Package ‘PhIPData’

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Description PhIPData defines an S4 class for phage-immunoprecipitation sequencing (PhIP-seq) experiments. Building upon the RangedSummarizedExperiment class, PhIPData enables users to coordinate metadata with experimental data in analyses. Additionally, PhIPData provides specialized methods to subset and identify beads-only samples, subset objects using virus aliases, and use existing peptide libraries to populate object parameters.
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## aliases

### Using aliases to subset virus data

#### Description

Rather than typing out full viruses names or repeating regexpressions, users can use aliases as a convenient tool to subset PhIPData objects by viral species.

#### Usage

- `getAlias(virus)`
- `setAlias(virus, pattern)`
- `deleteAlias(virus)`

#### Arguments

- `virus`: character vector of the alias
- `pattern`: character vector of regexpressions corresponding to the alias

#### Details

Aliases are cached to an rda file containing only a `data.frame` with two columns: `alias` and `pattern`. The `alias` column contains the alias while the `pattern` column contains the corresponding regexpression of interest.

Once an alias is added to the database, it can always be accessed once the package is loaded. It is recommended to use the functions `setAlias` and `deleteAlias`. If an alias already exists in the database, `setAlias` replaces the matched pattern. If an alias does not exist in the database, `getAlias` returns `NA_character_`. 
Value

getAlias() returns a vector of regexpressions corresponding to queried inputs. The returned vector is the same length as the input vector. Queries that do not exist in the database return NA_character_.

Functions

- getAlias: return a regexpression corresponding to the alias.
- setAlias: define/modify the regexpression for an alias.
- deleteAlias: remove an alias from the database.

Examples

```r
## Edit and modify aliases in the database
setAlias("test_virus", "test_pattern")
getAlias("test_virus")
setAlias("test_virus", "test_pattern2")
getAlias("test_virus")
deleteAlias("test_virus")

## Edit and modify multiple aliases at once.
setAlias(c("virus_1", "virus_2"), c("pattern_1", "pattern_2"))
getAlias(c("virus_1", "virus_2"))
deleteAlias(c("virus_1", "virus_2"))

## Example of how to subset HIV using `getAlias`
## Often, it is useful to set the `ignore.case` of `grep`/`grepl` to TRUE.
counts_dat <- matrix(1:10, nrow = 5)
peptide_meta <- data.frame(species = c(
  rep("Epstein-Barr virus", 2),
  rep("human immunodeficiency virus", 3)
))
phip_obj <- PhIPData(counts = counts_dat, peptideInfo = peptide_meta)
subset(phip_obj, grepl(getAlias("HIV"), species, ignore.case = TRUE))
```

Description

getBeadsName and setBeadsName are two function to get and set the string that encodes which samples are beads-only samples. Information about beads-only samples are stored in the groups column of sampleInfo.

Usage

getBeadsName()

setBeadsName(name)
librarySize

Arguments

name a string indicating how beads-only samples are encoded.

Details

If name is of length greater than one, only the first element of the vector is used. Non-character values of name are first coerced into strings.

Value

a string indicating how beads-only samples are encoded.

Functions

• getBeadsName: function that returns a string corresponding to how beads-only samples are encoded.
• setBeadsName: function to set the string that indicates which samples are beads-only samples in the groups column of sampleInfo.

Examples

## Returns the default string, "beads"
getBeadsName()

## Not run since it changes defaults/user settings
## Not run:
setBeadsName("beads-only")

## End(Not run)

librarySize Calculate total read counts for each sample.

Description

This function is a wrapper function for colSums on the counts assay.

Usage

librarySize(object, ..., withDimnames = TRUE)

Arguments

object PhIPData object
... arguments passed to colSums
withDimnames logical; if true, the vector names are the sample names; otherwise the vector is unnamed.
peptideLibraries

Value

a (named) numeric vector. The length of the vector is equal to the number of samples.

Examples

```r
example("PhIPData")
librarySize(phip_obj)
```

## Return an unnamed vector
librarySize(phip_obj, withDimnames = FALSE)

peptideLibraries     Peptide libraries

Description

PhIP-Seq experiments often use identical peptide libraries different cohorts. These functions enable the user to conveniently reuse tidied libraries.

Usage

- `getLibrary(name)`
- `makeLibrary(library, name)`
- `removeLibrary(name)`
- `listLibrary()`

Arguments

- `name` name of the library
- `library` a matrix, data.frame, or DataFrame with the peptide information for the specified library.

Details

Each library is stored as a DataFrame in .rds file. New libraries can be stored for future use with the makeLibrary function.

Value

`getLibrary` returns a DataFrame corresponding to the peptide information for the specified library.
Functions

- `getLibrary`: return a DataFrame with the peptide information corresponding to the library.
- `makeLibrary`: create and store a DataFrame with the specified peptide information.
- `removeLibrary`: delete stored libraries
- `listLibrary`: list all available libraries

Examples

```r
## Create a new library
pep_meta <- data.frame(species = c(
  rep("human immunodeficiency virus", 3),
  rep("Epstein-Barr virus", 2)
))
makeLibrary(pep_meta, "new_library")

## Use new library
counts_dat <- matrix(1:10, nrow = 5)
phip_obj <- PhIPData(
  counts = counts_dat,
  peptideInfo = getLibrary("new_library")
)

## List libraries
listLibrary()

## Delete created library
removeLibrary("new_library")
```

---

**PhIPData-class**  
*The PhIPData class*

**Description**

The PhIPData class is a matrix-like container designed to organize results from phage-immunoprecipitation (PhIP-Seq) experiments. Rows in PhIPData objects represent peptides and columns represent samples. Each object contains at least three assays:

- `counts`: a matrix of raw read counts,
- `logfc`: a matrix of log2 estimated fold-change in comparison to beads-only samples,
- `prob`: a matrix of probabilities associated with whether a sample has an enriched antibody response for a peptide.

The PhIPData class extends the RangedSummarizedExperiment class, so methods documented in RangedSummarizedExperiment and SummarizedExperiment also work on PhIPData objects.
PhIPData-class

Usage

PhIPData(
  counts = matrix(nrow = 0, ncol = 0),
  logfc = matrix(nrow = 0, ncol = 0),
  prob = matrix(nrow = 0, ncol = 0),
  peptideInfo = S4Vectors::DataFrame(),
  sampleInfo = S4Vectors::DataFrame(),
  metadata = list(),
  .defaultNames = "info"
)

Arguments

counts a matrix, data.frame, or DataFrame of integer read counts.

logfc a matrix, data.frame, or DataFrame of log2 estimated fold changes.

prob a matrix, data.frame, or DataFrame of probability values (p-values or posterior probabilities) for enrichment estimates.

peptideInfo a data.frame or DataFrame of peptide information.

sampleInfo a data.frame or DataFrame of additional sample information.

metadata a list object containing experiment-specific metadata.

.defaultNames vector of names to use when sample and peptide identifiers disagree across the metadata and the counts, logfc, and prob matrices. If .defaultNames is of length 1, the same source is used for both peptide and sample identifiers. If .defaultNames is longer than 2, the first and second elements correspond to the names for peptides and samples, respectively.

Valid options are:

  • "info": names should be taken from the SampleInfo or peptideInfo objects.
  • "counts": names should be taken from the row/column names of the counts object.
  • "logfc": names should be taken from the row/column names of the logfc object.
  • "prob": names should be taken from the row/column names of the prob object.

Details

Rows of PhIPData objects correspond to peptides of interest and are organized in GRanges or GRangesList objects. Though originally designed for genomic ranges, the sequence name and genomic range information in GRanges objects can be replaced with peptide names and amino acid positions, respectively. If no peptide names are given, peptides are given the names of pep_rownum. Peptide positions are specified by columns pos_start and pos_end in the peptideInfo argument of the constructor. Missing position information is set to 0. Additional peptide annotation can also be stored in GRanges objects and can be used to subset PhIPData objects as shown below.
Columns of PhIPData objects represent samples. Sample metadata are stored in a DataFrame and can be accessed as shown below. If no sample names are specified, samples are given default names of sample_colnum.

Unlike RangedSummarizedExperiment/SummarizedExperiment objects, PhIPData objects must contain counts, logfc, prob. If any of the three assays are missing when the constructor is called, an empty matrix of the same names and dimensions is initialized for that assay. Sample and peptide names are harmonized across assays and annotation during construction and replacement.

Though 'counts' typically contain integer values for the number of reads aligned to each peptide, 'PhIPData' only requires that stored values are non-negative numeric values. Pseudocounts or non-integer count values can also be stored in the 'counts' assay.

**Value**

A PhIPData object.

**Constructor**

PhIPData objects are constructed using the homonymous function and arguments as described above. Any PhIPData object can be created so long as peptide and sample identifiers (or lack thereof) are specified via any of the parameters.

**See Also**

PhIPData-methods for accessors and modifiers for PhIPData components. SummarizedExperiment

**Examples**

```r
## Construct a new PhIPData object
counts_dat <- matrix(sample(1:1e6, 25, replace = TRUE), nrow = 5)
logfc_dat <- matrix(rnorm(25, 0, 10), nrow = 5)
prob_dat <- matrix(rbeta(25, 1, 1), nrow = 5)

peptide_meta <- data.frame(
  pos_start = 1:5,
  pos_end = 6:10,
  species = c(rep("HIV", 3), rep("EBV", 2))
)

sample_meta <- data.frame(
  gender = sample(c("M", "F"), 5, TRUE),
  group = sample(c("ctrl", "trt", "beads"), 5, TRUE)
)

exp_meta <- list(
  date_run = as.Date("2021/01/20"),
  reads_per_sample = colSums(counts_dat)
)

rownames(counts_dat) <- rownames(logfc_dat) <-
                    rownames(prob_dat) <- rownames(peptide_meta) <-
                    paste0("pep_", 1:5)

colnames(counts_dat) <- colnames(logfc_dat) <-
```

PhIPData-methods

```r
colnames(prob_dat) <- rownames(sample_meta) <- paste0("sample_", 1:5)

phip_obj <- PhIPData(
  counts_dat, logfc_dat, prob_dat,
  peptide_meta, sample_meta, exp_meta
)
phip_obj
```

---

**PhIPData-methods**

**Accessing and Modifying Information in PhIPData objects**

**Description**

Methods to extract and modify assay(s) (including convenient functions for `counts`, `logfc`, and `prob`), `sampleInfo`, `peptideInfo`, and `metadata`.

**Usage**

```r
## S4 method for signature 'PhIPData'
counts(object, ...)

logfc(object, ...)

## S4 method for signature 'PhIPData'
logfc(object, ...)

prob(object, ...)

## S4 method for signature 'PhIPData'
prob(object, ...)

peptideInfo(object, ...)

## S4 method for signature 'PhIPData'
peptideInfo(object, ...)

sampleInfo(object, ...)

## S4 method for signature 'PhIPData'
sampleInfo(object, ...)

## S4 replacement method for signature 'PhIPData,list'
assays(x, withDimnames = TRUE, ...) <- value

## S4 replacement method for signature 'PhIPData,SimpleList'
assays(x, withDimnames = TRUE, ...) <- value
```
## S4 replacement method for signature 'PhIPData,missing'
assay(x, i, withDimnames = TRUE, ...) <- value

## S4 replacement method for signature 'PhIPData,numeric'
assay(x, i, withDimnames = TRUE, ...) <- value

## S4 replacement method for signature 'PhIPData,character'
assay(x, i, withDimnames = TRUE, ...) <- value

## S4 replacement method for signature 'PhIPData'
counts(object, ...) <- value
logfc(object, ...) <- value

## S4 replacement method for signature 'PhIPData'
logfc(object, ...) <- value
prob(object, ...) <- value

## S4 replacement method for signature 'PhIPData'
prob(object, ...) <- value
peptideInfo(object) <- value

## S4 replacement method for signature 'PhIPData'
peptideInfo(object) <- value
sampleInfo(object, ...) <- value

## S4 replacement method for signature 'PhIPData'
sampleInfo(object) <- value

### Arguments

- **object**: A PhIPData object
- **...**: parameters for assays, which are typically not needed.
- **x**: A PhIPData object
- **withDimnames**: Parameter for RangedSummarizedExperiment class functions. Overrided since row/column names are automatically synced within each object.
- **value**: A matrix, data.frame, or DataFrame of the same dimensions (not necessarily the same names)
- **i**: A numeric, character

### Details

In addition to the functions detailed in RangedSummarizedExperiment, the PhIPData class includes conveniently named functions to quickly access and modify frequently used components of PhIP-Data objects.
Replacement functions ensure that names of the replacement object are matched with the names of the PhIPData object.

Since packages for identifying differential expression in RNA-seq experiments are frequently used for estimating fold-changes for peptide enrichments, the class also includes coercion methods to and from DGELists.

**Value**

Accessors: a DataFrame object

Setters: a PhIPData object

**Available methods**

In the following code snippets, x is a PhIPData object, value is a matrix-like object with the same dimensions as x, and ... are further arguments passed to assay (for the getter) or assay<- (for the setter).

`counts(x, ...)`, `counts(x, ...) <- value`: Get or set a matrix of raw read counts

`logfc(x, ...), logfc(x, ...) <- value`: Get or set a matrix of log2 estimated fold changes (in comparison to beads-only samples)

`prob(x, ...), prob(x, ...) <- value`: Get or set a matrix of probabilities associated with whether a sample has an enriched antibody response for a peptide.

**See Also**

`assays` for SummarizedExperiment operations.

**Examples**

```r
example("PhIPData")

replacement_dat <- matrix(1L, nrow = 5, ncol = 5)

## SummarizedExperiment Accessors and Setters
assays(phip_obj)
assays(phip_obj)$counts <- replacement_dat
assay(phip_obj, "logfc")
assay(phip_obj, "logfc") <- replacement_dat

## counts
counts(phip_obj)
counts(phip_obj) <- counts_dat

## logfc
logfc(phip_obj)
logfc(phip_obj) <- logfc_dat

## prob
prob(phip_obj)
prob(phip_obj) <- replacement_dat
```
propReads  

**Proportion of sample reads**

### Description
This function calculates the proportion of total sample reads pulled by each peptide.

### Usage

```r
propReads(object, withDimnames = TRUE)
```

### Arguments

- `object`: PhIPData object
- `withDimnames`: logical; if true return a matrix with the same dimension names as the original object.

### Value
A (named) numeric matrix with the same dimensions as the function input. Matrix values are between 0 and 1.

### Examples

```r
example("PhIPData")
propReads(phip_obj)
```

## Return an unnamed matrix

```r
propReads(phip_obj, withDimnames = FALSE)
```

---

subsetBeads  

**Subset beads-only samples**

### Description
Function to subset PhIP-seq data for beads-only samples.

### Usage

```r
subsetBeads(object)
```
subsetBeads

Arguments

  object          PhIPData object

Value

  a PhIPData object.

Examples

  example("PhIPData")
  subsetBeads(phip_obj)
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