

# Package ‘rmimp’

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**Type** Package

**Title** Predicting the impact of mutations on kinase-substrate phosphorylation

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**Description** No description

**License** LGPL

## R topics documented:

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| mimp | <i>Predict the impact of single variants on phosphorylation.</i> |
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## Description

This function takes in mutation, sequence and phosphorylation data to predict the impact the mutation has on phosphorylation.

## Usage

```
mimp(muts, seqs, psites, perc.bg = 90, perc.fg = 10, thresh.log2 = 0,  
      display.results = T, include.cent = F, model.data = "hconf")
```

## Arguments

**muts** Mutation data file: a space delimited text file OR data frame containing two columns (1) gene and (1) mutation. Example:

```
TP53      R282W  
CTNNB1    S33C  
CTNNB1    S37F
```

|              |   |
|--------------|---|
| seqs         | Sequence data file containing protein sequences in FASTA format OR named list of sequences where each list element is the uppercase sequence and the name of each element is that of the protein. Example: list(TP53="ABCXYZ", CDK2="HJKEWR")   |
| psites       | Phosphorylation data file (optional): a space delimited text file OR data frame containing two columns (1) gene and (1) positions of phosphorylation sites. Example:<br><br><div style="text-align: center;"> TP53        280<br/> CTNNB1    29<br/> CTNNB1    44 </div>  |
| perc.bg      | Percentile value between 0 - 100. This value is used to compute a threshold, beta from the negative (background) distribution of scores. By default this is the 90th percentile of the background distribution of scores. Anything below the threshold is considered a negative hit.  |
| perc.fg      | Percentile value between 0 - 100. This value is used to compute a threshold, alpha from the positive (foreground) distribution of scores. By default this is the 10th percentile of the foreground distribution of scores. Anything above the threshold is considered a positive hit.   |
| thresh.log2  | Threshold for the absolute value of log ratio. Anything less than this value is discarded (default: 0).   |
| include.cent | If TRUE, gains and losses caused by mutation in the central STY residue are kept. Scores of peptides with a non-STY central residue is given a score of -1 (default: FALSE).  |
| model.data   | Name of specificity model data to use, can be "hconf" : individual experimental kinase specificity models used to scan for rewiring events. For experimental kinase specificity models, grouped by family, set to "hconf-fam". Both are considered high confidence. For lower confidence predicted specificity models, set to "lconf". NOTE: Predicted models are purely speculative and should be used with EXTREME CAUTION. |

## Value

The data is returned in a `data.frame` with the following columns:

|           |  |
|-----------|--|
| gene      | gene with the rewiring event   |
| mut       | mutation causing the rewiring event  |
| psite_pos | position of the central residue of the phosphosite   |
| mut_dist  | distance of the mutation from the central phosphosite  |
| wt        | sequence of the wildtype phosphosite (before the mutation)   |
| mt        | sequence of the mutated phosphosite (after the mutation)   |
| score_wt  | matrix similarity score of the wildtype phosphosite  |
| score_mt  | matrix similarity score of the mutated phosphosite   |
| log_ratio | Log2 ratio between mutant and wildtype scores. A high positive log ratio represents a high confidence gain-of-signaling event. A high negative log ratio represents a high confidence loss-of-signaling event. |
| pwm       | name of the kinase being rewired   |

|         |  |
|---------|--|
| pwm_fam | family/subfamily of kinase being rewired. If a kinase subfamily is available the family and subfamily will be separated by an underscore e.g. "DMPK_ROCK". If no subfamily is available, only the family is shown e.g. "GSK" |
| perc_wt | Percentile rank of the wt score  |
| perc_mt | Percentile rank of the mutant score  |

### Examples

```
# Get the path to example mutation data
mut.file = system.file("extdata", "mutation_data.txt", package = "rmimp")

# Get the path to example FASTA sequence data
seq.file = system.file("extdata", "sequence_data.txt", package = "rmimp")

# View the files in a text editor
browseURL(mut.file)
browseURL(seq.file)

# Run rewiring analysis
results = mimp(mut.file, seq.file, display.results=TRUE)

# Show head of results
head(results)
```

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|--------------|--|
| results2html | <i>Display MIMP results interactively in browser</i> |
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### Description

Display MIMP results interactively in browser

### Usage

```
results2html(x, max.rows = 5000)
```

### Arguments

|          |   |
|----------|---|
| x        | Data frame resulting from mimp call.                                    |
| max.rows | If data contains more rows than this value, results won't be displayed. |