

Package ‘vDiveR’

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Type Package

Title Visualization of Viral Protein Sequence Diversity Dynamics

Version 2.1.0

Description To ease the visualization of outputs from Diversity Motif Analyser ('DiMA'; <https://github.com/BVU-BILSAB/DiMA>). 'vDiveR' allows visualization of the diversity motifs (index and its variants – major, minor and unique) for elucidation of the underlying inherent dynamics. Please refer <https://vddiver-manual.readthedocs.io/en/latest/> for more information.

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concat_conserved_kmer	<i>k-mer sequences concatenation</i>
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Description

This function concatenates completely (index incidence = 100%) only or both both the completely and highly (90% <= index incidence < 100%) conserved k-mer positions that overlapped at least one k-mer position or are adjacent to each other and generate the CCS/HCS sequence in either CSV or FASTA format

Usage

```
concat_conserved_kmer(  
  data,  
  conservation_level = "HCS",  
  kmer = 9,  
  threshold_pct = NULL  
)
```

Arguments

- data DiMA JSON converted csv file data
- conservation_level CCS (completely conserved) / HCS (highly conserved)
- kmer size of the k-mer window
- threshold_pct manually set threshold of index.incidence for HCS

Value

A list with csv and fasta dataframes

Examples

```
csv<-concat_conserved_kmer(proteins_1host)$csv
csv_2hosts<-concat_conserved_kmer(protein_2hosts, conservation_level = "CCS")$csv
fasta <- concat_conserved_kmer(protein_2hosts, conservation_level = "HCS")$fasta
```

extract_from_GISAID	<i>Extract metadata via fasta file from GISAID</i>
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Description

This function gets the metadata from each header of GISAID fasta file

Usage

```
extract_from_GISAID(file_path)
```

Arguments

file_path	path of fasta file
-----------	--------------------

extract_from_NCBI	<i>Extract metadata via fasta file from ncbi</i>
-------------------	--

Description

This function gets the metadata from each head of fasta file

Usage

```
extract_from_NCBI(file_path)
```

Arguments

file_path	path of fasta file
-----------	--------------------

 json2csv

JSON2CSV

Description

This function converts DiMA (v5.0.9) JSON output file to a dataframe with 17 predefined columns which further acts as the input for other functions provided in this vDiveR package.

Usage

```
json2csv(
  json_data,
  host_name = "unknown host",
  protein_name = "unknown protein"
)
```

Arguments

json_data	DiMA JSON output dataframe
host_name	name of the host species
protein_name	name of the protein

Value

A dataframe which acts as input for the other functions in vDiveR package

Examples

```
inputdf<-json2csv(JSON_sample)
```

 JSON_sample

DiMA (v5.0.9) JSON Output File

Description

A sample DiMA JSON Output File which acts as the input for JSON2CSV()

Usage

```
JSON_sample
```

Format

A Diversity Motif Analyzer (DiMA) tool JSON file

`match_region_to_target`*Map data region to ggplot2 region*

Description

This function maps the provided data region to ggplot2 region.

Usage

```
match_region_to_target(region_column, region_list, max_dist = 3)
```

Arguments

<code>region_column</code>	a character vector of data region
<code>region_list</code>	a character vector of ggplot2 region
<code>max_dist</code>	maximum distance for string matching

`metadata`*Metadata Input Sample*

Description

A dummy dataset that acts as an input for `plot_world_map()` and `plot_time()`

Usage

```
metadata
```

Format

A data frame with 1000 rows and 3 variables:

ID unique identifier of the sequence

region geographical region of the sequence collection

date collection date of the sequence

metadata_extraction	<i>Metadata Extraction from NCBI/GISAID (Epi-Flu/EpiCoV/EpiPox/EpiArbo) FASTA file</i>
---------------------	--

Description

This function retrieves metadata (ID, region, date) from the input FASTA file, with the source of, either NCBI (with default FASTA header) or GISAID (with default FASTA header). The function will return a dataframe that has three columns consisting ID, collected region and collected date. Records that do not have region or date information will be excluded from the output dataframe.

Usage

```
metadata_extraction(file_path, source)
```

Arguments

- file_path path of fasta file
- source the source of fasta file, either "NCBI" or "GISAID"

Value

A dataframe that has three columns consisting ID, collected region and collected date

Examples

```
filepath <- system.file('extdata','GISAID_EpiCoV.faa', package = 'vDiver')
meta_gisaid <- metadata_extraction(filepath, 'GISAID')
```

plot_conservation_level	<i>Conservation Levels Distribution Plot</i>
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Description

This function plots conservation levels distribution of k-mer positions, which consists of completely conserved (black) (index incidence = 100%), highly conserved (blue) (90% <= index incidence < 100%), mixed variable (green) (20% < index incidence <= 90%), highly diverse (purple) (10% < index incidence <= 20%) and extremely diverse (pink) (index incidence <= 10%).

Usage

```
plot_conservation_level(
  df,
  protein_order = NULL,
  conservation_label = 1,
  host = 1,
  base_size = 11,
  line_dot_size = 2,
  label_size = 2.6,
  alpha = 0.6
)
```

Arguments

df	DiMA JSON converted csv file data
protein_order	order of proteins displayed in plot
conservation_label	0 (partial; show present conservation labels only) or 1 (full; show ALL conservation labels) in plot
host	number of host (1/2)
base_size	base font size in plot
line_dot_size	lines and dots size
label_size	conservation labels font size
alpha	any number from 0 (transparent) to 1 (opaque)

Value

A plot

Examples

```
plot_conservation_level(proteins_1host, conservation_label = 1, alpha=0.8, base_size = 15)
plot_conservation_level(protein_2hosts, conservation_label = 0, host=2)
```

plot_correlation	<i>Entropy and total variant incidence correlation plot</i>
------------------	---

Description

This function plots the correlation between entropy and total variant incidence of all the provided protein(s).

Usage

```
plot_correlation(
  df,
  host = 1,
  alpha = 1/3,
  line_dot_size = 3,
  base_size = 11,
  ylabel = "k-mer entropy (bits)\n",
  xlabel = "\nTotal variants (%)",
  ymax = ceiling(max(df$entropy)),
  ybreak = 0.5
)
```

Arguments

df	DiMA JSON converted csv file data
host	number of host (1/2)
alpha	any number from 0 (transparent) to 1 (opaque)
line_dot_size	dot size in scatter plot
base_size	base font size in plot
ylabel	y-axis label
xlabel	x-axis label
ymax	maximum y-axis
ybreak	y-axis breaks

Value

A scatter plot

Examples

```
plot_correlation(proteins_1host)
plot_correlation(protein_2hosts, base_size = 2, ybreak=1, ymax=10, host = 2)
```

plot_dynamics_protein *Dynamics of Diversity Motifs (Protein) Plot*

Description

This function compactly display the dynamics of diversity motifs (index and its variants: major, minor and unique) in the form of dot plot(s) as well as violin plots for all the provided individual protein(s).

Usage

```
plot_dynamics_protein(
  df,
  host = 1,
  protein_order = NULL,
  base_size = 8,
  alpha = 1/3,
  line_dot_size = 3,
  bw = "nrd0",
  adjust = 1
)
```

Arguments

df	DiMA JSON converted csv file data
host	number of host (1/2)
protein_order	order of proteins displayed in plot
base_size	base font size in plot
alpha	any number from 0 (transparent) to 1 (opaque)
line_dot_size	dot size in scatter plot
bw	smoothing bandwidth of violin plot (default: nrd0)
adjust	adjust the width of violin plot (default: 1)

Value

A plot

Examples

```
plot_dynamics_protein(proteins_1host)
```

plot_dynamics_proteome

Dynamics of Diversity Motifs (Proteome) Plot

Description

This function compactly display the dynamics of diversity motifs (index and its variants: major, minor and unique) in the form of dot plot as well as violin plot for all the provided proteins at proteome level.

Usage

```
plot_dynamics_proteome(  
  df,  
  host = 1,  
  line_dot_size = 2,  
  base_size = 10,  
  alpha = 1/3,  
  bw = "nrd0",  
  adjust = 1  
)
```

Arguments

df	DiMA JSON converted csv file data
host	number of host (1/2)
line_dot_size	size of dot in plot
base_size	word size in plot
alpha	any number from 0 (transparent) to 1 (opaque)
bw	smoothing bandwidth of violin plot (default: nrd0)
adjust	adjust the width of violin plot (default: 1)

Value

A plot

Examples

```
plot_dynamics_proteome(proteins_1host)
```

plot_entropy	<i>Entropy plot</i>
--------------	---------------------

Description

This function plot entropy (black) and total variant (red) incidence of each k-mer position across the studied proteins and highlight region(s) with zero entropy in yellow. k-mer position with low support is marked with a red triangle underneath the x-axis line.

Usage

```
plot_entropy(  
  df,  
  host = 1,  
  protein_order = "",  
  kmer_size = 9,
```

```

    ymax = 10,
    line_size = 2,
    base_size = 8,
    all = TRUE,
    highlight_zero_entropy = TRUE
)
```

Arguments

df	DiMA JSON converted csv file data
host	number of host (1/2)
protein_order	order of proteins displayed in plot
kmer_size	size of the k-mer window
ymax	maximum y-axis
line_size	size of the horizontal (reference) line in plot
base_size	word size in plot
all	plot both the entropy and total variants (pass FALSE in to plot only the entropy)
highlight_zero_entropy	highlight region with zero entropy (default: TRUE)

Value

A plot

Examples

```

plot_entropy(proteins_1host)
plot_entropy(protein_2hosts, host = 2)
```

plot_time	<i>Time Distribution of Sequences Plot</i>
-----------	--

Description

This function plots the time distribution of provided sequences in the form of bar plot with 'Month' as x-axis and 'Number of Sequences' as y-axis. Aside from the plot, this function also returns a dataframe with 2 columns: 'Date' and 'Number of sequences'. The input dataframe of this function is obtainable from `metadata_extraction()`, with NCBI Protein / GISAID (EpiFlu/EpiCoV/EpiPox/EpiArbo) FASTA file as input.

Usage

```
plot_time(
  metadata,
  date_format = "%Y-%m-%d",
  base_size = 8,
  date_break = "2 month",
  scale = "count"
)
```

Arguments

metadata	a dataframe with 3 columns, 'ID', 'region', and 'date'
date_format	date format of the input dataframe
base_size	word size in plot
date_break	date break for the scale_x_date
scale	plot counts or log scale the data

Value

A single plot or a list with 2 elements (a plot followed by a dataframe, default)

Examples

```
time_plot <- plot_time(metadata, date_format="%d/%m/%Y")$plot
time_df <- plot_time(metadata, date_format="%d/%m/%Y")$df
```

plot_world_map	<i>Geographical Distribution of Sequences Plot</i>
----------------	--

Description

This function plots a world map and color the affected geographical region(s) from light (lower) to dark (higher), depends on the cumulative number of sequences. Aside from the plot, this function also returns a dataframe with 2 columns: 'Region' and 'Number of Sequences'. The input dataframe of this function is obtainable from `metadata_extraction()`, with NCBI Protein / GISAID (EpiFlu/EpiCoV/EpiPox/EpiArbo) FASTA file as input.

Usage

```
plot_world_map(metadata, base_size = 8)
```

Arguments

metadata	a dataframe with 3 columns, 'ID', 'region', and 'date'
base_size	word size in plot

Value

A list with 2 elements (a plot followed by a dataframe)

Examples

```
geographical_plot <- plot_world_map(metadata)$plot
geographical_df <- plot_world_map(metadata)$df
```

proteins_1host

DiMA (v5.0.9) JSON converted-CSV Output Sample 1

Description

A dummy dataset with two proteins (A and B) from one host, human

Usage

```
proteins_1host
```

Format

A data frame with 806 rows and 17 variables:

proteinName name of the protein

position starting position of the aligned, overlapping k-mer window

count number of k-mer sequences at the given position

lowSupport k-mer position with sequences lesser than the minimum support threshold (TRUE) are considered of low support, in terms of sample size

entropy level of variability at the k-mer position, with zero representing completely conserved

indexSequence the predominant sequence (index motif) at the given k-mer position

index.incidence the fraction (in percentage) of the index sequences at the k-mer position

major.incidence the fraction (in percentage) of the major sequence (the predominant variant to the index) at the k-mer position

minor.incidence the fraction (in percentage) of minor sequences (of frequency lesser than the major variant, but not singletons) at the k-mer position

unique.incidence the fraction (in percentage) of unique sequences (singletons, observed only once) at the k-mer position

totalVariants.incidence the fraction (in percentage) of sequences at the k-mer position that are variants to the index (includes: major, minor and unique variants)

distinctVariant.incidence incidence of the distinct k-mer peptides at the k-mer position

multiIndex presence of more than one index sequence of equal incidence

host species name of the organism host to the virus

highestEntropy.position k-mer position that has the highest entropy value

highestEntropy highest entropy values observed in the studied protein

averageEntropy average entropy values across all the k-mer positions

protein_2hosts

DiMA (v5.0.9) JSON converted-CSV Output Sample 2

Description

A dummy dataset with 1 protein (Core) from two hosts, human and bat

Usage

protein_2hosts

Format

A data frame with 200 rows and 17 variables:

proteinName name of the protein

position starting position of the aligned, overlapping k-mer window

count number of k-mer sequences at the given position

lowSupport k-mer position with sequences lesser than the minimum support threshold (TRUE) are considered of low support, in terms of sample size

entropy level of variability at the k-mer position, with zero representing completely conserved

indexSequence the predominant sequence (index motif) at the given k-mer position

index.incidence the fraction (in percentage) of the index sequences at the k-mer position

major.incidence the fraction (in percentage) of the major sequence (the predominant variant to the index) at the k-mer position

minor.incidence the fraction (in percentage) of minor sequences (of frequency lesser than the major variant, but not singletons) at the k-mer position

unique.incidence the fraction (in percentage) of unique sequences (singletons, observed only once) at the k-mer position

totalVariants.incidence the fraction (in percentage) of sequences at the k-mer position that are variants to the index (includes: major, minor and unique variants)

distinctVariant.incidence incidence of the distinct k-mer peptides at the k-mer position

multiIndex presence of more than one index sequence of equal incidence

host species name of the organism host to the virus

highestEntropy.position k-mer position that has the highest entropy value

highestEntropy highest entropy values observed in the studied protein

averageEntropy average entropy values across all the k-mer positions

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