Package 'seqCAT'

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Title High Throughput Sequencing Cell Authentication Toolkit

Version 1.0.0

Description The seqCAT package uses variant calling data (in the form of VCF files) from high throughput sequencing technologies to authenticate and validate the source, function and characteristics of biological samples used in scientific endeavours.

Depends R (>= 3.4), GenomicRanges (>= 1.26.4), VariantAnnotation(>= 1.20.3)

Imports dplyr (>= 0.5.0), GenomeInfoDb (>= 1.13.4), ggplot2 (>= 2.2.1), IRanges (>= 2.8.2), lazyeval (>= 0.2.0), scales (>= 0.4.1), S4Vectors (>= 0.12.2), stats, SummarizedExperiment (>= 1.4.0), tidyr (>= 0.6.1), utils

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R topics documented:

`calculate_similarity` *SNV profile similarity calculations*

Description

Calculate the similarity statistics for SNV profile comparisons.

Usage

```
calculate_similarity(overlaps, similarity = NULL, a = 1, b = 5)
```

Arguments

`overlaps`	The input SNV overlaps dataframe.
`similarity`	Optional dataframe to add results to.
`a`	Similarity score parameter a (integer).
`b`	Similarity score parameter b (integer).

Details

This function calculates various summary statistics and sample similarities for a given profile comparison dataframe. It returns a small dataframe with the overall similarity score (whose parameters ‘a‘ and ‘b‘ can be adjusted in the function call), total SNV overlaps, the concordance of the overlaps and the sample names in question. This dataframe can also be given to the function, in which case it will simply add another row for the current samples, facilitating downstream aggregate analyses.

Value

A dataframe with summary statistics.

Examples

```
# Load test data
data(test_comparison)

# Calculate similarities
similarity <- calculate_similarity(test_comparison)

# Add another row of summary statistics
calculate_similarity(test_comparison, similarity = similarity)
```

compare_many *Comparisons of many SNV profiles*

Description

Overlap and compare genotypes in many SNV profiles.

Usage

```
compare_many(many, one = NULL, a = 1, b = 5)
```

Arguments

many	SNV profiles to be compared (list of GRanges objects).
one	SNV profile to be compared to all others (GRanges object).
a	Similarity score parameter a (integer).
b	Similarity score parameter b (integer).

Details

This is a function that compares all the combinations of the SNV profiles input to it, either in a one-to-many or many-to-many manner. It returns both a dataframe containing summary statistics for all unique combinations and a list of dataframes with all the performed comparisons, for easy re-use and downstream analyses of said comparisons.

Value

A list of summary statistics and comparisons.

Examples

```
# Load test data
data(test_profile_1)
data(test_profile_2)

# Perform many-to-many comparisons
profiles <- list(test_profile_1, test_profile_2)
comparisons <- compare_many(profiles)

# View aggregate similarities
## Not run: comparisons[[1]])

# View data of first comparison
## Not run: head(comparisons[[2]][[1]])
```

compare_profiles *Binary SNV profile comparisons*

Description

Overlap and compare genotypes in two SNV profiles.

Usage

```
compare_profiles(profile_1, profile_2)
```

Arguments

profile_1	The first SNV profile (GRanges object).
profile_2	The second SNV profile (GRanges object).

Details

This is a function for finding overlapping variants in two different SNV profiles (stored as GenomicRanges objects), followed by comparing the genotypes of the overlapping variants. The "compare_overlaps" function calls the "add_metadata" function twice in succession in order to merge the metadata for the two profiles (supplied as GRanges objects), returns the results as a dataframe, compares the genotypes of the overlapping variants using the "compare_genotypes" function and, finally, returns the final dataframe with all variant overlaps and their similarity.

Value

A dataframe.

Examples

```
# Load test data
data(test_profile_1)
data(test_profile_2)

# Compare the two profiles
comparison <- compare_profiles(test_profile_1, test_profile_2)
```

create_profile *SNV profile creation*

Description

Create an SNV profile from data in a VCF file.

Usage

```
create_profile(vcf_file, sample, output_file, filter_depth = 10,
  python = FALSE)
```

Arguments

vcf_file	The VCF file from which the profile will be created (path).
sample	The sample in the VCF for which a profile will be created (character).
output_file	The output file with the SNV profile (path).
filter_depth	Remove variants below this sequencing depth (integer).
python	Extract variants using Python instead of R (boolean).

Details

This function creates a SNV profile from a given VCF file by extracting the variants that pass the filtering criterias. It can either be performed using R, or by the create_profile.py function included (which requires that Python is installed, along with the PyVCF package). Profile creation is performed to facilitate and accelerate the cell authentication procedures, which is especially relevant when more than one pairwise comparison will be performed on the same sample.

Value

Does not return any data object, but outputs results to output_file (to save computational time from having to repeatedly create profiles).

Examples

```
# Path to the test VCF file
vcf_file = system.file("extdata", "test.vcf.gz", package = "seqCAT")

# Create SNV profiles
## Not run:
 create_profile(vcf_file, "sample1", "profile1.txt")
 create_profile(vcf_file, "sample1", "profile1.txt", filter_depth = 15)
 create_profile(vcf_file, "sample1", "profile1.txt", python = TRUE)

## End(Not run)
```

---

filter_variants *Variant filtering*

Description

Filter variants on sequencing depth.

Usage

```
filter_variants(overlaps, filter_depth = 10)
```

Arguments

overlaps	The dataframe containing the variant data to be filtered.
filter_depth	Threshold for variant depth (integer; default 10).

Details

This is a function for filtering variants on sequencing depth. Variants with a depth lower than 10 are removed by default, but can be changed in the function call.

Value

A data frame containing the filtered variants.

Examples

```
# Load test comparisons
data(test_comparison)

# Filter variants
filt_1 <- filter_variants(test_comparison)
filt_2 <- filter_variants(test_comparison, filter_depth = 20)
```

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plot_heatmap *Plot similarity heatmap*

Description

Plot a heatmap of similarities from many-to-many SNV profile comparisons.

Usage

```
plot_heatmap(similarities, annotate = TRUE, annotate_size = 5,
  legend = TRUE, cluster = TRUE, limits = c(0, 50, 90, 100),
  colour = "#1954A6")
```

Arguments

similarities	The long-format dataframe containing the data.
annotate	Annotate each cell with the score (boolean).
annotate_size	The size of the annotations (numeric).
legend	Show a legend for the colour gradient (boolean).
cluster	Cluster the samples based on similarity (boolean).
limits	The limits for the colour gradient (vector of four integers).
colour	The main colour to use for the gradient (character).

Details

This function creates publication-ready plots of heatmaps for many-to-many sample comparisons, taking a long-format dataframe containing the summary statistics of each comparison as input.

Value

A ggplot2 graphical object.

Examples

```
# Load test similarities
data(test_similarities)

# Plot a similarity heatmap
heatmap <- plot_heatmap(test_similarities)
```

`plot_impacts` *Plot SNV impact distribution*

Description

Plot SNV impact distributions for a binary SNV profile comparison.

Usage

```
plot_impacts(comparison, annotate = TRUE, legend = TRUE,
  palette = c("#0D2D59", "#1954A6"))
```

Arguments

`comparison`	The SNV profile comparison to be plotted.
`annotate`	Annotate each category (boolean).
`legend`	Show the legend (boolean).
`palette`	Colour palette for filling of bars (character vector).

Details

This function creates publication-ready plots of the impact distribution from a binary dataset comparison across the matched/mismatched SNVs.

Value

A ggplot2 graphical object.

Examples

```
# Load test comparison data
data(test_comparison)

# Plot the impact distribution
impacts <- plot_impacts(test_comparison)
```

`read_cosmic` *Read COSMIC SNV data*

Description

Read COSMIC cell line-specific mutational data.

This function lists the available cell lines in the provided CosmicCLP_MutantExport.tsv.gz file, and take about half the time it takes to read the full file with the read_cosmic function, making it useful for just seeing if your particular cell line is listed in COSMIC or not.

Usage

```
read_cosmic(file_path, cell_line)

list_cosmic(file_path)
```

Arguments

`file_path`	The CosmicCLP_MutantExport.tsv.gz file (path).
`cell_line`	The cell line to be investigated (character).

Details

This function reads the "CosmicCLP_MutantExport.tsv.gz" file obtained from COSMIC and returns a GRanges object with all the listed mutations for the specified cell line, which can then be use in downstream profile comparisons. Only non-duplicated (gene-level) SNVs are included in COSMIC profiles.

Value

A GRanges object with COSMIC SNVs.

A vector of cell line names

Examples

```
# Path to COSMIC test data
file <- system.file("extdata",
                    "subset_CosmicCLP_MutantExport.tsv.gz",
                    package = "seqCAT")

# Read COSMIC test data for HCT116 cell line
cosmic_hct116 <- read_cosmic(file, "HCT116")
file <- system.file("extdata",
                    "subset_CosmicCLP_MutantExport.tsv.gz",
                    package = "seqCAT")
cell_lines <- list_cosmic(file)
```

`read_profile` *Read SNV profile*

Description

Read SNV profiles for use in downstream comparisons.

Usage

```
read_profile(file, sample_name)
```

Arguments

`file`	The SNV profile to be read (path).
`sample_name`	The sample of the SNV profile (character).

Details

This is a function for reading SNV profiles created from VCF files. The data is returned as a GenomicRanges object, suitable for merging of metadata.

Value

A GRanges object.

Examples

```
# Path to test data
profile = system.file("extdata",
                      "test_profile_1.txt.gz",
                      package = "seqCAT")

# Read test profile
profile_1 <- read_profile(profile, "sample1")
```

`seqCAT` *seqCAT: High Throughput Sequencing Cell Authentication Toolkit*

Description

The *seqCAT* package provides a number of functions for performing evaluation, characterisation and authentication of biological samples through analysis of high throughput sequencing data.

`test_comparison` *Overlapping and compared SNVs*

Description

Overlapping and compared variants from "sample1" and "sample2" originating from the example.vcf file included in the inst/extdata directory, for use in unit tests.

Usage

```
data(test_comparison)
```

Format

A dataframe with 51 rows and 39 columns:

chr chromosome

pos SNV position

DP.sample_1 total variant depth, sample 1

AD1.sample_1 allelic depth, allele 1, sample 1

AD2.sample_1 allelic depth, allele 2, sample 1

A1.sample_1 allele 1, sample 1

A2.sample_1 allele 2, sample 1

warnings.sample_1 warnings from variant calling, sample 1

DP.sample_2 total variant depth, sample 2

AD1.sample_2 allelic depth, allele 1, sample 2

AD2.sample_2 allelic depth, allele 2, sample 2

A1.sample_2 allele 1, sample 2

A2.sample_2 allele 2, sample 2

warnings.sample_2 warnings from variant calling, sample 2

sample_1 name, sample 1

sample_2 name, sample 2

match status of genotype comparison

rsID mutation ID

gene associated gene

ENSGID ensembl gene ID

ENSTID ensembl transcript ID

REF reference allele

ALT alternative allele

impact putative variant impact

effect variant effect

feature transcript feature

biotype transcript biotype

`test_profile_1` *SNV profile 1*

Description

SNV profile in GRanges format from "sample1", originating from the test_profile_1.txt in the inst/extdata directory, for use in unit tests.

Usage

```
data(test_profile_1)
```

Format

A GRanges object with 383 elements and 17 metadata columns:

rsID mutation ID, if available

gene associated gene

ENSGID ensembl gene ID

ENSTID ensembl transcript ID

REF reference allele

ALT alternative allele

impact putative variant impact

effect variant effect

feature transcript feature

biotype transcript biotype

DP total variant depth

AD1 allelic depth, allele 1

AD2 allelic depth, allele 2

A1 allele 1

A2 allele 2

warnings warnings from variant calling

sample sample name

`test_profile_2` *SNV profile 2*

Description

SNV profile in GRanges format from "sample2", originating from the test_profile_2.txt in the inst/extdata directory, for use in unit tests.

Usage

```
data(test_profile_2)
```

Format

A GRanges object with 382 elements and 17 metadata columns:

rsID mutation ID, if available

gene associated gene

ENSGID ensembl gene ID

ENSTID ensembl transcript ID

REF reference allele

ALT alternative allele

impact putative variant impact

effect variant effect

feature transcript feature

biotype transcript biotype

DP total variant depth

AD1 allelic depth, allele 1

AD2 allelic depth, allele 2

A1 allele 1

A2 allele 2

warnings warnings from variant calling

sample sample name

`test_similarities` *Collated similarities object*

Description

Collated similarities of multiple sample comparisons from "sample1" and "sample" from the example.vcf file, for use in unit tests.

Usage

```
data(test_similarities)
```

Format

A dataframe with 3 rows and 6 columns:

sample_1 name of sample 1

sample_2 name of sample 2

overlaps the number of overlaps for the comparison

matches the number of matches for the comparison

concordance the concordance of the profiles

similarity_score the similarity score of the profiles

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