Package ‘atenas’

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Type Package
Title Analysis of Transposable Elements
Version 1.8.0
Description Quantify expression of transposable elements (TEs) from RNA-seq data through
different methods, including ERVmap, TEtranscripts and Telescope. A common interface
is provided to use each of these methods, which consists of building a parameter object,
calling the quantification function with this object and getting a SummarizedExperiment
object as output container of the quantified expression profiles. The implementation
allows one to quantify TEs and gene transcripts in an integrated manner.
License Artistic-2.0
Encoding UTF-8
Depends R (>= 4.3.0), SummarizedExperiment
Imports methods, stats, Matrix, BiocGenerics, BiocParallel, S4Vectors,
IRanges, GenomicRanges, GenomicAlignments, Rsamtools,
GenomeInfoDb, SQUAREM, sparseMatrixStats, AnnotationHub,
matrixStats
Suggests covr, BiocStyle, knitr, rmarkdown, RUnit,
TxDb.Dmelanogaster.UCSC.dm6.ensGene, RColorBrewer
biocViews Transcription, Transcriptomics, RNASeq, Sequencing,
Preprocessing, Software, GeneExpression, Coverage,
DifferentialExpression, FunctionalGenomics
VignetteBuilder knitr
URL https://github.com/functionalgenomics/atenas
BugReports https://github.com/functionalgenomics/atenas/issues
RoxygenNote 7.2.3
Collate 'AllGenerics.R' 'AllClasses.R' 'ERVmap.R' 'TEtranscripts.R'
'Telescope.R' 'annotations.R' 'atenas.R' 'atenasMethod.R'
'overlappingModes.R' 'qtex.R' 'utils.R' 'zzz.R'
git_url https://git.bioconductor.org/packages/atenas
git_branch RELEASE_3_18
The atena package provides a complete re-implementation in R of three existing methods for the quantification of transposable element (TE) expression in order to facilitate its integration into Bioconductor workflows for the analysis of RNA-seq data. The three methods are TEtranscripts (Jin et al. (2015)), ERVmap (Tokuyama et al. (2018)) and Telescope (Bendall et al. (2019)).
**Details**

The main functions are:

- **TEtranscriptsParam** - build parameter objects of the class `TEtranscriptsParam-class` for the TEtranscripts expression quantification method
- **ERVmapParam** - build parameter objects of the class `ERVmapParam-class` for the ERVmap expression quantification method
- **TelescopeParam** - build parameter objects of the class `TelescopeParam-class` for the Telescope expression quantification method
- **qtex** - call the TE expression quantification method using a previously built parameter object

For detailed information on usage, see the package vignette, by typing `vignette("atena")`.

All questions and bug reports should be posted to the Bioconductor Support Site:

[https://support.bioconductor.org](https://support.bioconductor.org)

The code of the development version of the package is available at the GitHub repository:

[https://github.com/functionalgenomics/atena](https://github.com/functionalgenomics/atena)

**References**


Tokuyama M et al. ERVmap analysis reveals genome-wide transcription of human endogenous retroviruses. PNAS. 2018;115(50):12565-12572. DOI: [https://doi.org/10.1073/pnas.1814589115](https://doi.org/10.1073/pnas.1814589115)

Bendall et al. Telescope: characterization of the retrotranscriptome by accurate estimation of transposable element expression. PLOS Comp. Biol. 2019;15(9):e1006453. DOI: [https://doi.org/10.1371/journal.pcbi.1006453](https://doi.org/10.1371/journal.pcbi.1006453)

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`annotaTEs` **Get RepeatMasker UCSC annotations**

**Description**

The `annotaTEs()` function fetches RepeatMasker UCSC transposable element (TE) annotations using `AnnotationHub` and parses them.

**Usage**

`annotaTEs(genome = "hg38", parsefun = rmskidentity, AHid = NULL, ...)"`
Arguments

- **genome**: The genome version of the desired RepeatMasker annotations (e.g. "hg38").
- **parsefun**: A function to parse the annotations:
  - Function `rmskidentity` returns RepeatMasker annotations as present in AnnotationHub, without processing them.
  - Function `rmskbasicparser` parses annotations by removing low complexity regions, simple repeats, satellites, rRNA, scRNA, snRNA, srpRNA and tRNA. Also removes TEs with a strand different than "+" or "-". Modifies "repFamily" and "repClass" columns when a "?" is present or when they are defined as "Unknown" or "Other". Finally, assigns a unique id to each TE instance by adding the suffix ".dup" plus a number at the end of the "repName".
  - Function `rmskatenaparser` parses RepeatMasker annotations reconstructing fragmented TEs by assembling together fragments from the same TE that are close enough. For LTR class TEs, it tries to reconstruct full-length and partial TEs following the LTR - internal region - LTR structure. Input is a GRanges object and output is a GRangesList object.
  - Function `OneCodeToFindThemAll` parses annotations following the 'One code to find them all' method by (Bailly-Bechet et al. 2014). Input is a GRanges object and output is a GRangesList object.
  - User-defined function. Input and output should be GRanges objects.
- **AHid**: AnnotationHub unique identifier, of the form AH12345, of an object with TE annotations. This is an optional argument to specify a concrete AnnotationHub resource, for instance when more than one RepeatMasker annotation available for a specific genome version. If AHid is not specified, the latest RepeatMasker annotation is be used.

Details

Given a specific genome version, the `annotaTEs()` function fetches RepeatMasker annotations from UCSC Genome Browser using the AnnotationHub package. Since RepeatMasker not only provides TE annotations but also low complexity DNA sequences and other types of repeats, a specific parsefun can be set to parse these annotations (e.g. `rmskbasicparser` or a user-defined function). If no parsing is required, parsefun can be set to `rmskidentity`.

Value

A GRanges object with transposable element annotations.

See Also

AnnotationHub
atenaParam-class

Examples

```r
rmsk_gr <- annotaTEs(genome = "hg19", parsefun = rmskidentity)
```

Description

This is a class for storing parameters to quantify TE (and gene) expression using the atena method. It is a subclass of the 'QuantifyParam-class'.

Build an object of the class atenaParam.

Usage

```r
atenaParam(
  bfl,
  teFeatures,
  aggregateby = character(0),
  ovMode = "ovUnion",
  geneFeatures = NULL,
  singleEnd = TRUE,
  strandMode = 1L,
  ignoreStrand = FALSE,
  fragments = TRUE,
  pi_prior = 0L,
  theta_prior = 0L,
  em_epsilon = 1e-07,
  maxIter = 100L,
  reassign_mode = "exclude",
  conf_prob = 0.9
)
```

## S4 method for signature 'atenaParam'
show(object)

Arguments

- **bfl**: A BamFile or BamFileList object, or a character string vector of BAM filenames.
- **teFeatures**: A GRanges or GRangesList object. Elements in this object should have names, which are used as a grouping factor for genomic ranges forming a common locus. This grouping is performed previous to TE expression quantification, unlike the aggregation of quantifications performed when the aggregateby parameter is specified, which is performed after individual TE instances are quantified.
aggregateby  Character vector with column names from the annotation to be used to aggregate quantifications. By default, this is an empty vector, which means that the names of the input GRanges or GRangesList object given in the teFeatures parameter are used to aggregate quantifications.

ovMode  Character vector indicating the overlapping mode. Available options are: "ovUnion" (default) and "ovIntersectionStrict", which implement the corresponding methods from HTSeq (https://htseq.readthedocs.io/en/release_0.11.1/count.html). Ambiguous alignments (alignments overlapping > 1 feature) are not counted.

geneFeatures  (Default NULL) A GRanges or GRangesList object with the gene annotated features to be quantified. Unique reads are first tallied with respect to these gene features whereas multi-mapping reads are preferentially assigned to TEs. Elements should have names indicating the gene name/id. In case that geneFeatures is a GRanges and contains a metadata column named type, only the elements with type = exon are considered for the analysis. Then, exon counts are summarized to the gene level. If NULL, gene expression is not quantified.

singleEnd  (Default TRUE) Logical value indicating if reads are single (TRUE) or paired-end (FALSE).

strandMode  (Default 1) Numeric vector which can take values 0, 1 or 2. The strand mode is a per-object switch on GAlignmentPairs objects that controls the behavior of the strand getter. See GAlignmentPairs class for further detail. If singleEnd = TRUE, then strandMode is ignored.

ignoreStrand  (Default FALSE) A logical which defines if the strand should be taken into consideration when computing the overlap between reads and annotated features. When ignoreStrand = FALSE, an aligned read is considered to be overlapping an annotated feature as long as they have a non-empty intersecting genomic range on the same strand, while when ignoreStrand = TRUE the strand is not considered.

fragments  (Default TRUE) A logical; applied to paired-end data only. When fragments=FALSE, the read-counting method only counts 'mated pairs' from opposite strands (non-ambiguous properly paired reads), while when fragments=TRUE same-strand pairs, singletons, reads with unmapped pairs and other ambiguous or not properly paired fragments are also counted (see "Pairing criteria" in readGAlignments()). For further details see summarizeOverlaps().

pi_prior  (Default 0) A positive numeric object indicating the prior on pi. The same prior can be specified for all features setting pi_prior as a scalar, or each feature can have a specific prior by setting pi_prior as a vector with names() corresponding to all feature names. Setting a pi prior is equivalent to adding n unique reads.

theta_prior  (Default 0) A positive numeric object indicating the prior on Q. The same prior can be specified for all features setting theta_prior as a scalar, or each feature can have a specific prior by setting theta_prior as a vector with names() corresponding to all feature names. Equivalent to adding n non-unique reads.

em_epsilon  (Default 1e-7) A numeric scalar indicating the EM Algorithm Epsilon cutoff.

maxIter  A positive integer scalar storing the maximum number of iterations of the EM SQUAREM algorithm (Du and Varadhan, 2020). Default is 100 and this value is passed to the maxiter parameter of the squarem() function.
**reassign_mode**  (Default 'exclude') Character vector indicating reassignment mode after EM step. Available methods are 'exclude' (reads with more than one best assignment are excluded from the final counts), 'choose' (when reads have more than one best assignment, one of them is randomly chosen), 'average' (the read count is divided evenly among the best assignments) and 'conf' (only assignments that exceed a certain threshold -defined by conf_prob parameter- are accepted, then the read count is proportionally divided among the assignments above conf_prob).

**conf_prob**  (Default 0.9) Minimum probability for high confidence assignment.

**object**  A atenaParam object.

**Details**

This is the constructor function for objects of the class atenaParam-class. This type of object is the input to the function qtex() for quantifying expression of transposable elements, which will call the atena method with this type of object. The atena method uses a multiple '__no_feature' approach in which as many '__no_feature' features as different overlapping patterns of multimapping reads in the overlapping matrix are used to represent alignments mapping outside annotations.

**Value**

A atenaParam object.

**Slots**

**singleEnd**  (Default TRUE) Logical value indicating if reads are single (TRUE) or paired-end (FALSE).

**strandMode**  (Default 1) Numeric vector which can take values 0, 1 or 2. The strand mode is a per-object switch on GAlignmentPairs objects that controls the behavior of the strand getter. See GAlignmentPairs class for further detail. If singleEnd = TRUE, then strandMode is ignored.

**ignoreStrand**  (Default FALSE) A logical which defines if the strand should be taken into consideration when computing the overlap between reads and annotated features. When ignoreStrand = FALSE, an aligned read is considered to be overlapping an annotated feature as long as they have a non-empty intersecting genomic range on the same strand, while when ignoreStrand = TRUE the strand is not considered.

**fragments**  (Default TRUE) A logical; applied to paired-end data only. When fragments=FALSE, the read-counting method only counts 'mated pairs' from opposite strands (non-ambiguous properly paired reads), while when fragments=TRUE same-strand pairs, singletons, reads with unmapped pairs and other ambiguous or not properly paired fragments are also counted (see "Pairing criteria" in readGAlignments()). For further details see summarizeOverlaps().

**pi_prior**  (Default 0) A positive numeric object indicating the prior on pi. The same prior can be specified for all features setting pi_prior as a scalar, or each feature can have a specific prior by setting pi_prior as a vector with names() corresponding to all feature names. Setting a pi prior is equivalent to adding n unique reads.

**theta_prior**  (Default 0) A positive numeric object indicating the prior on Q. The same prior can be specified for all features setting theta_prior as a scalar, or each feature can have a specific prior by setting theta_prior as a vector with names() corresponding to all feature names. Equivalent to adding n non-unique reads.
em_epsilon  (Default 1e-7) A numeric scalar indicating the EM Algorithm Epsilon cutoff.

maxIter A positive integer scalar storing the maximum number of iterations of the EM SQUAREM algorithm (Du and Varadhan, 2020). Default is 100 and this value is passed to the maxiter parameter of the squarem() function.

reassign_mode  (Default 'exclude') Character vector indicating reassignment mode after EM step. Available methods are 'exclude' (reads with more than one best assignment are excluded from the final counts), 'choose' (when reads have more than one best assignment, one of them is randomly chosen), 'average' (the read count is divided evenly among the best assignments) and 'conf' (only assignments that exceed a certain threshold -defined by conf_prob parameter- are accepted, then the read count is proportionally divided among the assignments above conf_prob).

conf_prob  (Default 0.9) Minimum probability for high confidence assignment.

Examples

```r
bamfiles <- list.files(system.file("extdata", package="atena"),
                      pattern="*.bam", full.names=TRUE)
rmskat <- annotaTEs(genome = "dm6", parsefun = rmskatenaparser,
                    strict = FALSE, insert = 500)
rmskLTR <- getLTRs(rmskat, relLength = 0.8,
                    full_length = TRUE,
                    partial = TRUE,
                    otherLTR = TRUE)
atpar <- atenaParam(bfl=bamfiles,
                    teFeatures=rmskLTR,
                    singleEnd = TRUE,
                    ignoreStrand=TRUE)
atpar
```

---

ERVmapParam-class  

ERVmap parameter class

Description

This is a class for storing parameters provided to the ERVmap algorithm. It is a subclass of the 'QuantifyParam-class'.

Build an object of the class ERVmapParam

Usage

```r
ERVmapParam(
  bfl,
  teFeatures,
  aggregateby = character(0),
  ovMode = "ovUnion",
```

---
geneFeatures = NULL,
singleEnd = TRUE,
ignoreStrand = TRUE,
strandMode = 1L,
fragments = !singleEnd,
maxMismatchRate = 0.02,
suboptimalAlignmentTag = "auto",
suboptimalAlignmentCutoff = 5,
geneCountMode = "all"
)

## S4 method for signature 'ERVmapParam'
show(object)

Arguments

- `bfl`: A `BamFile` or `BamFileList` object, or a character string vector of BAM filenames.
- `teFeatures`: A `GRanges` or `GRangesList` object with the transposable element (TE) annotated features to be quantified. Elements in this object should have names, which are used as a grouping factor for genomic ranges forming a common locus, unless other metadata column names are specified in the `aggregateby` parameter.
- `aggregateby`: Character vector with column names in the annotation to be used to aggregate quantifications. By default, this is an empty vector, which means that the names of the input `GRanges` or `GRangesList` object given in the `teFeatures` parameter are used to aggregate quantifications.
- `ovMode`: Character vector indicating the overlapping mode. Available options are: "ovUnion" (default) and "ovIntersectionStrict", which implement the corresponding methods from HTSeq (https://htseq.readthedocs.io/en/release_0.11.1/count.html). Ambiguous alignments (alignments overlapping > 1 feature) are addressed as in the original ERVmap algorithm.
- `geneFeatures`: (Default NULL) A `GRanges` or `GRangesList` object with the gene annotated features to be quantified. Overlaps with unique reads are first tallied with respect to these gene features. Elements should have names indicating the gene name/id. In case that `geneFeatures` is a `GRanges` and contains a metadata column named `type`, only the elements with `type = exon` are considered for the analysis. Then, exon counts are summarized to the gene level. If NULL, gene expression is not quantified.
- `singleEnd`: (Default TRUE) Logical value indicating if reads are single (`TRUE`) or paired-end (`FALSE`).
- `ignoreStrand`: (Default TRUE) A logical which defines if the strand should be taken into consideration when computing the overlap between reads and annotated features. When `ignore_strand = FALSE`, an aligned read is considered to be overlapping an annotated feature as long as they have a non-empty intersecting genomic range on the same strand, while when `ignoreStrand = TRUE` the strand is not considered.
strandMode  (Default 1) Numeric vector which can take values 0, 1 or 2. The strand mode is a per-object switch on \texttt{GAlignmentPairs} objects that controls the behavior of the strand getter. See \texttt{GAlignmentPairs} class for further detail. If \texttt{singleEnd = TRUE}, then \texttt{strandMode} is ignored.

fragments  (Default not \texttt{singleEnd}) A logical; applied to paired-end data only. When \texttt{fragments=TRUE}, the read-counting method in the original ERVmap algorithm is applied: each mate of a paired-end read is counted (including ambiguous and not properly paired reads). When \texttt{fragments=FALSE}, if the two mates of a paired-end read map to the same element, they are counted as a single hit and singletons, reads with unmapped pairs and other ambiguous or not properly paired fragments are not counted (see "Pairing criteria" in \texttt{readGAlignments()}).

maxMismatchRate  (Default 0.02) Numeric value storing the maximum mismatch rate employed by the ERVmap algorithm to discard aligned reads whose rate of sum of hard and soft clipping or whose rate of the edit distance over the genome reference to the length of the read is above this threshold.

suboptimalAlignmentTag  (Default "auto") Character string storing the tag name in the BAM files that stores the suboptimal alignment score used in the third filter of ERVmap; see Tokuyama et al. (2018). The default, \texttt{suboptimalAlignmentTag="auto"}, first extracts the name of the read mapper software from one or more BAM files. If BAM files were generated by BWA, the suboptimal alignment scores are obtained from a tag called \texttt{XS}. For other read mappers, the suboptimal alignment score is considered to be missing since, except from BWA, no other aligner provides a tag with suboptimal alignment scores. In this case, the available secondary alignments are used to implement an analogous approach to that of the third ERVmap filter. When \texttt{suboptimalAlignmentTag="none"}, it also performs the latter approach even when the tag \texttt{XS} is available. When this parameter is different from "auto" and "none", a tag with the given name is used to extract the suboptimal alignment score.

suboptimalAlignmentCutoff  (Default 5) Numeric value storing the cutoff above which the difference between the alignment score and the suboptimal alignment score is considered sufficiently large to retain the alignment. When this value is set to \texttt{NA}, the filtering step based on suboptimal alignment scores is skipped.

geneCountMode  (Default "all") Character string indicating if the ERVmap read filters applied to quantify TEs expression should also be applied when quantifying gene expression ("ervmap") or not ("all"), in which case all primary alignments mapping to genes are counted.

Details

This is the constructor function for objects of the class \texttt{ERVmapParam-class}. This type of object is the input to the function \texttt{qtex()} for quantifying expression of transposable elements using the ERVmap method Tokuyama et al. (2018). The ERVmap algorithm processes reads following conservative filtering criteria to provide reliable raw count data for each TE.
**ERVmapParam-class**

**Value**

A `ERVmapParam` object.

**Slots**

readMapper The name of the software used to align reads, obtained from the BAM file header.

singleEnd (Default FALSE) Logical value indicating if reads are single (TRUE) or paired-end (FALSE).

strandMode (Default 1) Numeric vector which can take values 0, 1 or 2. The strand mode is a per-object switch on `GAlignmentPairs` objects that controls the behavior of the strand getter. See `GAlignmentPairs` class for further detail. If `singleEnd = TRUE`, then `strandMode #` is ignored.

ignoreStrand (Default TRUE) A logical which defines if the strand should be taken into consideration when computing the overlap between reads and TEs in the annotations. When `ignore_strand = FALSE`, only those reads which overlap the TE and are on the same strand are counted. On the contrary, when `ignore_strand = TRUE`, any read overlapping an element in `teFeatures` is counted regardless of the strand.

fragments (Default not singleEnd) A logical; applied to paired-end data only. When `fragments=TRUE`, the read-counting method in the original ERVmap algorithm is applied: each mate of a paired-end read is counted (including ambiguous and not properly paired reads). When `fragments=FALSE`, if the two mates of a paired-end read map to the same element, they are counted as a single hit and singletons, reads with unmapped pairs and other ambiguous or not properly paired fragments are not counted (see "Pairing criteria" in `readGAlignments()`).

maxMismatchRate (Default 0.02) Numeric value storing the maximum mismatch rate employed by the ERVmap algorithm to discard aligned reads whose rate of sum of hard and soft clipping, or of the edit distance over the genome reference, to the length of the read is above this threshold.

suboptimalAlignmentTag (Default "auto") Character string storing the tag name in the BAM files that stores the suboptimal alignment score used in the third filter of ERVmap; see Tokuyama et al. (2018). The default, suboptimalAlignmentTag="auto", assumes that either the BAM files were generated by BWA and include a tag called XS that stores the suboptimal alignment score or, if the XS tag is not available, then it uses the available secondary alignments to implement an analogous approach to that of the third ERVmap filter. When suboptimalAlignmentTag="none", it also performs the latter approach even when the tag XS is available. When this parameter is different from "auto" and "none", a tag with the given name is used to extract the suboptimal alignment score. The absence of that tag will prompt an error.

suboptimalAlignmentCutoff (Default 5) Numeric value storing the cutoff above which the difference between the alignment score and the suboptimal alignment score is considered sufficiently large to retain the alignment. When this value is set to NA, then the filtering step based on suboptimal alignment scores is skipped.

geneCountMode (Default "all") Character string indicating if the ERVmap read filters applied to quantify TEs expression should also be applied when quantifying gene expression ("ervmap") or not ("all"), in which case all primary alignments mapping to genes are counted.

**References**

Tokuyama M et al. ERVmap analysis reveals genome-wide transcription of human endogenous retroviruses. PNAS. 2018;115(50):12565-12572. DOI: [https://doi.org/10.1073/pnas.1814589115](https://doi.org/10.1073/pnas.1814589115)
Tokuyama M et al. ERVmap analysis reveals genome-wide transcription of human endogenous retroviruses. PNAS. 2018;115(50):12565-12572. DOI: https://doi.org/10.1073/pnas.1814589115

**Examples**

```r
bamfiles <- list.files(system.file("extdata", package="atena"),
  pattern="*.bam", full.names=TRUE)
rmskat <- annotaTEs(genome = "dm6", parsefun = rmskatenaparser,
  strict = FALSE, insert = 500)
rmskLTR <- getLTRs(rmskat, relLength = 0.8,
  full_length = TRUE,
  partial = TRUE,
  otherLTR = TRUE)
empar <- ERVmapParam(bamfiles,
  teFeatures = rmskLTR,
  singleEnd = TRUE,
  ignoreStrand = TRUE,
  suboptimalAlignmentCutoff=NA)
empar
```

---

**getDNAtransposons**

*Getter of DNA class TEs from parsed RepeatMasker annotations*

**Description**

Getter of DNA class TEs from parsed RepeatMasker annotations

**Usage**

```r
getDNAtransposons(parsed_ann, relLength = 0.9)
```

**Arguments**

- `parsed_ann` A GRangesList object obtained from parsing RepeatMasker annotations with OneCodeToFindThemAll() or rmskatenaparser() function.
- `relLength` (Default 0.9) Numeric value that can take values between 0 to 1. Sets the minimum relative length required for features. Elements with a lower relative length than `relLength` will be filtered. The relative length used is the one obtained by OneCodeToFindThemAll() or rmskatenaparser() (length of the reconstructed TE / length of the reference).

**Details**

Retrieves DNA class TEs from RepeatMasker annotations after parsing using the OneCodeToFindThemAll() or rmskatenaparser() function. The `relLength` parameter can be used to filter out elements with a lower relative length.
getLINEs

Value

A GRangesList object with annotations from DNA transposons.

Examples

rmsk_gr <- annotateTES(genome = "dm6", parsefun = rmskatenaparser,
                        strict = FALSE)

rmsk_gr_DNAtrans <- getDNAtransposons(rmsk_gr, relLength = 0.95)

getLINEs

Getter of LINE class TEs from parsed RepeatMasker annotations

Description

Getter of LINE class TEs from parsed RepeatMasker annotations

Usage

getLINEs(parsed_ann, relLength = 0.9)

Arguments

parsed_ann A GRangesList object obtained from parsing RepeatMasker annotations with
            OneCodeToFindThemAll() or rmskatenaparser() function.
relLength (Default 0.9) Numeric value that can take values between 0 to 1. Sets the mini-
            mum relative length required for features. Elements with a lower relative length
            than relLength will be filtered. The relative length used is the one obtained
            by OneCodeToFindThemAll() or rmskatenaparser(). (length of the recon-
            structed TE / length of the reference).

Details

Retrieves LINE class TEs from RepeatMasker annotations after parsing using the OneCodeToFindThemAll() or
rmskatenaparser() function. The relLength parameter can be used to filter out elements with
a lower relative length.

Value

A GRangesList object with annotations from LINEs.

Examples

rmsk_gr <- annotateTES(genome = "dm6", parsefun = rmskatenaparser,
                        strict = FALSE)

rmsk_gr_LINE <- getLINEs(rmsk_gr, relLength = 0.95)
getLTRs  

*Getter of LTR class TEs from parsed RepeatMasker annotations*

**Description**

Getter of LTR class TEs from parsed RepeatMasker annotations

**Usage**

```r
getLTRs(
  parsed_ann,
  relLength = 0.9,
  full_length = TRUE,
  partial = FALSE,
  soloLTR = FALSE,
  otherLTR = FALSE
)
```

**Arguments**

- `parsed_ann`: A GRangesList object obtained from parsing RepeatMasker annotations with `OneCodeToFindThemAll()` or `rmskatenaparser()` function.
- `relLength`: (Default 0.9) Numeric value that can take values between 0 to 1. Sets the minimum relative length required for features. Elements with a lower relative length than `relLength` will be filtered. The relative length used is the one obtained by `OneCodeToFindThemAll()` or `rmskatenaparser()`. (length of the reconstructed TE / length of the reference).
- `full_length`: (Default TRUE) A logical. Should reconstructed full-length LTR TEs (elements with structure LTR - internal region - LTR) be reported?
- `partial`: (Default FALSE) A logical. Should partially reconstructed LTR TEs be reported (structure LTR - internal region or internal region - LTR)?
- `soloLTR`: (Default FALSE) A logical. Should solo LTRs be reported? Note that only fragments unambiguously identified as LTRs thanks to the identification of their equivalent internal region are considered as LTRs.
- `otherLTR`: (Default FALSE) A logical. Should other TEs from the LTR class, not included in any of the previous three categories, be reported? These include TEs from LTR class that cannot be unambiguously identified as LTR or internal region, and thus cannot be reconstructed into partial or full-length elements; as well as solo internal regions.

**Details**

Retrieves LTR class TEs from RepeatMasker annotations after parsing using the `OneCodeToFindThemAll()` or `rmskatenaparser()` function. The `relLength` parameter can be used to filter out elements with a lower relative length. The other parameters can be used to fine-tune the type of elements to be reported.
getSINEs

Value

A GRangesList object with annotations from LTR.

Examples

rmsk_gr <- annotateTEs(genome = "dm6", parsefun = rmskatenaparser,
                        strict = FALSE)
rmsk_gr_ltr <- getLTRs(rmsk_gr, relLength = 0.95, full_length = TRUE,
                       partial = TRUE)

getSINEs

Getter of SINE class TEs from parsed RepeatMasker annotations

Description

Getter of SINE class TEs from parsed RepeatMasker annotations

Usage

getSINEs(parsed_ann, relLength = 0.9)

Arguments

parsed_ann A GRangesList object obtained from parsing RepeatMasker annotations with OneCodeToFindThemAll() or rmskatenaparser() function.
relLength (Default 0.9) Numeric value that can take values between 0 to 1. Sets the minimum relative length required for features. Elements with a lower relative length than relLength will be filtered. The relative length used is the one obtained by OneCodeToFindThemAll() or rmskatenaparser() (length of the reconstructed TE / length of the reference).

Details

Retrieves SINE class TEs from RepeatMasker annotations after parsing using the OneCodeToFindThemAll() or rmskatenaparser() function. The relLength parameter can be used to filter out elements with a lower relative length.

Value

A GRangesList object with annotations from SINEs.

Examples

rmsk_gr <- annotateTEs(genome = "dm6", parsefun = rmskatenaparser,
                        strict = FALSE)
rmsk_gr_sine <- getSINEs(rmsk_gr, relLength = 0.95)
OneCodeToFindThemAll parser of RepeatMasker annotations

Description

OneCodeToFindThemAll parser of RepeatMasker annotations

Usage

OneCodeToFindThemAll(
  gr,  
  dictionary = NULL,  
  fuzzy = FALSE,  
  strict = FALSE,  
  insert = -1,  
  BPPARAM = SerialParam(progressbar = TRUE)
)

Arguments

- **gr**: A GRanges object with RepeatMasker annotations from AnnotationHub
- **dictionary** (Default NULL) When NULL, a dictionary is built based on names of repeats. If not, a data.frame with equivalences LTR - internal regions created by the user, where first column should be the name of the internal region and the second column should be the LTR(s). When more than one LTR, these should be separated by ":".
- **fuzzy** (Default FALSE) A logical; if TRUE, the search for equivalences between internal parts and LTRs to reconstruct LTR class transposable elements is less stringent, allowing more matches between corresponding subparts. This option can increase the proportion of false positives (incorrectly reconstructed LTR class TEs).
- **strict** (Default FALSE) A logical; if TRUE, the 80-80 rule is applied, i.e. only copies with more than 80 and more than 80 bp long are reported.
- **insert** (Default -1) An integer. When insert < 0, two fragments are assembled if the distance separating their furthest extremities is less than twice the reference length of the element. When insert > 0, fragments are assembled if the distance between their closest extremities is equal or less than insert. When insert = 0, two fragments are assembled if they are in contact next to each other.
- **BPPARAM** See ?bplapply in the BiocParallel package. Can be used to run function in parallel.

Details

Implementation of One code to find them all (Bailly-Bechet et al. 2014). Parses RepeatMasker annotations from UCSC by assembling together fragments from the same transposable element (TE) that are close enough (determined by the insert parameter). For TEs from the LTR class,
the parser tries to reconstruct full-length, when possible, or partial TEs following the LTR - internal region - LTR structure. Equivalences between internal regions and flanking LTRs can be set by the user with the dictionary parameter or can be obtained by the parser. In this last case, the fuzzy parameter determines the level of stringency when searching for LTR - internal region equivalences.

Value

A GRangesList object.

References


Examples

```r
## Not run:
rmrk_gr <- annotaTEs(genome = "dm6", parsefun = OneCodeToFindThemAll,
                      fuzzy = FALSE, strict = FALSE)

## End(Not run)
```

ovUnion

Pre-defined overlapping mode functions

Description

The following functions control the way in which overlaps between aligned reads and annotated features are resolved when an aligned read overlaps more than one feature on the same locus:

Usage

```r
ovUnion(reads, features, ignoreStrand, inter.feature = TRUE)

ovIntersectionStrict(reads, features, ignoreStrand, inter.feature = TRUE)
```

Arguments

- **reads**: A GAlignments, GAlignmentList or a GAlignmentPairs object.
- **features**: A GRanges object with annotated features.
- **ignoreStrand**: (Default FALSE) A logical which defines if the strand should be taken into consideration when computing the overlap between reads and annotated features. When ignoreStrand = FALSE, an aligned read will be considered to be overlapping an annotated feature as long as they have a non-empty intersecting genomic ranges on the same strand, while when ignoreStrand = TRUE the strand will not be considered.
inter.feature  When TRUE, ambiguous alignments (alignments overlapping > 1 features) are removed and not counted. When inter.feature is set to FALSE, these ambiguous overlaps are taken into account and addressed differently depending on the TE quantification.

Details

- ovUnion(): (default)
- ovIntersectionStrict():
- User supplied: a function taking the same parameters as the previous three functions and returning a Hits object.

They take the following parameters:

These functions are given to the mode parameter of the qtex() function and are similar to the functions Union() and IntersectionStrict() from the GenomicAlignments package, with the difference that instead of returning counts of reads overlapping annotated features, they return the actual overlaps, because the counting is deferred to other algorithms that follow some specific strategy when a read maps to more than one feature. For this same reason, these functions lack the inter.feature argument found in the corresponding functions from the GenomicAlignments package.

Value

A Hits object; see the Hits-class manual page.

Examples

```r
bamfiles <- list.files(system.file("extdata", package="atena"),
                      pattern="*.bam", full.names=TRUE)
rm-skat <- annotaTEs(genome = "dm6", parsefun = rmskatenaparser,
                     strict = FALSE, insert = 500)
rm-skLRT <- getLTRs(rmskat, relLength = 0.8,
                    full_length = TRUE,
                    partial = TRUE,
                    otherLTR = TRUE)
tspar <- TelescopeParam(bfl=bamfiles,
                        teFeatures=rmskLTR,
                        singleEnd = TRUE,
                        ignoreStrand=TRUE)
tsquant <- qtex(tspar, mode=ovIntersectionStrict)
```

qtex,ERVmapParam-method

Quantify transposable element expression

Description

The qtex() method quantifies transposable element expression.
Usage

## S4 method for signature 'ERVmapParam'
qtex(
  x,
  phenodata = NULL,
  mode = ovUnion,
  yieldSize = 1000000L,
  verbose = 1,
  BPPARAM = SerialParam(progressbar = ifelse(verbose == 1, TRUE, FALSE))
)

## S4 method for signature 'TEtranscriptsParam'
qtex(
  x,
  phenodata = NULL,
  mode = ovUnion,
  yieldSize = 1000000L,
  BPPARAM = SerialParam(progressbar = TRUE)
)

## S4 method for signature 'TelescopeParam'
qtex(
  x,
  phenodata = NULL,
  mode = ovUnion,
  yieldSize = 1000000L,
  BPPARAM = SerialParam(progressbar = TRUE)
)

## S4 method for signature 'atenaParam'
qtex(
  x,
  phenodata = NULL,
  mode = ovUnion,
  yieldSize = 1000000L,
  BPPARAM = SerialParam(progressbar = TRUE)
)

Arguments

x An AtenaParam object of one of the following subclasses:
  • A ERVMapParam object built using the constructor function ERVMapParam().
    This object will trigger qtex() to use the algorithm by Tokuyama et al.
    (2018).
  • A TelescopeParam object built using the constructor function TelescopeParam().
    This object will trigger qtex() to use the algorithm by Bendall et al. (2019).

phenodata A data.frame or DataFrame object storing phenotypic data to include in the
resulting `SummarizedExperiment` object. If phenodata is set, its row names will become the column names of the resulting `SummarizedExperiment` object.

- **mode**
  One of the pre-defined overlapping methods such as `ovUnion()`, `ovIntersectionStrict` or a user-supplied overlapping function. For a user-supplied overlapping function, the input parameters must match those of the pre-defined methods and the function must return a `Hits` object with subject hits matching the annotated features. This parameter is analogous to the `mode` parameter of the `summarizeOverlaps()` function from the GenomicAlignments package.

- **yieldSize**
  Field inherited from `BamFile`. The method for signature `ERVmapParam()` reads the BAM file by chunks. `yieldSize` represents the number of records (chunk size) to yield each time the file is read.

- **verbose**
  (Default 1). When `verbose > 1`, detailed information on the quantification steps is provided. Warnings are always present regardless of the value of `verbose`.

- **BPPARAM**
  An object of a `BiocParallelParam` subclass to configure the parallel execution of the code. By default, a `SerialParam` object is used, which does not use any parallelization, with the flag `progress=TRUE` to show progress through the calculations.

**Details**

Giving some `AtenaParam` object sub-class as input, the `qtex()` method quantifies the expression of transposable elements (TEs). The particular algorithm to perform the quantification will be selected depending on the specific sub-class of input `AtenaParam` object, see argument `x` above.

**Value**

A `SummarizedExperiment` object.

**References**


**See Also**

`ERVmapParam` `TelescopeParam`

**Examples**

```r
bamfiles <- list.files(system.file("extdata", package="atena"),
  pattern="*.bam", full.names=TRUE)
rmskat <- annotaTEs(genome = "dm6", parsefun = rmskatenaparser,
  strict = FALSE, insert = 500)
rmskLTR <- getLTRs(rmskat, relLength = 0.8,
  full_length = TRUE,
  partial = TRUE,
```
QuantifyParam-class

otherLTR = TRUE)

```r
tspar <- TelescopeParam(bfl=bamfiles,
  teFeatures=rmskLTR,
  singleEnd = TRUE,
  ignoreStrand=TRUE)

tquant <- qtex(tspar)
```

---

QuantifyParam-class  
**Atena parameter class**

**Description**

This is a virtual class from which other classes are derived for storing parameters provided to quantification methods of transposable elements from RNA-seq data.

**Usage**

### S4 method for signature 'QuantifyParam'

```r
path(object)
```

### S4 method for signature 'QuantifyParam'

```r
features(object)
```

**Arguments**

- `object`  
  A `QuantifyParam` object.

**Value**

- `path()`: Filesystem paths to the BAM files in the input parameter object.
- `features()`: The `GenomicRanges` or `GenomicRangesList` object with the features in the input parameter object.

**Slots**

- `bfl`  
  A `BamFileList` object.
- `features`  
  A `GRanges` object.
- `aggregateby`  
  Character vector with column names in the annotation to be used to aggregate quantifications.
- `ovMode`  
  Character vector indicating the overlapping mode. Available options are: "ovUnion" (default) and "ovIntersectionStrict", which implement the corresponding methods from HTSeq ([https://htseq.readthedocs.io/en/release_0.11.1/count.html](https://htseq.readthedocs.io/en/release_0.11.1/count.html)). In the TEtranscripts, ERVmap and Telescope methods ambiguous alignments (alignments overlapping > 1 feature) are addressed differently depending on the method. In the atena method, those overlaps are not counted.
See Also

ERVmapParam-class TelescopeParam-class TEtranscriptsParam-class atenaParam-class

Examples

```r
bamfiles <- list.files(system.file("extdata", package="atena"), pattern="*.bam", full.names=TRUE)
rmmskat <- annotaTEs(genome = "dm6", parsefun = rmskatenaparser, strict = FALSE, insert = 500)
rmmskLTR <- getLTRs(rmmskat, relLength = 0.8, full_length = TRUE, partial = TRUE)
ttpar <- TEtranscriptsParam(bamfiles, teFeatures = rmmskLTR, singleEnd = TRUE, ignoreStrand=TRUE)
path(ttpar)
```

---

**rmskatenaparser**

*atena annotation parser of RepeatMasker annotations*

**Description**

*atena annotation parser of RepeatMasker annotations*

**Usage**

```r
rmskatenaparser(gr, strict = FALSE, insert = 1000)
```

**Arguments**

- `gr` A GRanges object with RepeatMasker annotations from AnnotationHub
- `strict` (Default FALSE) A logical; if TRUE, the 80-80 rule is applied, i.e. only copies with more than 80 and more than 80 bp long are reported.
- `insert` (Default 1000L) An integer > 0. Fragments are assembled together if the distance between their closest extremities is equal or less than `insert`. When `insert = 0`, two fragments are assembled if they are in contact next to each other.

**Details**

*atena annotation parser of RepeatMasker annotations. Parses RepeatMasker annotations from UCSC by assembling together fragments from the same transposable element (TE) that are close enough (determined by the `insert` parameter). For TEs from the LTR class, the parser tries to reconstruct full-length, when possible, or partial TEs following the LTR - internal region - LTR structure. Equivalences between LTR and internal regions are found by, first, identifying LTR regions (those
with the "LTR" substring in their name) and internal regions (those with a suffix such as "-int", "-I", etc.). Then, LTR are assigned to internal regions for which the comparison of the two names are has a higher number of equal consecutive characters.

Value

A GRangesList object.

Examples

rmsk_gr <- annotaTEs(genome = "dm6", parsefun = rmskatenaparser,
                      strict = FALSE)

rmskbasicparser

Parser of RepeatMasker annotations

Description

Parser of RepeatMasker annotations

Usage

rmskbasicparser(gr)

Arguments

gr A GRanges object with RepeatMasker annotations from AnnotationHub

Details

Parses annotations by removing low complexity regions, simple repeats, satellites, rRNA, scRNA, snRNA, srpRNA and tRNA. Also removes TEs with a strand different than "+" or ".". Modifies "repFamily" and "repClass" columns when a "?" is present or when they are defined as "Unknown" or "Other". Finally, assigns a unique id to each TE instance by adding the suffix "_dup" plus a number at the end of the "repName".

Value

A GRanges object.

Examples

rmsk_gr <- annotaTEs(genome = "dm6", parsefun = rmskbasicparser)
riskyIdentity

Description

Identity function for parsefun

Usage

riskyIdentity(gr)

Arguments

gr A GRanges object.

Details

Identity function: returns the GRanges object without any modification.

Value

A GRanges object.

Examples

risky_gr <- annotate(genome = "dm6", parsefun = riskyIdentity)

TelescopeParam-class

Telescope parameter class

Description

This is a class for storing parameters provided to the Telescope algorithm.

Build an object of the class TelescopeParam.

Usage

TelescopeParam(
  bfl,
  teFeatures,
  aggregateby = character(0),
  ovMode = "ovUnion",
  geneFeatures = NULL,
  singleEnd = TRUE,
  strandMode = 1L,
ignoreStrand = FALSE,
fragments = FALSE,
minOverlFract = 0.2,
p1_prior = 0L,
theta_prior = 0L,
em_epsilon = 1e-07,
maxIter = 100L,
reassign_mode = "exclude",
conf_prob = 0.9
)

## S4 method for signature 'TelescopeParam'
show(object)

### Arguments

- **bfl**: A BamFile or BamFileList object, or a character string vector of BAM file- names.
- **teFeatures**: A GRanges or GRangesList object. Elements in this object should have names, which are used as a grouping factor for genomic ranges forming a common locus (equivalent to "locus" column in Telescope). This grouping is performed previous to TE expression quantification, unlike the aggregation of quantifications performed when the aggregateby parameter is specified, which is performed after individual TE instances are quantified.
- **aggregateby**: Character vector with column names from the annotation to be used to aggregate quantifications. By default, this is an empty vector, which means that the names of the input GRanges or GRangesList object given in the teFeatures parameter are used to aggregate quantifications.
- **ovMode**: Character vector indicating the overlapping mode. Available options are: "ovUnion" (default) and "ovIntersectionStrict", which implement the corresponding methods from HTSeq ([https://htseq.readthedocs.io/en/release_0.11.1/count.html](https://htseq.readthedocs.io/en/release_0.11.1/count.html)). Ambiguous alignments (alignments overlapping > 1 feature) are addressed as in the original Telescope method: the overlap with the longest overlapping length is kept.
- **geneFeatures**: (Default NULL) A GRanges or GRangesList object with the gene annotated features to be quantified. The TEtranscripts approach for gene expression quantification is used, in which overlaps with multi-mapping reads are preferentially assigned to TEs. Elements should have names indicating the gene name/id. In case that geneFeatures is a GRanges and contains a metadata column named type, only the elements with type = exon are considered for the analysis. Then, exon counts are summarized to the gene level. If NULL, gene expression is not quantified.
- **singleEnd**: (Default TRUE) Logical value indicating if reads are single (TRUE) or paired-end (FALSE).
- **strandMode**: (Default 1) Numeric vector which can take values 0, 1 or 2. The strand mode is a per-object switch on GAlignmentPairs objects that controls the behavior of the strand getter. See GAlignmentPairs class for further detail. If singleEnd = TRUE, then strandMode is ignored.
ignoreStrand (Default FALSE) A logical which defines if the strand should be taken into consideration when computing the overlap between reads and annotated features. When ignoreStrand = FALSE, an aligned read is considered to be overlapping an annotated feature as long as they have a non-empty intersecting genomic range on the same strand, while when ignoreStrand = TRUE the strand is not considered.

fragments (Default FALSE) A logical; applied to paired-end data only. When fragments=FALSE, the read-counting method only counts `mated pairs` from opposite strands (non-ambiguous properly paired reads), while when fragments=TRUE same-strand pairs, singletons, reads with unmapped pairs and other ambiguous or not properly paired fragments are also counted (see "Pairing criteria" in readGAlignments()). fragments=TRUE is equivalent to the original Telescope algorithm. For further details see summarizeOverlaps().

minOverlFract (Default 0.2) A numeric scalar. minOverlFract is multiplied by the read length and the resulting value is used to discard alignments for which the overlapping length (number of base pairs the alignment and the feature overlap) is lower. When no minimum overlap is required, set minOverlFract = 0.

pi_prior (Default 0) A positive integer scalar indicating the prior on pi. This is equivalent to adding n unique reads.

theta_prior (Default 0) A positive integer scalar storing the prior on Q. Equivalent to adding n non-unique reads.

em_epsilon (Default 1e-7) A numeric scalar indicating the EM Algorithm Epsilon cutoff.

maxIter A positive integer scalar storing the maximum number of iterations of the EM SQUAREM algorithm (Du and Varadhan, 2020). Default is 100 and this value is passed to the maxiter parameter of the squarem() function.

reassign_mode (Default 'exclude') Character vector indicating reassignment mode after EM step. Available methods are 'exclude' (reads with more than one best assignment are excluded from the final counts), 'choose' (when reads have more than one best assignment, one of them is randomly chosen), 'average' (the read count is divided evenly among the best assignments) and 'conf' (only assignments that exceed a certain threshold -defined by conf_prob parameter- are accepted, then the read count is proportionally divided among the assignments above conf_prob).

conf_prob (Default 0.9) Minimum probability for high confidence assignment.

object A TelescopeParam object.

Details

This is the constructor function for objects of the class TelescopeParam-class. This type of object is the input to the function qtex() for quantifying expression of transposable elements, which will call the Telescope algorithm Bendall et al. (2019) with this type of object.

Value

A TelescopeParam object.
Slots

singleEnd (Default TRUE) Logical value indicating if reads are single (TRUE) or paired-end (FALSE).

strandMode (Default 1) Numeric vector which can take values 0, 1 or 2. The strand mode is a per-
object switch on GAlignmentPairs objects that controls the behavior of the strand getter. See
GAlignmentPairs class for further detail. If singleEnd = TRUE, then strandMode is ignored.

ignoreStrand (Default FALSE) A logical which defines if the strand should be taken into consid-
eration when computing the overlap between reads and annotated features. When ignoreStrand
= FALSE, an aligned read is considered to be overlapping an annotated feature as long as they
have a non-empty intersecting genomic range on the same strand, while when ignoreStrand
= TRUE the strand is not considered.

fragments (Default FALSE) A logical; applied to paired-end data only. When fragments=FALSE,
the read-counting method only counts ‘mated pairs’ from opposite strands (non-ambiguous
properly paired reads), while when fragments=TRUE same-strand pairs, singletons, reads with
unmapped pairs and other ambiguous or not properly paired fragments are also counted (see
“Pairing criteria” in readGAlignments()). fragments=TRUE is equivalent to the original Tele-
scope algorithm. For further details see summarizeOverlaps().

minOverlFract (Default 0.2) A numeric scalar. minOverlFract is multiplied by the read length
and the resulting value is used to discard alignments for which the overlapping length (number
of base pairs the alignment and the feature overlap) is lower. When no minimum overlap is
required, set minOverlFract = 0.

pi_prior (Default 0) A positive integer scalar indicating the prior on pi. This is equivalent to
adding n unique reads.

theta_prior (Default 0) A positive integer scalar storing the prior on Q. Equivalent to adding n
unique reads.

em_epsilon (Default 1e-7) A numeric scalar indicating the EM Algorithm Epsilon cutoff.

maxIter A positive integer scalar storing the maximum number of iterations of the EM SQUAREM
algorithm (Du and Varadhan, 2020). Default is 100 and this value is passed to the maxiter
parameter of the squarem() function.

reassign_mode (Default ‘exclude’) Character vector indicating reassignment mode after EM step.
Available methods are ‘exclude’ (reads with more than one best assignment are excluded
from the final counts), ‘choose’ (when reads have more than one best assignment, one of
them is randomly chosen), ‘average’ (the read count is divided evenly among the best assign-
ments) and ‘conf’ (only assignments that exceed a certain threshold -defined by conf_prob
parameter- are accepted, then the read count is proportionally divided among the assignments
above conf_prob).

conf_prob (Default 0.9) Minimum probability for high confidence assignment.

References

Bendall et al. Telescope: characterization of the retrotranscriptome by accurate estimation of trans-
posable element expression. PLOS Comp. Biol. 2019;15(9):e1006453. DOI: https://doi.org/
10.1371/journal.pcbi.1006453

Bendall et al. Telescope: characterization of the retrotranscriptome by accurate estimation of trans-
posable element expression. PLOS Comp. Biol. 2019;15(9):e1006453. DOI: https://doi.org/
10.1371/journal.pcbi.1006453
Examples

bamfiles <- list.files(system.file("extdata", package="atena"),
    pattern="*.bam", full.names=TRUE)
rmstk <- annotaTEs(genome = "dm6", parsefun = rmskatenaparser,
    strict = FALSE, insert = 500)
rmstLTR <- getLTRs(rmstkat, relLength = 0.8,
    full_length = TRUE,
    partial = TRUE,
    otherLTR = TRUE)
tspar <- TelescopeParam(bfl=bamfiles,
    teFeatures=rmstLTR,
    singleEnd = TRUE,
    ignoreStrand=TRUE)

tspar

TEtranscriptsParam-class

 TEtranscripts parameter class

Description

This is a class for storing parameters provided to the TEtranscripts algorithm. It is a subclass of the 'QuantifyParam-class'.

Usage

TEtranscriptsParam(
    bfl,
    teFeatures,
    aggregateby = character(0),
    ovMode = "ovUnion",
    geneFeatures = NULL,
    singleEnd = TRUE,
    ignoreStrand = FALSE,
    strandMode = 1L,
    fragments = TRUE,
    tolerance = 1e-04,
    maxIter = 100L
)

## S4 method for signature 'TEtranscriptsParam'
show(object)
Arguments

**bfl**
a character string vector of BAM file names.

**teFeatures**
A GRanges or GRangesList object with the TE annotated features to be quantified. Elements in this object should have names, which are used as a grouping factor for genomic ranges forming a common locus, unless other metadata column names are specified in the aggregateby parameter.

**aggregateby**
Character vector with column names from the annotation to be used to aggregate quantifications. By default, this is an empty vector, which means that the names of the input GRanges or GRangesList object given in the teFeatures parameter are used to aggregate quantifications.

**ovMode**
Character vector indicating the overlapping mode. Available options are: "ovUnion" (default) and "ovIntersectionStrict", which implement the corresponding methods from HTSeq ([https://htseq.readthedocs.io/en/release_0.11.1/count.html](https://htseq.readthedocs.io/en/release_0.11.1/count.html)). Ambiguous alignments (alignments overlapping > 1 feature) are addressed as in the original TEtranscripts method.

**geneFeatures**
(Default NULL) A GRanges or GRangesList object with the gene annotated features to be quantified. Following the TEtranscripts algorithm, overlaps with unique reads are first tallied with respect to these gene features. Elements should have names indicating the gene name/id. In case that geneFeatures is a GRanges and contains a metadata column named type, only the elements with type = exon are considered for the analysis. Then, exon counts are summarized to the gene level. If NULL, gene expression is not quantified.

**singleEnd**
(Default TRUE) Logical value indicating if reads are single (TRUE) or paired-end (FALSE).

**ignoreStrand**
(Default FALSE) A logical which defines if the strand should be taken into consideration when computing the overlap between reads and annotated features. When ignoreStrand = FALSE, an aligned read is considered to be overlapping an annotated feature as long as they have a non-empty intersecting genomic range on the same strand, while when ignoreStrand = TRUE the strand is not considered.

**strandMode**
(Default 1) Numeric vector which can take values 0, 1 or 2. The strand mode is a per-object switch on GAlignmentPairs objects that controls the behavior of the strand getter. See GAlignmentPairs class for further detail. If singleEnd = TRUE, then strandMode is ignored.

**fragments**
(Default TRUE) A logical; applied to paired-end data only. In both cases (fragments=FALSE and fragments=TRUE), the read-counting method discards not properly paired reads. Moreover, when fragments=FALSE, only non-ambiguous properly paired reads are counted. When fragments=TRUE, ambiguous reads are also counted (see "Pairing criteria" in readGAlignments()). fragments=True is equivalent to the behavior of the TEtranscripts algorithm. For further details see summarizeOverlaps().

**tolerance**
A positive numeric scalar storing the minimum tolerance above which the SQUAREM algorithm (Du and Varadhan, 2020) keeps iterating. Default is 1e-4 and this value is passed to the tol parameter of the squarem() function.
maxIter  
A positive integer scalar storing the maximum number of iterations of the SQUAREM algorithm (Du and Varadhan, 2020). Default is 100 and this value is passed to the maxiter parameter of the squarem() function.

object  
A TEtranscriptsParam object.

Details

This is the constructor function for objects of the class TEtranscriptsParam-class. This type of object is the input to the function qtx() for quantifying expression of transposable elements using the TEtranscripts method Jin et al. (2015). The TEtranscripts algorithm quantifies TE expression by using an EM algorithm to optimally distribute ambiguously mapped reads.

Value

A TEtranscriptsParam object.

Slots

singleEnd (Default FALSE) Logical value indicating if reads are single (TRUE) or paired-end (FALSE).

ignoreStrand (Default FALSE) A logical which defines if the strand should be taken into consideration when computing the overlap between reads and annotated features. When ignoreStrand = FALSE, an aligned read will be considered to be overlapping an annotated feature as long as they have a non-empty intersecting genomic ranges on the same strand, while when ignoreStrand = TRUE the strand will not be considered.

strandMode (Default 1) Numeric vector which can take values 0, 1 or 2. The strand mode is a per-object switch on GAlignmentPairs objects that controls the behavior of the strand getter. See GAlignmentPairs class for further detail. If singleEnd = TRUE, then use either strandMode = NULL or do not specify the strandMode parameter.

fragments (Default TRUE) A logical; applied to paired-end data only. In both cases (fragments=FALSE and fragments=TRUE), the read-counting method discards not properly paired reads. Moreover, when fragments=FALSE, only non-ambiguous properly paired reads are counted. When fragments=TRUE, ambiguous reads are also counted (see "Pairing criteria" in readGAlignments()). fragments=TRUE is equivalent to the behavior of the TEtranscripts algorithm. For further details see summarizeOverlaps().

tolerance  A positive numeric scalar storing the minimum tolerance above which the SQUAREM algorithm (Du and Varadhan, 2020) keeps iterating. Default is 1e-4 and this value is passed to the tol parameter of the squarem() function.

maxIter  A positive integer scalar storing the maximum number of iterations of the SQUAREM algorithm (Du and Varadhan, 2020). Default is 100 and this value is passed to the maxiter parameter of the squarem() function.

References


**Examples**

```r
bamfiles <- list.files(system.file("extdata", package="atena"),
  pattern="*.bam", full.names=TRUE)
rmskat <- annotaTEs(genome = "dm6", parsefun = rmskatenaparser,
  strict = FALSE, insert = 500)
rmskLTR <- getLTRs(rmskat, relLength = 0.8,
  full_length = TRUE,
  partial = TRUE,
  otherLTR = TRUE)
library(TxDb.Dmelanogaster.UCSC.dm6.ensGene)
txdb <- TxDb.Dmelanogaster.UCSC.dm6.ensGene
txdb_genes <- genes(txdb)
tpar <- TEtranscriptsParam(bamfiles,
  teFeatures = rmskLTR,
  geneFeatures = txdb_genes,
  singleEnd = TRUE,
  ignoreStrand=TRUE,
  aggregateby = c("repName"))

ttpar
```
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