Package ‘knowYourCG’

May 29, 2024

Type Package
Title Functional analysis of DNA methylome datasets
Version 1.0.0
Description knowYourCG automates the functional analysis of DNA methylation data. The package tests the enrichment of discrete CpG probes across thousands of curated biological and technical features. GSEA-like analysis can be performed on continuous methylation data query sets. knowYourCG can also take beta matrices as input to perform feature aggregation over the curated database sets.

Depends R (>= 4.4.0)

URL https://github.com/zhou-lab/knowYourCG

BugReports https://github.com/zhou-lab/knowYourCG/issues

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VignetteBuilder knitr

Encoding UTF-8

LazyData false

RoxygenNote 7.2.3

Imports sesameData, dplyr, methods, rlang, GenomicRanges, IRanges, reshape2, S4Vectors, stats, stringr, utils

biocViews Epigenetics, DNAMethylation, MethylationArray

Suggests testthat (>= 3.0.0), SummarizedExperiment, rmarkdown, knitr, sesame, gprofiler2

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aggregateTestEnrichments

Description

Aggregate test enrichment results

Usage

aggregateTestEnrichments(result_list, column = "estimate", return_df = FALSE)

Arguments

result_list a list of results from testEnrichment
column the column name to aggregate (Default: estimate)
return_df whether to return a merged data frame

Value

a matrix for all results
Examples

```r
## pick some big TFBS-overlapping CpG groups
sesameData::sesameDataCache(data_titles =
  c("KYCG.MM285.TFBSconsensus.20220116","KYCG.MM285.chromHMM.20210210",
    "probeIDSignature", "MM285.address"))

queries <- cg_lists[(sapply(cg_lists, length) > 40000)]
result_list <- lapply(queries, testEnrichment, "MM285.chromHMM")
mtx <- aggregateTestEnrichments(result_list)
```

annoProbes

Annotate Probe IDs using KYCG databases

Description

see sesameData_annoProbes if you’d like to annotate by genomic coordinates (in GRanges)

Usage

```r
annoProbes(
  probeIDs,
  databases,
  db_names = NULL,
  platform = NULL,
  sep = ",",
  indicator = FALSE,
  silent = FALSE
)
```

Arguments

- `probeIDs`: probe IDs in a character vector
- `databases`: character or actual database (i.e. list of probe IDs)
- `db_names`: specific database (default to all databases)
- `platform`: EPIC, MM285 etc. will infer from probe IDs if not given
- `sep`: delimiter used in paste
- `indicator`: return the indicator matrix instead of a concatenated annotation (in the case of have multiple annotations)
- `silent`: suppress message

Value

named annotation vector, or indicator matrix
Examples

```r
sesameData::sesameDataCache(data_titles=
c("MM285.address","probeIDSignature","KYCG.MM285.designGroup.20210210"))
probes <- names(sesameData::sesameData_getManifestGRanges("MM285"))
anno <- annoProbes(probeIDs=probes, "designGroup", silent = TRUE)
```


description

build gene-probe association database

Usage

```r
buildGeneDBs(
  probeIDs = NULL,
  platform = NULL,
  genome = NULL,
  max_distance = 10000,
  silent = FALSE
)
```

Arguments

- `probeIDs` the query probe list. If NULL, use all the probes on the platform
- `platform` HM450, EPIC, MM285, Mammal40, will infer from query if not given
- `genome` hg38, mm10, ..., will infer if not given.
- `max_distance` probe-gene distance for association
- `silent` suppress messages

Value

gene databases

Examples

```r
sesameData::sesameDataCache(data_titles=
c("EPIC.address","genomeInfo.hg38","probeIDSignature"))
query <- c("cg04707299", "cg13380562", "cg00480749")
dbs <- buildGeneDBs(query, platform = "EPIC")
testEnrichment(query, dbs, platform = "EPIC")
```
dbStats aggregates methylation of a given betas matrix over specified database set features

Description

dbStats aggregates methylation of a given betas matrix over specified database set features

Usage

dbStats(betas, databases, fun = mean, na.rm = TRUE, n_min = NULL, f_min = 0.1)

Arguments

betas matrix of beta values where probes are on the rows and samples are on the columns
databases List of vectors corresponding to probe locations for which the features will be extracted
fun aggregation function, default to mean
na.rm whether to remove NA
n_min min number of non-NA for aggregation function to apply, overrides f_min
f_min min fraction of non-NA for aggregation function to apply

Value

matrix with samples on the rows and database set on the columns

Examples

library(SummarizedExperiment)
sesameData::sesameDataCache(data_titles=
c("MM285.467.SE.tissue20Kprobes","KYCG.MM285.probeType.20210630"))
se <- sesameData::sesameDataGet("MM285.467.SE.tissue20Kprobes")
head(dbStats(assay(se), "MM285.probeType")[,1:3])
sesameData::sesameDataGet_resetEnv()
getDBs  

Get databases by full or partial names of the database group(s)

Description

Get databases by full or partial names of the database group(s)

Usage

getDBs(  
group_nms,  
db_names = NULL,  
platform = NULL,  
summary = FALSE,  
allow_multi = FALSE,  
type = NULL,  
silent = FALSE  
)

Arguments

group_nms  database group names
db_names  name of the database, fetch only the given databases
platform  EPIC, HM450, MM285, ... If given, will restrict to that platform.
summary  return a summary of database instead of db itself
allow_multi  allow multiple groups to be returned for
type  numerical, categorical, default: all
silent  no messages each query.

Value

a list of databases, return NULL if no database is found

Examples

sesameData::sesameDataCache(data_titles=  
c("KYCG.MM285.chromHMM.20210210","KYCG.MM285.probeType.20210630"))  
dbs <- getDBs("MM285.chromHMM")  
dbs <- getDBs(c("MM285.chromHMM", "MM285.probeType"))
listDBGroups

List database group names

Description
List database group names

Usage
listDBGroups(filter = NULL, path = NULL, type = NULL)

Arguments
- filter: keywords for filtering
- path: file path to downloaded knowledgebase sets
- type: categorical, numerical (default: all)

Value
a list of db group names

Examples
head(listDBGroups("chromHMM"))
## or listDBGroups(path = "~/Downloads")

testEnrichment
testEnrichment tests for the enrichment of a set of probes (query set) in a number of features (database sets).

Description
testEnrichment tests for the enrichment of a set of probes (query set) in a number of features (database sets).

Usage
testEnrichment(
  probeIDs,
  databases = NULL,
  universe = NULL,
  alternative = "greater",
  include_genes = FALSE,
  platform = NULL,
  silent = FALSE
)
testEnrichmentFisher

Arguments

probeIDs  Vector of probes of interest (e.g., significant probes)
databases List of vectors corresponding to the database sets of interest with associated meta data as an attribute to each element. Optional. (Default: NA)
universe  Vector of probes in the universe set containing all of the probes to be considered in the test. If it is not provided, it will be inferred from the provided platform. (Default: NA).
alternative "two.sided", "greater", or "less"
include_genes include gene link enrichment testing
platform  String corresponding to the type of platform to use. Either MM285, EPIC, HM450, or HM27. If it is not provided, it will be inferred from the query set probeIDs (Default: NA).
silent output message? (Default: FALSE)

Value

A data frame containing features corresponding to the test estimate, p-value, and type of test.

Examples

library(SummarizedExperiment)
sesameData::sesameDataCache(data_titles=
c("MM285.tissueSignature","KYCG.MM285.chromHMM.20210210","MM285.address"))
df <- rowData(sesameData::sesameDataGet("MM285.tissueSignature"))
probes <- df$Probe_ID[df$branch == "B_cell"]
res <- testEnrichment(probes, "chromHMM", platform="MM285")
sesameData::sesameDataGet_resetEnv()

Description

Estimates log2 Odds ratio

Usage

testEnrichmentFisher(query, database, universe, alternative = "greater")
**Arguments**

- **query**: Vector of probes of interest (e.g., significant probes)
- **database**: Vectors corresponding to the database set of interest with associated meta data as an attribute to each element.
- **universe**: Vector of probes in the universe set containing all of
- **alternative**: greater or two.sided (default: greater) the probes to be considered in the test. (Default: NULL)

**Value**

A DataFrame with the estimate/statistic, p-value, and name of test for the given results.

**testEnrichmentSEA** uses the GSEA-like test to estimate the association of a categorical variable against a continuous variable.

**Description**

estimate represent enrichment score and negative estimate indicate a test for depletion

**Usage**

```r
testEnrichmentSEA(
  query,
  database,
  platform = NULL,
  silent = FALSE,
  precise = FALSE,
  prepPlot = FALSE
)
```

**Arguments**

- **query**: query, if numerical, expect categorical database, if categorical expect numerical database
- **databases**: database, numerical or categorical, but needs to be different from query
- **platform**: EPIC, MM285, ..., infer if not given
- **silent**: suppress message (default: FALSE)
- **precise**: whether to compute precise p-value (up to numerical limit) of interest.
- **prepPlot**: return the raw enrichment scores and presence vectors for plotting

**Value**

A DataFrame with the estimate/statistic, p-value, and name of test for the given results.
Examples

```r
sesameData::sesameDataCache(data_titles=c("KYCG.MM285.designGroup.20210210","KYCG.MM285.seqContextN.20210630","probeIDSsignature"))
query <- getDBs("KYCG.MM285.designGroup")["TSS"]
res <- testEnrichmentSEA(query, "MM285.seqContextN")
```

testEnrichmentSpearman

`testEnrichmentSpearman` uses the Spearman statistical test to estimate the association between two continuous variables.

Description

testEnrichmentSpearman uses the Spearman statistical test to estimate the association between two continuous variables.

Usage

```r
testEnrichmentSpearman(num_query, num_db)
```

Arguments

- **num_query**: named numeric vector of probes of interest where names are probe IDs (e.g. significant probes)
- **num_db**: List of vectors corresponding to the database set of interest with associated metadata as an attribute to each element.

Value

A DataFrame with the estimate/statistic, p-value, and name of test for the given results.

testGO

`testGO` tests Gene Ontology of genes overlapping CpG query

Description

estimate represent enrichment score and negative estimate indicate a test for depletion

Usage

```r
testGO(probeIDs, platform = NULL, organism = "hsapiens", gene_name = TRUE, ...)
```
testProbeProximity

testProbeProximity tests if a query set of probes share closer genomic proximity than if randomly distributed

Arguments

- `probeIDs`: Vector of CpG probes IDs or a data frame with gene_name column, usually the output of testEnrichment() function
- `platform`: EPIC, MM285, ..., infer if not given
- `organism`: The organism corresponding to the CpG platform or genes in gene_name column
- `gene_name`: If query is data frame from testEnrichment output, whether to use the gene_name column. If set to FALSE, TFBS will be used (default: FALSE)
- `...`: Additional arguments to sesameData_getGenesByProbes and gost()

Value

A list of enriched terms and meta data from gprofiler2 output

Examples

```r
library(SummarizedExperiment)
sesameData::sesameDataCache(data_titles = c("MM285.tissueSignature","probeIDSignature", "MM285.address","genomeInfo.mm10"))
df <- rowData(sesameData::sesameDataGet('MM285.tissueSignature'))
query <- df$Probe_ID[df$branch == "fetal_liver" & df$type == "Hypo"]
res <- testGO(query, platform="MM285")
```

testProbeProximity

testProbeProximity tests if a query set of probes share closer genomic proximity than if randomly distributed

Description

testProbeProximity tests if a query set of probes share closer genomic proximity than if randomly distributed

Usage

```r
testProbeProximity(
  probeIDs,       # Vector of CpG probes IDs or a data frame with gene_name column, usually the output of testEnrichment() function
  gr = NULL,      # If query is data frame from testEnrichment output, whether to use the gene_name column. If set to FALSE, TFBS will be used (default: FALSE)
  platform = NULL, # The organism corresponding to the CpG platform or genes in gene_name column
  organism = NULL, # EPIC, MM285, ..., infer if not given
  iterations = 100, # Additional arguments to sesameData_getGenesByProbes and gost()
  bin_size = 1500  # Integer
)"`
Arguments

probeIDs  Vector of probes of interest (e.g., significant probes)
gr        GRanges to draw samples and compute genomic distances
platform  String corresponding to the type of platform to use. Either MM285, EPIC, HM450, or HM27. If it is not provided, it will be inferred from the query set probeIDs (Default: NA).
iterations Number of random samples to generate null distribution (Default: 100).
bin_size   the poisson interval size for computing neighboring hits

Value

list containing a dataframe for the poisson statistics and a data frame for the probes in close proximity

Examples

sesameData::sesameDataCache(data_titles=
c(“MM285.tissueSignature”, “MM285.address”, “probeIDssignature”))
library(SummarizedExperiment)
df <- rowData(sesameData::sesameDataGet("MM285.tissueSignature"))
probes <- df$Probe_ID[df$branch == "B_cell"]
res <- testProbeProximity(probeIDs=probes,platform="MM285")
esameData::sesameDataGet_resetEnv()
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