

Package ‘Uniquorn’

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Title Identification of cancer cell lines based on their weighted mutational/ variational fingerprint

Version 2.16.0

Description

This packages enables users to identify cancer cell lines. Cancer cell line misidentification and cross-contamination reprints a significant challenge for cancer researchers.

The identification is vital and in the frame of this package based on the locations/ loci of somatic and germline mutations/ variations.

The input format is vcf/ vcf.gz and the files have to contain a single cancer cell line sample (i.e. a single member/genotype/gt column in the vcf file).

The implemented method is optimized for the Next-generation whole exome and whole genome DNA-sequencing technology. RNA-

seq data is very likely to work as well but hasn't been rigiously tested yet. Panel-seq will require manual adjustment of thresholds

Imports stringr, R.utils, WriteXLS, stats, doParallel, foreach, GenomicRanges, IRanges, VariantAnnotation

Depends R (>= 3.5)

License Artistic-2.0

LazyData TRUE

Type Package

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add_custom_vcf_to_database

add_custom_vcf_to_database This function adds the variants of parsed custom CCLs to a monet DB instance

Description

add_custom_vcf_to_database This function adds the variants of parsed custom CCLs to a monet DB instance

Usage

```

add_custom_vcf_to_database(
  vcf_input_files,
  ref_gen = "GRCH37",
  library_name = "CUSTOM",
  n_threads = 1,
  test_mode = FALSE
)

```

Arguments

vcf_input_files	a character vector containing the input vcf files. This may be one or many vcf files.
ref_gen	a character string specifying the reference genome version. All training sets are associated with a reference genome version. Default is "GRCH37".
library_name	a character string giving the name of the library to add the cancer cell lines to. Default is "CUSTOM". Library name will be automatically added as a suffix to the identifier.
n_threads	an integer specifying the number of threads to be used.
test_mode	Is this a test? Just for internal use

Value

Message wheather the adding was successful

Examples

```

HT29_vcf_file = system.file("extdata/HT29_TEST.vcf", package = "Uniquorn");
add_custom_vcf_to_database(
  vcf_input_files = HT29_vcf_file,
  library_name = "CELLMINER",
  ref_gen = "GRCH37",
  n_threads = 1,
  test_mode = TRUE
)

```

add_missing_cls	<i>add_missing_cls</i>
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Description

add_missing_cls

Usage

```
add_missing_cls(res_table, dif_cls)
```

Arguments

<code>res_table</code>	Table that contains the identification results
<code>dif_cls</code>	Missing CLs

Value

Results table with added missing cls

```
add_penalty_statistics
```

```
add_penalty_statistics
```

Description

Add penalty statistics to results

Usage

```
add_penalty_statistics(match_t, minimum_matching_mutations)
```

Arguments

<code>match_t</code>	object that contains the matching variants
<code>minimum_matching_mutations</code>	a numerical giving the minimum amount of mutations that has to match between query and training sample for a positive prediction

Value

The updated statistics

```
add_p_q_values_statistics
      add_p_q_values_statistics
```

Description

A hypergeometric distribution-assumption allows to calculate the p-values for a significant or non-significant overlap in this function

Usage

```
add_p_q_values_statistics(g_query, match_t, p_value, ref_gen,
      minimum_matching_mutations, top_hits_per_library)
```

Arguments

g_query	IRanges object that contains the query variants
match_t	A table that contains the nubmber of matching variants
p_value	Threshold for the significance p-value
ref_gen	Reference genome version
minimum_matching_mutations	Manual lower amount of matching mutations require for a significant match between a query and a reference
top_hits_per_library	limits significant similarities to the first n hits

Details

add_p_q_values_statistics Calculates the p-values

Value

R table with a statistic

```
create_bed_file      create_bed_file
```

Description

Creates BED files from the found and not found annotated mutations

Usage

```

create_bed_file(
  match_t,
  vcf_fingerprint,
  output_file,
  ref_gen,
  manual_identifier
)

```

Arguments

match_t	R table which contains the mutations from the training database for the cancer cell lines
vcf_fingerprint	contains the mutations that are present in the query cancer cell line's vcf file
output_file	Path to output file
ref_gen	Reference genome version
manual_identifier	Manually enter a vector of CL name(s) whose bed files should be created, independently from them passing the detection threshold

Value

Returns a message which indicates if the BED file creation has succeeded

identify_vcf_file	<i>identify_VCF_file</i>
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Description

Identifies a cancer cell lines contained in a vcf file based on the pattern (start & length) of all contained mutations/ variations.

Usage

```

identify_vcf_file(
  vcf_file,
  output_file,
  ref_gen,
  minimum_matching_mutations,
  mutational_weight_inclusion_threshold,
  write_xls,
  output_bed_file,
  top_hits_per_library,
  manual_identifier,
)

```

```

    verbose,
    p_value,
    confidence_score,
    n_threads,
    write_results
)

```

Arguments

vcf_file	Input vcf file. Only one sample column allowed.
output_file	Path of the output file. If blank, autogenerated as name of input file plus '_uniquorn_ident.tab' suffix.
ref_gen	Reference genome version. All training sets are associated with a reference genome version. Default: GRCH37
minimum_matching_mutations	The minimum amount of mutations that has to match between query and training sample for a positive prediction
mutational_weight_inclusion_threshold	Include only mutations with a weight of at least x. Range: 0.0 to 1.0. 1= unique to CL. ~0 = found in many CL samples.
write_xls	Create identification results additionally as xls file for easier reading
output_bed_file	If BED files for IGV visualization should be created for the Cancer Cell lines that pass the threshold
top_hits_per_library	Limit the number of significant similarities per library to n (default 3) many hits. Is particularly used in contexts when heterogeneous query and reference CCLs are being compared.
manual_identifier	Manually enter a vector of CL name(s) whose bed files should be created, independently from them passing the detection threshold
verbose	Print additional information
p_value	Required p-value for identification. Note that if you set the confidence score, the confidence score overrides the p-value
confidence_score	Cutoff for positive prediction between 0 and 100. Calculated by transforming the p-value by $-1 * \log(p\text{-value})$ Note that if you set the confidence score, the confidence score overrides the p-value
n_threads	Number of threads to be used
write_results	Write identification results to file

Details

identify_vcf_file parses the vcf file and predicts the identity of the sample

Value

R table with a statistic of the identification result

Examples

```
HT29_vcf_file = system.file("extdata/HT29.vcf", package = "Uniquorn");
```

```
identification = identify_vcf_file(  
  vcf_file = HT29_vcf_file,  
  verbose = FALSE,  
  write_results = FALSE  
)
```

```
initiate_canonical_databases  
  initiate_canonical_databases
```

Description

Parses data into r list variable

Usage

```
initiate_canonical_databases(  
  cosmic_file = "CosmicCLP_MutantExport.tsv",  
  ccle_file = "CCLE_hybrid_capture1650_hg19_NoCommonSNPs_CDS_2012.05.07.maf",  
  ref_gen = "GRCH37"  
)
```

Arguments

cosmic_file	The path to the cosmic DNA genotype data file. Ensure that the right reference genome is used
ccle_file	The path to the ccle DNA genotype data file. Ensure that the right reference genome is used
ref_gen	Reference genome version

Value

Returns message if parsing process has succeeded

Examples

```
initiate_canonical_databases(  
  cosmic_file = "CosmicCLP_MutantExport.tsv",  
  ccle_file = "CCLE_hybrid_capture1650_hg19_NoCommonSNPs_CDS_2012.05.07.maf",  
  ref_gen = "GRCH37"  
)
```

```
init_and_load_identification
    init_and_load_identification
```

Description

Initiate the analysis Output basic information

Usage

```
init_and_load_identification(  
    verbose,  
    ref_gen,  
    vcf_file,  
    output_dir  
)
```

Arguments

verbose	Print additional information
ref_gen	Reference genome version. All training sets are associated with a reference genome version. Default: GRCH37
vcf_file	Path to vcf_file
output_dir	Output directory for identification results

Details

init_and_load_identification parses vcf file and output basic information

Value

Three file path instances and the fingerprint

```
match_query_ccl_to_database
    match_query_ccl_to_database
```

Description

Matches query ccl to the database

Usage

```
match_query_ccl_to_database(g_query, ref_gen = "GRCH37", library_name,  
    mutational_weight_inclusion_threshold)
```

Arguments

<code>g_query</code>	IRanges object that contains the variants
<code>ref_gen</code>	Reference genome version. All training sets are associated with a reference genome version. Default: GRCH37
<code>library_name</code>	a character string giving the name of the library
<code>mutational_weight_inclusion_threshold</code>	a numerical giving the lower bound for mutational weight to be included

Value

The R Table `sim_list` which contains the CoSMIC CLP fingerprints

`parse_ccle_genotype_data`
parse_ccle_genotype_data

Description

Parses ccle genotype data

Usage

```
parse_ccle_genotype_data(ccle_file, ref_gen = "GRCH37")
```

Arguments

<code>ccle_file</code>	Path to CCLE file on hard disk
<code>ref_gen</code>	Reference genome version

Value

The R Table `sim_list` which contains the CCLE fingerprints

```
parse_cosmic_genotype_data
      parse_cosmic_genotype_data
```

Description

Parses cosmic genotype data

Usage

```
parse_cosmic_genotype_data(cosmic_file, ref_gen = "GRCH37")
```

Arguments

cosmic_file	Path to cosmic clp file in hard disk
ref_gen	Reference genome version

Value

The R Table sim_list which contains the CoSMIC CLP fingerprints

```
parse_vcf_file      Filter Parsed VCF Files
```

Description

Intern utility function. Filters the parsed VCF file for all informations except for the start and length of variations/mutations.

Usage

```
parse_vcf_file(
  vcf_file,
  ref_gen,
  library_name
)
```

Arguments

vcf_file	character string giving the path to the vcf file on the operating system.
ref_gen	Reference genome version
library_name	Name of the reference library

Value

Loci-based DNA-mutational fingerprint of the cancer cell line as found in the input VCF file.

parse_vcf_query_into_db

parse_vcf_query_into_db This function adds the variants of parsed custom CCLs to a monet DB instance

Description

parse_vcf_query_into_db This function adds the variants of parsed custom CCLs to a monet DB instance

Usage

```
parse_vcf_query_into_db(g_query, ref_gen = "GRCH37", library_name,
                        test_mode = FALSE)
```

Arguments

g_query	a GenomicRanges object
ref_gen	a character string specifying the reference genome version. All training sets are associated with a reference genome version. Default is "GRCH37".
library_name	a character string giving the name of the library to add the cancer cell lines to. Default is "CUSTOM". Library name will be automatically added as a suffix to the identifier.
test_mode	Is this a test? Just for internal use

Value

Message wheather the adding was successful

read_library_names *Library Name Reader*

Description

This function procides information on the reference library names

Usage

```
read_library_names(ref_gen)
```

Arguments

ref_gen	a character vector specifying the reference genome version. All training sets are associated with a reference genome version. Default is "GRCH37".
---------	----------------------------------------------------------------------------------------------------------------------------------------------------

Arguments

ccl_names	A character vector giving the names of the cancer cell line identifiers to be removed. Can be one or many
ref_gen	A character vector specifying the reference genome version. All training sets are associated with a reference genome version. Default is "GRCH37".
library_name	Name of the library from which ccls are to be removed
test_mode	Signifies if this is a test run

Value

Message that indicates whether the removal was succesful.

Examples

```
remove_ccls_from_database(
  ccl_names = "HT29",
  ref_gen = "GRCH37",
  library_name = "CELLMINER",
  test_mode = TRUE
)
```

```
remove_library_from_database
```

Remove entire Library from Database

Description

This function removes a entire library from the database by removing all associated cancer cell line fingerprints from the database.

Usage

```
remove_library_from_database(library, ref_gen = "GRCH37", test_mode = FALSE)
```

Arguments

library	a character vector giving the names of the library to be removed.
ref_gen	a character vector specifying the reference genome version. All training sets are associated with a reference genome version. Default is "GRCH37".
test_mode	is this a test? Just for internal use.

Value

Message that indicates whether the removal was succesful.

Examples

```
remove_library_from_database(library = "CELLMINER",
                             ref_gen = "GRCH37",
                             test_mode = TRUE)
```

```
show_contained_ccls    show_contained_ccls
```

Description

This function displays the names, amount of mutations and the overall weight of the mutations of all contained cancer cell line fingerprints for a chosen reference genome and optional library.

Usage

```
show_contained_ccls(ref_gen, verbose)
```

Arguments

ref_gen	a character vector specifying the reference genome version. All training sets are associated with a reference genome version. Default is "GRCH37".
verbose	Should DB informations be printed?

Value

R table which contains identifiers of all cancer cell line samples which match the specified parameters (reference genome and library).

Examples

```
## Show all contained cancer cell lines for reference GRCH37:
show_contained_ccls(ref_gen = "GRCH37", verbose = TRUE)
```

```
show_contained_variants_for_ccl
      Variants In Cancer Cell Line
```

Description

This function shows all mutations present in the database for a selected cancer cell line and reference genome.

Usage

```
show_contained_variants_for_ccl(
  name_ccl,
  ref_gen,
  library_name,
  mutational_weight_inclusion_threshold
)
```

Arguments

name_ccl	a character vector giving the identifier of the cancer cell line for which mutations will be shown.
ref_gen	a character vector specifying the reference genome version. All training sets are associated with a reference genome version. Default is "GRCH37".
library_name	Name of the reference library
mutational_weight_inclusion_threshold	Include only mutations with a weight of at least x. Range: 0.0 to 1.0. 1= unique to CL. ~0 = found in many CCL samples.

Value

GenomicRanges object that contains the ccl's variants

Examples

```
## Show all mutations for Cancer Cell Line 'SK_OV_3'
show_contained_variants_for_ccl(
  name_ccl = "SK_OV_3",
  ref_gen = "GRCH37",
  library_name = "CELLMINER",
  mutational_weight_inclusion_threshold = 0
)
```

```
show_contained_variants_in_library
```

All variants contained in reference library

Description

This function shows all variants contained in a reference library for a given inclusion weight. Default inclusion weight is 0 (all variants).

Usage

```
show_contained_variants_in_library(
  ref_gen,
  library_name,
  mutational_weight_inclusion_threshold
)
```


Arguments

ref_gen a character vector specifying the reference genome version. All training sets are associated with a reference genome version. Default is "GRCH37".

library_name Name of the reference library.

mutational_weight_inclusion_threshold Include only mutations with a weight of at least x. Range: 0.0 to 1.0. 1 = unique to CL. ~0 = found in many CL samples.

Value

Returns a GenomicRanges object that contains the variants

Examples

```
## Show all variants contained in reference library CELLMINER
show_contained_variants_in_library(
  ref_gen = "GRCH37",
  library_name = "CELLMINER",
  mutational_weight_inclusion_threshold = 0
)
```

show_which_ccls_contain_variant

Cancer cell lines with specific variant

Description

This function displays all cancer cell lines in the database which contain a specified variant. Utilizes closed interval coordinates.

Usage

```
show_which_ccls_contain_variant(
  start,
  end,
  chromosome,
  ref_gen,
  library_name,
  mutational_weight_inclusion_threshold
)
```

Arguments

start Start coordinate

end Stop coordinate

chromosome Chromosome, 'chr' prefixes are ignored

`ref_gen` a character vector specifying the reference genome version. All training sets are associated with a reference genome version. Default is "GRCH37".

`library_name` Name of the reference library

`mutational_weight_inclusion_threshold`
Include only mutations with a weight of at least x. Range: 0.0 to 1.0. 1= unique to CL. ~0 = found in many CCL samples.

Value

Returns a `GenomicRanges` object that contains the variant if present. Member `ccls` can be found in the `$Member_ccl` vector

Examples

```
show_which_ccls_contain_variant(  
  start = 92030762,  
  end = 92030762,  
  chromosome = 8,  
  ref_gen = "GRCH37",  
  library_name = "CELLMINER",  
  mutational_weight_inclusion_threshold = 0  
)
```

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