

# Package ‘CrispRVariants’

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**Type** Package

**Title** Tools for counting and visualising mutations in a target location

**Version** 1.39.0

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**Description** CrispRVariants provides tools for analysing the results of a CRISPR-Cas9 mutagenesis sequencing experiment, or other sequencing experiments where variants within a given region are of interest. These tools allow users to localize variant allele combinations with respect to any genomic location (e.g. the Cas9 cut site), plot allele combinations and calculate mutation rates with flexible filtering of unrelated variants.

**biocViews** ImmunoOncology, CRISPR, GenomicVariation, VariantDetection, GeneticVariability, DataRepresentation, Visualization, Sequencing

**Depends** R (>= 4.3.0), ggplot2 (>= 2.2.0)

**Encoding** UTF-8

**License** GPL-2

**Imports** AnnotationDbi, BiocParallel, Biostrings, methods, GenomeInfoDb, GenomicAlignments, GenomicRanges, grDevices, grid, gridExtra, IRanges, reshape2, Rsamtools, S4Vectors (>= 0.9.38), utils

**Suggests** BiocStyle, GenomicFeatures, knitr, rmarkdown, readxl, rtracklayer, sangerseqR, testthat, VariantAnnotation

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<i>.explodeCigarOpCombs</i>	<i>.explodeCigarOpCombs</i>
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**Description**

Breaks cigar strings into individual operations

**Usage**

`.explodeCigarOpCombs(cigar, ops = GenomicAlignments::CIGAR_OPS)`

**Arguments**

- cigar                      character(m) A vector of cigar strings
- ops                        character(n) Which operations should be kept?

**Value**

- The operations, as a CharacterList
- Exploded cigar operations with operation widths

**Author(s)**

Helen Lindsay

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<code>.formatVarLabels</code>	<i>formatVarLabels</i>
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---

## Description

Internal CrispRVariants function for creating allele labels given variants and positions

Assume that the reference may be on the negative strand and regions are given with respect to the reference sequence.

## Usage

```
.formatVarLabels(
  grl,
  labels,
  position = c("start", "end"),
  genome.to.pos = NULL,
  pos.to.lab = ":",
  as.string = TRUE
)

.findMismatches(
  alns,
  ref.seq,
  ref.start,
  regions = NULL,
  strand = "+",
  min.pct = 0
)
```

## Arguments

<code>grl</code>	(GRangesList) A GRangesList of variants. The position of the variants is used in labels
<code>labels</code>	(character(n)) A vector of labels for each variant. In CrispRVariants, this is the size and type of the variant, e.g. "9D" for a 9 bp deletion.
<code>position</code>	One of "start" and "end". Determines whether the start or the end coordinate is used when labeling variants.
<code>genome.to.pos</code>	Optional named vector for transforming variant coordinates into another coordinate system (Default: NULL)
<code>pos.to.lab</code>	(character(1)) Character to join positions and labels (Default: ":", e.g. -3:9D)
<code>as.string</code>	Should individual variant labels be pasted into a single comma separated string when one alignment has multiple variants? (Default: TRUE)
<code>alns</code>	A GAlignments object, where the aligned sequences should span the reference sequence

<code>ref.seq</code>	A DNASTring object, the sequence for comparison when checking for mismatches. The sequence does not necessarily have to match the mapping reference sequence. Must span all regions if regions are provided.
<code>ref.start</code>	(numeric(1)) The genomic start position of the reference sequence
<code>regions</code>	A GRanges object, regions to check for mismatches with coordinates relative to the reference sequence
<code>strand</code>	One of "+", "-"
<code>min.pct</code>	(numeric(1), between 0 and 100) Only return SNVs that occur at in least min.pct change, not any change at a position.

**Value**

A data frame of sequence indices, genomic position of mismatch and mismatch base

---

<code>.getAxisCoords</code>	<i>.getAxisCoords</i>
-----------------------------	-----------------------

---

**Description**

Manually specify x-tick locations and labels, as sometimes ggplot defaults are too dense. Used internally by CrispRVariants for creating alignment plot with `plotAlignments`.

**Usage**

```
.getAxisCoords(
  locations,
  labels = NULL,
  loc.boundaries = NULL,
  lab.boundaries = c(-1, 1),
  label.at = 5,
  min.tick.sep = 1
)
```

**Arguments**

<code>locations</code>	character(n) Actual x coordinates, or the desired range of the x coordinates. If labels are provided, all tick locations must be in locations and have a matching label.
<code>labels</code>	character(n) labels for the x axis ticks. Should be the same length as locations if provided. Note that if not all tick locations are included in locations, it must be possible to extrapolate labels from locations (Default: NULL)
<code>loc.boundaries</code>	numeric(i) Locations that must be included. (Default: NULL)
<code>lab.boundaries</code>	numeric(j) Labels that must be included. (Default: c(-1,1), for showing the cut sites). Boundaries must be in labels and have a matching tick location.
<code>label.at</code>	numeric(1) Add ticks when label modulo label.at is zero (Default = 5)
<code>min.tick.sep</code>	numeric(1) Minimum distance between ticks, excluding boundary ticks. (Default: 1)

**Value**

A list containing vectors named `tick_locs` and `tick_labs`

**Author(s)**

Helen Lindsay

---

`.intersperse`*.intersperse*

---

**Description**

create a vector of elements in `outer` interspersed with elements in `inner`. Similar to python `zip`. No element checking.

**Usage**

```
.intersperse(outer, inner)
```

**Arguments**

<code>outer</code>	vector that will be the first and last elements
<code>inner</code>	vector that will join elements of <code>outer</code>

**Value**

A vector interspersing elements of `outer` and `inner`. If `outer` is `c(a,b,c)` and `inner` is `c(d,e)`, returns `c(a,d,b,e,c)`

**Author(s)**

Helen Lindsay

**Examples**

```
CrispRVariants:::.intersperse(c(1:10), c(1:9)*10)
```

---

*.invertKeepRanges*      *Helper functions for selectAlnRegions*

---

**Description**

(.invertKeepRanges) Internal CrispRVariants function used by seqsToPartialAlns for checking arguments and getting region to delete. Returns FALSE if no region to delete found, or region to be deleted is entire target.

(.checkRelativeLocs) Shift keep to start at 1 if it is within the target

(.adjustRelativeInsLocs) Internal CrispRVariants function for shifting insertion locations relative to the target region when removing a segment of the alignments. Note that this function does not do input checking but assumes this has been done upstream. Insertions at the left border of a gap region are removed.

(.offsetIndices) Get indices of a vector grouped by "x", cumulatively adding offsets to each group according to "offset"

**Usage**

```
.invertKeepRanges(target, keep)

.checkRelativeLocs(target, keep)

.adjustRelativeInsLocs(target, keep, starts, gap_nchars)

.offsetIndices(x, offset)
```

**Arguments**

target	The complete region spanned by the alignments (GRanges)
keep	Region to display, relative to the target region, i.e. not genomic coords (IRanges or GRanges)
starts	numeric(n) Insertion locations. When plotting, the insertion symbol appears at the left border of the start location.
gap_nchars	character(n) Number of letters added to when joining segments before each region in keep. If first base of keep is 1, the first entry of gap_nchars should be 0.
x	A vector of group lengths
offset	A vector of offset lengths matching x

**Value**

Gaps between keep (IRanges), or FALSE if no gap ranges found  
insertion\_sites (data.frame) with modified start column

**Author(s)**

Helen Lindsay

**Examples**

```
CrispRVariants:::.offsetIndices(rep(2,5), c(0:4)*10)
```

---

abifToFastq

---

*Read a file in ab1 (Sanger) format and convert to fastq*


---

**Description**

This is an R implementation of Wibowo Arindrarto's abifpy.py trimming module, which itself implements Richard Mott's trimming algorithm See <https://github.com/bow/abifpy> for more details.

**Usage**

```
abifToFastq(
  seqname,
  fname,
  outfname,
  trim = TRUE,
  cutoff = 0.05,
  min_seq_len = 20,
  offset = 33,
  recall = FALSE
)
```

**Arguments**

seqname	name of sequence, to appear in fastq file
fname	filename of sequence in ab1 format
outfname	filename to append the fastq output to
trim	should low quality bases be trimmed from the ends? TRUE or FALSE
cutoff	probability cutoff
min_seq_len	minimum number of sequenced bases required in order to trim the read
offset	phred offset for quality scores
recall	Use sangerseqR to resolve the primary sequence if two sequences are present. May cause quality scores to be ignored. (Default: FALSE)

**Details**

Requires Bioconductor package SangerseqR



**Value**

None. Sequences are appended to the outfname.

**Author(s)**

Helen Lindsay

**Examples**

```
# Running this code will write the fastq file to "IM2033.fastq"
ab1_fname <- system.file("extdata", "IM2033.ab1", package = "CrispRVariants")
abifToFastq("IM2033", ab1_fname, "IM2033.fastq")
```

---

addClipped

*Extrapolates mapping location from clipped, aligned reads*

---

**Description**

Extrapolates the mapping location of a read by assuming that the clipped regions should map adjacent to the mapped locations. This is not always a good assumption, particularly in the case of chimeric reads!

**Usage**

```
addClipped(bam, ...)

## S4 method for signature 'GAlignments'
addClipped(bam, ...)
```

**Arguments**

bam	A GAlignments object
...	additional arguments

**Value**

A [GRanges](#) representation of the extended mapping locations

**Author(s)**

Helen Lindsay

---

addCodonFrame	<i>Internal CrispRVariants function for indicating codon frame on an alignment tile plot</i>
---------------	--

---

### Description

Adds vertical dotted lines in intervals of three nucleotides. Codon frame is supplied, alignments are assumed not to span an intron-exon junction.

### Usage

```
addCodonFrame(p, width, codon.frame)
```

### Arguments

p	A ggplot object, typically from CrispRVariants:::makeAlignmentTilePlot
width	The number of nucleotides in the alignments
codon.frame	The leftmost starting location of the next codon - 1,2,or 3

### Value

A ggplot object with added vertical lines indicating the frame

### Author(s)

Helen Lindsay

---

alleles	<i>Get allele names</i>
---------	-------------------------

---

### Description

Function to access allele names

### Usage

```
alleles(obj, ...)

## S4 method for signature 'CrisprSet'
alleles(obj, ...)
```

### Arguments

obj	An object containing variant alleles
...	additional arguments

**Value**

A data frame relating CIGAR strings to variant labels

**Author(s)**

Helen Lindsay

**Examples**

```
data("gol_clutch1")
alleles <- alleles(gol)
```

---

alns

*Get alignments*

---

**Description**

Return alignments from an object that contains them. For a `CrisprSet` object, these are truncated, non-chimeric alignments

**Usage**

```
alns(obj, ...)
```

## S4 method for signature 'CrisprSet'

```
alns(obj, ...)
```

**Arguments**

obj	An object containing aligned sequences
...	additional arguments

**Value**

A `GAlignmentsList` of consensus sequences on the positive strand.

**Author(s)**

Helen Lindsay

**Examples**

```
data("gol_clutch1")
alns <- alns(gol)
```

---

annotateGenePlot	<i>Plots and annotates transcripts</i>
------------------	--

---

## Description

Plots the gene structure, annotates this with the target location

## Usage

```
annotateGenePlot(
  txdb,
  target,
  target.colour = "red",
  target.size = 1,
  gene.text.size = 10,
  panel.spacing = grid::unit(c(0.1, 0.1, 0.1, 0.1), "lines"),
  plot.title = NULL,
  all.transcripts = TRUE
)
```

## Arguments

<code>txdb</code>	A GenomicFeatures:TxDb object
<code>target</code>	Location of target (GRanges)
<code>target.colour</code>	Colour of box indicating targt region
<code>target.size</code>	Thickness of box indicating target region
<code>gene.text.size</code>	Size for figure label
<code>panel.spacing</code>	Unit object, margin size
<code>plot.title</code>	A title for the plot. If no plot.title is supplied, the title is the list of gene ids shown (default). If plot.title == FALSE, the plot will not have a title.
<code>all.transcripts</code>	If TRUE (default), all transcripts of genes overlapping the target are shown, including transcripts that do not themselves overlap the target. If FALSE, only the transcripts that overlap the target are shown.

## Value

A ggplot2 plot of the transcript structures

---

arrangePlots	<i>Arrange plots for plotVariants:CrisprSet</i>
--------------	---

---

## Description

Arranges 3 plots in two rows. The vertical margins of the left.plot and right.plot constrained to be equal

## Usage

```
arrangePlots(
  top.plot,
  left.plot,
  right.plot,
  fig.height = NULL,
  col.width.ratio = c(2, 1),
  row.ht.ratio = c(1, 6),
  left.plot.margin = grid::unit(c(0.1, 0.2, 3, 0.2), "lines")
)
```

## Arguments

top.plot	ggplot grob, placed on top of the figure, spanning the figure width
left.plot	ggplot, placed in the second row on the left
right.plot	ggplot, placed in the second row on the right. y-axis labels are removed.
fig.height	Actual height for the figure. If not provided, figure height is the sum of the row.ht.ratio (Default: NULL)
col.width.ratio	Vector specifying column width ratio (Default: c(2, 1))
row.ht.ratio	Vector specifying row height ratio (Default: c(1,6))
left.plot.margin	Unit object specifying margins of left.plot. Margins of right.plot are constrained by the left.plot.

## Value

The arranged plots

---

barplotAlleleFreqs      *Plots barplots of the spectrum of variants for a sample set*


---

### Description

For signature "matrix", this function optionally does a very naive classification of variants by size. Frameshift variant combinations are those whose sum is not divisible by three. Intron boundaries are *\*NOT\** considered, use with caution! For signature "CrisprSet", the function uses the VariantAnnotation package to localize variant alleles with respect to annotated transcripts. Variants are annotated as "coding" when they are coding in any transcript.

(signature("CrisprSet")) Groups variants by size and type and produces a barplot showing the variant spectrum for each sample. Accepts all arguments accepted by barplotAlleleFreqs for signature("matrix"). Requires package "VariantAnnotation"

signature("matrix") Accepts a matrix of allele counts, with rownames being alleles and column names samples.

### Usage

```
barplotAlleleFreqs(obj, ...)

## S4 method for signature 'CrisprSet'
barplotAlleleFreqs(
  obj,
  ...,
  txdb,
  min.freq = 0,
  include.chimeras = TRUE,
  group = NULL,
  palette = c("rainbow", "bluered")
)

## S4 method for signature 'matrix'
barplotAlleleFreqs(
  obj,
  category.labels = NULL,
  group = NULL,
  bar.colours = NULL,
  group.colours = NULL,
  legend.text.size = 10,
  axis.text.size = 10,
  legend.symbol.size = 1,
  snv.label = "SNV",
  novar.label = "no variant",
  chimera.label = "Other",
  include.table = TRUE,
  classify = TRUE
)
```

**Arguments**

<code>obj</code>	The object to be plotted
<code>...</code>	additional arguments
<code>txdb</code>	A transcript database object
<code>min.freq</code>	Include variants with at frequency least <code>min.freq</code> in at least one sample. (Default: 0, i.e. no cutoff)
<code>include.chimeras</code>	Should chimeric reads be included in results? (Default: TRUE)
<code>group</code>	A grouping factor for the columns in <code>obj</code> . Columns in the same group will be displayed in the same text colour (Default: NULL)
<code>palette</code>	Colour palette. Options are "rainbow", a quantitative palette (default) or "bluered", a gradient palette.
<code>category.labels</code>	Labels for each category, corresponding to the rows of <code>obj</code> . Only applicable when categories are provided, i.e. "classify" is FALSE. (Default: NULL)
<code>bar.colours</code>	Colours for the categories in the barplot. Colours must be provided if there are more than 6 different categories.
<code>group.colours</code>	Colours for the text labels for the experimental groups A set of 15 different colours is provided.
<code>legend.text.size</code>	The size of the legend text, in points.
<code>axis.text.size</code>	The size of the axis text, in points
<code>legend.symbol.size</code>	The size of the symbols in the legend
<code>snv.label</code>	The row label for single nucleotide variants
<code>novar.label</code>	The row label for non-variant sequences
<code>chimera.label</code>	The row label for chimeric (non-linearly aligned) variant alleles
<code>include.table</code>	Should a table of allele (variant combination) counts and total sequences be plotted? (Default: TRUE)
<code>classify</code>	If TRUE, performs a naive classification by size (Default: TRUE)

**Value**

A ggplot2 barplot of the allele distribution and optionally a table of allele counts

**Author(s)**

Helen Lindsay

**Examples**

```
data("gol_clutch1")
barplotAlleleFreqs(variantCounts(gol))

# Just show the barplot without the counts table:
barplotAlleleFreqs(variantCounts(gol), include.table = FALSE)
```

---

collapsePairs	<i>Internal CrispRVariants function for collapsing pairs with concordant indels</i>
---------------	---

---

## Description

Given a set of alignments to a target region, finds read pairs. Compares insertion/deletion locations within pairs using the cigar string. Pairs with non-identical indels are excluded. Pairs with identical indels are collapsed to a single read, taking the consensus sequence of the pairs.

## Usage

```
collapsePairs(
  alns,
  use.consensus = TRUE,
  keep.unpaired = TRUE,
  verbose = TRUE,
  ...
)
```

## Arguments

alns	A GAlignments object. We do not use GAlignmentPairs because amplicon-seq can result in pairs in non-standard pairing orientation. Must include BAM flag, must not include unmapped reads.
use.consensus	Should the consensus sequence be used if pairs have a mismatch? Setting this to be TRUE makes this function much slower (Default: TRUE)
keep.unpaired	Should unpaired and chimeric reads be included? (Default: TRUE)
verbose	Report statistics on reads kept and excluded
...	Additional items with the same length as alns, that should be filtered to match alns.

## Value

The alignments, with non-concordant pairs removed and concordant pairs represented by a single read.

## Author(s)

Helen Lindsay



---

consensusSeqs	<i>Get consensus sequences for variant alleles</i>
---------------	--

---

**Description**

Return consensus sequences of variant alleles. At present, chimeric alignments are not included.

**Usage**

```
consensusSeqs(obj, ...)  
  
## S4 method for signature 'CrisprSet'  
consensusSeqs(obj, ..., top.n = NULL, min.freq = 0, min.count = 1)
```

**Arguments**

obj	An object containing aligned sequences
...	additional arguments
top.n	(Integer n) If specified, return variants ranked at least n according to frequency across all samples (Default: 0, i.e. no cutoff)
min.freq	(Float n least one sample (Default: 0)
min.count	(Integer n) Return variants with count greater than n in at least one sample (Default: 0)

**Value**

A DNASTringSet of consensus sequences on the positive strand.

**Author(s)**

Helen Lindsay

**Examples**

```
data("gol_clutch1")  
seqs <- consensusSeqs(gol, sample = 2)
```

---

countDeletions	<i>Count the number of reads containing an insertion or deletion</i>
----------------	--

---

### Description

Counts the number of reads containing a deletion or insertion (indel) of any size in a set of aligned reads. For countDeletions and countInsertions Reads may be filtered according to whether they contain more than one indel of the same or different types.

### Usage

```
countDeletions(alns, ...)

## S4 method for signature 'GAlignments'
countDeletions(
  alns,
  ...,
  multi.del = FALSE,
  del.and.ins = FALSE,
  del.ops = c("D")
)

countInsertions(alns, ...)

## S4 method for signature 'GAlignments'
countInsertions(
  alns,
  ...,
  ins.and.del = FALSE,
  multi.ins = FALSE,
  del.ops = c("D")
)

countIndels(alns)

## S4 method for signature 'GAlignments'
countIndels(alns)

indelPercent(alns)

## S4 method for signature 'GAlignments'
indelPercent(alns)
```

### Arguments

alns	The aligned reads
...	extra arguments

<code>multi.del</code>	If TRUE, returns the exact number of deletions, i.e., if one read contains 2 deletions, it contributes 2 to the total count (default: FALSE)
<code>del.and.ins</code>	If TRUE, counts deletions regardless of whether reads also contain insertions. If FALSE, counts reads that contain deletions but not insertions (default: FALSE)
<code>del.ops</code>	Cigar operations counted as deletions. Default: <code>c("D")</code>
<code>ins.and.del</code>	If TRUE, counts insertions regardless of whether reads also contain deletions. If FALSE, counts reads that contain insertions but not deletions (default: FALSE)
<code>multi.ins</code>	If TRUE, returns the exact number of insertions, i.e., if one read contains 2 insertions, it contributes 2 to the total count (default: FALSE)

**Value**

`countDeletions`: The number of reads containing a deletion (integer)  
`countInsertions`: The number of reads containing an insertion (integer)  
`countIndels`: The number of reads containing at least one insertion  
`indelPercent`: The percentage of reads containing an insertion or deletion (numeric)

**Author(s)**

Helen Lindsay

**Examples**

```
bam_fname <- system.file("extdata", "gol_F1_clutch_2_embryo_4_s.bam",
  package = "CrisprVariants")
bam <- GenomicAlignments::readGAlignments(bam_fname, use.names = TRUE)
countDeletions(bam)
countInsertions(bam)
countIndels(bam)
indelPercent(bam)
```

---

CrisprRun-class

*CrisprRun class*


---

**Description**

A ReferenceClass container for a single sample of alignments narrowed to a target region. Typically `CrisprRun` objects will not be accessed directly, but if necessary via a `CrisprSet` class which contains a list of `CrisprRun` objects. Note that the `CrisprVariants` plotting functions don't work on `CrisprRun` objects.

**Arguments**

<code>bam</code>	a GAlignments object containing (narrowed) alignments to the target region. Filtering of the bam should generally be done before initialising a CrisprRun object
<code>target</code>	The target location, a GRanges object
<code>genome.ranges</code>	A GRangesList of genomic coordinates for the cigar operations. If bam is a standard GAlignments object, this is equivalent to <code>cigarRangesAlongReferenceSpace + start(bam)</code>
<code>rc</code>	(reverse complement) Should the alignments be reverse complemented, i.e. displayed with respect to the negative strand? (Default: FALSE)
<code>name</code>	A name for this set of reads, used in plots if present (Default: NULL)
<code>chimeras</code>	Off-target chimeric alignments not in bam. (Default: empty)
<code>verbose</code>	Print information about initialisation progress (Default: FALSE)

**Fields**

<code>aln</code> s	A GAlignments object containing the narrowed reads. Note that if the alignments are represented with respect to the reverse strand, the "start" remains with respect to the forward strand, whilst the cigar and the sequence are reverse complemented.
<code>name</code>	The name of the sample
<code>cigar_labels</code>	A vector of labels for the reads, based on the cigar strings, optionally renumbered with respect to a new zero point (e.g. the cut site) and shortened to only insertion and deletion locations. Set at initialisation of a CrisprSet object, but not at initialisation of a CrisprRun object.
<code>chimeras</code>	Chimeric, off-target alignments corresponding to alignments in <code>aln</code> s

**Methods**

<code>getCigarLabels( target, target.loc, genome_to_target, ref, separate.snv, rc, match.label, mismatch.label)</code>	Description: Sets the "cig_labels" field, returns the cigar labels. Input parameters: <code>target</code> : (GRanges) the counting region. <code>target.loc</code> : The location of the cut site with respect to the target <code>genome_to_target</code> : A vector with names being genomic locations and values being locations with respect to the cut site <code>separate.snv</code> : Should single nucleotide variants be called? (Default: TRUE) <code>match.label</code> : Label for non-variant reads (Default: no variant) <code>mismatch.label</code> : Label for single nucleotide variants (Default: SNV) <code>rc</code> : Should the variants be displayed with respect to the negative strand? (Default: FALSE) <code>keep.ops</code> : CIGAR operations to remain in the variant label (usually indels) <code>upstream</code> : distance upstream of the cut site to call SNVs <code>downstream</code> : distance downstream of the cut site to call SNVs <code>regions</code> : IRanges(k) Regions for counting insertions and deletions. Insertions on the right border are not counted. <code>snv.regions</code> Regions for calling SNVs
<code>getInsertionSeqs(target)</code>	Description: Return a table relating insertion sequences to alignment indices Input parameters:

**Author(s)**

Helen Lindsay

**See Also**[CrisprSet](#)**Examples**

```
# readsToTarget with signature("GAlignments", "GRanges") returns a
# CrisprRun object

bam_fname <- system.file("extdata", "gol_F1_clutch_1_embryo_1_s.bam",
                          package = "CrisprVariants")
param <- Rsamtools::ScanBamParam(what = c("seq", "flag"))
alns <- GenomicAlignments::readGAlignments(bam_fname, param = param,
                                           use.names = TRUE)

reference <- Biostrings::DNAString("GGTCTCTCGCAGGATGTTGCTGG")
gd <- GenomicRanges::GRanges("18", IRanges::IRanges(4647377, 4647399), strand = "+")

crispr_run <- readsToTarget(alns, target = gd, reference = reference,
                           name = "Sample name", target.loc = 17)

# Alternatively, CrisprRun objects can be accessed from a CrisprSet object
# e.g. crispr_set$crispr_runs[[1]]
```

CrisprSet-class

*CrisprSet class***Description**

A ReferenceClass container for holding a set of narrowed alignments, each corresponding to the same target region. Individual samples are represented as CrisprRun objects. CrisprRun objects with no on-target reads are excluded. CrisprSet objects are constructed with [readsToTarget](#) or [readsToTargets](#). For most use cases, a CrisprSet object should not be initialized directly.

**Arguments**

<code>crispr.runs</code>	A list of CrisprRun objects, typically representing individual samples within an experiment
<code>reference</code>	The reference sequence, must be the same length as the target region
<code>target</code>	The target location (GRanges). Variants will be counted over this region. Need not correspond to the guide sequence.
<code>rc</code>	Should the alignments be reverse complemented, i.e. displayed w.r.t the reverse strand? (default: FALSE)
<code>names</code>	A list of names for each of the samples, e.g. for displaying in plots. If not supplied, the names of the crispr.runs are used, which default to the filenames of the bam files if available (Default: NULL)

<code>renumbered</code>	Should the variants be renumbered using <code>target.loc</code> as the zero point? If TRUE, variants are described by the location of their 5'-most base with respect to the <code>target.loc</code> . A 3bp deletion starting 5bp 5' of the cut site would be labelled as -5:3D (Default: TRUE)
<code>target.loc</code>	The location of the Cas9 cut site with respect to the supplied target. (Or some other central location). Can be displayed on plots and used as the zero point for renumbering variants. For a target region with the PAM location from bases 21-23, the <code>target.loc</code> is base 17 (default: 17)
<code>match.label</code>	Label for sequences with no variants (default: "no variant")
<code>mismatch.label</code>	Label for sequences with only single nucleotide variants (default: "SNV")
<code>bpparam</code>	A BiocParallel parameter object specifying how many cores to use. The parallelisable step is calling SNVs. Parallelisation is by sample. (default: SerialParam, i.e. no parallelization)
<code>verbose</code>	If true, prints information about initialisation progress (default: TRUE)

## Fields

`crispr_runs` A list of `CrisprRun` objects, typically corresponding to samples of an experiment.

`ref` The reference sequence for the target region, as a `Biostrings::DNAString` object

`cigar_freqs` A matrix of counts for each variant

`target` The target location, as a `GRanges` object

## Methods

`classifyCodingBySize(var_type, cutoff = 10)` Description: This is a naive classification of variants as frameshift or in-frame Coding indels are summed, and indels with sum divisible by 3 are considered frameshift. Note that this may not be correct for variants that span an intron-exon boundary Input paramters: `var_type`: A vector of `var_type`. Only variants with `var_type == "coding"` are considered. Intended to work with `classifyVariantsByLoc` cutoff: Variants are divided into those less than and greater than "cutoff" (Default: 10) Result: A character vector with a classification for each variant allele

`classifyVariantsByLoc(txdb, add_chr = TRUE, verbose = TRUE, ...)` Description: Uses the `VariantAnnotation` package to look up the location of the variants. `VariantAnnotation` allows multiple classification tags per variant, this function returns a single tag. The following preference order is used: `spliceSite > coding > intron > fiveUTR > threeUTR > promoter > intergenic`

Input parameters: `txdb`: A `BSgenome` transcription database `add_chr`: Add "chr" to chromosome names to make compatible with UCSC (default: TRUE) `verbose`: Print progress (default: TRUE) ...: Filtering arguments for `variantCounts`

Return value: A vector of classification tags, matching the rownames of `.self$cigar_freqs` (the variant count table)

`classifyVariantsByType(...)` Description: Classifies variants as insertions, deletions, or complex (combinations). In development Input parameters: ... Optional arguments to "variantCounts" for filtering variants before classification Return value: A named vector classifying variant alleles as insertions, deletions, etc

`consensusAlleles( cig_freqs = .self$cigar_freqs, return_nms = TRUE, match.ops = c("M", "X", "=") )`  
 Description: Get variants by their cigar string, make the pairwise alignments for the consensus sequence for each variant allele

Input parameters: `cig_freqs`: A table of variant allele frequencies (by default: `.self$cigar_freqs`, but could also be filtered) `return_nms`: If true, return a list of sequences and labels (Default: FALSE) `match.ops`: CIGAR operations for 1-1 alignments

Return: A `DNASet` of the consensus sequences for the specified alleles, or a list containing the consensus sequences and names for the labels if `return_nms = TRUE`

`filterUniqueLowQual(min_count = 2, max_n = 0, verbose = TRUE)` Description: Deletes reads containing rare variant combinations and more than a minimum number of ambiguity characters within the target region. These are assumed to be alignment errors.

Input parameters: `min_count`: the number of times a variant combination must occur across all samples to keep (default: 2, i.e. a variant must occur at least twice in one or more samples to keep) `max_n`: maximum number of ambiguity ("N") bases a read with a rare variant combination may contain. (default: 0) `verbose`: If TRUE, print the number of sequences removed (default: TRUE)

`filterVariants( cig_freqs = NULL, names = NULL, columns = NULL, include.chimeras = TRUE )`

Description: Relabels specified variants in a table of variant allele counts as non-variant, e.g. variants known to exist in control samples. Accepts either a size, e.g. "1D", or a specific mutation, e.g. "-4:3D". For alleles that include one variant to be filtered and one other variant, the other variant will be retained. If SNVs are included, these will be removed entirely, but note that SNVs are only called in reads that do not contain an insertion/deletion variant

Input parameters: `cig_freqs`: A table of variant allele counts (Default: NULL, i.e. `.self$cigar_freqs`) `names`: Labels of variants alleles to remove (Default: NULL) `columns`: Indices or names of control samples. Remove all variants that occur in these columns. (Default: NULL) `include.chimeras`: Should chimeric reads be included? (Default: TRUE)

`heatmapCigarFreqs( as.percent = TRUE, x.size = 8, y.size = 8, x.axis.title = NULL, x.angle = 90, min.freq =`

Description: Internal method for `CrisprVariants::plotFreqHeatmap`, optionally filters the table of variants, then a table of variant counts, coloured by counts or proportions.

Input parameters: `as.percent`: Should colours represent the percentage of reads per sample (TRUE) or the actual counts (FALSE)? (Default: TRUE) `x.size`: Font size for x axis labels (Default: 8) `y.size`: Font size for y axis labels (Default: 8) `x.axis.title`: Title for x axis `min.freq`: Include only variants with frequency at least `min.freq` in at least one sample `min.count`: Include only variants with count at least `min.count` in at least one sample `top.n`: Include only the n most common variants `type`: Should labels show counts or proportions? (Default: counts) `header`: What should be displayed in the header of the heatmap. Default: total count for type = "counts" or proportion of reads shown in the matrix for type = "proportions". If "counts" is selected, total counts will be shown for both types. "efficiency" shows the mutation efficiency (calculated with default settings) `order`: Reorder the columns according to this order (Default: NULL) `alleles`: Names of alleles to include. Selection of alleles takes place after filtering (Default: NULL). `exclude`: Names of alleles to exclude (Default: NULL) `create.plot`: Should the plot be created (TRUE, default), or the data used in the plot returned. ...: Extra filtering or plotting options

Return value: A `ggplot2` plot object. Call `"print(obj)"` to display

See also: `CrisprVariants::plotFreqHeatmap`

`makePairwiseAlns(cig_freqs = .self$cigar_freqs, ...)` Description: Get variants by their cigar string, make the pairwise alignments for the consensus sequence for each variant allele

Input parameters: `cig_freqs`: A table of variant allele frequencies (by default: `.self$cigar_freqs`, but could also be filtered) ...: Extra arguments for `CrisprVariants::seqsToAln`, e.g. which symbol should be used for representing deleted bases

```
mutationEfficiency(snv = c("non_variant", "include", "exclude"), include.chimeras = TRUE, exclude.cols =
```

Description: Calculates summary statistics for the mutation efficiency, i.e. the percentage of reads that contain a variant. Reads that do not contain an insertion or deletion, but do contain a single nucleotide variant (snv) can be considered as mutated, non-mutated, or not included in efficiency calculations as they are ambiguous. Note: `mutationEfficiency` does not treat partial alignments differently

Input parameters: `snv`: One of "include" (consider reads with mismatches to be mutated), "exclude" (do not include reads with snvs in efficiency calculations), and "non\_variant" (consider reads with mismatches to be non-mutated). `include.chimeras`: Should chimeras be counted as variants? (Default: TRUE) `exclude.cols`: A list of column names to exclude from calculation, e.g. if one sample is a control (default: NULL, i.e. include all columns) `group`: A grouping variable. Efficiency will be calculated per group, instead of for individual. Cannot be used with `exclude.cols`. `filter.vars`: Variants that should not be counted as mutations. `filter.cols`: Column names to be considered controls. Variants occurring in a control sample will not be counted as mutations. `count.alleles`: If TRUE, also report statistics about the number of alleles per sample/per group. (Default: FALSE) `per.sample`: Return efficiencies for each sample (Default: TRUE) `min.freq`: Minimum frequency for counting alleles. Does not apply to calculating efficiency. To filter when calculating efficiency, first use "variantCounts". (Default: 0, i.e. no filtering) Return value: A vector of efficiency statistics per sample and overall, or a matrix if a group is supplied.

```
plotVariants(min.freq = 0, min.count = 0, top.n = nrow(.self$cigar_freqs), alleles = NULL, renumbered = .s
```

Description: Internal method for `CrisprVariants:plotAlignments`, optionally filters the table of variants, then plots variants with respect to the reference sequence, collapsing insertions and displaying insertion sequences below the plot.

Input parameters: `min.freq`: i (in at least one sample) `min.count`: i (integer) include variants that occur at least i times in at least one sample `top.n`: n (integer) Plot only the n most frequent variants (default: plot all) Note that if there are ties in variant ranks, `top.n` only includes ties with all members ranking  $\leq$  `top.n` `alleles`: Alleles to include after filtering. Default NULL means use all alleles that pass filtering. `renumbered`: If TRUE, the x-axis is numbered with respect to the target (cut) site. If FALSE, x-axis shows genomic locations. (default: TRUE) `add.other`: Add a blank row named "Other" for chimeric alignments, if there are any (Default: TRUE) `create.plot`: Data is plotted if TRUE and returned without if FALSE. (Default: TRUE) `plot.regions`: Subregion of the target to plot (Default: NULL) `allow.partial`: Should partial alignments be allowed? (Default: TRUE) ... additional arguments for `plotAlignments`

Return value: A ggplot2 plot object. Call "print(obj)" to display

### Author(s)

Helen Lindsay

### See Also

[readsToTarget](#) and [readsToTargets](#) for initialising a `CrisprSet`, [CrisprRun](#) container for sample data.



**Examples**

```
# Load the metadata table
md_fname <- system.file("extdata", "gol_F1_metadata_small.txt", package = "CrispRVariants")
md <- read.table(md_fname, sep = "\t", stringsAsFactors = FALSE)

# Get bam filenames and their full paths
bam_fnames <- sapply(md$bam.filename, function(fn){
  system.file("extdata", fn, package = "CrispRVariants")})

reference <- Biostrings::DNAString("GGTCTCTCGCAGGATGTTGCTGG")
gd <- GenomicRanges::GRanges("18", IRanges::IRanges(4647377, 4647399), strand = "+")

crispr_set <- readsToTarget(bam_fnames, target = gd, reference = reference,
                           names = md$experiment.name, target.loc = 17)
```

---

dispatchDots

*dispatchDots*


---

**Description**

Update default values for func with values from dot args

**Usage**

```
dispatchDots(func, ..., call = FALSE)
```

**Arguments**

func	Function to call
...	dot args to pass to function
call	If TRUE, call the function with the argument list and return this result (Default: FALSE)

**Value**

A list of arguments to pass to func, or if call is TRUE, the result of calling func with these arguments.

**Author(s)**

Helen Lindsay

**Examples**

```
# Set up a function to dispatch dot arguments to:
f <- function(a=1, b=2, c=3){
  print(c(a,b,c))
}
# Set up a function for passing dots:
```

```

g <- function(...){
  CrispRVariants:::dispatchDots(f, ...)
}

g(a = 5)
g(a = 5, call = TRUE)
# Unrelated arguments will not be passed on
g(a = 5, d = 6)

```

---

excludeFromBam	<i>Removes reads from a bam file</i>
----------------	--------------------------------------

---

### Description

Returns a GAlignments excluding reads based on either name and/or location

### Usage

```
excludeFromBam(bam, exclude.ranges = GRanges(), exclude.names = NA)
```

### Arguments

bam                    a GAlignments object  
 exclude.ranges    Regions to exclude, as [GRanges](#).  
 exclude.names    A character vector of alignments names to exclude

### Value

The bam minus the excluded regions

### Author(s)

Helen Lindsay

---

findChimeras	<i>Find chimeric reads</i>
--------------	----------------------------

---

### Description

Find chimeric reads, assuming that the GAlignments object does not contain multimapping reads. That is, read names that appear more than ones in the file are considered chimeras. Chimeric reads are reads that cannot be mapped as a single, linear alignment. Reads from structural rearrangements such as inversions can be mapped as chimeras. Note that the indices of all chimeric reads are returned, these are not separated into individual chimeric sets.

**Usage**

```
findChimeras(bam, by.flag = FALSE)
```

**Arguments**

bam	A GAlignments object, must include names
by.flag	Can the chimeras be detected just using the supplementary alignment flag? (Default: FALSE). If TRUE, detects supplementary alignments and returns reads with the same name as a supplementary alignment (quicker). If FALSE, all alignments with duplicated names are returned.

**Value**

A vector of indices of chimeric sequences within the original bam

**Author(s)**

Helen Lindsay

**See Also**

[plotChimeras](#) for plotting chimeric alignment sets.

**Examples**

```
bam_fname <- system.file("extdata", "gol_F1_clutch_2_embryo_4_s.bam",
                          package = "CrisprVariants")
bam <- GenomicAlignments::readGAlignments(bam_fname, use.names = TRUE)
chimera_indices <- findChimeras(bam)
chimeras <- bam[chimera_indices]
```

---

findSNVs

*Find frequent SNVs*


---

**Description**

Find single nucleotide variants (SNVs) above a specified frequency in a table of variants.

**Usage**

```
findSNVs(obj, ...)

## S4 method for signature 'CrisprSet'
findSNVs(obj, ..., freq = 0.25, include.chimeras = TRUE)
```

**Arguments**

obj	An object containing variant counts
...	additional arguments
freq	minimum frequency snv to return (Default: 0.25)
include.chimeras	include chimeric reads when calculating SNV frequencies (Default: TRUE)

**Value**

A vector of SNVs and their frequencies

**Author(s)**

Helen Lindsay

---

getChimeras	<i>Get chimeric alignments</i>
-------------	--------------------------------

---

**Description**

Return chimeric alignments from a collection of aligned sequences

**Usage**

```
getChimeras(obj, ...)

## S4 method for signature 'CrisprSet'
getChimeras(obj, ..., sample)
```

**Arguments**

obj	An object containing aligned sequences
...	additional arguments
sample	The sample name or sample index to return

**Value**

A GAlignment object containing the chimeric read groups

**Author(s)**

Helen Lindsay

**Examples**

```
data("gol_clutch1")
chimeras <- getChimeras(gol, sample = 2)
```

---

getInsertionsTable	<i>getInsertionsTable</i>
--------------------	---------------------------

---

**Description**

Returns a table of insertion sequences present in a GAlignments object. This table is aggregated and used by plotAlignments.

**Usage**

```
getInsertionsTable(alns, pos = 1L)
```

**Arguments**

alns	A GAlignments object
pos	(Integer(1)) The amount by which to shift genomic coordinates upstream to get coordinates relative to a display region

**Value**

A data frame of insertion sequences, genomic and relative locations

**Author(s)**

Helen Lindsay

---

gol_clutch1	<i>Variant sequences from golden clutch 1 (Burger et al)</i>
-------------	--

---

**Description**

This dataset is a subset of the crispant data for the golden gene used by Burger et al (submitted).

**Usage**

```
data(gol_clutch1)
```

**Format**

A CrisprSet object countaining 8 samples

**Details**

- gol The variants as a CrisprSet object

**Value**

A CrisprSet object named "gol"

indelLabels

*indelLabels*

---

**Description**

Makes allele labels for insertion / deletion variants

**Usage**

```
indelLabels(  
  alns,  
  rc = FALSE,  
  genome.to.pos = NULL,  
  keep.ops = c("I", "D", "N"),  
  regions = NULL,  
  as.string = TRUE,  
  ...  
)
```

**Arguments**

<code>alns</code>	(GAlignments) aligned reads for finding variants
<code>rc</code>	Should the variants be displayed with respect to the negative strand? (Default: FALSE)
<code>genome.to.pos</code>	A vector with names being genomic locations and values being positions to use in labels (Default: NULL)
<code>keep.ops</code>	CIGAR operations to remain in the variant label (usually indels)
<code>regions</code>	IRanges(k) Regions for counting insertions and deletions. Insertions on the right border are not counted.
<code>as.string</code>	Return labels as strings (Default: TRUE)
<code>...</code>	extra formatting arguments

**Value**

A vector of labels for alns

**Author(s)**

Helen Lindsay

---

makeAlignmentTilePlot *Internal CrispRVariants function for creating the plotAlignments background*

---

## Description

Takes a matrix of characters, x and y locations and colours, creates a ggplot geom\_tile plot with tiles labelled by the characters.

## Usage

```
makeAlignmentTilePlot(  
  m,  
  ref,  
  xlab,  
  plot.text.size,  
  axis.text.size,  
  xtick.labs,  
  xtick.breaks,  
  tile.height  
)
```

## Arguments

m	A matrix with column headings Var1: y location, Var2: x location, cols: tile fill colour, isref: transparency value text_cols: text colour
ref	The reference sequence, only used for checking the number of x-tick labels when x-tick breaks are not supplied
xlab	Label for the x axis
plot.text.size	Size for text within plot
axis.text.size	Size for text on axes
xtick.labs	x axis labels
xtick.breaks	Locations of x labels
tile.height	Controls whitespace between tiles

## Value

A ggplot object

## Author(s)

Helen Lindsay

mergeChimeras

*mergeChimeras***Description**

Merges chimeric alignments where the individual segments border an unmapped region (a long deletion). If bases of the read are mapped to both ends of the gap, the multimapped reads are only included in the leftmost genomic segment. If there are more than `max_unmapped` unmapped bases between the mapped bases, the read is not considered mergeable. Currently experimental and only tested with reads mapped by bwa mem.

**Usage**

```
mergeChimeras(
  bam,
  chimera_idx = NULL,
  verbose = TRUE,
  max_read_overlap = 10,
  max_unmapped = 4,
  name = NULL
)
```

**Arguments**

<code>bam</code>	A <code>GenomicAlignments::GAlignments</code> object
<code>chimera_idx</code>	Indices of chimeric reads within <code>bam</code>
<code>verbose</code>	Should information about the number of mergeable alignments be printed? (Default: TRUE)
<code>max_read_overlap</code>	Maximum number of bases in a mergeable read that are aligned to two genomic locations (Default: 10)
<code>max_unmapped</code>	Maximum number of bases in a mergeable read that are unmapped and located between two mapped segments (Default: 4)
<code>name</code>	Name of the sample, used when reporting verbose output.

**Value**

A list of the merged and unmerged chimeric alignments

**Author(s)**

Helen Lindsay



---

mergeCrisprSets	<i>Merge two CrisprSets</i>
-----------------	-----------------------------

---

**Description**

Merge two CrisprSet objects sharing a reference and target location

**Usage**

```
mergeCrisprSets(x, y, ...)  
  
## S4 method for signature 'CrisprSet,CrisprSet'  
mergeCrisprSets(  
  x,  
  y,  
  ...,  
  x.samples = NULL,  
  y.samples = NULL,  
  names = NULL,  
  order = NULL  
)
```

**Arguments**

x	A CrisprSet object
y	A second CrisprSet object
...	extra arguments
x.samples	A subset of column names or indices to keep from CrisprSet x (Default: NULL, i.e. keep all)
y.samples	A subset of column names or indices to keep from CrisprSet y (Default: NULL, i.e. keep all)
names	New names for the merged CrisprSet object (Default: NULL)
order	A list of sample names, matching the names in x and y, specifying the order of the samples in the new CrisprSet. (Not implemented yet)

**Value**

A merged CrisprSet object

**Author(s)**

Helen Lindsay

## Examples

```
# Load the metadata table
md_fname <- system.file("extdata", "gol_F1_metadata_small.txt", package = "CrispRVariants")
md <- read.table(md_fname, sep = "\t", stringsAsFactors = FALSE)

# Get bam filenames and their full paths
bam_fnames <- sapply(md$bam.filename, function(fn){
  system.file("extdata", fn, package = "CrispRVariants")})

reference <- Biostrings::DNAString("GGTCTCTCGCAGGATGTTGCTGG")
gd <- GenomicRanges::GRanges("18", IRanges::IRanges(4647377, 4647399),
  strand = "+")

crispr_set1 <- readsToTarget(bam_fnames[c(1:4)], target = gd,
  reference = reference, names = md$experiment.name[1:4], target.loc = 17)
crispr_set2 <- readsToTarget(bam_fnames[c(5:8)], target = gd,
  reference = reference, names = md$experiment.name[5:8], target.loc = 17)
mergeCrisprSets(crispr_set1,crispr_set2)
```

---

mismatchLabels

nonindelLabels

---

## Description

Make variant labels for variants without an insertion or deletion

## Usage

```
mismatchLabels(
  alns,
  target,
  ref.seq,
  regions = NULL,
  min.pct = 0,
  mismatch.label = "SNV",
  genome.to.pos = NULL,
  as.string = TRUE
)
```

## Arguments

alns	A GAlignments object, where the aligned sequences should span the reference sequence
target	(GRanges(1)) The region for counting mismatches
ref.seq	A DNAString object, the sequence for comparison when checking for mismatches. The sequence does not necessarily have to match the mapping reference sequence. Must span all regions if regions are provided.

regions	A GRanges object, regions to check for mismatches with coordinates relative to the reference sequence
min.pct	(numeric(1), between 0 and 100) Only return SNVs that occur at in least min.pct change, not any change at a position.
mismatch.label	(character(1)) Label to append to the start of mismatch strings, if returning as a single string (Default: "SNV:")
genome.to.pos	Optional named vector for transforming variant coordinates into another coordinate system (Default: NULL)
as.string	Should individual variant labels be pasted into a single comma separated string when one alignment has multiple variants? (Default: TRUE)

**Value**

A data frame of sequence indices, genomic position of mismatch and mismatch base

---

mutationEfficiency	<i>Get mutation efficiency</i>
--------------------	--------------------------------

---

**Description**

Returns the percentage of sequences that contain at least one mutation.

**Usage**

```
mutationEfficiency(obj, ...)

## S4 method for signature 'CrisprSet'
mutationEfficiency(
  obj,
  ...,
  snv = c("non_variant", "include", "exclude"),
  include.chimeras = TRUE,
  exclude.cols = NULL,
  filter.vars = NULL,
  filter.cols = NULL,
  group = NULL
)
```

**Arguments**

obj	An object containing variant counts
...	additional arguments
snv	Single nucleotide variants (SNVs) may be considered as mutations ("include"), treated as ambiguous sequences and not counted at all ("exclude"), or treated as non-mutations, e.g. sequencing errors or pre-existing SNVs ("non_variant", default)

<code>include.chimeras</code>	Should chimeric alignments be counted as variants when calculating mutation efficiency (Default: TRUE)
<code>exclude.cols</code>	A vector of names of columns in the variant counts table that will not be considered when counting mutation efficiency
<code>filter.vars</code>	Variants to remove before calculating efficiency. May be either a variant size, e.g. "1D", or a particular variant/variant combination, e.g. -5:3D
<code>filter.cols</code>	A vector of control sample names. Any variants present in the control samples will be counted as non-variant, unless they also contain another indel. Note that this is not compatible with counting snvs as variants.
<code>group</code>	A grouping vector. If provided, efficiency will be calculated per group (Default: NULL)

**Value**

A vector of efficiency statistics per sample and overall, or a matrix of efficiency statistics per group if a group is provided

**Author(s)**

Helen Lindsay

**Examples**

```
data("gol_clutch1")
mutationEfficiency(gol)
```

---

narrowAlignments	<i>Narrow a set of aligned reads to a target region</i>
------------------	---

---

**Description**

Aligned reads are narrowed to the target region. In the case of reads with deletions spanning the boundaries of the target, reads are narrowed to the start of the deletion,

**Usage**

```
narrowAlignments(alns, target, ...)

## S4 method for signature 'GAlignments,GRanges'
narrowAlignments(
  alns,
  target,
  ...,
  reverse.complement,
  minoverlap = NULL,
  verbose = FALSE,
```

```

    clipping.ops = c("S", "H"),
    match.ops = c("M", "X", "=")
  )

```

### Arguments

<code>alns</code>	A GAlignments object including a metadata column "seq" containing the sequence
<code>target</code>	A GRanges object
<code>...</code>	additional arguments
<code>reverse.complement</code>	Should the aligned reads be reverse complemented?
<code>minoverlap</code>	Minimum overlapping region between alignments and target. If not specified, alignments must span the entire target region. (Default: NULL)
<code>verbose</code>	(Default: FALSE)
<code>clipping.ops</code>	CIGAR operations corresponding to clipping (Default: c("S","H"))
<code>match.ops</code>	CIGAR operations corresponding to a match, i.e. a non-indel position (Default: c("M","X","="))

### Value

The narrowed alignments (GAlignments)

### Author(s)

Helen Lindsay

### Examples

```

bam_fname <- system.file("extdata", "gol_F1_clutch_2_embryo_4_s.bam",
                          package = "CrispRVariants")
bam <- GenomicAlignments::readGAlignments(bam_fname, use.names = TRUE)
target <- GenomicRanges::GRanges("18", IRanges::IRanges(4647377, 4647399),
                                strand = "+")
narrowAlignments(bam, target, reverse.complement = FALSE)

```

## Description

(signature("CrisprSet")) Wrapper for `CrisprSet$plotVariants`. Optionally filters a `CrisprSet` frequency table, then plots variants. More information in [CrisprSet](#)

(signature("DNAString")) Plots a set of pairwise alignments to a reference sequence. Alignments should all be the same length as the reference sequences. This is achieved by removing insertions with respect to the reference, see [seqsToAln](#). Insertions are indicated by symbols in the plot and a table showing the inserted sequences below the plot. The default options are intended for a figure 6-8 inches wide, with figure height best chosen according to the number of different variants and insertions to be displayed.

## Usage

```
plotAlignments(obj, ...)

## S4 method for signature 'CrisprSet'
plotAlignments(
  obj,
  ...,
  min.freq = 0,
  min.count = 1,
  top.n = 50,
  renumbered = obj$pars[["renumbered"]],
  add.other = TRUE,
  create.plot = TRUE
)

## S4 method for signature 'character'
plotAlignments(
  obj,
  ...,
  alns,
  ins.sites,
  highlight.pam = TRUE,
  show.plot = FALSE,
  target.loc = 17,
  pam.start = NA,
  pam.end = NA,
  ins.size = 2,
  legend.cols = 3,
  xlab = NULL,
  xtick.labs = NULL,
  xtick.breaks = NULL,
  plot.text.size = 2,
  axis.text.size = 8,
  legend.text.size = 6,
  highlight.guide = TRUE,
  guide.loc = NULL,
  tile.height = 0.55,
```

```

    max.insertion.size = 20,
    min.insertion.freq = 5,
    line.weight = 1,
    legend.symbol.size = ins.size,
    add.other = FALSE,
    codon.frame = NULL,
    style = c("all", "mismatches")
)

## S4 method for signature 'DNAStrng'
plotAlignments(
  obj,
  ...,
  alns,
  ins.sites,
  highlight.pam = TRUE,
  show.plot = FALSE,
  target.loc = 17,
  pam.start = NA,
  pam.end = NA,
  ins.size = 2,
  legend.cols = 3,
  xlab = NULL,
  xtick.labs = NULL,
  xtick.breaks = NULL,
  plot.text.size = 2,
  axis.text.size = 8,
  legend.text.size = 6,
  highlight.guide = TRUE,
  guide.loc = NULL,
  tile.height = 0.55,
  max.insertion.size = 20,
  min.insertion.freq = 5,
  line.weight = 1,
  legend.symbol.size = ins.size,
  add.other = FALSE,
  codon.frame = NULL
)

```

### Arguments

<code>obj</code>	The object to be plotted
<code>...</code>	Additional arguments
<code>min.freq</code>	<code>i</code> (one sample (default: 0, i.e no frequency cutoff))
<code>min.count</code>	<code>i</code> (integer) only plot variants with count $\geq i$ in at least one sample (default: 0, i.e no count cutoff)
<code>top.n</code>	(integer) Plot only the <code>n</code> most frequent variants (default: 50)

<code>renumbered</code>	If TRUE, the x-axis is numbered with respect to the target (default: TRUE)
<code>add.other</code>	Add a blank row labelled "Other" to the plot, for combining with <code>plotFreqHeatmap</code> (default: TRUE (signature "CrisprSet") FALSE (signature "matrix"))
<code>create.plot</code>	Should the data be plotted? If false, returns the data used for plotting (Default: TRUE)
<code>alns</code>	A named character vector of aligned sequences, with insertions removed
<code>ins.sites</code>	A table of insertion_sites, which must include cols named "start", "cigar", "seq" and "count" for the start of the insertion in the corresponding sequence
<code>highlight.pam</code>	should location of PAM with respect to the target site be indicated by a box? (Default: TRUE) If TRUE, and <code>pam.start</code> and <code>pam.end</code> are not supplied, PAM is inferred from <code>target.loc</code>
<code>show.plot</code>	Should the plot be displayed (TRUE) or just returned as a ggplot object (FALSE). (Default: FALSE)
<code>target.loc</code>	The location of the zero point / cleavage location. Base n, where the zero point is between bases n and n+1
<code>pam.start</code>	The first location of the PAM with respect to the reference.
<code>pam.end</code>	The last location of the PAM with respect to the reference. Default is two bases after the <code>pam.start</code>
<code>ins.size</code>	The size of the symbols representing insertions within the plot.
<code>legend.cols</code>	The number of columns in the legend. (Default:3)
<code>xlab</code>	A title for the x-axis (Default: NULL)
<code>xtick.labs</code>	Labels for the x-axis ticks (Default: NULL)
<code>xtick.breaks</code>	Locations for x-axis tick breaks (Default: NULL)
<code>plot.text.size</code>	The size of the text inside the plot
<code>axis.text.size</code>	The size of the axis labels
<code>legend.text.size</code>	The size of the legend labels
<code>highlight.guide</code>	Should the guide be indicated by a box in the reference sequence? (Default: TRUE)
<code>guide.loc</code>	The location of the guide region to be highlighted, as an IRanges object. Will be inferred from <code>target.loc</code> if <code>highlight.guide</code> = TRUE and no <code>guide.loc</code> is supplied, assuming the guide plus PAM is 23bp (Default: NULL)
<code>tile.height</code>	The height of the tiles within the plot. (Default: 0.55)
<code>max.insertion.size</code>	The maximum length of an insertion to be shown in the legend. If <code>max.insertion.size</code> = n, an insertion of length m > n will be annotated as "mI" in the figure. (Default: 20)
<code>min.insertion.freq</code>	Display inserted sequences with frequency at least x amongst the sequences with an insertion of this size and length (Default: 5)
<code>line.weight</code>	The line thickness for the vertical line indicating the zero point (cleavage site) and the boxes for the guide and PAM. (Default: 1)



legend.symbol.size	The size of the symbols indicating insertions in the legend. (Default: ins.size)
codon.frame	Codon position of the leftmost nucleotide. If provided, codon positions in the specified frame are indicated. (Default: NULL)
style	One of "all" (colour all tiles) and "mismatches" (colour only mismatch positions)

**Value**

A ggplot2 figure

**Author(s)**

Helen Lindsay

**See Also**

[seqsToAln](#), [ggplot](#)

**Examples**

```
#Load a CrisprSet object and plot
data("gol_clutch1")
plotAlignments(gol)
```

---

plotChimeras

---

*Display a dot plot of chimeric alignments*


---

**Description**

Produces a dot plot of a set of chimeric alignments. For chimeric alignments, a single read is split into several, possibly overlapping aligned blocks. Aligned sections of chimeric reads can be separated by large genomic distances, or on separate chromosomes. plotChimeras produces a dot plot, each aligned block highlighted, and chromosomes shown in different colours. Large gaps between aligned segments are collapsed and indicated on the plot with horizontal lines. The X-axis shows each base of the entire read. Note that the mapping to the fwd strand is shown if all strands agree. The chimeric alignments must be sorted!

**Usage**

```
plotChimeras(
  chimeric.alns,
  max.gap = 10,
  tick.sep = 20,
  text.size = 10,
  title.size = 16,
  gap.pad = 20,
  legend.title = "Chromosome",
  xangle = 90,
```

```
wrt.forward = FALSE,
annotate.within = 20,
annotations = GenomicRanges::GRanges()
)
```

### Arguments

chimeric.alns	A GAlignments object containing only the chimeric reads to be plotted
max.gap	If aligned segments are separated by more than max.gap,
tick.sep	How many bases should separate tick labels on plot. Default 20.
text.size	Size of X and Y tick labels on plot. Default 12
title.size	Size of X and Y axis labels on plot. Default 16
gap.pad	How much should aligned blocks be separated by? (Default: 20)
legend.title	Title for the legend. Default "Chromosome"
xangle	Angle for x axis text (Default 90, i.e vertical)
wrt.forward	Should chimeric alignments where all members map to the negative strand be displayed with respect to the forward strand, i.e. as the cigar strand is written (TRUE), or the negative strand (FALSE) (Default: FALSE)
annotate.within	annot_aln ranges in "annotations" within n bases of a chimeric alignment (Default 50)
annotations	A list of GRanges. Any that overlap with the chimeric alignments are highlighted in the plot.

### Value

A ggplot2 dotplot of the chimeric alignments versus the reference sequence

### Author(s)

Helen Lindsay

### See Also

[findChimeras](#) for finding chimeric alignment sets.

### Examples

```
bam_fname <- system.file("extdata", "gol_F1_clutch_2_embryo_4_s.bam",
                          package = "CrisprVariants")
bam <- GenomicAlignments::readGAlignments(bam_fname, use.names = TRUE)
# Choose a single chimeric read set to plot:
chimeras <- bam[names(bam) == "AB3092"]

# This read aligns in 3 pieces, all on chromosome 18.
# The plot shows the alignment annot_alns a small duplication and
# a long gap.
plotChimeras(chimeras)
```

---

plotFreqHeatmap*Plot a table of counts with colours indicating frequency*

---

**Description**

Creates a heatmap from a matrix of counts or proportions, where tiles are coloured by the proportion and labeled with the value.

**Usage**

```
plotFreqHeatmap(obj, ...)  
  
## S4 method for signature 'matrix'  
plotFreqHeatmap(  
  obj,  
  ...,  
  col.sums = NULL,  
  header = NA,  
  header.name = "Total",  
  group = NULL,  
  group.colours = NULL,  
  as.percent = TRUE,  
  x.axis.title = NULL,  
  x.size = 6,  
  y.size = 8,  
  x.angle = 90,  
  legend.text.size = 6,  
  plot.text.size = 3,  
  line.width = 1,  
  x.hjust = 1,  
  legend.position = "right",  
  x.labels = NULL,  
  legend.key.height = grid::unit(1, "lines")  
)  
  
## S4 method for signature 'CrisprSet'  
plotFreqHeatmap(  
  obj,  
  ...,  
  top.n = 50,  
  min.freq = 0,  
  min.count = 1,  
  type = c("counts", "proportions"),  
  order = NULL,  
  alleles = NULL  
)
```

**Arguments**

<code>obj</code>	A matrix of counts with rows = feature, columns = sample
<code>...</code>	additional arguments
<code>col.sums</code>	Alternative column sums to be used for calculating the tile colours if <code>as.percent = TRUE</code> , e.g. if "obj" is a subset of a larger data set. If "NULL" (default), the column sums of "obj" are used.
<code>header</code>	Alternative column titles, e.g. column sums for the unfiltered data set when obj is a subset. If set to "NA", column sums of obj are displayed. If "NULL", no header is displayed (Default: NA).
<code>header.name</code>	Label for the header row (Default: "Total")
<code>group</code>	Grouping factor for columns. If supplied, columns are ordered to match the levels (Default: NULL)
<code>group.colours</code>	Colours for column groups, should match levels of "group". If "NULL", groups are coloured differently (Default: NULL)
<code>as.percent</code>	Should colours represent the percentage of reads per sample (TRUE) or the actual counts (FALSE)? (Default: TRUE)
<code>x.axis.title</code>	A title for the x-axis. (Default: NULL)
<code>x.size</code>	Font size for x-labels (Default: 16)
<code>y.size</code>	Font size for y-labels (Default: 16)
<code>x.angle</code>	Angle for x-labels (Default: 90, i.e. vertical)
<code>legend.text.size</code>	Font size for legend (Default: 16)
<code>plot.text.size</code>	Font size counts within plot (Default: 3)
<code>line.width</code>	Line thickness of title box'
<code>x.hjust</code>	Horizontal justification of x axis labels (Default: 1)
<code>legend.position</code>	The position of the legend (Default: right)
<code>x.labels</code>	X-axis labels (Default: NULL, column.names of the matrix, doesn't do anything at the moment)
<code>legend.key.height</code>	The height of the legend key, as a "unit" object. (See <a href="#">unit</a> ).
<code>top.n</code>	Show the n top ranked variants. Note that if the nth and n+1th variants have equal rank, they will not be shown. (Default: 50)
<code>min.freq</code>	i ( one sample (Default: 0, i.e no frequency cutoff)
<code>min.count</code>	i (integer) only plot variants with count >= i in at least one sample (default: 0, i.e no count cutoff)
<code>type</code>	Plot either "counts" or "proportions"
<code>order</code>	A list of column names or indices specifying the order of the columns in the plot
<code>alleles</code>	A list of alleles to include. Can be used to display only alleles of interest or to order the alleles. The default value NULL means all alleles passing the frequency cut offs will be included.

**Value**

The ggplot2 plot of the variant frequencies

**Examples**

```
#Load a CrisprSet object for plotting
data("gol_clutch1")

# Plot the frequency heatmap
plotFreqHeatmap(gol)
```

---

plotVariants	<i>Plot alignments, frequencies and location of target sequence</i>
--------------	---

---

**Description**

Combines a plot of transcript structure, alleles aligned with respect to a reference genome and a heatmap of counts or proportions of each allele in a set of data.

**Usage**

```
plotVariants(obj, ...)

## S4 method for signature 'CrisprSet'
plotVariants(
  obj,
  ...,
  txdb = NULL,
  add.chr = TRUE,
  plotAlignments.args = list(),
  plotFreqHeatmap.args = list()
)
```

**Arguments**

obj	The object to be plotted
...	extra arguments for plot layout
txdb	GenomicFeatures:TxDb object (default: NULL)
add.chr	If target chromosome does not start with "chr", e.g. "chr5", add the "chr" prefix. (Default:TRUE)
plotAlignments.args	Extra arguments for plotAlignments
plotFreqHeatmap.args	Extra arguments for plotFreqHeatmap

**Value**

A ggplot2 plot of the variants

**See Also**

[arrangePlots](#) for general layout options and [annotateGenePlot](#) for options relating to the transcript plot.

**Examples**

```
#Load a CrisprSet object for plotting
data("gol_clutch1")

#Load the transcript db. This is a subset of the Ensembl Danio Rerio v73 gtf
# for the region 18:4640000-4650000 which includes the targeted gol gene

library(GenomicFeatures)
fn <- system.file("extdata", "Danio_rerio.Zv9.73.gol.sqlite",
                  package = "CrisprVariants")
txdb <- loadDb(fn)

# Plot the variants
p <- plotVariants(gol, txdb = txdb)

#In the above plot, the bottom margin is too large, the legend is
#cut off, and the text within the plots should be larger.
#These issues can be fixed with some adjustments:
p <- plotVariants(gol, txdb = txdb,
                  plotAlignments.args = list(plot.text.size = 4, legend.cols = 2),
                  plotFreqHeatmap.args = list(plot.text.size = 4),
                  left.plot.margin = grid::unit(c(0.1,0.2,0.5,1), "lines"))
```

---

rcAlns

---

*Internal CrisprVariants function for determining read orientation*


---

**Description**

Function for determining whether reads should be oriented to the target strand, always displayed on the positive strand, or oriented to

**Usage**

```
rcAlns(target.strand, orientation)
```

**Arguments**

target.strand	The target strand (one of "+", "-", "*")
orientation	One of "target", "opposite" and "positive" (Default: "target")

**Value**

A logical value indicating whether reads should be reverse complemented

**Author(s)**

Helen Lindsay

---

readsByPCRPrimer

*Finds overlaps between aligned reads and PCR primers*


---

**Description**

Short reads amplified with PCR primers should start and end at defined positions. However, the ends of an aligned read may be clipped as sequencing technologies are prone to making errors at the start and end. readsByPCRPrimer extrapolates the genomic location of entire reads from their aligned sections by adding clipped sections, then finds near exact matches to a set of PCR primers. Note that this is not always a good assumption, and is misleading in the case of chimeric reads where sections clipped in one part of a chimera are aligned in another.

**Usage**

```
readsByPCRPrimer(bam, primers, ...)
```

```
## S4 method for signature 'GAlignments,GRanges'
```

```
readsByPCRPrimer(
  bam,
  primers,
  ...,
  tolerance = 0,
  verbose = TRUE,
  ignore.strand = TRUE,
  allow.partial = TRUE,
  chimera.idx = NULL
)
```

```
## S4 method for signature 'GRanges,GRanges'
```

```
readsByPCRPrimer(
  bam,
  primers,
  ...,
  tolerance = 0,
  verbose = TRUE,
  ignore.strand = TRUE,
  allow.partial = TRUE,
  chimera.idx = NULL
)
```

**Arguments**

bam	A set of aligned reads
primers	A set of ranges that the unclipped reads may map to
...	Additional arguments
tolerance	Number of bases by which reads and primers may differ at each end (Default: 0)
verbose	Print number of full and partial matches (Default: TRUE)
ignore.strand	Passed to <a href="#">findOverlaps</a> and <a href="#">disjoin</a> . Should strand be ignored when finding overlaps. (Default: TRUE)
allow.partial	Should reads that do not match the PCR boundaries, but map to a region covered by only one primer be considered matches? (Default: TRUE)
chimera.idx	Indices of chimeric reads within the bam. If specified, chimeras overlapping multiple pcr primers will be removed.

**Value**

A [Hits](#) object where "query" is the index with respect to bam and "subject" is the index with respect to the primers.

**Author(s)**

Helen Lindsay

**See Also**

[GRanges](#), [GAlignments](#)

---

readsToTarget	<i>Trims reads to a target region.</i>
---------------	--

---

**Description**

Trims aligned reads to one or several target regions, optionally reverse complementing the alignments.

**Usage**

```
readsToTarget(reads, target, ...)

## S4 method for signature 'GAlignments,GRanges'
readsToTarget(
  reads,
  target,
  ...,
  reverse.complement = TRUE,
```



```

    chimeras = NULL,
    collapse.pairs = FALSE,
    use.consensus = FALSE,
    store.chimeras = FALSE,
    verbose = TRUE,
    name = NULL,
    minoverlap = NULL,
    orientation = c("target", "opposite", "positive")
)

## S4 method for signature 'GAlignmentsList,GRanges'
readsToTarget(
  reads,
  target,
  ...,
  reference = reference,
  names = NULL,
  reverse.complement = TRUE,
  target.loc = 17,
  chimeras = NULL,
  collapse.pairs = FALSE,
  use.consensus = FALSE,
  orientation = c("target", "opposite", "positive"),
  minoverlap = NULL,
  verbose = TRUE
)

## S4 method for signature 'character,GRanges'
readsToTarget(
  reads,
  target,
  ...,
  reference,
  reverse.complement = TRUE,
  target.loc = 17,
  exclude.ranges = GRanges(),
  exclude.names = NA,
  chimeras = c("count", "exclude", "ignore", "merge"),
  collapse.pairs = FALSE,
  use.consensus = FALSE,
  orientation = c("target", "opposite", "positive"),
  names = NULL,
  minoverlap = NULL,
  verbose = TRUE
)

readsToTargets(reads, targets, ...)

```

```
## S4 method for signature 'character,GRanges'
readsToTargets(
  reads,
  targets,
  ...,
  references,
  primer.ranges = NULL,
  target.loc = 17,
  reverse.complement = TRUE,
  collapse.pairs = FALSE,
  use.consensus = FALSE,
  ignore.strand = TRUE,
  names = NULL,
  bpparam = BiocParallel::SerialParam(),
  orientation = c("target", "opposite", "positive"),
  chimera.to.target = 5,
  verbose = TRUE
)

## S4 method for signature 'GAlignmentsList,GRanges'
readsToTargets(
  reads,
  targets,
  ...,
  references,
  primer.ranges = NULL,
  target.loc = 17,
  reverse.complement = TRUE,
  collapse.pairs = FALSE,
  use.consensus = FALSE,
  ignore.strand = TRUE,
  names = NULL,
  bpparam = BiocParallel::SerialParam(),
  chimera.to.target = 5,
  orientation = c("target", "opposite", "positive"),
  verbose = TRUE
)
```

## Arguments

<code>reads</code>	A GAlignments object, or a character vector of the filenames
<code>target</code>	A GRanges object specifying the range to narrow alignments to
<code>...</code>	Extra arguments for initialising CrisprSet
<code>reverse.complement</code>	(Default: TRUE) Should the alignments be oriented to match the strand of the target? If TRUE, targets located strand and targets on the negative strand with respect to the negative strand. If FALSE, the parameter 'orientation' must be set

	to determine the orientation. 'reverse.complement' will be replaced by 'orientation' in a later release.
chimeras	Flag to determine how chimeric reads are treated. One of "ignore", "exclude", and "merge". Default "count", "merge" not implemented yet
collapse.pairs	If reads are paired, should pairs be collapsed? (Default: FALSE) Note: only collapses primary alignments, and assumes that there is only one primary alignment per read.
use.consensus	Take the consensus sequence for non-matching pairs? If FALSE, the sequence of the first read is used. Can be very slow. (Default: FALSE)
store.chimeras	Should chimeric reads be stored? (Default: FALSE)
verbose	Print progress and statistics (Default: TRUE)
name	An experiment name for the reads. (Default: NULL)
minoverlap	Minimum number of bases the aligned read must share with the target site. If not specified, the aligned read must completely span the target region. (Default: NULL)
orientation	One of "target" (reads are displayed on the same strand as the target) "opposite" (reads are displayed on the opposite) strand from the target or "positive" (reads are displayed on the forward strand regardless of the strand of the target) (Default:"target")
reference	The reference sequence
names	Experiment names for each bam file. If not supplied, filenames are used.
target.loc	The zero point for renumbering (Default: 17)
exclude.ranges	Ranges to exclude from consideration, e.g. homologous to a pcr primer.
exclude.names	Alignment names to exclude
targets	A set of targets to narrow reads to
references	A set of reference sequences matching the targets. References for negative strand targets should be on the negative strand.
primer.ranges	A set of GRanges, corresponding to the targets. Read lengths are typically greater than target regions, and it can be that reads span multiple targets. If primer.ranges are available, they can be used to assign such reads to the correct target.
ignore.strand	Should strand be considered when finding overlaps? (See <a href="#">findOverlaps</a> )
bpparam	A BiocParallel parameter for parallelising across reads. Default: no parallelisation. (See <a href="#">bpparam</a> )
chimera.to.target	Number of bases that may separate a chimeric read set from the target.loc for it to be assigned to the target. (Default: 5)

## Value

(signature("GAlignments", "GRanges")) A [CrisprRun](#) object

(signature("character", "GRanges")) A [CrisprSet](#) object

**Author(s)**

Helen Lindsay

**Examples**

```
# Load the metadata table
md_fname <- system.file("extdata", "gol_F1_metadata_small.txt", package = "CrispRVariants")
md <- read.table(md_fname, sep = "\t", stringsAsFactors = FALSE)

# Get bam filenames and their full paths
bam_fnames <- sapply(md$bam.filename, function(fn){
  system.file("extdata", fn, package = "CrispRVariants")})

reference <- Biostrings::DNAString("GGTCTCTCGCAGGATGTTGCTGG")
gd <- GenomicRanges::GRanges("18", IRanges::IRanges(4647377, 4647399),
  strand = "+")

crispr_set <- readsToTarget(bam_fnames, target = gd, reference = reference,
  names = md$experiment.name, target.loc = 17)
```

readTargetBam

*Internal CrispRVariants function for reading and filtering a bam file***Description**

Includes options for excluding reads either by name or range. The latter is useful if chimeras are excluded. Reads are excluded before chimeras are detected, thus a chimeric read consisting of two sections, one of which overlaps an excluded region, will not be considered chimeric. Chimeric reads can be ignored, excluded, which means that all sections of a chimeric read will be removed, or merged, which means that chimeras will be collapsed into a single read where possible. (Not implemented yet) If chimeras = "merge", chimeric reads are merged if all segments

**Usage**

```
readTargetBam(
  file,
  target,
  exclude.ranges = GRanges(),
  exclude.names = NA,
  chimera.to.target = 5,
  chimeras = c("count", "ignore", "exclude", "merge"),
  max.read.overlap = 10,
  max.unmapped = 4,
  by.flag = TRUE,
  verbose = TRUE
)
```

**Arguments**

file	The name of a bam file to read in
target	A GRanges object containing a single target range
exclude.ranges	A GRanges object of regions that should not be counted, e.g. primer or cloning vector sequences that have a match in the genome
exclude.names	A vector of read names to exclude.
chimera.to.target	Maximum distance between endpoints of chimeras and target.loc for assigning chimeras to targets (default: 5)
chimeras	Flag to determine how chimeric reads are treated. One of "ignore", "exclude", "count" and "merge". Default "ignore".
max.read.overlap	Maximum number of bases mapped to two positions for chimeras to be merged (Default: 10)
max.unmapped	Maximum number of bases that are unmapped for chimeras to be merged (Default: 4)
by.flag	Is the supplementary alignment flag set? Used for identifying chimeric alignments, function is much faster if TRUE. Not all aligners set this flag. If FALSE, chimeric alignments are identified using read names (Default: TRUE)
verbose	Print stats about number of alignments read and filtered. (Default: TRUE)

**Value**

A GenomicAlignments::GAlignment obj

---

refFromAlns	<i>refFromAlns</i>
-------------	--------------------

---

**Description**

Reconstruct the reference sequence from alignments reads using the CIGAR

**Usage**

```
refFromAlns(alns, location, ...)

## S4 method for signature 'GAlignments,ANY'
refFromAlns(alns, location, ..., keep.names = FALSE)

## S4 method for signature 'GAlignments,GRanges'
refFromAlns(alns, location, ...)
```

**Arguments**

<code>alns</code>	Alignments to use for inferring the reference sequence
<code>location</code>	The location to infer the reference for.
<code>...</code>	additional arguments
<code>keep.names</code>	Should read names be added to the result if present? (Default: FALSE)

**Value**

The reference sequences corresponding to the provided alignments

A DNASTringSet (signature = c("GAlignments", "ANY"))

A DNASTring (signature = c("GAlignments", "GRanges"))

**Author(s)**

Helen Lindsay

**Examples**

```
bam <- system.file("extdata", "bam/ab1_ptena_wildtype_looking_embryo_1_s.bam",
  package = "CrispRVariants")
alns <- GenomicAlignments::readGAlignments(bam,
  param = Rsamtools::ScanBamParam(tag = "MD", what = "seq"))
# To get the reference sequence from the a given location:
location <- GenomicRanges::GRanges("chr17", IRanges::IRanges(23648420,23648430))
refFromAlns(alns, location = location)
```

---

reverseCigar

*Reverses the order of operations in a cigar string*

---

**Description**

For example, the string "20M5D15M" would become "15M5D20M"

**Usage**

```
reverseCigar(cigars)
```

**Arguments**

<code>cigars</code>	the cigar strings.
---------------------	--------------------

**Value**

The reversed cigar string

---

rmMultiPCRChimera	<i>Remove chimeric reads overlapping multiple primers</i>
-------------------	---

---

**Description**

Finds and removes sets of chimeric read alignments that overlap more than one guide, i.e. that cannot be unambiguously assigned to a single guide.

**Usage**

```
rmMultiPCRChimera(readnames, pcrhits, chimera_idx, ...)  
  
## S4 method for signature 'character,Hits,integer'  
rmMultiPCRChimera(readnames, pcrhits, chimera_idx, ..., verbose = TRUE)
```

**Arguments**

readnames	A set of read names, used for identifying chimeric read sets
pcrhits	A mapping between indices of reads and a set of pcr primers
chimera_idx	location of chimeric reads within the bam
...	Additional arguments
verbose	Display information about the chimeras (Default: TRUE)

**Value**

pcrhits, with chimeric reads mapping to different primers omitted.

**Author(s)**

Helen Lindsay

---

selectOps	<i>selectOps</i>
-----------	------------------

---

**Description**

select CIGAR operations in a region of interest.

**Usage**

```
selectOps(cigar, ...)

## S4 method for signature 'character'
selectOps(
  cigar,
  ...,
  ops = GenomicAlignments::CIGAR_OPS,
  op.regions = NULL,
  pos = 1L
)
```

**Arguments**

cigar	CIGAR strings
...	Extra arguments (Not currently used)
ops	CIGAR operations to consider (Default: all)
op.regions	(GRanges) Return operations only in these regions
pos	An offset for the cigar ranges

**Value**

A GRanges list of operation locations in reference space with a metadata column for the operation width in query space.

**Author(s)**

Helen Lindsay

---

seqsToAln

---

*Creates a text alignment from a set of cigar strings*


---

**Description**

Creates a one-to-one text alignment of a set of cigar strings with respect to the reference sequence by collapsing insertions and introducing gaps across deletions.

When genomic coordinates for the alignment start and the target region are provided, aligned sequences are cropped to the target region

Given a character vector of pairwise alignments and a region to display, trims alignments to the display regions, joined by a separator "join". Alignments should be equal length, e.g. created by seqsToAln



**Usage**

```
seqsToAIn(
  cigar,
  dnaseq,
  target,
  del_char = "-",
  aln_start = NULL,
  reverse_complement = FALSE,
  allow_partial = FALSE
)

selectAInRegions(
  alns,
  reference,
  target,
  keep,
  join = "/ %s /",
  border.gaps = FALSE
)
```

**Arguments**

<code>cigar</code>	A list of cigar strings to align
<code>dnaseq</code>	The set of sequences corresponding to the cigars, as <code>Biostrings::DNAStrings</code>
<code>target</code>	The target region to return, as <code>GRanges</code> . Sequences overlapping the target region are trimmed to exactly match it.
<code>del_char</code>	The character to represent deleted bases. Default "-"
<code>aln_start</code>	Genomic start locations of aligned sequences. Should be used in conjunction with <code>target_start</code> and <code>target_end</code> .
<code>reverse_complement</code>	(Default: FALSE)
<code>allow_partial</code>	Are alignments that do not span the target region allowed? (Default: FALSE)
<code>alns</code>	Character vector of pairwise alignments, with insertions removed
<code>reference</code>	Reference sequence
<code>keep</code>	Region to display, relative to the target region, i.e. not genomic coords ( <code>IRanges</code> or <code>GRanges</code> )
<code>join</code>	<code>character(1)</code> String used for joining alignment segments. Can accept a placeholder to fill in the number of bases deleted with " deleted
<code>border.gaps</code>	( <code>logical(1)</code> ) Should bases deleted from the borders be shown? (Default: FALSE)

**Value**

The sequences with insertions collapsed and deletions padded

A list of the truncated alignments (`alns`) and reference (`ref`)

**Author(s)**

Helen Lindsay

---

setDNATileColours	<i>Sets colours for plotting aligned DNA sequences.</i>
-------------------	---

---

**Description**Sets tile colours for [plotAlignments](#) with a DNA alphabet**Usage**

setDNATileColours(m)

**Arguments**

m	A matrix with a column named "value" of the characters at each tile position.
---	---

**Value**

A matrix with additional columns specifying tile and text colours

**Author(s)**

Helen Lindsay

---

setMismatchTileColours	<i>Sets colours for plotting mismatches in aligned DNA sequences.</i>
------------------------	---

---

**Description**Sets tile colours for [plotAlignments](#) with a DNA alphabet.**Usage**

setMismatchTileColours(m)

**Arguments**

m	A data frame of nucleotides and plotting locations, e.g. created by <a href="#">transformAlnsToLong</a>
---	---

**Value**

A matrix with additional columns specifying tile and text colours

**Author(s)**

Helen Lindsay

---

transformAlnsToLong	<i>Transform data for plotting</i>
---------------------	------------------------------------

---

**Description**

Orders and transforms a reference sequence and a set of aligned sequences into long format, i.e. one observation (tile position) per row. Used internally by [plotAlignments](#).

**Usage**

```
transformAlnsToLong(ref, alns, add.other = FALSE)
```

**Arguments**

ref	The reference sequence
alns	Character vector of aligned sequences
add.other	Add a blank row labelled "Other" (Default: FALSE)

**Value**

A matrix of characters and plotting locations

**Author(s)**

Helen Lindsay

---

variantCounts	<i>Get variant counts</i>
---------------	---------------------------

---

**Description**

Returns a matrix of counts where rows are sequence variants and columns are samples

**Usage**

```
variantCounts(obj, ...)  
  
## S4 method for signature 'CrisprSet'  
variantCounts(  
  obj,  
  ...,  
  top.n = NULL,  
  min.freq = 0,  
  min.count = 1,  
  include.chimeras = TRUE,
```

```
include.nonindel = TRUE,  
result = "counts",  
filter.vars = NULL  
)
```

Arguments

obj	An object containing variant counts
...	Additional arguments
top.n	(Integer n) If specified, return variants ranked at least n according to frequency across all samples (Default: 0, i.e. no cutoff)
min.freq	(Float n least one sample (Default: 0)
min.count	(Integer n) Return variants with count greater than n in at least one sample (Default: 0)
include.chimeras	Should chimeric reads be included in the counts table? (Default: TRUE)
include.nonindel	Should sequences without indels be returned? (Default:TRUE)
result	Return variants as either counts ("counts", default) or proportions ("proportions")
filter.vars	Labels of variants alleles to remove (Default: NULL)

Value

A matrix of counts where rows are variants and columns are samples

Author(s)

Helen Lindsay

Examples

```
data("gol_clutch1")  
  
#Return a matrix of the 5 most frequent variants  
variantCounts(gol, top.n = 5)
```

---

writeFastq	<i>Append a sequence to a fastq file</i>
------------	--

---

Description

Used by abifToFastq to write sanger sequences to fastq format As abifToFastq appends output to files, writeFastq checks that sequence names are unique. This function is faster with checking switched off.

**Usage**

```
writeFastq(outf, vals, allow_spaces = FALSE, check = TRUE)
```

**Arguments**

<code>outf</code>	Name of fastq file to append sequence
<code>vals</code>	A list containing entries named "seq" (sequence) and "quals" (quality scores, in ASCII format)
<code>allow_spaces</code>	Should spaces in the sequence name be substituted with underscores? TRUE or FALSE
<code>check</code>	Check whether reads with the same name already exist in the output fastq. (Default: TRUE)

**Value**

None. The sequences in "vals" are written to outf

**Author(s)**

Helen Lindsay

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