Package ‘Pbase’

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Type Package

Title Manipulating and exploring protein and proteomics data

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Suggests testthat (>= 0.8), ggplot2, BSgenome.Hsapiens.NCBI.GRCh38, TxDb.Hsapiens.UCSC.hg38.knownGene, AnnotationHub, knitr, rmarkdown, BiocStyle

Description A set of classes and functions to investigate and understand protein sequence data in the context of a proteomics experiment.

License GPL-3

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VignetteBuilder knitr

URL https://github.com/ComputationalProteomicsUnit/Pbase

BugReports https://github.com/ComputationalProteomicsUnit/Pbase/issues

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NeedsCompilation no

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calculateHeavyLabels

Description

A function to calculate heavy labeled peptides for proteins stored in a Proteins object.

Usage

`calculateHeavyLabels(proteins, peptides, maxN = 20L, nN = 4L, nC = 3L, endsWith = c("K", "R", "G"), ...)`

Arguments

- `proteins`: A Proteins object.
- `peptides`: A named character vector containing the peptides of interest. The names must match the UniProt accession numbers of the proteins in object.
- `maxN`: An integer, maximal length of the heavy labeled peptide.
- `nN`: An integer, minimal number of amino acids at the N terminus.
- `nC`: An integer, minimal number of amino acids at the C terminus.
- `endsWith`: A character vector containing the allowed amino acids at the end of the resulting sequence (every peptide that doesn’t end with one of these amino acids has to be one amino acid shorter as `maxN`).
- `...`: Additional parameters passed to .addOverhangs.

Details

The digestion efficiency with enzymes like trypsin is below 100%. That’s why spiked-in peptides for labeled quantitation have to follow the same digestion rules as the peptides of interest. Therefore it is necessary to extend the peptides of interest by a few amino acids on the N- and C-terminus. These extensions should not be a cleavage point of the used enzyme. This method provides an easy interface to find the sequences for heavy labeled peptides that could be used as spike-ins for the peptides of interest. Please see the references for a more detailed discussion.

TODO: There should be a function to find the best labels for a given protein automatically.

Value

A data.frame with 6 columns:

- `Protein`: The Protein accession number.
- `Peptide`: The peptide of interest.
- `N_overhang`: The added sequence of the N-terminus.
- `C_overhang`: The added sequence of the C-terminus.
- `spikeTideResult`: A short description of the used creation rule.
- `spikeTide`: The heavy labeled peptide that represents the peptide of interest best.
etrid2grl

Author(s)
Sebastian Gibb <mail@sebastiangibb.de> and Pavel Shliaha

References
The complete description of the issue: https://github.com/sgibb/cleaver/issues/5

Examples
## example protein database
data(p, package = "Pbase")

## digest proteins into peptides
cleavedProteins <- cleave(p)

## find spike-ins for the peptides of interest
calculateHeavyLabels(cleavedProteins,
  peptides = c(A4UGR9 = "MEGFHIK",
               A4UGR9 = "QGNMYTLSK",
               A6H8Y1 = "GASTNPQR"))

etrid2grl  From a transcript identifier to GRanges object

Description
This function takes on or more Ensembl transcript identifiers, queries Biomart and constructs a GRangesList object as would Gviz::BiomartGeneRegionTrack for a genomic region (in fact, currently most of the code has been taken from Gviz:::fetchBMData and Gviz:::chrName is used to validate chromosome names).

Usage
etrid2grl(etrid, ens, use.names = FALSE)

Arguments

etrid  A vector of Ensembl transcript identifiers.

ens  A instance of class Mart from biomaRt. If missing, useMart("ensembl", "hsapiens_gene_ensembl") is used.

use.names  If set to TRUE and etrid has names, then the latter are used to name the output.

Value
A GRangesList object of length length(etrid).

Author(s)
Laurent Gatto
Examples

id <- c("ENST00000612959", "ENST00000317091")
gr1 <- etrid2grl(id[1])
gr1
gr <- etrid2grl(id)
stopifnot(all.equal(id, names(grl)))

isReverse

Are all the ranges on the same strand

Description

Are all the ranges on the same strand

Usage

isReverse(gr)

isForward(gr)

Arguments

gr

A GRanges object.

Value

A logical if all the ranges in the gr object are on the "-" (or "+") strand.

Author(s)

Laurent Gatto

mapToGenome-methods

Map range coordinates between proteins and genome space

Description

Map range coordinates between peptide features along proteins and genome (reference) space.

Usage

## S4 method for signature 'Proteins,GRangesList'
mapToGenome(x, genome, drop.empty.ranges = TRUE, ...)
## S4 method for signature 'Proteins,GRangesList'
pmapToGenome(x, genome, drop.empty.ranges = TRUE, ...)
Arguments

x  
Proteins object containing peptides pranges to be mapped.

genome  
A GRangesList object used to map between x and the result. The ranges are typically created by the etrid2grl function.

drop.empty.ranges  
TRUE (default) or FALSE. Should empty ranges be dropped?

...  
Additional parameters passed to inner functions. Currently ignored.

Details

• mapToGenome maps the pranges(x) to the ranges of genome. Unless x and genome are of length 1, both must be named and items of x are matched to items of genome using their respective names. Names that do not co-occur in x and genome are ignored. If we have seqnames(x): "A", "B" and "C" and names(genome): "C", "A", "a", "z", "A" and "A".

the names of the output will be "A", "A", "A" and "C".

The output is ordered by (1) seqnames(x) and (2) the order of the elements in genome. In case less than length(x) are mapped, as for p["B"] above, a message informs the user.

• pmapToGenome is the element-wise (aka 'parallel') version of mapToGenome. The i-th pranges(x) is mapped to the i-th range in genome. x and genome must have the same length and do not need to be named (names are ignored).

Value

A named GRangesList object, with names matching names(genome). For pmapToGenome, the return value will have the same length as the inputs.

Author(s)

Laurent Gatto

See Also

• See ?mapToAlignments in the GenomicAlignments package for mapping coordinates between reads (local) and genome (reference) space using a CIGAR alignment.

• See ?mapToTranscripts in the GenomicRanges package for mapping coordinates between features in the transcriptome and genome space.

• The proteinCoding function to remove non-protein coding ranges before mapping peptides to their genomic coordinates.

• The mapping vignette for examples and visualisations.

See plotAsAnnotationTrack and plotAsAnnotationTrack for more details about the two plotting functions.
Examples

data(p)
grl <- etrid2grl(acols(p)$ENST)
pgcr1 <- proteinCoding(grl)
plotAsGeneRegionTrack(grl[[1]],
                      pgcr1[[1]])

mp <- mapToGenome(p[4], pgcr[4])
plotAsAnnotationTrack(mp[[1]], pgcr[4])

Data accompanying the Pbase package

Description

A small example Proteins test instance. This object is likely to change on a regular basis. It will be described more thoroughly when it becomes stable. The MSMS spectra that were searched against the database are available in the pms MSnExp object.

Usage

data(p)
data(pms)

See Also

The Pbase-data vignette.

Examples

data(p)

p
data(pms)
pms
**plotAsAnnotationTrack**  
*Plot gene region and annotation tracks*

**Description**

These functions convert ranges of peptides or exons to AnnotationTrack or GeneRegionTrack objects from the Gviz package and produces the corresponding plot. The genome argument controls whether additional ideogram and axis tracks are to be plotted. plotAsAnnotationTrack plots peptides that span multiple exons in red and connects them with a grey line. See pmapToGenome for example code.

**Usage**

```r
plotAsAnnotationTrack(x, ..., genome = "hg38", plot = TRUE)
plotAsGeneRegionTrack(..., genome = "hg38", plot = TRUE)
```

**Arguments**

- `x`  
  A Granges object containing peptides genomics coordinates, typically generated by pmapToGenome. These ranges are converted to an AnnotationTrack.

- `...`  
  One or more GRanges instances, typically resulting from calling etrid2grl, or, a single GRangesList. These ranges are converted to GeneRegionTrack instances.

- `genome`  
  A character of length 1, giving the name of the genome. Default is "hg38". If NULL, no chromosome and axis tracks are displayed.

- `plot`  
  A logical defining if the figure should be plotted. Default is TRUE.

**Value**

Used for its plotting side effects. Invisible returns a list of tracks.

**Author(s)**

Laurent Gatto

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**Pparams-class**  
*Class "Pparams"*

**Description**

Pbase parametrisation infrastructure.

**Objects from the Class**

New Pbase parameters can be generated with the Pparams() constructor. Pparams instances control various aspects of Pbase functions, as described in the Slots section below. If no parameters are passed to the respective functions, default values from Pparams() are used.
Slots

DbFormat: The format of the protein sequence fasta database used to generate the Proteins object. Currently only "UniProt" is supported. "RefSeq" will be added as well as a mechanism to support arbitrary and custom fasta header.

IdFormat: The format of the identification data files used to add pfeatures to Protein instances. Currently, "UniProt" is supported. "RefSeq" will be added as well as a mechanism to support arbitrary and custom fasta header.

IdReader: Package to be used to load the identification data. Currently one of mzR (via the openIDfile and psms functions) or mzID (via the mzID and flatten functions). Differences between these two architectures include the metadata available in the Proteins’ pfeatures, speed and stability (mzR is much faster but less mature and currently susceptible to crashes).

verbose: A logical defining if the various functions display messages (default) or remain silent.

Methods

show signature(object = "Pparams"):

Author(s)

Laurent Gatto <lg390@cam.ac.uk>

Examples

Pparams()
Pparams(IdReader = "mzID")

try(Pparams(IdReader = "mzid"))

proteinCoding-methods Only keep protein coding ranges

Description

Removed all the ranges that are not protein coding. Typically used on the output of etrid2grl before mapToGenome.

Methods

signature(object="GRanges", mcol="character", coding="character") Removes all the ranges that are not annotated as protein coding ranges, i.e. ranges whose mcols()$mcol is different from coding. The default values are mcols()$feature and "protein_coding". The method return the GRanges trimmed from all non-matching ranges.

signature(object="GRangesList", mcol="character", coding="character") As above but for GRanges in a GrangesList.
**Proteins-class**

---

**The Proteins Class for Proteomics Data And Meta-Data**

---

### Description

The Proteins class encapsulates data and meta-data for proteomics experiments. The class stores the protein sequences as well as specific subsets of interest, typically peptides, as ranges. The Proteins instances, the sequence and peptide slots are described by their respective metadata attributes.

### Objects from the Class

Objects can be created using its constructor Proteins. The constructor either takes a fasta file name as first argument or, alternatively a named uniprotIds argument with valid UniProt accession numbers (not yet implemented).

### Details

An instance of class Proteins is characterised by one or multiple protein sequences that can be accessed as AAStringSet with the aa accessor. Sequence-specific annotation, such as accession numbers, protein and gene names, ... is available with the ametadata or acols methods. General metadata such as the data of creation of the instance are stored as a list returned by the metadata accessor, which would typically contain a created character that documents when the object was created, a reference genome descriptor, a UniProtRelease with the release data of the UniProt database and possibly others.

Each sequence of a Proteins instance can also be characterised by a set of specific ranges describing peptides of interest. These peptide features can be extracted as an AAStringSetList, where each protein sequence contains 0 or more peptide features. These peptides features are encode as ranges along the original proteins sequences (a list of IRanges) that can be extracted with the pranges function. These peptide features have their own metadata describing for example peptide identification scores, number of missed cleavages, ... available with the pmetadata or pcols methods.

See also the Pbase-data vignette.

### Development notes

Since version 0.2.0, addIdentificationData supports multiple identification file names to be added to a Proteins instance (argument renamed filenames) using either mzID or mzR. Added new Pparams parametrisation infrastructure.

See news(package = "Pbase") for a description of all changes.

Other possible metadata fields: Uniprot.sw, biomaRt instances.

### Slots

- **metadata**: Object of class "list" containing global metadata, accessed with metadata.
- **aa**: Object of class "AAStringSet" storing the protein sequences, accessed with aa.
- **pranges**: Object of class "CompressedIRangesList" containing protein feature ranges such as theoretical (obtain by in silico cleavage) or observed peptides. Accessed as an IRangesList with pranges or and AAStringSetList with pfeatures.
- **__classVersion__**: Object of class "Versions" documenting the class versions. Intended for developer use and debugging.
Extends

Class "Versioned", directly.

Methods

**aa** signature(x = "Proteins"): Returns an AStringSet instance representing the sequences of the proteins.

**pfeatures** signature(x = "Proteins"): ...

**pranges** signature(x = "Proteins"): ...

**metadata** signature(x = "Proteins"): Returns a list of global metadata of the instance x, including data of instance creation or, if created from a set of UniProt identifiers (see constructors above), the UniProt version and UniProt.WS version number.

**ametadata** signature(x = "Proteins"): Returns a DataFrame of protein metadata.

**acols** signature(x = "Proteins"): See ametadata.

**pmetadata** signature(x = "Proteins"): Returns a list of feature metadata.

**pcols** signature(x = "Proteins"): See pmetadata.

**avarLabels** signature(x = "Proteins"): Returns the names of the sequences metadata.

**pvarLabels** signature(x = "Proteins"): Returns the names of the peptide feature metadata.

**seqnames** signature(x = "Proteins"): Returns the protein sequence names defined as UniProt accession numbers.

**length** signature(x = "Proteins"): Returns the number of proteins.

**[]** signature(x = "Proteins", i = "ANY", j = "missing"): Creates a subset of the Proteins instance.

**[[** signature(x = "Proteins", i = "ANY", j = "missing"): Returns an AString instance representing the sequence of the selected protein.

**pfilter** signature(x = "Proteins", mass = "numeric", len = "numeric", ...): ... Cleaves all proteins using the enzym rule while allowing missedCleavages missing cleavages. Please see cleave for details.

**addIdentificationData** signature(object = "Proteins", id = "character", rmEmptyRanges = "logical", par = "Pparams"): Adds identification data from an IdemMzMI file (id) to the Proteins object. If rmEmptyRanges is TRUE proteins without any identification data are removed. See Pparams for further settings.

**addPeptideFragments** signature(object = "Proteins", filenames = "character", rmEmptyRanges = "logical", par = "Pparams"): Adds identification data from a fasta file (filenames) to the Proteins object. Please note that both fasta files (the origin of the Proteins object and the ones given in filenames) must share the same Uniprot accession numbers. If rmEmptyRanges is TRUE proteins without any identification data are removed. See Pparams for further settings.

**plot** signature(x = "Proteins", y = "missing"): Plots all proteins and associated peptides using the Gviz/Pviz infrastructure.

**show** signature(object = "Proteins"): Displays object summary as text.
Proteins-class

Functions

- **rmEmptyRanges** signature(x = "Proteins"): removes proteins with empty peptide ranges.
- **proteotypic** signature(x = "Proteins"): returns a modified Proteins object. pcols(x) gains a "Proteotypic" logical column, indicating of the peptide is proteotypic or not.
- **proteinCoverage** signature(pattern = "Proteins"): calculates the coverage of proteins. pcols(x) gains a "Coverage" numeric column.
- **isCleaver** signature(x = "Proteins", missedCleavages = "numeric"): Tests whether a Protein object was cleaved already.

Author(s)

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References

Definition of the UniProt fasta comment format: [http://www.uniprot.org/help/fasta-headers](http://www.uniprot.org/help/fasta-headers)

See Also

- calculateHeavyLabels

Examples

```r
## Create a Protein object reading all proteins from a fasta file.
fastaFiles <- list.files(system.file("extdata", package = "Pbase"), pattern = "fasta", full.names = TRUE)
p <- Proteins(fastaFiles)
p
metadata(p)

## Adding custom metadata
metadata(p, "Comment") <- "I love R"
metadata(p)

## Plotting
plot(p[1:5], from = 1, to = 30)

## Cleaving
pp <- cleave(p[1:100])
pp <- proteotypic(pp)
pp
pcols(pp[1:2])

plot(pp[1:2], from = 20, to = 30)

## Protein coverage
pp <- proteinCoverage(pp)
avarLabels(pp)
acols(pp)$Coverage
pp

## Add identification data
idfile <- system.file("extdata/Thermo_Hela_PRTC_selected.mzid",
```
Proteins-class

package = "Pbase")
p <- addIdentificationData(p, idfile)
pranges(p)
pfeatures(p)

plot(p[1])
plot(p[1], # the first protein has 36 peptides
     fill = c(rep("orange", 13), rep("steelblue", 13)))
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