Package ‘DominoEffect’

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Title Identification and Annotation of Protein Hotspot Residues
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Description The functions support identification and annotation of hotspot residues in proteins. These are individual amino acids that accumulate mutations at a much higher rate than their surrounding regions.
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align_to_unip

Align protein segment around the hotspot to the UniProt/Swiss-Prot KB sequence.

Description

This function alignes the Ensembl protein region with a hotspot to the UniProt sequence. The Ensembl region encompasses 15 amino acids where the hotspot is in the middle. If the hotspot was at the protein start or end the region is still 15 amino acids long, but the hotspot position is shifted.

Usage

```r
align_to_unip(ens.seq, uni.seq, ensembl_mut_position)
```

Arguments

- `ens.seq`: AAString object with the Ensembl protein sequence corresponding to the representative transcript.
- `uni.seq`: AAString with the UniProt sequence for the identifier matching the Ensembl gene name.
- `ensembl_mut_position`: Numeric vector defining the hotspot position in the Ensembl sequence, i.e. in the ens.seq

Value

Returns a list where the first element is a character vector defining whether there was a significant alignment and the second element provides the hotspot position in the UniProt sequence.

Author(s)

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Examples

library(Biostrings)

ens.seq <- AAString("MDLSALREEVQVINAMQKILECPICLELIKEPVSTKCDHIFCKFCMLK")
uni.seq <- AAString("MDLSALRVEEVQVINAMQKILECPICLELIKEPVSTKCDHIFCKFCMLA")
ensembl_mut_position <- 25

align_to_unip(ens.seq, uni.seq, ensembl_mut_position)

descritpion

The function calculates boundaries of sequence windows around the mutation. It is possible to define up to two window lengths. If the mutation is close to the start or end of the protein, the region is extended into the other direction to keep the indicated size.

Usage

calculate_boundary(mut_pos_numeric, length_aa, flanking_region)

Arguments

mut_pos_numeric

Amino acid position of mutation

length_aa

Length of transcript in amino acids

flanking_region

Vector containing two flanking regions

Value

returns a list with the boundaries for the two regions

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Examples

calculate_boundary(250, 500, c(200, 300))
calculate_boundary(250, 500, 300)
DominoData  

Sample data

Description

Sample Data

Author(s)

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DominoEffect  

Identification of significant mutation hotspot residues.

Description

The function identifies individual amino acid residues, which accumulate a high fraction of the overall mutation load within a protein. After detecting mutation hotspots, the function obtains UniProt/Swiss-Prot KB functional and structural annotations for the affected protein regions and checks for the sequence agreement.

Usage

```R
DominoEffect(mutation_dataset, gene_data, snp_data, 
min_n_muts = 5, MAF_thresh = 0.01, 
flanking_region = c(200, 300), 
poisson.thr = 0.01, percentage.thr = 0.15, 
ratio.thr = 45, approach = "percentage", write_to_file = "NO", 
ens_release = "https://feb2023.archive.ensembl.org")
```

Arguments

- **mutation_dataset**: Object containing a table with the mutation data (e.g. TCGA point mutations mapped to protein level).
- **gene_data**: DominoData object containing information about Ensembl gene annotations: gene identifiers and representative transcript cDNA length.
- **snp_data**: Object containing a table with information on population SNPs.
- **min_n_muts**: Numeric vector defining a minimum number of mutations that need to occur at the same residue. Default: 5
- **MAF_thresh**: Numeric vector that defines Minor allele frequency threshold for considering reported mutations as population SNPs.
- **flanking_region**: Numeric vector that defines size of a window around the mutation that will be considered. Up to two window sizes are allowed.
**GPo_of_hotspots**

**poisson.thr**  
Numeric vector that defines a threshold for the adjusted p-value. Residues with an associated p-value that is lower than the defined value are reported. Default: 0.01

**percentage.thr**  
Number defining the fraction of mutations within the window that need to fall on a single residue in order for it to be classified as a hotspot. Default: 0.15

**ratio.thr**  
Number defining a requirement that a number of mutations on a single residue should exceed what would be expected by chance given a background mutation rate in the window (i.e. the surrounding region). Default: 45

**approach**  
Option to define selection criteria to use percentage.thr or ratio.thr as criterion for finding single residue mutation clusters. Options: "both", "percentage" or "ratio". Default: "percentage"

**write_to_file**  
Option if the identified and annotated hotspots should be written to a file (YES or NO). Default: NO

**ens_release**  
Release of ensembl to be used. Default: https://feb2023.archive.ensembl.org

**Value**

Results table

**Author(s)**

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**Examples**

```r
data("SnpData", package = "DominoEffect")
data("TestData", package = "DominoEffect")
data("DominoData", package = "DominoEffect")

hotspot_mutations <- DominoEffect(mutation_dataset = TestData,
gene_data = DominoData, snp_data = SnpData)
```

---

**GPo_of_hotspots**  
*Converts hotspot mutation table into a GPo object*

**Description**

This function converts the genomic information on hotspot mutations into a GPo object.

**Usage**

```r
GPo_of_hotspots(hotspot_mutations)
```
identify_hotspots

**Arguments**

- `hotspot_mutations`  
  Data frame with information on hotspot mutations generated by the DominoEffect package.

**Value**

GPo object that contains the genomic information on hotspot mutations.

**Author(s)**

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**Examples**

```r
data("SnpData", package = "DominoEffect")
data("TestData", package = "DominoEffect")
data("DominoData", package = "DominoEffect")

hotspot_mutations <- DominoEffect(mutation_dataset = TestData,
gene_data = DominoData, snp_data = SnpData)
GPo_of_hotspots(hotspot_mutations)
```

---

**identify_hotspots**  
Identify hotspots

**Description**

The function identify protein hotspot mutation residues

**Usage**

```r
identify_hotspots(mutation_dataset, gene_data,
snp_data, min_n_muts = 5, MAF_thres = 0.01, flanking_region = c(200, 300),
poisson.thr = 0.01, percentage.thr = 0.15, ratio.thr = 45, approach = "percentage")
```

**Arguments**

- `mutation_dataset`  
  Object containing a table with the mutation data (e.g. TCGA point mutations mapped to protein level).  
- `gene_data`  
  Data frame or Txdb object containing information about Ensembl gene annotations: gene identifiers and representative transcript cDNA length.  
- `snp_data`  
  Object containing a table or vcf object with information on population SNPs.  
- `min_n_muts`  
  Numeric vector defining a minimum number of mutations that need to occur at the same residue. Default: 5
import_txdb

**MAF_thresh**  Numeric vector that defines Minor allele frequency threshold for considering reported mutations as population SNPs.

**flanking_region**  Numeric vector that defines size of a window around the mutation that will be considered. Up to two window sizes are allowed.

**poisson.thr**  Numeric vector that defines a threshold for the adjusted p-value. Residues with an associated p-value that is lower than the defined value are reported. Default: 0.01

**percentage.thr**  Number defining the fraction of mutations within the window that need to fall on a single residue in order for it to be classified as a hotspot. Default: 0.15

**ratio.thr**  Number defining a requirement that a number of mutations on a single residue should exceed what would be expected by chance given a background mutation rate in the window (i.e. the surrounding region). Default: 45

**approach**  Option to define selection criteria to use percentage.thr or ratio.thr as criterion for finding single residue mutation clusters. Options: "both", "percentage" or "ratio". Default = "percentage"

**Value**

An object containing information on the significant hotspots, associated Gene and protein identifiers, number of mutations, percentage of mutations within defined windows that map to the same residue and associated p-values.

**Author(s)**

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**Examples**

data("SnpData", package = "DominoEffect")
data("TestData", package = "DominoEffect")
data("DominoData", package = "DominoEffect")
hotspot_mutations <- identify_hotspots(mutation_dataset = TestData, gene_data = DominoData, snp_data = SnpData)

---

**import_txdb**  *Imports txdb data and converts it into format required for DominoEffect package*

**Description**

This function imports txdb data and converts into a data frame used in the DominoEffect package only extracting the required information from the txdb object.

**Usage**

import_txdb(txdb_object)
map_to_func_elem

Arguments

  txdb_object  TxDB Object, e.g. from makeTxDbFromEnsembl

Value

  Data frame that can be used in DominoEffect package.

Author(s)

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Examples

#EnsTxDB <- makeTxDbFromEnsembl(organism="Homo sapiens", release=73, 
  #   server="ensembldb.ensembl.org")
#DominoData <- import_txdb(EnsTxDB)
#head(DominoData)

map_to_func_elem  Functional annotation of significant hotspot residues.

Description

  This function retrieves Uniprot annotations for the functional elements in the proteins with significant hotspots and overlaps and maps the hotspot residues to these.

Usage

  map_to_func_elem(hotspot_results, write_to_file = "NO", ens_release = "109")

Arguments

  hotspot_results  Object containing information about the hotspot residues identified using the function identify_hotspots().

  write_to_file  A character vector defining whether the resulting annotated hotspots should be saved in a file (YES or NO).

  ens_release  A character vector defining whether default gene annotations are used, i.e. Ensembl release 109, or if the gene_data correspond to a different Ensembl release. For the current Ensembl version this should be set to ens_release="www.ensembl.org". For the archive versions to the corresponding archive website.

Value

  Updated results file containing an additional columns with the information on the annotated functional and structural region within which the mutation is mapped.
Author(s)

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Examples

data("TestData", package = "DominoEffect")
data("DominoData", package = "DominoEffect")
data("SnpData", package = "DominoEffect")

hotspot_mutations <- identify_hotspots(TestData, DominoData, SnpData)
hotspot_mutations <- map_to_func_elem(hotspot_mutations,
write_to_file = "NO", ens_release = "109")

head(hotspot_mutations)
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