Package ‘recountmethylation’

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Version  1.12.0
Title  Access and analyze public DNA methylation array data compilations
Description  Resources for cross-study analyses of public DNAm array data from NCBI GEO repo, produced using Illumina's Infinium HumanMethylation450K (HM450K) and MethylationEPIC (EPIC) platforms.
Provided functions enable download, summary, and filtering of large compilation files. Vignettes detail background about file formats, example analyses, and more. Note the disclaimer on package load and consult the main manuscripts for further info.
License  Artistic-2.0
Encoding  UTF-8
URL  https://github.com/metamaden/recountmethylation
BugReports  https://github.com/metamaden/recountmethylation/issues
LazyData  FALSE
Depends  R (>= 4.1)
Imports  minfi, HDF5Array, rhdf5, S4Vectors, utils, methods, RCurl, R.utils, BiocFileCache, basilisk, reticulate, DelayedMatrixStats
Suggests  minfiData, minfiDataEPIC, knitr, testthat, ggplot2, gridExtra, rmarkdown, BiocStyle, GenomicRanges, limma, ExperimentHub, AnnotationHub
VignetteBuilder  knitr
biocViews  DNAMethylation, Epigenetics, Microarray, MethylationArray, ExperimentHub
RoxygenNote  7.2.0
git_url  https://git.bioconductor.org/packages/recountmethylation
git_branch  RELEASE_3_18
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Description

Get background signal for BeadArray metric calculations.
Get the Biotin staining Red BeadArray metric.
Get the Biotin staining Green BeadArray metric.
Get the Non-polymorphic Red BeadArray metric.
Get the Non-polymorphic Green BeadArray metric.
Get the Bisulfite Conversion I Red BeadArray metric.
Get the Restoration BeadArray metric.
Get the Specificity I Red BeadArray metric.
Get the Specificity I Green BeadArray metric.
Get the Specificity II Green BeadArray metric.
Get the Extension Red BeadArray metric.
Get the Extension Green BeadArray metric.
Get the Hybridication (high vs. medium) BeadArray metric.
Get the Hybridication (medium vs. low) BeadArray metric.
Get the Target Removal I BeadArray metric.
Get the Target Removal 2 BeadArray metric.
Get the Bisulfite Conversion I Green BeadArray metric.
Get the Bisulfite Conversion II BeadArray metric.

**Usage**

```r
ba.background(cdf, ct = "Extension")

ba.biotinstaining.red(
  rs,
  biotin.baseline,
  rm,
  cdf,
  mnum = 1,
  mtot = 17,
  cnamei = "biotin.stain.red",
  ct = "Biotin|DNP"
)

ba.biotinstaining.grn(
  gs,
  biotin.baseline,
  rm,
  cdf,
  mnum = 2,
  mtot = 17,
  cnamei = "biotin.stain.grn",
  ct = "Biotin|DNP"
)

ba.nonpolymorphic.red(
  rs,
  rm,
  cdf,
```
mnum = 3,
mtot = 17,
cnamei = "nonpolymorphic.red",
c = "NP"
)

ba.nonpolymorphic.grn(
  gs,
  rm,
  cdf,
  mnum = 4,
  mtot = 17,
  cnamei = "nonpolymorphic.grn",
  c = "NP"
)

ba.bisulfiteconv1.red(
  rs,
  rm,
  cdf,
  mnum = 5,
  mtot = 17,
  cnamei = "bisulfite.conv1.red",
  c = "Conversion I-"
)

ba.restoration(
  gs,
  rm,
  cdf,
  addr.bkg,
  baseline = 3000,
  mnum = 6,
  mtot = 17,
  cnamei = "restoration.grn",
  c = "RESTORATION"
)

ba.specificity1.red(
  rs,
  rm,
  cdf,
  mnum = 7,
  mtot = 17,
  cnamei = "specificityI.red"
)

ba.specificity1.grn(

ba.background

    gs,
    rm,
    cdf,
    mnum = 8,
    mtot = 17,
    cnamei = "specificityI.grn"
)

ba.specificity2( 
    rs,
    gs,
    rm,
    cdf,
    mnum = 9,
    mtot = 17,
    cnamei = "specificityII",
    ct = "Specificity 2"
)

ba.extension.red(
    rs,
    rm,
    cdf,
    mnum = 10,
    mtot = 17,
    cnamei = "extension.red",
    ct = "Extension.*"
)

ba.extension.grn(
    gs,
    rm,
    cdf,
    mnum = 11,
    mtot = 17,
    cnamei = "extension.grn",
    ct = "Extension.*"
)

ba.hybridization.hi.vs.med( 
    gs,
    rm,
    cdf,
    mnum = 12,
    mtot = 17,
    cnamei = "hyb.hi.med",
    ct = "High|Medium"
)
ba.hybridization.med.vs.low(
    gs,
    rm,
    cdf,
    mnum = 13,
    mtot = 17,
    cnamei = "hyb.med.low",
    ct = "Low|Medium"
)

ba.targetremoval1(
    gs,
    rm,
    cdf,
    baseline = 3000,
    mnum = 14,
    mtot = 17,
    cnamei = "target.removal1",
    ct = "Extension|Target Removal 1"
)

ba.targetremoval2(
    gs,
    rm,
    cdf,
    baseline = 3000,
    mnum = 15,
    mtot = 17,
    cnamei = "target.removal2",
    ct = "Extension|Target Removal 2"
)

ba.bisulfiteconv1.grn(
    gs,
    rm,
    cdf,
    mnum = 16,
    mtot = 17,
    cnamei = "bisulfite.conv1.grn",
    ct = "Conversion I-
"
)

ba.bisulfiteconv2(
    rs,
    gs,
    rm,
    cdf,
Arguments

cdf  Control probe data frame (required).
ct  Column name/metric title (character, required).
rs  Red signal matrix (required).
biotin.baseline  Baseline signal for the biotin stain assay (required).
rm  Results matrix containing control metrics (required).
mnum  Metric number out of total (numeric).
mtot  Total metrics to be calculated (numeric, 17).
cnamei  Column name (character, required).
gs  Green signal matrix (required).
addr.bkg  Background signal probe address (required).
baseline  Baseline measure for signals (integer, 3000).

description

Get the BeadArray control metrics from HM450K platform red/grn signals.

Usage

bactrl(
    cdf = NULL,
    baset = "reduced",
    rg = NULL,
    rs = NULL,
    gs = NULL,
    baseline = 3000,
    biotin.baseline = 1
)
Arguments

CDF    Control probe annotations (optional, NULL, data.frame, cols = properties, rows = probes).

baset  Either the most informative BeadArray metrics (a.k.a. "reduced" set, 5 metrics, recommended), or the full set of 17 metrics ("all").

rg     An RGChannelSet object (optional, NULL).

rs     Red signal data (optional, NULL, data.frame, columns are probes, rows are samples, column names are addresses, rownames are samples/GSM IDs).

gs     Green signal data (optional, NULL, data.frame, columns are probes, rows are samples, column names are addresses, rownames are samples/GSM IDs).

baseline Baseline measure for signals (integer, 3000).

biotin.baseline Baseline to use for biotin controls (integer, 1).

Details

This function calculates the BeadArray quality metrics based on Illumina’s documentation and previous work (see references). Based on previous work, this function can calculate the full set of 17 metrics (e.g. baset = 'all'), or the reduced set of 5 metrics (e.g. baset = 'reduced'), where the latter is recommended for most purposes. For additional details, consult the BeadArray metrics vignette in this package.

Value

Matrix of BeadArray signal values

References


See Also

bathresh

Examples

dir <- system.file("extdata", "bactrl", package = "recountmethylation")
rgf <- get(load(file.path(dir, "rgf-cgctrl-test_hm450k-minfidata.rda")))
mba <- bactrl(rg = rgf)
bathresh

Get BeadArray control outcomes from a matrix of metric signals

Description
Get Illumina’s prescribed minimum quality thresholds for BeadArray metrics.

Usage
bathresh()

Value
Data frame of minimum BeadArray quality thresholds.

References

See Also
bactrl

Examples
dfthresh <- bathresh()

data_mdpost
Retrieve all available sample metadata from an HDF5 database.

Description
Retrieve all available sample metadata in a dataset from an HDF5 database. Returns data in metadata dataset "dsn" contained in an h5 file located at path "dbn."

Usage
data_mdpost(dbn = "remethdb2.h5", dsn = "mdpost")

Arguments
dbn Path to h5 HDF5 database file.
dsn Name or group path to HDF5 dataset containing the sample metadata and learned annotations.
gds_idat2rg

Get IDATs as an RGChannelSet from GEO/GDS

Description
Queries and downloads GSM IDAT files in GEO Data Sets db, then returns the assay data as an "RGChannelSet", calling gds_idatquery() then minfi::read.metharray().

Usage

gds_idat2rg(
gsmvi,
rmdl = TRUE,
ext = "gz",
dfp = ".idats/",

silent = TRUE
)

Arguments

gsmvi
A vector of GSM IDs (alphanumeric character strings).

rmdl
Whether to remove downloaded IDAT files when finished (default TRUE).

ext
Extension for downloaded files (default "gz").

dfp
Destination for IDAT downloads.

burl

silent
Whether to suppress warnings on download removal (default TRUE).

Value
An RGChannelSet object
**gds_idatquery**

Query and download IDATs from GEO Data Sets

### Description

Queries GEO Data Sets for IDATs, and downloads available IDATs. This uses anticipated string pattern to construct the URL path for the query. IDATs are detected from the supplement for a GSE record.

### Usage

```r
gds_idatquery(
  gsmvi,
  ext = "gz",
  expand = TRUE,
  verbose = FALSE,
  dfp = "idats",
)
)
```

### Arguments

- `gsmvi` Vector of valid GSM IDs.
- `ext` Filename extension.
- `expand` Whether to expand compressed files.
- `verbose` Whether to show verbose messages (default FALSE).
- `dfp` Destination directory for downloads.
- `burl` Base URL string for RCurl query.

### Value

Lists the basename paths and filenames of IDATs downloaded.

### Examples

```r
query <- try(gds_idatquery(gsmvi = c("GSM2465267", "GSM2814572")))
```
getdb  

Access database files.

Description

Combines download and load functions for databases. If the "namematch" argument isn't provided, the latest available file is downloaded. All files include metadata for the available samples.

There are 6 functions. Functions with "h5se" access HDF5-SummarizedExperiment files, and "h5" functions access HDF5 databases. The 4 h5se functions are "rg" (RGChannelSet), "gm" (MethylSet), "gr" (GenomicRatioSet), and "test" (data for 2 samples from "gr"). The 2 h5 functions are "rg" (red and green signal datasets), and "test" (data for 2 samples from "rg"). See vignette for details about file types and classes.

Usage

getdb_h5se_test(
  platform = NULL,
  dfp = NULL,
  namematch = "remethdb-h5se_gr-test.*",
  verbose = FALSE
)

getdb_h5_test(
  platform = NULL,
  namematch = "remethdb-h5_rg-test_.*",
  dfp = NULL,
  verbose = FALSE
)

getdb_h5se_gr(
  platform = c("hm450k", "epic"),
  dfp = NULL,
  namematch = "remethdb_h5se-gr_.*",
  verbose = FALSE
)

getdb_h5se_gm(
  platform = c("hm450k", "epic"),
  dfp = NULL,
  namematch = "remethdb_h5se-gm_.*",
  verbose = FALSE
)

getdb_h5se_rg(
  platform = c("hm450k", "epic"),
  dfp = NULL,
getrg

```r
namematch = "remethdb-h5se_rg_.*",
verbose = FALSE
)

getdb_h5_rg(
    platform = c("hm450k", "epic"),
    dfp = NULL,
    namematch = "remethdb-h5_rg_.*",
    verbose = FALSE
)
```

Arguments

- **platform**: Valid supported DNAm array platform type. Currently either "epic" for EPIC/HM850K, or "hm450k" for HM450K.
- **dfp**: Folder to search for database file (optional, if NULL then searches cache dir specified by BiocFileCache).
- **namematch**: Filename pattern to match when searching for database (see defaults).
- **verbose**: Whether to return verbose messages (default FALSE).

Value

Either a SummarizedExperiment object for h5se functions, or a file path for h5 functions.

See Also

- `get_rmdl()`

Examples

```r
h5 <- getdb_h5_test(dfp = tempdir())
```

Description

Queries signal datasets in an h5 HDF5 database file. Handles identity queries to rows (GSM IDs) or columns (bead addresses). Returns query matches either as a list of datasets or a single RGChannelSet, with option to include sample metadata.
Usage

```r
getrg(
  dbn,
  gsmv = NULL,
  cgv = NULL,
  data.type = c("se"),
  dsv = c("redsignal", "greensignal"),
  all.gsm = FALSE,
  all.cg = TRUE,
  metadata = TRUE,
  md.dsn = "mdpost",
  verbose = FALSE
)
```

Arguments

- **dbn**: Name of the HDF5 database file.
- **gsmv**: Vector valid GSM IDs (rows) to query, either NULL or vector of length > 2 valid GSM IDs, or "all.gsm" should be TRUE.
- **cgv**: Vector of valid bead addresses (columns) to query in the signal datasets (default NULL).
- **data.type**: Format for returned query matches, either as datasets "df" or RGChannelSet "se" object.
- **dsv**: Vector of raw signal datasets or group paths to query, including both the red channel 'redsignal' and green channel 'greensignal' datasets.
- **all.gsm**: Whether to query all available GSM IDs.
- **all.cg**: Whether to query all available CpG probe addresses.
- **metadata**: Whether to access available postprocessed metadata for queried samples.
- **md.dsn**: Name of metadata dataset in h5 file.
- **verbose**: Whether to post status messages.

Value

Returns either an RGChannelSet or list of data.frame objects from dataset query matches.

See Also

- `rgse()`

Examples

```r
path <- system.file("extdata", "h5test", package = "recountmethylation")
fn <- list.files(path)
dopath <- file.path(path, fn)
rg <- getrg(dbn = doopath, all.gsm = TRUE, metadata = FALSE)
dim(rg) # [1] 11162 2
```
get_crossreactive_cpgs

Description
Get cross-reactive CpG probe IDs for Illumina BeadArray platforms.

Usage
get_crossreactive_cpgs(probeset = "all")

Arguments
probeset Specify the set of probes to filter ("all", "hm450k", "epic", "chen", "pidsley", "illumina").

Details
Prior work showed significant cross-reactivity at subsets of CpG probes on Illumina’s BeadArray platforms, including HM450K and EPIC. This was primarily due to the probe sequence, as the targeted 50-bp sequence can be either too short or too degenerate to bind a particular DNA region with high specificity. This can cause cross-reaction with off-target DNA locations, including at entirely different chromosomes than the target sequence. Consult the individual publication sources for details about the identification and consequences of cross-reactive CpG probes.

You can retrieve a cross-reactive probe set in a variety of ways. For instance, declare the publication source with either "chen" (for Chen et al 2013), "pidsley" (for Pidsley et al 2016), or "illumina" (for official Illumina documentation), or declare the platform category as either "all" (both HM450K and EPIC), "hm450k", or "epic.".

Value
Vector of cross-reactive CpG probe IDs.

References
See Also

bactrl, get_qcsignal

Examples

```r
length(get_crossreactive_cpgs("all")) # 46324
length(get_crossreactive_cpgs("hm450k")) # 30540
length(get_crossreactive_cpgs("epic")) # 43410
length(get_crossreactive_cpgs("chen")) # 29233
length(get_crossreactive_cpgs("pidsley")) # 43254
length(get_crossreactive_cpgs("illumina")) # 1031
```

Description

Get the hashed features for a data table. Uses reticulate package to call the Python script to do feature hashing on a table of data. It is assumed the input table has sample data in rows, with probe data in columns. The input data table should have row names but not column names.

Usage

```r
get_fh(csv_savepath, csv_openpath, ndim = 1000, lstart = 1)
```

Arguments

- `csv_savepath`: Name/path of hashed features table to write (required, string, writes new csv where rows = samples, cols = hashed features).
- `csv_openpath`: Name/path of table to hash (required, string, assumes a csv where rows = samples, cols = probes).
- `ndim`: Number of hashed features (integer, 1000).
- `lstart`: Line index to start on (0-based for Python, required, int, 0).

Value

Path to new hashed features table.

Examples

```r
# get example bval csv
# of_fpath <- system.file("extdata", "fhtest",
# package = "recountmethylation")
# of_fpath <- file.path(of_fpath, "tbval_test.csv")
# write new hashed features results
# get_fh(csv_savepath = "bval_fn.csv", csv_openpath = of_fpath, ndim = 100)
```
Description

Get the medians of the log2-transformed M and U signals. This function uses the DelayedMatrixStats implementations of colMedians for rapid calculations on DelayedArray-formatted matrices.

Usage

get_qcsignal(se = NULL, mm = NULL, mu = NULL, sample_idv = NULL)

Arguments

- se: Valid SummarizedExperiment object, such as a MethylSet or similar object for which getMeth() and getUnmeth() methods are defined (optional).
- mm: Matrix of methylated/M signals (optional, not required if se provided).
- mu: Matrix of unmethylated/U signals (optional, not required if se provided).
- sample_idv: Vector of sample IDs to label rows in the returned data frame (optional, uses mm colnames instead if not provided).

Details

Calculates the log2 of median signal for methylated/M and unmethylated/U signals separately.

Value

Data frame of signal summaries.

See Also

bactrl

Examples

library(minfiData)
data(MsetEx)
se <- MsetEx
class(se)
  # [1] "MethylSet"
  # attr("package")
  # [1] "minfi"
ms <- get_qcsignal(se)
get_rmdl

Get DNAm assay data.

Description
Uses RCurl to download the latest HDF5-SummarizedExperiment or HDF5 database compilation files objects from the server. Calls servermatrix and performs various quality checks to validate files and downloads. This function is wrapped in the getdb() set of functions (type "?getdb" for details).

Usage
get_rmdl(
  which.class = c("rg", "gm", "gr", "test"),
  which.type = c("h5se", "h5"),
  which.platform = c("hm450k", "epic"),
  fn = NULL,
  dfp = "downloads",
  url = "https://methylation.recount.bio/",
  show.files = FALSE,
  download = TRUE,
  sslver = FALSE,
  verbose = TRUE
)

Arguments

which.class
   Either "rg", "gm", "gr", or "test" for RGChannelSet, MethylSet, GenomicRatioSet, or 2-sample subset.

which.type
   Either "h5se" for an HDF5-SummarizedExperiment or "h5" for an HDF5 database.

which.platform
   Supported DNAm array platform type. Currently supports either "epic" for EPIC/HM850K, or "hm450k" for HM450K.

fn
   Name of file on server to download (optional, default NULL).

dfp
   Download destination directory (default "downloads").

url
   The server URL to locate files for download.

show.files
   Whether to print server file data to console (default FALSE).

download
   Whether to download (TRUE) or return queried filename (FALSE).

sslver
   Whether to use server certificate check (default FALSE).

verbose
   Whether to return verbose messages (default TRUE).

Value
New filepath to dir containing the downloaded data.
**get_servermatrix**

See Also

servermatrix(), getURL(), loadHDF5SummarizedExperiment(), h5ls()

Examples

```
# prints file info from server:
path <- try(get_rmdl(which.class = "test", which.type = "h5se",
show.files = TRUE, download = FALSE))
```

Description

Get a matrix of server files. If the RCurl call fails, a matrix is loaded from the stored package files at `sm_path`.

Usage

```
get_servermatrix(  
  dn = NULL,  
  sslver = FALSE,  
  printmatrix = TRUE,  
  url = "https://methylation.recount.bio/",  
  verbose = FALSE,  
  sm_path = system.file("extdata", "servermatrix_rda", package = "recountmethylation")  
)
```

Arguments

- **dn**: Server data returned from RCurl (default NULL).
- **sslver**: Whether to use SSL certificate authentication for server connection (default FALSE).
- **printmatrix**: Whether to print the data matrix to console (default TRUE).
- **url**: Server website url (default "https://methylation.recount.bio/").
- **verbose**: Whether to show verbose messages (default FALSE).
- **sm_path**: Path to the servermatrix_rda dir containing the stored servermatrix files (default: system.file...).

Value

Matrix of server files and file metadata

See Also

servermatrix, get_rmdl, smfilt
Examples

```r
sm <- get_servermatrix(url = "")
```

**hread**

*Query and store an HDF5 dataset on row and column indices.*

**Description**

Connect to an HDF5 database h5 file with rhdf5::h5read(). Returns the subsetted data.

**Usage**

```r
hread(ri, ci, dsn = "redsignal", dbn = "remethdb2.h5")
```

**Arguments**

- `ri`: Row indices in dataset.
- `ci`: Column indices in dataset.
- `dsn`: Name of dataset or group of dataset to connect with.
- `dbn`: Path to h5 database file.

**Value**

HDF5 database connection object.

**Examples**

```r
# Get tests data pointer
path <- system.file("extdata", "h5test", package = "recountmethylation")
fn <- list.files(path)
dbpath <- file.path(path, fn)
# red signal, first 2 assay addr, 3 samples
reds <- hread(1:2, 1:3, d = "redsignal", dbn = dbpath)
dim(reds) # [1] 2 3
```
Description

Make search index from table of hashed features. Additional details about the hnswlib search index parameters (e.g. 'space_val', 'efc_val', 'm_val', and 'ef_val') can be found in the Python package docstrings and ReadMe.

Usage

```r
make_si(
    fh_csv_fpath,
    si_fname = "new_search_index.pickle",
    si_dict_fname = "new_index_dict.pickle",
    threads = 4,
    space_val = "l2",
    efc_val = 2000,
    m_val = 1000,
    ef_val = 2000
)
```

Arguments

- `fh_csv_fpath`: Name/path of csv (e.g. a table of hashed features) containing data for the index (required, string, "bvaltest.csv", where rows = samples, cols = features).
- `si_fname`: Name of new search index file to save (required, string, "new_search_index.pickle").
- `si_dict_fname`: Name of new index dictionary, with string labels, to save (required, string, "new_index_dict.pickle").
- `threads`: Number of threads for processing new index (required, int, 4).
- `space_val`: Space value for new search index (required, valid string, "l2").
- `efc_val`: EFC value for the index (required, int, 2000).
- `m_val`: M value for the index (required, int, 1000).
- `ef_val`: EF value for the index (required, int, 2000).

Value

Boolean, TRUE if new search index and dictionary created, FALSE if creating the new search index and dictionary files failed, otherwise NULL.

Examples

```r
# fh_csv_fpath <- system.file("extdata", "fhtest",
# package = "recountmethylation")
# fh_csv_fpath <- file.path(fh_csv_fpath, "bval_fn.csv")
# make_si(fh_csv_fpath)
```
matchds_1to2

Match two datasets on rows and columns

Description

Match 2 datasets using the character vectors of row or column names. This is used to assemble an "RGChannelSet" from a query to an h5 dataset.

Usage

matchds_1to2(
  ds1,
  ds2,
  mi1 = c("rows", "columns"),
  mi2 = c("rows", "columns"),
  subset.match = FALSE
)

Arguments

ds1 First dataset to match
ds2 Second dataset to match
mi1 Match index of ds1 (either "rows" or "columns")
mi2 Match index of ds2 (either "rows" or "columns")
subset.match If index lengths don’t match, match on the common subset instead

Value

A list of the matched datasets.

Examples

# get 2 data matrices
ds1 <- matrix(seq(1, 10, 1), nrow = 5)
ds2 <- matrix(seq(11, 20, 1), nrow = 5)
rownames(ds1) <- rownames(ds2) <- paste0("row", seq(1, 5, 1))
colnames(ds1) <- colnames(ds2) <- paste0("col", c(1, 2))
ds2 <- ds2[rev(seq(1, 5, 1)), c(2, 1)]
# match row and column names
lmatched <- matchds_1to2(ds1, ds2, mi1 = "rows", mi2 = "rows")
lmatched <- matchds_1to2(lmatched[[1]], lmatched[[2]], mi1 = "columns",
  mi2 = "columns")
# check matches
ds1m <- lmatched[[1]]
ds2m <- lmatched[[2]]
identical(rownames(ds1m), rownames(ds2m))
identical(colnames(ds1m), colnames(ds2m))
Description

Query an HNSW search index. Does K Nearest Neighbors lookup on a previously saved search index object, returning the K nearest neighbors of the queried sample(s). The `query_si()` function returns verbose output, which can be silenced with `suppressMessages()`.

Usage

```r
query_si(
  sample_idv,
  fh_csv_fpath,
  si_fname = "new_search_index",
  si_fpath = ".",
  lkval = c(1, 2)
)
```

Arguments

- `sample_idv`: Vector of valid sample IDs, or GSM IDs, which are included in the rownames of the hashed features table at `fh_csv_fpath` (required, vector of char strings).
- `fh_csv_fpath`: Path to the hashed features table, which includes rownames corresponding to sample ID strings in the `sample_idv` vector (required, char).
- `si_fname`: Base filename of the search index object, used to find the search index and index dict files, which are expected to be located at `si_fpath` (required, char).
- `si_fpath`: Path to the directory containing the search index and index dict files (required, char).
- `lkval`: Vector of K nearest neighbors to return per query (optional, int, `c(1, 2)`).

Value

Nearest neighbors results of search index query.

Examples

```r
# file paths
# fh table
# fh_csv_fname <- system.file("extdata", "fhtest",
#   package = "recountmethylation")
# fh_csv_fname <- file.path(fh_csv_fname, "bval_fh10.csv")
# si dict
# index_dict_fname <- system.file("extdata", "sitest",
#   package = "recountmethylation")
# index_dict_fname <- file.path(index_dict_fname, "new_index_dict.pickle")
```
# set sample ids to query
# sample_idv <- c("GSM1038308.1548799666.hlink.GSM1038308_5958154021_R01C01",
# "GSM1038309.1548799666.hlink.GSM1038309_5958154021_R02C01")
# set a list of k nearest neighbors to query
# lkval <- c(1,2,3)

# get query results as a data frame (with verbose results messaging)
# dfk <- query_si(sample_idv = sample_idv, lkval = lkval,
# fh_csv_fname = "bval_fn.csv",
# index_dict_fname = "new_index_dict.pickle")
# returns:

## Starting basilisk process...
## Defining the virtual env dependencies...
## Running virtual environment setup...
## Sourcing Python functions...
## Querying the search index...
## Getting hashed features data for samples...
## Getting index data for sample:
## GSM1038308.1548799666.hlink.GSM1038308_5958154021_R01C01'
## Getting index data for sample:
## GSM1038309.1548799666.hlink.GSM1038309_5958154021_R02C01'
## Beginning queries of k neighbors from lk...
## ii = 0 , ki = 1
## Loading search index...
## Querying 2 elements in data with k = 1 nearest neighbors...
## Query completed, time: 0.0007359981536865234
## Applying labels to query results...
## Returning data (sample id, k index, and distance)...  
## ii = 1 , ki = 2
## Loading search index...
## Querying 2 elements in data with k = 2 nearest neighbors...
## Query completed, time: 0.0006208419799804688
## Applying labels to query results...
## Returning data (sample id, k index, and distance)...  
## ii = 2 , ki = 3
## Provided k '3' > n si samples, skipping...
## Returning query results...

---

**rgse**

*Form an RGChannelSet from a list containing signal data matrices*

**Description**

Forms an RGChannelSet from signal data list. This is called by certain queries to h5 files.

**Usage**

```
rgse(ldat, verbose = FALSE)
```
servermatrix

Arguments

  ldat      List of raw signal data query results. Must include 2 data.frame objects named "redsignal" and "greensignal."
  verbose  Whether to post status messages.

Value

Returns a RGChannelSet object from raw signal dataset queries.

See Also

getrg(), RGChannelSet()

Examples

path <- system.file("extdata", "h5test", package = "recountmethylation")
fn <- list.files(path)
dbpath <- file.path(path, fn)
rg <- getrg(dbn = dbpath, all.gsm = TRUE, metadata = FALSE)
dim(rg) # [1] 11162 2
class(rg)
# [1] "RGChannelSet"
# attr(,"package")
# [1] "minfi"

Description

Called by get_rmdl() to get a matrix of database files and file info from the server. Verifies valid versions and timestamps in filenames, and that h5se directories contain both an assays and an se.rds file.

Usage

servermatrix(
  dn = NULL,
  sslver = FALSE,
  printmatrix = TRUE,
  url = "https://methylation.recount.bio/",
  verbose = FALSE
)

setup_sienv

Arguments

- **dn**: Server data returned from RCurl (default NULL).
- **sslver**: Whether to use SSL certificate authentication for server connection (default FALSE).
- **printmatrix**: Whether to print the data matrix to console (default TRUE).
- **url**: Server website url (default "https://methylation.recount.bio/").
- **verbose**: Whether to show verbose messages (default FALSE).

Value

Matrix of server files and file metadata

See Also

get_rmdl, smfilt

Examples

```r
dn <- "remethdb-h5se_gr-test_0-0-1_1590090412 29-May-2020 07:28 -"
sm <- get_servermatrix(dn = dn)
```

Description

Set up a new virtual environment for search index construction using the basilisk package.

Usage

```r
setup_sienv(
  env.name = "dnam_si_hnswlib",
  pkgv = c("python==3.7.1", "hnswlib==0.5.1", "pandas==1.2.2", "numpy==1.20.1",
            "mmh3==3.0.0", "h5py==3.2.1")
)
```

Arguments

- **env.name**: Name of the new virtual environment (required, "dnam_si-hnswlib")
- **pkgv**: Vector of the dependencies and their versions for the new virtual environment (required, format: "packagename==versionnum").

Value

New basilisk environment object.
smfilt

Description
Filters the data matrix returned from servermatrix().

Usage

smfilt(sm, typesdf = NULL)

Arguments

- **sm**: Data matrix returned from servermatrix().
- **typesdf**: Data.frame containing database file info for dm filters.

Value
Filtered data matrix of server file info.

See Also

get_rmdl, servermatrix

Examples

dm <- matrix(c("remethdb_h5-rg_epic_0-0-2_1589820348.h5","08-Jan-2021",
   "09:46","66751358297"), nrow = 1)
smfilt(dm)
Index

ba.background, 2
ba.biotinstaining.grn (ba.background), 2
ba.biotinstaining.red (ba.background), 2
ba.bisulfiteconv1.grn (ba.background), 2
ba.bisulfiteconv1.red (ba.background), 2
ba.bisulfiteconv2 (ba.background), 2
ba.extension.grn (ba.background), 2
ba.extension.red (ba.background), 2
ba.hybridization.hi.vs.med (ba.background), 2
ba.hybridization.med.vs.low (ba.background), 2
ba.nonpolymorphic.grn (ba.background), 2
ba.nonpolymorphic.red (ba.background), 2
ba.restoration (ba.background), 2
ba.specificity1.grn (ba.background), 2
ba.specificity1.red (ba.background), 2
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ba.targetremoval2 (ba.background), 2
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