

Package ‘TCGArretriever’

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Type Package

Title Retrieve Genomic and Clinical Data from CBioPortal Including
TCGA Data

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Author Damiano Fantini

Maintainer Damiano Fantini <damiano.fantini@gmail.com>

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Imports httr, reshape2, jsonlite

Suggests stats, utils, knitr, rmarkdown, ggplot2, dplyr

VignetteBuilder knitr

Description The Cancer Genome Atlas (TCGA) is a program aimed at improving our understanding of Cancer Biology. Several TCGA Datasets are available online. 'TCGArretriever' helps accessing and downloading TCGA data hosted on 'cBioPortal' via its Web Interface (see <<https://www.cbioportal.org/>> for more information).

License GPL-3

LazyData true

URL https://www.data-pulse.com/dev_site/TCGArretriever/

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<i>blcaOutputExamples</i>	<i>TCGAretriever Examples</i>
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Description

A list of objects including examples of the output returned by different ‘TCGAretriever’ functions. The objects were obtained from the ‘“blca_tcga”’ study (bladder cancer).

Usage

```
data(blcaOutputExamples)
```

Format

A list including 7 elements.

- exmpl_1** data.frame (dimensions: 10 by 13). Sample output of the ‘get_cancer_studies()‘ function.
- exmpl_2** data.frame (dimensions: 10 by 5). Sample output of the ‘get_cancer_types()‘ function.
- exmpl_3** data.frame (dimensions: 9 by 5). Sample output of the ‘get_case_lists()‘ function.
- exmpl_4** list including 9 elements. Sample output of the ‘expand_cases()‘ function.
- exmpl_5** data.frame (dimensions: 10 by 94). Sample output of the ‘get_clinical_data()‘ function.
- exmpl_6** data.frame (dimensions: 9 by 8). Sample output of the ‘get_genetic_profiles()‘ function.
- exmpl_7** data.frame (dimensions: 10 by 3). Sample output of the ‘get_gene_identifiers()‘ function.
- exmpl_8** data.frame (dimensions: 2 by 10). Sample output of the ‘get_molecular_data()‘ function.
- exmpl_9** data.frame (dimensions: 6 by 27). Sample output of the ‘get_mutation_data()‘ function.

Details

The object was built using the following lines of code.

```
blcaOutputExamples <- list( exmpl_1 = head(get_cancer_studies(), 10), exmpl_2 = head(get_cancer_types(), 10), exmpl_3 = head(get_case_lists("blca_tcga"), 10), exmpl_4 = expand_cases("blca_tcga"), exmpl_5 = head(get_clinical_data("blca_tcga"), 10), exmpl_6 = head(get_genetic_profiles("blca_tcga"), 10), exmpl_7 = head(get_gene_identifiers(), 10), exmpl_8 = get_molecular_data(case_list_id = 'blca_tcga_3way_complete', gprofile_id = 'blca_tcga_rna_seq_v2_mrna', glist = c("TP53", "E2F1"))[, 1:10], exmpl_9 = head(get_mutation_data(case_list_id = 'blca_tcga_sequenced', gprofile_id = 'blca_tcga_mutations', glist = c('TP53', 'PTEN'))))
```

Examples

```
data(blcaOutputExamples)
blcaOutputExamples$exmpl_1
```

expand_cases

Samples Included in a List of Samples.

Description

Each study includes one or more "case lists". Each case list is a collection of samples that were analyzed using one or more platforms/assays. It is possible to obtain a list of all sample identifiers for each case list of interest.

Usage

```
expand_cases(csid, dryrun = FALSE)
```

Arguments

- | | |
|--------|--|
| csid | String corresponding to a TCGA Cancer Study identifier. |
| dryrun | Logical. If TRUE, all other arguments (if any) are ignored and a representative example is returned as output. No Internet connection is required for executing the operation when 'dryrun' is TRUE. |

Value

list containing as many elements as TCGA case lists available for a given TCGA Study. Each element is named after a case list identifier. Also, each element is a character vector including all sample identifiers (case ids) corresponding to the corresponding case list identifier.

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
# Set `dryrun = FALSE` (default option) in production!
x <- expand_cases("blca_tcga", dryrun = TRUE)
lapply(x, utils::head)
```

`fetch_all_tcga` *Fetch All Molecular Data for a Cancer Profile of Interest.*

Description

Recursively query cbioportal to retrieve data corresponding to all available genes. Data are returned as a ‘`data.frame`’ that can be easily manipulated for downstream analyses.

Usage

```
fetch_all_tcгадаda(case_list_id, gprofile_id, mutations = FALSE)
```

Arguments

<code>case_list_id</code>	string corresponding to the identifier of the TCGA Case List of interest
<code>gprofile_id</code>	string corresponding to the identifier of the TCGA Profile of interest
<code>mutations</code>	logical. If TRUE, extended mutation data are fetched instead of the standard TCGA data

Value

A `data.frame` is returned, including the desired TCGA data. Typically, rows are genes and columns are cases. If "extended mutation" data are retrieved (`mutations = TRUE`), rows correspond to individual mutations while columns are populated with mutation features

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
get_cancer_studies
```

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```
## End(Not run)
```

get_cancer_studies *Retrieve the List of Cancer Studies Available at cbioportal.*

Description

Retrieve information about the studies or datasets available at cbioportal.org. Information include a ‘studyId’, description, references, and more.

Usage

```
get_cancer_studies(dryrun = FALSE)
```

Arguments

dryrun Logical. If TRUE, all other arguments (if any) are ignored and a representative example is returned as output. No Internet connection is required for executing the operation when ‘dryrun’ is TRUE.

Value

Data Frame including cancer study information.

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
# Set `dryrun = FALSE` (default option) in production!
all_studies <- get_cancer_studies(dryrun = TRUE)
utils::head(all_studies)
```

`get_cancer_types` *Retrieve Cancer Types.*

Description

Retrieve information about cancer types and corresponding abbreviations from cbioportal.org. Information include identifiers, names, and parental cancer type.

Usage

```
get_cancer_types(dryrun = FALSE)
```

Arguments

`dryrun` Logical. If TRUE, all other arguments (if any) are ignored and a representative example is returned as output. No Internet connection is required for executing the operation when ‘dryrun’ is TRUE.

Value

A data.frame including cancer type information.

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
# Set `dryrun = FALSE` (default option) in production!
all_canc <- get_cancer_types(dryrun = TRUE)
utils::head(all_canc)
```

get_case_lists

Retrieve Case List Available for a Specific Cancer Study.

Description

Each study includes one or more "case lists". Each case list is a collection of samples that were analyzed using one or more platforms/assays. It is possible to obtain a list of case list identifiers from cbioportal.org for a cancer study of interest. Identifier, name, description and category are returned for each entry.

Usage

```
get_case_lists(csid, dryrun = FALSE)
```

Arguments

csid	String corresponding to the Identifier of the Study of Interest
dryrun	Logical. If TRUE, all other arguments (if any) are ignored and a representative example is returned as output. No Internet connection is required for executing the operation when 'dryrun' is TRUE.

Value

Data Frame including Case List information.

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
# Set `dryrun = FALSE` (default option) in production!
blca_case_lists <- get_case_lists("blca_tcga", dryrun = TRUE)
blca_case_lists
```

get_clinical_data *Retrieve Clinical Information for a Cancer Study.*

Description

Retrieve Clinical Information about the samples included in a cancer study of interest. For each sample/case, information about the corresponding cancer patient are returned. These may include sex, age, therapeutic regimen, tumor stage, survival status, as well as other information.

Usage

```
get_clinical_data(csid, case_list_id = NULL, dryrun = FALSE)
```

Arguments

<code>csid</code>	String corresponding to a TCGA Cancer Study identifier.
<code>case_list_id</code>	String corresponding to the case_list identifier of interest. This Can be NULL.
<code>dryrun</code>	Logical. If TRUE, all other arguments (if any) are ignored and a representative example is returned as output. No Internet connection is required for executing the operation when ‘dryrun’ is TRUE.

Value

`data.frame` including clinical information of a list of samples/cases of interest.

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
# Set `dryrun = FALSE` (default option) in production!
clinic_data <- get_clinical_data("blca_tcga", dryrun = TRUE)
utils::head(clinic_data[, 1:7])
```

get_genetic_profiles *Retrieve Genetic Profiles for a TCGA Study of Interest*

Description

Retrieve Information about all genetic profiles associated with a cancer study of interest. Each cancer study includes one or more types of molecular analyses. The corresponding assays or platforms are referred to as genetic profiles. A genetic profile identifier is required to download molecular data.

Usage

```
get_genetic_profiles(csid = NULL, dryrun = FALSE)
```

Arguments

csid	String corresponding to the cancer study id of interest
dryrun	Logical. If TRUE, all other arguments (if any) are ignored and a representative example is returned as output. No Internet connection is required for executing the operation when 'dryrun' is TRUE.

Value

data.frame including information about genetic profiles.

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
# Set `dryrun = FALSE` (default option) in production!
get_genetic_profiles("blca_tcga", dryrun = TRUE)
```

get_gene_identifiers Retrieve All Gene Identifiers

Description

Obtain all valid gene identifiers, including ENTREZ gene identifiers and HUGO gene symbols. Genes are classified according to the gene type (*e.g.* , 'protein-coding', 'pseudogene', 'miRNA', ...). Note that miRNA and phosphoprotein genes are associated with a negative entrezGeneId.

Usage

```
get_gene_identifiers(dryrun = FALSE)
```

Arguments

dryrun Logical. If TRUE, all other arguments (if any) are ignored and a representative example is returned as output. No Internet connection is required for executing the operation when 'dryrun' is TRUE.

Value

Data Frame including gene identifiers.

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
# Set `dryrun = FALSE` (default option) in production!
x <- get_gene_identifiers(dryrun = TRUE)
```

get_molecular_data *Retrieve Molecular Data corresponding to a Genetic Profile of Interest.*

Description

Retrieve Data corresponding to a Genetic Profile of interest from a cancer study of interest. This function is the workhorse of the TCGAretriever package and can be used to fetch data concerning several genes at once. For retrieving mutation data, please use the ‘get_mutation_data()‘ function. For large queries (more than 500 genes), please use the ‘fetch_all_tcгадata()‘ function.

Usage

```
get_molecular_data(  
  case_list_id,  
  gprofile_id,  
  glist = c("TP53", "E2F1"),  
  dryrun = FALSE  
)
```

Arguments

case_list_id	String corresponding to the Identifier of a list of cases.
gprofile_id	String corresponding to the Identifier of a genetic Profile of interest.
glist	Vector including one or more gene identifiers (ENTREZID or OFFICIAL_SYMBOL). ENTREZID gene identifiers should be passed as numeric.
dryrun	Logical. If TRUE, all other arguments (if any) are ignored and a representative example is returned as output. No Internet connection is required for executing the operation when ‘dryrun‘ is TRUE.

Value

data.frame including the molecular data of interest. Rows are genes, columns are samples.

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
# Set `dryrun = FALSE` (default option) in production!
x <- get_molecular_data(case_list_id = 'blca_tcga_3way_complete',
                        gprofile_id = 'blca_tcga_rna_seq_v2_mrna',
                        glist = c("TP53", "E2F1"), dryrun = TRUE)
x[, 1:10]
```

get_mutation_data

Retrieve Mutation Data corresponding to a Genetic Profile of Interest.

Description

Retrieve DNA Sequence Variations (Mutations) identified by exome sequencing projects. This function is the workhorse of the TCGAretriever package for mutation data and can be used to fetch data concerning several genes at once. For retrieving non-mutation data, please use the ‘get_molecular_data()’ function. For large queries (more than 500 genes), please use the ‘fetch_all_tcgadata()’ function.

Usage

```
get_mutation_data(
  case_list_id,
  gprofile_id,
  glist = c("TP53", "E2F1"),
  dryrun = FALSE
)
```

Arguments

<code>case_list_id</code>	String corresponding to the Identifier of a list of cases.
<code>gprofile_id</code>	String corresponding to the Identifier of a genetic Profile of interest.
<code>glist</code>	Vector including one or more gene identifiers (ENTREZID or OFFICIAL_SYMBOL). ENTREZID gene identifiers should be passed as numeric.
<code>dryrun</code>	Logical. If TRUE, all other arguments (if any) are ignored and a representative example is returned as output. No Internet connection is required for executing the operation when ‘dryrun’ is TRUE.

Value

data Frame inclusing one row per mutation

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
# Set `dryrun = FALSE` (default option) in production!
x <- get_mutation_data(case_list_id = 'blca_tcga_sequenced',
                      gprofile_id = 'blca_tcga_mutations',
                      glist = c('TP53', 'PTEN'), dryrun = TRUE)
utils::head(x[, c(4, 7, 23, 15, 16, 17, 24, 18, 21)])
```

make_groups

Split Numeric Vectors in Groups.

Description

Assign each element of a numeric vector to a group. Grouping is based on ranks: numeric values are sorted and then split in 2 or more groups. Values may be sorted in an increasing or decreasing fashion. The vector is returned in the original order. Labels may be assigned to each group.

Usage

```
make_groups(num_vector, groups, group_labels = NULL, desc = FALSE)
```

Arguments

num_vector	numeric vector. It includes the values to be assigned to the different groups
groups	integer. The number of groups that will be generated
group_labels	character vector. Labels for each group. Note that the length of group_labels has to be equal to the number of groups
desc	logical. If TRUE, the sorting is applied in a decreasing fashion

Value

data.frame including the vector provided as argument in the original order ("value") and the grouping vector ("rank"). If labels are provided as an argument, group labels are also included in the data.frame ("labels").

Author(s)

Damiano Fantini, <damiano.fantini@gmail.com>

References

https://www.data-pulse.com/dev_site/TCGAretriever/

Examples

```
exprs_geneX <- c(19.1,18.4,22.4,15.5,20.2,17.4,9.4,12.4,31.2,33.2,18.4,22.1)
groups_num <- 3
groups_labels <- c("high", "med", "low")
make_groups(exprs_geneX, groups_num, groups_labels, desc = TRUE)
```

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