Package 'crisprBase'

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Title Base functions and classes for CRISPR gRNA design

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Imports BiocGenerics, Biostrings, GenomicRanges, IRanges, S4Vectors, stringr

Suggests knitr, rmarkdown, testthat

biocViews CRISPR, FunctionalGenomics

Description Provides S4 classes for general nucleases, CRISPR nucleases and base editors. Several CRISPR-specific genome arithmetic functions are implemented to help extract genomic coordinates of spacer and protospacer sequences. Commonly-used CRISPR nuclease objects are provided that can be readily used in other packages. Both DNA- and RNA-targeting nucleases are supported.

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Encoding UTF-8

RoxygenNote 7.1.2

VignetteBuilder knitr

BugReports https://github.com/Jfortin1/crisprBase/issues

URL https://github.com/Jfortin1/crisprBase

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R topics documented:

`annotateMismatches` *Annotate mismatches between spacer and protospacer sequences*

Description

Annotate mismatches between spacer and protospacer sequences.

Usage

```
annotateMismatches(spacers, protospacers, rnase = FALSE)
```

Arguments

`spacers`	A character vector specifying spacer sequences (gRNA).
`protospacers`	A character vector specifying protospacer sequences (target DNA).
`rnase`	Is it for an RNAse? FALSE by default. If TRUE, spacers and protospacers are expected to be the reverse complement of each other.

Value

A data.frame storing spacer and protospacer columns, as well as number of mismatches, and positions for the different mismatches, if any. Positions are relative to the 5' end of the spacer sequences. For RNAses (e.g. CasRx), this means that a mismatch at position 1 corresponds to the last nucleotide of the protospacer sequence.

Author(s)

Jean-Philippe Fortin

Examples

```
spacers <- c("CCGGAGCGAGTTGCAGTAAGCAG",
    "GCCGGAGCGAGTTGCAGTAAGCA",
    "GGCCGGAGCGAGTTGCAGTAAGC")

protospacers=c("CTGCTTACTGCAACTCGCTCTGG",
    "TGCTTAATGCAACCCGCTCCGGC",
    "GCTTACTGCAACTCGCTCCGGCC")

ann <- annotateMismatches(spacers,
    protospacers,
    rnase=TRUE)
```

AsCas12a *AsCas12a CrisprNuclease object*

Description

CrisprNuclease object for the Wildtype Acidaminococcus Cas12a (AsCas12a) nuclease.

Usage

```
data(AsCas12a, package="crisprBase")
```

Format

CrisprNuclease object.

Details

The AsCas12a nuclease recognizes TTTV PAM sequences. Spacer sequences must be located downstream of PAM sequences.

baseEditorName *An S4 class to represent a base editor*

Description

An S4 class to represent a base editor

Usage

```
baseEditorName(object)

baseEditorName(object) <- value

editingWeights(object, ...)

editingWeights(object) <- value

editingStrand(object)

editingStrand(object) <- value

BaseEditor(
  CrisprNuclease,
  baseEditorName = NA_character_,
  editingStrand = c("original", "opposite"),
  editingWeights = NULL
)

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

## S4 method for signature 'BaseEditor'
baseEditorName(object)

## S4 replacement method for signature 'BaseEditor'
baseEditorName(object) <- value

## S4 method for signature 'BaseEditor'
editingWeights(object, substitutions = NULL)

## S4 replacement method for signature 'BaseEditor'
editingWeights(object) <- value

## S4 method for signature 'BaseEditor'
editingStrand(object)

## S4 replacement method for signature 'BaseEditor'
```

```
editingStrand(object) <- value
```

Arguments

object	BaseEditor object.
value	Value to replaced with.
...	Additional arguments for class-specific methods
CrisprNuclease	A CrisprNuclease object.
baseEditorName	String specifying base editor name.
editingStrand	String indicating which strand with respect to the target protospacer sequence will be edited. Must be either "original" or "opposite". "original" by default.
editingWeights	Numeric matrix of editing weights. Column names must be indicating relative position to the PAM site. Row names must be of the form "X2Y" where "X" represents the origin base, and "Y" represents the subtituted base. For instance, "C2T" indicates the row corresponding to C to T editing.
substitutions	Character vector indicating which substitutions should be returned.

Value

A BaseEditor object

Functions

- BaseEditor: Create a BaseEditor object

Slots

baseEditorName Name of the base editor.

editingWeights Matrix of editing weights.

editingStrand String indicating which strand with respect to the target protospacer sequence will be edited. Must be either "original" or "opposite". "original" by default.

Constructors

Use the constructor link{BaseEditor} to create a BaseEditor object.

Accessors

baseEditorName: To get the name of the base editor.

editingWeights: To return the matrix of editing weights.

editingStrand: To return the editing strand.

Setters

baseEditorName<-: To change the name of the base editor.

editingWeights<-: To change the matrix of editing weights.

editingStrand<-: To change the editing strand.

Examples

```
# Creating an object for BE4max (C to T editor)
# based on experimental weights

ws <- c(0.7, 0.7, 0.8, 1.8, 1, 2, 1.4, 1.2, 2.3, 1.3, 2.4, 2.2, 3.4,
      2.2, 2.1, 3.5, 5.8, 16.2, 31.8, 63.2, 90.3, 100, 87, 62, 31.4,
      16.3, 10, 5.6, 3.3, 1.9, 1.8, 2.4, 1.7, 0.5, 0.2, 0.1)
ws <- matrix(ws, nrow=1, ncol=length(ws))
rownames(ws) <- "C2T"
colnames(ws) <- -36:-1
data(SpCas9, package="crisprBase")
BE4max <- BaseEditor(SpCas9,
                     baseEditorName="BE4max",
                     editingStrand="original",
                     editingWeights=ws)
metadata(BE4max)$description_base_editor <- "BE4max cytosine base editor."
```

BE4max *BE4max BaseEditor object*

Description

BaseEditor for the cytosine base editor CRISPR/Cas9 system BE4max. Editing weights were obtained from https://doi.org/10.1016/j.cell.2020.05.037

Usage

```
data(BE4max, package="crisprBase")
```

Format

BaseEditor object.

Details

BaseEditor for the cytosine base editor CRISPR/Cas9 system BE4max. Editing weights were obtained from https://doi.org/10.1016/j.cell.2020.05.037.

CasRx *CasRx CrisprNuclease object*

Description

CrisprNuclease object for the Cas13d-NLS from Ruminococcus flavefaciens strain XPD3002 nuclease (RNase).

Usage

```
data(CasRx, package="crisprBase")
```

Format

CrisprNuclease object.

Details

The CasRx nuclease was derived from Cas13d Ruminococcus flavefaciens string XPD3002. See 10.1016/j.cell.2018.02.033.

enAsCas12a *enAsCas12a CrisprNuclease object*

Description

CrisprNuclease object for the Enhanced Acidaminococcus Cas12a (AsCas12a) nuclease.

Usage

```
data(enAsCas12a, package="crisprBase")
```

Format

CrisprNuclease object.

Details

The enAsCas12a nuclease recognizes an extended set of PAM sequences beyong the canonical TTTV sequence for AsCas12a. Spacer sequences must be located downstream of PAM sequences.

`extractPamFromTarget` *Extract PAM sequences from target sequences*

Description

Extract PAM sequences from target sequences (protospacer + PAM) using information stored in a CrisprNuclease object.

Usage

```
extractPamFromTarget(targets, object)
```

Arguments

`targets`	Character vector of target sequences.
`object`	`CrisprNuclease` corresponding to the target sequences.

Value

Character vector of PAM sequences of length equal to that of the `targets` character vector.

Author(s)

Jean-Philippe Fortin

Examples

```
data(SpCas9, AsCas12a, package="crisprBase")
# Extracting PAM sequences from Cas9 protospacers:
targets <- c("AGGTGCTGATTGTAGTGCTGCGG",
             "AGGTGCTGATTGTAGTGCTGAGG")
extractPamFromTarget(targets, SpCas9)
# Extracting PAM sequences from Cas12a targets:
targets <- c("TTTAAGGTGCTGATTGTAGTGCTGTGT",
            "TTTCAGGTGCTGATTGTAGTGCTGAAA")
extractPamFromTarget(targets, AsCas12a)
```

`extractProtospacerFromTarget` *Extract protospacer sequences from target sequences*

Description

Extract protospacer sequences from target sequences (protospacer + PAM) using information stored in a CrisprNuclease object.

Usage

```
extractProtospacerFromTarget(targets, object)
```

Arguments

`targets`	Character vector of targets sequences.
`object`	`CrisprNuclease` corresponding to the targets sequences.

Value

Character vector of protospacer sequences of length equal to that of the `targets` character vector.

Author(s)

Jean-Philippe Fortin

Examples

```
data(SpCas9, AsCas12a, package="crisprBase")
# Extracting protospacer sequences from Cas9 targets:
targets <- c("AGGTGCTGATTGTAGTGCTGCGG",
             "AGGTGCTGATTGTAGTGCTGAGG")
extractProtospacerFromTarget(targets, SpCas9)
# Extracting protospacer sequences from Cas12a targets:
targets <- c("TTTAAGGTGCTGATTGTAGTGCTGTGT",
             "TTTCAGGTGCTGATTGTAGTGCTGAAA")
extractProtospacerFromTarget(targets, AsCas12a)
```

getAvailableCrisprNucleases *Return list of available CrisprNuclease objects in crisprBase*

Description

Return list of available CrisprNuclease objects in crisprBase.

Usage

```
getAvailableCrisprNucleases()
```

Value

Character vector of available `CrisprNuclease` objects found in `crisprBase`.

Author(s)

Jean-Philippe Fortin

Examples

```
getAvailableCrisprNucleases()
```

getPamRanges *Construct a PAM GRanges from a list of PAM sites*

Description

Construct a PAM GRanges from a list of PAM sites using information stored in a CrisprNuclease object.

Usage

```
getPamRanges(
  gr = NULL,
  seqnames = NULL,
  pam_site = NULL,
  strand = NULL,
  nuclease = NULL
)
```

Arguments

gr	GRanges object of width 1 specifying the coordinates of the first nucleotide of the PAM sequences.
seqnames	Character vector of genomic sequence names. Ignored if gr is not NULL.
pam_site	Numeric vector specifying the coordinates of the first nucleotide of the PAM sequences corresponding to the PAM sequences. Ignored if gr is not NULL.
strand	Character vector specifying the strand of the PAM. Ignored if gr is not NULL.
nuclease	CrisprNuclease object.

Value

GRanges object representing genomic coordinates of PAM sequences.

Author(s)

Jean-Philippe Fortin

Examples

```
data(SpCas9, AsCas12a, package="crisprBase")
if (require(GenomicRanges)){
gr <- GRanges("chr10",
              IRanges(start=c(100,120), width=1),
              strand=c("+","-"))
getPamRanges(gr, nuclease=SpCas9)
getPamRanges(gr, nuclease=AsCas12a)
}
```

---

getProtospacerRanges *Construct a protospacer GRanges from a list of PAM sites*

Description

Construct a protospacer GRanges from a list of PAM sites using information stored in a CrisprNuclease object.

Usage

```
getProtospacerRanges(
  gr = NULL,
  seqnames = NULL,
  pam_site = NULL,
  strand = NULL,
  nuclease = NULL,
  spacer_len = NULL
)
```

Arguments

gr	GRanges object of width 1 specifying the coordinates of the first nucleotide of the PAM sequences.
seqnames	Character vector of genomic sequence names. Ignored if gr is not NULL.
pam_site	Numeric vector specifying the coordinates of the first nucleotide of the PAM sequences corresponding to the protospacers. Ignored if gr is not NULL.
strand	Character vector specifying the strand of the protospacer. Ignored if gr is not NULL.
nuclease	CrisprNuclease object.
spacer_len	Non-negative integer to overwrite the default spacer length stored in the CrisprNuclease object. s

Value

GRanges object representing genomic coordinates of protospacer sequences.

Author(s)

Jean-Philippe Fortin

Examples

```
data(SpCas9, AsCas12a, package="crisprBase")
if (require(GenomicRanges)){
gr <- GRanges("chr10",
              IRanges(start=c(100,120), width=1),
              strand=c("+","-"))
getProtospacerRanges(gr, nuclease=SpCas9)
getProtospacerRanges(gr, nuclease=AsCas12a)
}
```

---

getTargetRanges *Construct a target GRanges from a list of PAM sites*

Description

Construct a target (protospacer + PAM) GRanges from a list of PAM sites using information stored in a CrisprNuclease object.

Usage

```
getTargetRanges(
  gr = NULL,
  seqnames = NULL,
  pam_site = NULL,
  strand = NULL,
  nuclease = NULL,
  spacer_len = NULL
)
```

Arguments

gr	GRanges object of width 1 specifying the coordinates of the first nucleotide of the PAM sequences.
seqnames	Character vector of genomic sequence names. Ignored if gr is not NULL.
pam_site	Numeric vector specifying the coordinates of the first nucleotide of the PAM sequences corresponding to the targets. Ignored if gr is not NULL.
strand	Character vector specifying the strand of the target. Ignored if gr is not NULL.
nuclease	CrisprNuclease object.
spacer_len	Non-negative integer to overwrite the default spacer length stored in the CrisprNuclease object.

Value

GRanges object representing genomic coordinates of the target sequences.

Author(s)

Jean-Philippe Fortin

Examples

```
data(SpCas9, AsCas12a, package="crisprBase")
library(GenomicRanges)
gr <- GRanges("chr10",
              IRanges(start=c(100,120), width=1),
              strand=c("+","-"))
getTargetRanges(gr, nuclease=SpCas9)
getTargetRanges(gr, nuclease=AsCas12a)
```

motifs *An S4 class to represent a nuclease.*

Description

Return motif string representations of recognition sites.

Return length of the recognition sites sequences.

Usage

```
motifs(object, ...)

motifLength(object, ...)

nucleaseName(object)

targetType(object)

weights(object, ...)

nucleaseName(object) <- value

targetType(object) <- value

weights(object) <- value

cutSites(object, ...)

isCutting(object)

isRnase(object)

isDnase(object)

Nuclease(
  nucleaseName,
  targetType = c("DNA", "RNA"),
  motifs = NULL,
  cutSites = NULL,
  weights = rep(1, length(motifs)),
  metadata = list()
)

## S4 method for signature 'Nuclease'
show(object)
```

```
## S4 method for signature 'Nuclease'
nucleaseName(object)

## S4 replacement method for signature 'Nuclease'
nucleaseName(object) <- value

## S4 method for signature 'Nuclease'
targetType(object)

## S4 replacement method for signature 'Nuclease'
targetType(object) <- value

## S4 method for signature 'Nuclease'
weights(object, expand = FALSE)

## S4 replacement method for signature 'Nuclease'
weights(object) <- value

## S4 method for signature 'Nuclease'
isCutting(object)

## S4 method for signature 'Nuclease'
isRnase(object)

## S4 method for signature 'Nuclease'
isDnase(object)

## S4 method for signature 'Nuclease'
motifs(
  object,
  primary = FALSE,
  strand = c("+", "-"),
  expand = FALSE,
  as.character = FALSE
)

## S4 method for signature 'Nuclease'
motifLength(object)

## S4 method for signature 'Nuclease'
cutSites(object, strand = c("+", "-", "both"), combine = TRUE, middle = FALSE)
```

Arguments

object	Nuclease object.
...	Additional arguments for class-specific methods
value	New value to pass to the setter functions.
nucleaseName	Name of the nuclease.

targetType	String specifying target type ("DNA" or "RNA").
motifs	Character vector of recognition sequence motifs written from 5' to 3' written in Rebase convention. If the point of cleavage has been determined, the precise site is marked with ^. Only letters in the IUPAC code are accepted. For nucleases that cleave away from their recognition sequence, the cleavage sites are indicated in parentheses. See details for more information.
cutSites	Matrix with 2 rows (+ and - strand, respectively) specifying the cleavage coordinates relative to the first nucleotide of the motif sequence. Each column corresponds to a motif specified in the motifs slot.
weights	Optional numeric vector specifying relative weights for the recognition motifs to specify cleavage probabilities.
metadata	Optional list providing global metadata information.
expand	Should sequences be expanded to only contain ATCG nucleotides? FALSE by default.
primary	Should only the motif with the highest weight be returned? FALSE by default. Only relevant if weights are stored in the Nuclease object.
strand	Strand to allow reverse complementation of the motif. "+" by default.
as.character	Should the motif sequences be returned as a character vector? FALSE by default.
combine	Should only unique values be considered? TRUE by default.
middle	For staggered cuts, should the middle point between the cut on the forward strand and the cut on the reverse strand be considered as the cut site? FALSE by default.

Value

A Nuclease object

Functions

- Nuclease: Create a Nuclease object

Slots

nucleaseName Name of the nuclease.

targetType Character string indicationg target type ("DNA" or "RNA").

motifs DNAStringSet of recognition sequence motifs written from 5' to 3'.

cutSites Matrix with 2 rows (+ and - strand, respectively) specifying the cleavage coordinates relative to the first nucleotide of the motif sequence. Each column corresponds to a motif specified in the motifs slot.

weights Optional numeric vector specifying relative weights for the motifs corresponding to cleavage probabilities.

metadata Optional string providing a description of the nuclease.

Constructors

Use the constructor `link{Nuclease}` to create a Nuclease object.

Accessors

`nucleaseName`: To get the name of the nuclease.

`targetType`: To get the target type ("DNA" or "RNA").

`metadata`: To get the metadata list of the nuclease.

`motifs`: To get the recognition mofif nucleotide sequences.

`weights`: To get nuclease weights.

`cutSites`: To get nuclease cut sites.

See Also

See the CrisprNuclease for CRISPR-specific nucleases.

Examples

```
EcoRI <- Nuclease("EcoRI",
                  motifs=c("G^AATTC"),
                  metadata=list(description="EcoRI restriction enzyme"))
```

---

plotEditingWeights *Quick plot to visualize editing weights*

Description

Quick plot to visualize editing weights from a BaseEditor object.

Usage

```
plotEditingWeights(
  baseEditor,
  discardEmptyRows = TRUE,
  substitutions = NULL,
  ...
)
```

Arguments

`baseEditor`	A `BaseEditor` object.
`discardEmptyRows`	Should rows that have all weight equal to 0 be discarded? TRUE by default.
`substitutions`	Character vector specifying substitutions to be plotted. If NULL (default), all substitutions are shown.
`...`	Additional arguments to be passed to `plot`

Value

Nothing. A plot is generated as a side effect.

Examples

```
if (interactive()){
    data(BE4max, package="crisprBase")
    plotEditingWeights(BE4max)
}
```

`restrictionEnzymes` *List of Nuclease objects representing common restriction enzymes*

Description

List of Nuclease objects representing common restriction enzymes from REBASE database.

Usage

```
data(restrictionEnzymes, package="crisprBase")
```

Format

List of Nuclease objects.

Details

List of Nuclease objects representing common restriction enzymes from REBASE database.

`SaCas9` *SaCas9 CrisprNuclease object*

Description

CrisprNuclease object for the wildtype Staphylococcus aureus Cas9 (SaCas9) nuclease.

Usage

```
data(SaCas9, package="crisprBase")
```

Format

CrisprNuclease object.

Details

The AsCas9 nuclease recognizes NNGRRT PAM sequences. Spacer sequences must be located upstream of PAM sequences.

spacerLength *An S4 class to represent a CRISPR nuclease.*

Description

An S4 class to represent a CRISPR nuclease.

Usage

```
spacerLength(object)

targetLength(object)

pamLength(object)

spacerGap(object)

hasSpacerGap(object)

spacerGap(object) <- value

spacerLength(object) <- value

pamSide(object)

pamSide(object) <- value

pams(object, ...)

pamIndices(object, ...)

spacerIndices(object, ...)

prototypeSequence(object, ...)

CrisprNuclease(
  nucleaseName,
  targetType = c("DNA", "RNA"),
  pams = NA_character_,
  weights = rep(1, length(pams)),
  metadata = list(),
  pam_side = NA_character_,
  spacer_gap = 0L,
  spacer_length = NA_integer_
)

## S4 method for signature 'CrisprNuclease'
```

```
show(object)

## S4 method for signature 'CrisprNuclease'
pamLength(object)

## S4 method for signature 'CrisprNuclease'
spacerLength(object)

## S4 replacement method for signature 'CrisprNuclease'
spacerLength(object) <- value

## S4 method for signature 'CrisprNuclease'
pamSide(object)

## S4 replacement method for signature 'CrisprNuclease'
pamSide(object) <- value

## S4 method for signature 'CrisprNuclease'
spacerGap(object)

## S4 replacement method for signature 'CrisprNuclease'
spacerGap(object) <- value

## S4 method for signature 'CrisprNuclease'
hasSpacerGap(object)

## S4 method for signature 'CrisprNuclease'
targetLength(object)

## S4 method for signature 'CrisprNuclease'
pams(object, primary = TRUE, ignore_pam = FALSE, as.character = FALSE)

## S4 method for signature 'CrisprNuclease'
pamIndices(object)

## S4 method for signature 'CrisprNuclease'
spacerIndices(object)

## S4 method for signature 'CrisprNuclease'
prototypeSequence(object, primary = TRUE)
```

Arguments

object	CrisprNuclease object.
value	For spacerLength<- and gapLength<-, must be a non-negative integer. For pamSide, must be either '5prime' or '3prime'.
...	Additional arguments for class-specific methods
nucleaseName	Name of the CRISPR nuclease.

targetType	String specifying target type ("DNA" or "RNA").
pams	Character vector of PAM sequence motifs written from 5' to 3. If the point of cleavage has been determined, the precise site is marked with ^. Only letters in the IUPAC code are accepted. For nucleases that cleave away from their recognition sequence, the cleavage sites are indicated in parentheses. See details for more information.
weights	Optional numeric vector specifying relative weights of the PAM sequences to specify cleavage probabilities.
metadata	Optional list providing global metadata information.
pam_side	String specifying the side of the PAM sequence sequence with respect to the protospacer sequence. Must be either '3prime' (e.g. Cas9) or '5prime' (e.g. Cas12a)
spacer_gap	Integer specifying the length (in nucleotides) between the spacer sequence and the PAM sequence (e.g. 0 for Cas9 and Cas12a).
spacer_length	Integer specifying the length of the spacer sequence
primary	Should only the PAM sequence with the heighest weight be returned? If no cleavage weights are stored in the CrisprNuclease object, all sequences are returned. TRUE by default.
ignore_pam	Should all possible k-mer sequences for a given PAM length be returned, irrespetively of the PAM sequence motifs stored in the CrisprNuclease object? FALSE by default.
as.character	Should the PAM sequences be returned as a character vector? FALSE by default.

Value

A CrisprNuclease object

Functions

- `CrisprNuclease`: Create a CrisprNuclease object

Slots

`pam_side` String specifying the side of the PAM sequence with respect to the protospacer sequence. Must be either '3prime' (e.g. SpCas9) or '5prime' (e.g. AsCas12a)

`spacer_length` Integer specifying the length of the spacer sequence

`spacer_gap` Integer specifying the length (in nucleotides) between the spacer sequence and the PAM sequence (e.g. 0 for SpCas9 and AsCas12a).

Constructors

Use the constructor `link{CrisprNuclease}` to create a CrisprNuclease object.

Accessors

`nucleaseName`: To get the name of the CRISPR nuclease.

`spacerLength`: To return the length of the spacer sequence.

`targetLength`: To return the length of the target sequence (protospacer + pam).

`pamLength`: To return the length of the PAM sequence.

`pamSide`: To return the side of the PAM sequence with respect to the spacer sequence.

`spacerGap`: To return the length of the gap between the PAM and spacer sequences.

`pams`: To return the list of PAM sequences.

Setters

`spacerGap<-`: To change the length of the gap between the PAM and spacer sequences.

`pamSide<-`: To change the side of the PAM sequence with respect to the protospacer sequence.

`spacerLength<-`: To change the length of the spacer sequence.

Utility functions for genomic arithmetics

`pamIndices`: To return the relative coordinates of the PAM sequence within the target sequence.

`spacerIndices`: To return the relatiive coordinates of the spacer sequence within the target sequence.

Examples

```
SpCas9 <- CrisprNuclease("SpCas9",
                         pams=c("(3/3)NGG", "(3/3)NAG", "(3/3)NGA"),
                         weights=c(1, 0.2593, 0.0694),
                         metadata=list(description="Wildtype Streptococcus
                                       pyogenes Cas9 (SpCas9) nuclease"),
                         pam_side="3prime",
                         spacer_length=20)
```

SpCas9 *SpCas9 CrisprNuclease object*

Description

CrisprNuclease object for the wildtype Streptococcus pyogenes Cas9 (SpCas9) nuclease.

Usage

```
data(SpCas9, package="crisprBase")
```

Format

CrisprNuclease object.

Details

The SpCas9 nuclease recognizes NGG PAM sequences. Spacer sequences must be located upstream of PAM sequences.

`SpGCas9` *SpGCas9 CrisprNuclease object*

Description

CrisprNuclease object for the engineered Streptococcus pyogenes Cas9 SpG nuclease.

Usage

```
data(SpGCas9, package="crisprBase")
```

Format

CrisprNuclease object.

Details

The SpGCas9 nuclease recognizes NGN PAM sequences. Spacer sequences must be located upstream of PAM sequences.

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