinate points. The position value is corrected so that it afterwards points to the same amino-acid as before.

Usage

```
silic_tryp(seq,pos,id)
```

Arguments

| | |
|-----|---|
| seq | Character. Amino-acid sequences which are to be truncated. |
| pos | Numeric. Points to an amino-acid inside the sequence. |
| id | Numeric. An identifier which is copied to the result table. |

Details

The routine implements the "Keil"-rule, where sites are described by the regex "[RK](?!P)". The cut position is between [RK] and the following character. The sequence fragment which contains the exon-intron boundary (depicted by position) is returned. Dependent numeric values are recalculated.

Value

data.frame

Author(s)

Wolfgang Kaisers

Examples

```
silic_tryp(seq="AXKUEMRFG",pos=4)
```

| | |
|-------------------|---|
| sortTable-methods | <i>Sorting contained data with sortTable.</i> |
|-------------------|---|

Description

Sorting tables by key columns.

Methods

signature(x = "cRanges") Method for 'cRanges'. Key columns: seqid, start, end
signature(x = "gapSites") Method for 'gapSites'. Key columns: seqid, lend, rstart

```
SpliceCountSet-class  Class "SpliceCountSet"
```

Description

Directly inherits from ExpressionSet

Objects from the Class

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

Author(s)

Wolfgang Kaisers

Examples

```
# scs<-new("SpliceCountSet")
```

```
trim          trim and resize methods: trim_left, trim_right, resize_left, resize_right
```

Description

The trim and resize functions change number of nucleotides contained in align-gap features (exonic). Trim functions cut feature sizes down to `maxlen`. Resize functions reset all sizes to a fixed value. The functions operate directly on the passed objects. There is no return value.

Usage

```
trim_left(x,maxlen)
```

Arguments

| | |
|---------------------|--|
| <code>x</code> | gapSites. Object from which the lJunc values are calculated. |
| <code>maxlen</code> | Numeric. Maximum number of nucleotides on feature (exon) side of boundary. |

Value

None.

Author(s)

Wolfgang Kaisers

Examples

```

# A) Create gapSites object
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam[1],idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
ga
# B) Trim
trim_left(ga,3)
trim_right(ga,2)
ga
# C) Resize
resize_left(ga,5)
resize_right(ga,6)
ga

```

truncateSeq

truncateSeq method

Description

truncateSeqs amino acid sequences at positions depicted by '*' (stop-codon).

Usage

```
truncateSeq(x, rme=TRUE, trunc=42L)
```

Arguments

| | |
|-------|--|
| x | caRanges. Object in which amino-acid sequences are to be truncated. |
| rme | Logical. Default is TRUE. When TRUE, sites with resulting empty sequence (i.e. stop-codon upstream of the splice position) are removed from dataset. |
| trunc | Integer. ASCII code for character at which truncation should occur. Default value is 42='*' (stop-codon). |

Details

The function truncateSeqs the contained amino acid sequences. When the stop-codon is found on the left side of position, the function returns an empty sequence for that site. The position values for these records are also set to 0.

Value

Object of same class as input.

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read gap-sites from BAM-file
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
# B) Load DNA sequence
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
ucf<-system.file("extdata","uc_small.RData",package="spliceSites")
uc<-loadGenome(ucf)
# C) Calculate codon frame data and add DNA
lj<-lJunc(ga,featlen=21,gaplen=21,strand='+')
ljc<-lCodons(lj,frame=1,keepStrand=TRUE)
cdr<-dnaRanges(ljc,dna_small)
# D) Translate DNA to amino acid and truncate
ar<-translate(cdr)
tra<-truncateSeq(ar)
```

truncate_seq

*truncate_seq function***Description**

truncateSeqs amino acid sequences at positions depicted by '*' (stop-codon).

Usage

```
truncate_seq(seq,pos,id,rme=TRUE,trunc=42L)
```

Arguments

| | |
|-------|---|
| seq | Character. Amino-acid sequences which are to be truncated. |
| pos | Numeric. Points to an amino-acid inside the sequence. |
| id | Numeric. An identifier which is copied to the result table. |
| rme | Logical. Empty sequences are removed when set to TRUE. |
| trunc | Integer. ASCII code for character at which truncation should occur. Default value is 42='*' (stop-codon). |

Details

The function truncateSeqs the contained amino acid sequences. When the stop-codon is found on the left side of position, the function returns an empty sequence for that site. The position values for these records are also set to 0.

Value

data.frame

Author(s)

Wolfgang Kaisers

Examples

```
truncate_seq(seq="ARPX*QR", pos=3)
```

| | |
|---------------|-----------------------------|
| trypsinCleave | <i>trypsinCleave method</i> |
|---------------|-----------------------------|

Description

trypsinCleave cleaves amino acid sequences and returns the fragment which contains the position described by position entry in data.frame.

Usage

```
trypsinCleave(x, minLen=5, ...)
```

Arguments

| | |
|--------|---|
| x | caRanges (aaGapSites). Object in which amino-acid sequences are to be truncated. |
| minLen | Numeric. Default is 5. Data sets where the remaining sequence fragment is shorter than minLen are excluded. |
| ... | Additional arguments which may be passed to the routine (currently unused). |

Details

The routine implements the "Keil"-rule, where sites are described by the regex "[RK](?!P)". The cut position is between [RK] and the following character. The sequence fragment which contains the exon-intron boundary (depicted by position) is returned. Dependent numeric values are recalculated. The returned sequence ends on "[RK]" unless the returned fragment is a sequence suffix.

Value

Same class as given object.

Author(s)

Wolfgang Kaisers

Examples

```

bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
ga
lj<-lJunc(ga,featlen=21,gaplen=21,strand='+')
ljc<-lCodons(lj,frame=1,keepStrand=TRUE)
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
cdr<-dnaRanges(ljc,dna_small)
ar<-translate(cdr)
tra<-truncateSeq(ar)
tyc<-trypsinCleave(tra)

```

uniqueJuncAnn

uniqueJuncAnn method for ExpressionSet

Description

uniqueJuncAnns adds annotation data to ExpressionSet and removes not-matching sites.

Usage

```
uniqueJuncAnn(object,junc,ann=TRUE,...)
```

Arguments

| | |
|--------|---|
| object | ExpressionSet. Object containing gap-site expression data. |
| junc | refJunctions. Object containing splice-junction sites. |
| ann | logical. Default: TRUE. When TRUE the unannotated sites are removed, otherwise the annotated sites are removed. |
| ... | Unused. |

Value

ExpressionSet

Author(s)

Wolfgang Kaisers

Examples

```

# A) Names of BAM-files
bam<-character(2)
bam[1]<-system.file("extdata","rna_fem.bam",package="spliceSites")
bam[2]<-system.file("extdata","rna_mal.bam",package="spliceSites")

# B) Experiment Profile
prof<-data.frame(gender=c("f","m"))
meta<-data.frame(labelDescription=names(prof),row.names=names(prof))
pd<-new("AnnotatedDataFrame",data=prof,varMetadata=meta)

# C) Read ExpressionSet
es<-readExpSet(bam,phenoData=pd)

# D) Annotate ExpressionSet
ucf<-system.file("extdata","uc_small.RData",package="spliceSites")
uc<-loadGenome(ucf)
ucj<-getSpliceTable(uc)

# E) Extract unique annotated junction sites.
uja<-uniqueJuncAnn(es,ucj)

```

write.files

write.files

Description

Writes table data and sequence in separate files.

Usage

```
write.files(x, path, filename,...)
```

Arguments

| | |
|----------|--|
| x | caRanges or aaGapSites object for which data is written. |
| path | Path for writing files. |
| filename | Basic filename to which suffixes are added. |
| ... | Other arguments passed to "write.table". |

Details

There are two files written: A text file with tabulated values from data.frame (separated by ";") and a fasta file which contains the stored dna sequence.

Value

None.

Note

The function tries to create directory 'path' when it does not exist.

Author(s)

Wolfgang Kaisers

Examples

```
# A) Read gap-sites from BAM-files
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
# B) Load DNA sequence
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
# C) Add DNA sequence
lj<-lJunc(ga,featlen=21,gaplen=21,strand='+')
ljc<-lCodons(lj,frame=1,keepStrand=TRUE)
cdr<-dnaRanges(ljc,dna_small)
# D) Translate DNA to amino-acid
ar<-translate(cdr)
# E) Write "ar.csv" and "ar.fa"
# write.files(ar, ".", "ar")
```

xCodons

xCodon methods

Description

The xCodon functions work on cRanges objects. On the contained ranges the function can have two effects: an upstream frame-shift of 0 to 2 positions and a downstream trim to full codons (i.e. (end-start+1)%3==0). The lCodon function acts in '+' strand mode (left frame-shift, right truncation) and the rCodon function acts in '-' strand mode (right frame-shift, left truncation).

Usage

```
lCodons(x, frame=1, keepStrand=TRUE)
```

Arguments

| | |
|------------|--|
| x | cRanges. Object in which codon positions are calculated |
| frame | Numeric. Default is 1. Accepted values are 1,2 or 3. The value causes a frame-shift of size (frame-1). |
| keepStrand | Logical. Default is TRUE. When FALSE, lCodons overwrites strand entries by '+' and rCodons overwrites strand entries by '-'. |

Details

The function causes an upstream frameshift and a downstream truncation. lCodon works with '+'-strand view (left-to-right) and rCodon works with '-'-strand view (right to left). The underlying rationale is: The cRanges object contains ranges around exon-intron boundaries. The boundary itself is marked by the position value. The functions calculate genomic ranges which can be supplemented by the reference DNA-sequence which then can readily be translated into amino-acid sequences. The different values for frame and keepStrand are used to produce all six putative amino-acid sequences for this exon-intron boundary.

Author(s)

Wolfgang Kaisers

Examples

```
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam,idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
dnafile<-system.file("extdata","dna_small.RData",package="spliceSites")
load(dnafile)
ucf<-system.file("extdata","uc_small.RData",package="spliceSites")
uc<-loadGenome(ucf)

lj<-lJunc(ga,featlen=21,gaplen=21,strand='+')
ljc<-lCodons(lj,frame=1,keepStrand=TRUE)
rj<-rJunc(ga,featlen=21,gaplen=21,strand='-')
rjc<-rCodons(rj,frame=1,keepStrand=TRUE)
```

xJunc

xJunc methods: lJunc, rJunc, lrJunc

Description

The term 'xJunc' envelopes three functions: lJunc, rJunc and lrJunc. All three functions take a gapSites object and return ranges which are restricted around align-gap (exon-intron) boundaries. The functions lJunc and rJunc return cRanges objects, the lrJunc function returns a gapSites object.

Usage

```
lJunc(x,featlen,gaplen,unique=FALSE,strand,...)
```

Arguments

| | |
|---------|--|
| x | gapSites. Object from which the lJunc values are calculated. |
| featlen | Numeric. Number of nucleotides on feature (exon) side of boundary. |
| gaplen | Numeric. Number of nucleotides on gap (intron) side of boundary. |

| | |
|--------|---|
| unique | Logical. Default is 'FALSE'. When 'TRUE', the function removes duplicate entries which can be due to alternative splice events. |
| strand | Character. Mandatory. All strand entries are set to the given value. |
| ... | Optional arguments passed additionally to the function (currently unused). |

Details

The functions are intended to provide position information which crosses exon-intron boundaries. Added DNA sequences can be used to produce seqlogs. The functions are intended to be used in advance of xCodons functions. Later on added AA sequences can be used to search for proteins where intronic sequences are retained.

Value

cRanges

Author(s)

Wolfgang Kaisers

Examples

```
# A) Create gapSites object
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam[1],idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
ga

# B) Extract junction data
lj<-lJunc(ga,featlen=6,gaplen=6,strand='+')
ljm<-lJunc(ga,featlen=6,gaplen=6,strand='-')
rj<-rJunc(ga,featlen=6,gaplen=6,strand='+')
rjm<-rJunc(ga,featlen=6,gaplen=6,strand='-')
lrj<-lrJunc(ga,lfeatlen=6,rfeatlen=6,strand='+')
lrjm<-lrJunc(ga,lfeatlen=6,rfeatlen=6,strand='-')
```

xJuncStrand

xJuncStrand methods: lJuncStrand, rJuncStrand, lrJuncStrand

Description

The term 'xJuncStrand' envelopes three functions: lJuncStrand, rJuncStrand and lrJuncStrand. All three functions take a gapSites object and return ranges which are restricted around align-gap (exon-intron) boundaries. The functions lJuncStrand and rJuncStrand return cRanges objects, the lrJuncStrand function returns a gapSites object. The resulting objects contain strand information which is copied from the input objects.

Usage

```
lJuncStrand(x, featlen, gaplen, ...)
```

Arguments

| | |
|---------|--|
| x | gapSites. Object from which the lJuncStrand values are calculated. |
| featlen | Numeric. Number of nucleotides on feature (exon) side of boundary. |
| gaplen | Numeric. Number of nucleotides on gap (intron) side of boundary. |
| ... | Optional arguments passed additionally to the function (currently unused). |

Details

The functions are intended to provide position information which crosses exon-intron boundaries. Added DNA sequences can be used to produce seqlogos. The functions are intended to be used in advance of xCodons functions. Later on added AA sequences can be used to search for proteins where intronic sequences are retained.

Value

cRanges

Author(s)

Wolfgang Kaisers

Examples

```
# A) Create gapSites object
bam<-system.file("extdata","rna_fem.bam",package="spliceSites")
reader<-bamReader(bam[1],idx=TRUE)
ga<-alignGapList(reader)
bamClose(reader)
ga

# B) Extract JuncStrandtion data
lj<-lJuncStrand(ga,featlen=6,gaplen=6)
ljm<-lJuncStrand(ga,featlen=6,gaplen=6)
rj<-rJuncStrand(ga,featlen=6,gaplen=6)
rjm<-rJuncStrand(ga,featlen=6,gaplen=6)
lrj<-lrJuncStrand(ga,lfeatlen=6,rfeatlen=6)
lrjm<-lrJuncStrand(ga,lfeatlen=6,rfeatlen=6)
```

[-methods

Methods for Function [.

Description

Methods for function [

Methods

signature(x = "cRanges", i="ANY", j="ANY", drop="ANY") Method for 'cRanges'.

signature(x = "gapSites", i="ANY", j="ANY", drop="ANY") Method for 'gapSites.'

Index

- *Topic **ExpressionSet**
 - annotate-ExpressionSet, 11
 - readCuffGeneFpkm, 36
 - readExpSet, 37
- *Topic **SpliceCountSet**
 - SpliceCountSet-class, 42
- *Topic **addGeneAligns**
 - addGeneAligns, 5
- *Topic **addMaxEnt**
 - addMaxEnt, 7
- *Topic **alignGapList**
 - getGapSites, 27
- *Topic **annGapSites**
 - annGapSites-class, 9
- *Topic **annotate**
 - annotate-ExpressionSet, 11
 - annotation, 12
- *Topic **annotation**
 - annotation, 12
- *Topic **as.data.frame**
 - as.data.frame-methods, 13
- *Topic **bamCount**
 - countByGeneName, 16
- *Topic **bamRange**
 - rangeByGeneName, 35
- *Topic **bamReader**
 - getGapSites, 27
- *Topic **cRanges**
 - cRanges-class, 17
 - extractByGeneName, 21
 - extractRange, 22
- *Topic **caRanges**
 - caRanges-class, 13
 - extractByGeneName, 21
 - extractRange, 22
- *Topic **cdRanges**
 - cdRanges-class, 15
 - extractByGeneName, 21
 - extractRange, 22
- *Topic **classes**
 - aaGapSites-class, 4
 - caRanges-class, 13
 - cdRanges-class, 15
 - cRanges-class, 17
 - dnaGapSites-class, 19
 - gapSites-class, 25
 - hbond-class, 28
 - maxEnt-class, 32
- *Topic **c**
 - c-methods, 13
- *Topic **dim**
 - dim-methods, 19
- *Topic **dnaRanges**
 - dnaRanges, 20
- *Topic **extractByGeneName**
 - extractByGeneName, 21
- *Topic **extractRange**
 - extractRange, 22
- *Topic **gapAligns**
 - addGeneAligns, 5
- *Topic **gapSites**
 - alt_X_ranks, 8
 - annotation, 12
 - countByGeneName, 16
 - extractByGeneName, 21
 - extractRange, 22
 - gapSites, 23
 - gapSites-class, 25
 - getGapSites, 27
 - readMergedBamGaps, 38
- *Topic **hbond**
 - addHbond, 6
- *Topic **head**
 - head-methods, 29
- *Topic **initialize**
 - initialize-methods, 29
- *Topic **keyProfiler**
 - keyProfiler-class, 30

- readTabledBamGaps, 39
- *Topic **lCodons**
 - trim, 42
 - xCodons, 48
 - xJunc, 49
 - xJuncStrand, 50
- *Topic **lrcodons**
 - lrcodons, 31
- *Topic **maxEnt**
 - addMaxEnt, 7
- *Topic **merge**
 - merge-methods, 33
- *Topic **methods**
 - [-methods, 52
 - as.data.frame-methods, 13
 - c-methods, 13
 - dim-methods, 19
 - head-methods, 29
 - initialize-methods, 29
 - merge-methods, 33
 - sortTable-methods, 41
- *Topic **package**
 - spliceSites-package, 2
- *Topic **plotGeneAlignDepth**
 - plotGeneAlignDepth, 33
- *Topic **rCodons**
 - trim, 42
 - xCodons, 48
 - xJunc, 49
 - xJuncStrand, 50
- *Topic **rangeByGeneName**
 - rangeByGeneName, 35
- *Topic **readCuffGeneFpkm**
 - readCuffGeneFpkm, 36
- *Topic **readExpSet**
 - readExpSet, 37
- *Topic **readMergedBamGaps**
 - readMergedBamGaps, 38
- *Topic **readTabledBamGaps**
 - readTabledBamGaps, 39
- *Topic **refGenome**
 - annotation, 12
 - rangeByGeneName, 35
- *Topic **seqlogo**
 - seqlogo, 40
- *Topic **silic_tryp**
 - silic_tryp, 41
- *Topic **sortTables**
 - sortTable-methods, 41
- *Topic **stop-codon**
 - trim, 42
 - truncateSeq, 43
 - uniqueJuncAnn, 46
 - xCodons, 48
 - xJunc, 49
 - xJuncStrand, 50
- *Topic **truncateSeq**
 - truncateSeq, 43
- *Topic **truncate_seq**
 - truncate_seq, 44
- *Topic **trypsinCleave**
 - trypsinCleave, 45
- *Topic **uniqueJuncAnn**
 - uniqueJuncAnn, 46
- *Topic **write.files**
 - write.files, 47
- [-methods, 52
- aaGapSites-class, 4
- addGeneAligns, 5
- addGeneAligns.gapSites-method
 - (addGeneAligns), 5
- addGeneAligns-method (addGeneAligns), 5
- addHbond, 6
- addHbond.cdRanges-method (addHbond), 6
- addHbond.gapSites-method (addHbond), 6
- addHbond-methods (addHbond), 6
- addKeyTable (keyProfiler-class), 30
- addKeyTable.keyProfiler-method
 - (keyProfiler-class), 30
- addKeyTable-methods
 - (keyProfiler-class), 30
- addMaxEnt, 7
- addMaxEnt.gapSites-method (addMaxEnt), 7
- addMaxEnt-methods (addMaxEnt), 7
- alignGapList (getGapSites), 27
- alt_left_ranks (alt_X_ranks), 8
- alt_left_ranks.gapSites-method
 - (alt_X_ranks), 8
- alt_left_ranks-methods (alt_X_ranks), 8
- alt_ranks (alt_X_ranks), 8
- alt_ranks.gapSites-method
 - (alt_X_ranks), 8
- alt_ranks-methods (alt_X_ranks), 8
- alt_right_ranks (alt_X_ranks), 8
- alt_right_ranks.gapSites-method
 - (alt_X_ranks), 8

- alt_right_ranks-method (alt_X_ranks), 8
- alt_X_ranks, 8
- annGapSites-class, 9
- annotate (annotation), 12
- annotate, ExpressionSet, refJunctions-method (annotate-ExpressionSet), 11
- annotate, gapSites, refJunctions-method (annotation), 12
- annotate-ExpressionSet, 11
- annotate-methods (annotation), 12
- annotation, 12
- annotation, gapSites-method (annotation), 12
- annotation-methods (annotation), 12
- annotation<- (annotation), 12
- annotation<-, gapSites, ANY-method (annotation), 12
- annotation<-, gapSites, refJunctions-method (annotation), 12
- annotation<--methods (annotation), 12
- appendKeyTable (keyProfiler-class), 30
- appendKeyTable, keyProfiler-method (keyProfiler-class), 30
- appendKeyTable-methods (keyProfiler-class), 30
- as.data.frame-methods, 13
- as.data.frame.cRanges (cRanges-class), 17
- as.data.frame.gapSites (gapSites-class), 25

- basedir, hbond-method (hbond-class), 28
- basedir, maxEnt-method (maxEnt-class), 32
- basedir<-, hbond-method (hbond-class), 28
- basedir<-, maxEnt-method (maxEnt-class), 32

- c, caRanges-method (caRanges-class), 13
- c, cdRanges-method (cdRanges-class), 15
- c, cRanges-method (cRanges-class), 17
- c, gapSites-method (gapSites-class), 25
- c-methods, 13
- caRanges-class, 13
- cdRanges-class, 15
- count (cRanges-class), 17
- count, cRanges-method (cRanges-class), 17
- count-methods (cRanges-class), 17
- countByGeneName, 16
- cRanges, 13, 15
- cRanges-class, 17
- dim, cRanges-method (cRanges-class), 17
- dim, gapSites-method (gapSites-class), 25
- dim-methods, 19
- dnaGapSites (gapSites), 23
- dnaGapSites, gapSites, DNASTringSet-method (gapSites-class), 25
- dnaGapSites-class, 19
- dnaGapSites-methods (gapSites), 23
- dnaRanges, 20
- dnaRanges, cRanges, DNASTringSet, logical-method (dnaRanges), 20
- dnaRanges, cRanges, DNASTringSet, missing-method (dnaRanges), 20
- dnaRanges-methods (dnaRanges), 20
- do_group_align_data (gapSites-class), 25

- end, cRanges-method (cRanges-class), 17
- extractByGeneName, 21
- extractByGeneName, cRanges, ANY-method (extractByGeneName), 21
- extractByGeneName, cRanges-method (extractByGeneName), 21
- extractByGeneName, gapSites, ANY-method (extractByGeneName), 21
- extractByGeneName, gapSites-method (extractByGeneName), 21
- extractByGeneName-methods (extractByGeneName), 21
- extractRange, 22
- extractRange, cRanges-method (extractRange), 22
- extractRange, gapSites-method (extractRange), 22
- extractRange-methods (extractRange), 22

- gapSites, 4, 10, 19, 23
- gapSites-class, 25
- getAnnStrand (gapSites-class), 25
- getAnnStrand, gapSites-method (gapSites-class), 25
- getAnnStrand-methods (gapSites-class), 25
- getGapSites, 27
- getKeyTable (keyProfiler-class), 30
- getKeyTable, keyProfiler-method (keyProfiler-class), 30

- getKeyTable-methods
 - (keyProfiler-class), 30
- getMeStrand (addMaxEnt), 7
- getMeStrand, gapSites-method
 - (addMaxEnt), 7
- getMeStrand-methods (addMaxEnt), 7
- getProfile (gapSites-class), 25
- getProfile, gapSites-method
 - (gapSites-class), 25
- getProfile-methods (gapSites-class), 25
- getSequence (cdRanges-class), 15
- getSequence, caRanges-method
 - (caRanges-class), 13
- getSequence, cdRanges-method
 - (cdRanges-class), 15
- getSequence-methods (cdRanges-class), 15
- gptm (gapSites-class), 25
- gptm, gapSites-method (gapSites-class), 25
- gptm-methods (gapSites-class), 25
- hbond (hbond-class), 28
- hbond, hbond-method (hbond-class), 28
- hbond-class, 28
- hbond-methods (hbond-class), 28
- head, aaGapSites-method
 - (aaGapSites-class), 4
- head, caRanges-method (caRanges-class), 13
- head, cdRanges-method (cdRanges-class), 15
- head, cRanges-method (cRanges-class), 17
- head, dnaGapSites-method
 - (dnaGapSites-class), 19
- head, gapSites-method (gapSites-class), 25
- head-methods, 29
- id (cRanges-class), 17
- id, cRanges-method (cRanges-class), 17
- id-methods (cRanges-class), 17
- initialize, cdRanges-method
 - (cdRanges-class), 15
- initialize, cRanges-method
 - (cRanges-class), 17
- initialize, keyProfiler-method
 - (keyProfiler-class), 30
- initialize, SpliceCountSet-method
 - (SpliceCountSet-class), 42
- initialize-methods, 29
- keyProfiler-class, 30
- lCodons (xCodons), 48
- lCodons, cRanges-method (xCodons), 48
- lCodons-methods (xCodons), 48
- lend (gapSites-class), 25
- lend, gapSites-method (gapSites-class), 25
- lend-methods (gapSites-class), 25
- lJunc (xJunc), 49
- lJunc, gapSites-method (xJunc), 49
- lJunc-methods (xJunc), 49
- lJuncStrand (xJuncStrand), 50
- lJuncStrand, gapSites-method
 - (xJuncStrand), 50
- lJuncStrand-methods (xJuncStrand), 50
- load.hbond (hbond-class), 28
- load.maxEnt (maxEnt-class), 32
- lrcodons, 31
- lrcodons, gapSites-method (lrcodons), 31
- lrcodons-methods (lrcodons), 31
- lrJunc (xJunc), 49
- lrJunc, gapSites-method (xJunc), 49
- lrJunc-methods (xJunc), 49
- lrJuncStrand (xJuncStrand), 50
- lrJuncStrand, gapSites-method
 - (xJuncStrand), 50
- lrJuncStrand-methods (xJuncStrand), 50
- lstart (gapSites-class), 25
- lstart, gapSites-method
 - (gapSites-class), 25
- lstart-methods (gapSites-class), 25
- maxEnt-class, 32
- merge-methods, 33
- merge.gapSites (gapSites-class), 25
- nAlignGaps (gapSites-class), 25
- nAlignGaps, gapSites-method
 - (gapSites-class), 25
- nAlignGaps-methods (gapSites-class), 25
- nAligns (gapSites-class), 25
- nAligns, gapSites-method
 - (gapSites-class), 25
- nAligns-methods (gapSites-class), 25
- overlap_genome (gapSites-class), 25

- plot_diff (annGapSites-class), 9
- plot_diff, annGapSites-method (annGapSites-class), 9
- plot_diff-methods (annGapSites-class), 9
- plot_diff_ranks (alt_X_ranks), 8
- plot_diff_ranks, gapSites-method (alt_X_ranks), 8
- plot_diff_ranks-methods (alt_X_ranks), 8
- plotGeneAlignDepth, 33
- plotGeneAlignDepth, bamReader-method (plotGeneAlignDepth), 33
- plotGeneAlignDepth-methods (plotGeneAlignDepth), 33

- rangeByGeneName, 35
- rbamtools, 3
- rCodons (xCodons), 48
- rCodons, cRanges-method (xCodons), 48
- rCodons-methods (xCodons), 48
- readCuffGeneFpkm, 36
- readExpSet, 37
- readMergedBamGaps, 38
- readTabledBamGaps, 39
- refGenome, 3
- rend (gapSites-class), 25
- rend, gapSites-method (gapSites-class), 25
- rend-methods (gapSites-class), 25
- resize_left (trim), 42
- resize_left, gapSites-method (trim), 42
- resize_left-methods (trim), 42
- resize_right (trim), 42
- resize_right, gapSites-method (trim), 42
- resize_right-methods (trim), 42
- rJunc (xJunc), 49
- rJunc, gapSites-method (xJunc), 49
- rJunc-methods (xJunc), 49
- rJuncStrand (xJuncStrand), 50
- rJuncStrand, gapSites-method (xJuncStrand), 50
- rJuncStrand-methods (xJuncStrand), 50
- rpmg (gapSites-class), 25
- rpmg, gapSites-method (gapSites-class), 25
- rpmg-methods (gapSites-class), 25
- rstart (gapSites-class), 25
- rstart, gapSites-method (gapSites-class), 25
- rstart-methods (gapSites-class), 25

- saveMaxEnt (maxEnt-class), 32
- saveMaxEnt, maxEnt-method (maxEnt-class), 32
- saveMaxEnt-methods (maxEnt-class), 32
- score3 (maxEnt-class), 32
- score3, maxEnt-method (maxEnt-class), 32
- score3-methods (maxEnt-class), 32
- score5 (maxEnt-class), 32
- score5, maxEnt-method (maxEnt-class), 32
- score5-methods (maxEnt-class), 32
- scoreSeq3 (maxEnt-class), 32
- scoreSeq3, maxEnt-method (maxEnt-class), 32
- scoreSeq3-methods (maxEnt-class), 32
- scoreSeq5 (maxEnt-class), 32
- scoreSeq5, maxEnt-method (maxEnt-class), 32
- scoreSeq5-methods (maxEnt-class), 32
- seqid (gapSites-class), 25
- seqid, cRanges-method (cRanges-class), 17
- seqid, gapSites-method (gapSites-class), 25
- seqid-methods (gapSites-class), 25
- seqlogo, 40
- seqlogo, cdRanges-method (seqlogo), 40
- seqlogo, dnaGapSites-method (dnaGapSites-class), 19
- seqlogo-methods (seqlogo), 40
- setMeStrand (addMaxEnt), 7
- setMeStrand, gapSites-method (addMaxEnt), 7
- setMeStrand-methods (addMaxEnt), 7
- show, aaGapSites-method (aaGapSites-class), 4
- show, cRanges-method (cRanges-class), 17
- show, dnaGapSites-method (dnaGapSites-class), 19
- show, gapSites-method (gapSites-class), 25
- silic_tryp, 41
- sortTable (cRanges-class), 17
- sortTable, cRanges-method (cRanges-class), 17
- sortTable, gapSites-method (gapSites-class), 25
- sortTable-methods, 41
- SpliceCountSet-class, 42
- spliceSites (spliceSites-package), 2

- spliceSites-package, [2](#)
- start, cRanges-method (cRanges-class), [17](#)
- strand (cRanges-class), [17](#)
- strand, cRanges-method (cRanges-class), [17](#)
- strand, gapSites-method (gapSites-class), [25](#)
- strand-methods (cRanges-class), [17](#)
- strand<-, gapSites-method (gapSites-class), [25](#)

- translate (cdRanges-class), [15](#)
- translate, cdRanges-method (cdRanges-class), [15](#)
- translate, dnaGapSites-method (dnaGapSites-class), [19](#)
- translate-methods (cdRanges-class), [15](#)
- trim, [42](#)
- trim_left (trim), [42](#)
- trim_left, gapSites-method (trim), [42](#)
- trim_left-methods (trim), [42](#)
- trim_right (trim), [42](#)
- trim_right, gapSites-method (trim), [42](#)
- trim_right-methods (trim), [42](#)
- truncate_seq, [44](#)
- truncateSeq, [43](#)
- truncateSeq, aaGapSites-method (truncateSeq), [43](#)
- truncateSeq, caRanges-method (truncateSeq), [43](#)
- truncateSeq-methods (truncateSeq), [43](#)
- trypsinCleave, [45](#)
- trypsinCleave, aaGapSites-method (trypsinCleave), [45](#)
- trypsinCleave, caRanges-method (trypsinCleave), [45](#)
- trypsinCleave-methods (trypsinCleave), [45](#)

- uniqueJuncAnn, [46](#)
- uniqueJuncAnn, ExpressionSet, refJunctions-method (uniqueJuncAnn), [46](#)
- uniqueJuncAnn-methods (uniqueJuncAnn), [46](#)

- width, cRanges-method (cRanges-class), [17](#)
- write.annDNA.tables (gapSites-class), [25](#)
- write.annDNA.tables, gapSites, DNASTringSet, character-method (gapSites-class), [25](#)
- write.annDNA.tables-methods (gapSites-class), [25](#)
- write.files, [47](#)
- write.files, aaGapSites-method (aaGapSites-class), [4](#)
- write.files, caRanges-method (caRanges-class), [13](#)
- write.files-methods (write.files), [47](#)

- xCodons, [48](#)
- xJunc, [49](#)
- xJuncStrand, [50](#)