

# Package ‘g3viz’

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**Type** Package

**Title** Interactively Visualize Genetic Mutation Data using a Lollipop-Diagram

**Version** 1.1.2

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**Description** R interface for 'g3-lollipop' JavaScript library.  
Visualize genetic mutation data using an interactive lollipop diagram in RStudio or your browser.

**License** MIT + file LICENSE

**Encoding** UTF-8

**LazyData** true

**Depends** R (>= 3.0.0)

**Imports** jsonlite, cgdSr, stringr, htmlwidgets

**Suggests** shiny (>= 1.0.0), knitr, rmarkdown, kableExtra

**URL** <https://github.com/G3viz/g3viz>

**BugReports** <https://github.com/G3viz/g3viz/issues>

**RoxygenNote** 6.1.1

**VignetteBuilder** knitr

**NeedsCompilation** no

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## R topics documented:

g3Lollipop . . . . .	2
g3Lollipop-shiny . . . . .	3
g3Lollipop.options . . . . .	4
g3Lollipop.theme . . . . .	7
getMutationsFromCbiportal . . . . .	8

guessMAFColumnName . . . . .	9
hgnc2pfam . . . . .	10
hgnc2pfam.df . . . . .	11
hgnc2uniprot . . . . .	11
mapMutationTypeToMutationClass . . . . .	12
mutation.table.df . . . . .	14
parseProteinChange . . . . .	14
readMAF . . . . .	15
uniprot2pfam . . . . .	16

<b>Index</b>	<b>17</b>
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g3Lollipop	<i>Render g3lollipop diagram for the given mutation data</i>
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---

## Description

Render g3lollipop diagram for the given mutation data

## Usage

```
g3Lollipop(mutation.dat, gene.symbol, uniprot.id = NA,
  gene.symbol.col = "Hugo_Symbol", aa.pos.col = "AA_Position",
  protein.change.col = c("Protein_Change", "HGVS_Short"),
  factor.col = "Mutation_Class", plot.options = g3Lollipop.options(),
  save.png.btn = TRUE, save.svg.btn = TRUE, btn.style = NA,
  output.filename = "output")
```

## Arguments

mutation.dat	Input genomic mutation data frame
gene.symbol	HGNC primary gene symbol
uniprot.id	UniProt ID, in case that the specified gene symbol links to multiple UniProt entries (isoforms). For example, <i>AKAP7</i> gene has two isoforms in <b>UniProt</b> , <b>O43687</b> and <b>Q9P0M2</b> .
gene.symbol.col	Column name of Hugo gene symbols (e.g., TP53). Default <i>Hugo_Symbol</i> .
aa.pos.col	Column name of the parsed amino-acid change position. Default <i>AA_Position</i> .
protein.change.col	Column name of protein change information (e.g., p.K960R, G658S, L14Sfs*15). Default is a list of <i>Protein_Change</i> , <i>HGVS_Short</i> .
factor.col	column of classes in the plot legend. IF NA, use parsed <i>Mutation_Class</i> column, otherwise, use specified. Default NA.
plot.options	g3lollipop diagram options in list format. Check <a href="#">g3Lollipop.options</a>
save.png.btn	If add <i>save-as-png</i> button to the diagram. Default TRUE.
save.svg.btn	If add <i>save-as-svg</i> button to the diagram. Default TRUE.

`btn.style` button style, including browser default button style, and two built-in styles, *blue* or *gray*. Default NA, indicating browser default.

`output.filename` Specify output file name.

## Examples

```
# system mutation data
maf.file <- system.file("extdata", "TCGA.BRCA.varscan.somatic.maf.gz", package = "g3viz")
# read in MAF file
mutation.dat <- readMAF(maf.file)

# use built-in chart theme
chart.options <- g3Lollipop.theme(theme.name = "default",
                                  title.text = "PIK3CA gene (default theme)")

# generate chart
g3Lollipop(mutation.dat,
            gene.symbol = "PIK3CA",
            plot.options = chart.options,
            btn.style = "blue",
            output.filename = "default_theme")
```

---

g3Lollipop-shiny      *Shiny bindings for g3Lollipop*

---

## Description

Output and render functions for using g3viz lollipop diagram within Shiny applications and interactive Rmd documents.

## Usage

```
g3LollipopOutput(outputId, width = "100%", height = "520px")

renderG3Lollipop(expr, env = parent.frame(), quoted = FALSE)
```

## Arguments

`outputId` output variable to read from

`width, height` Must be a valid CSS unit (like '100%', '400px', 'auto') or a number, which will be coerced to a string and have 'px' appended.

`expr` An expression that generates a g3-lollipop

`env` The environment in which to evaluate `expr`.

`quoted` Is `expr` a quoted expression (with `quote()`)? This is useful if you want to save an expression in a variable.

---

g3Lollipop.options      *G3Lollipop plot options*

---

## Description

G3Lollipop plot options

## Usage

```
g3Lollipop.options(chart.width = 800, chart.type = "circle",
  chart.margin = list(left = 40, right = 20, top = 15, bottom = 25),
  chart.background = "transparent", transition.time = 600,
  y.axis.label = "# of mutations",
  axis.label.font = "normal 12px Arial", axis.label.color = "#4f4f4f",
  axis.label.alignment = "middle", axis.label.dy = "-2em",
  y.axis.line.color = "#c4c8ca", y.axis.line.style = "dash",
  y.axis.line.width = 1, y.max.range.ratio = 1.1,
  legend.margin = list(left = 10, right = 0, top = 5, bottom = 5),
  legend.interactive = TRUE, legend.title = NA,
  lollipop.track.height = 420,
  lollipop.track.background = "rgb(233,233,233)",
  lollipop.pop.min.size = 2, lollipop.pop.max.size = 12,
  lollipop.pop.info.limit = 8, lollipop.pop.info.color = "#EEE",
  lollipop.pop.info.dy = "0.35em",
  lollipop.line.color = "rgb(42,42,42)", lollipop.line.width = 0.5,
  lollipop.circle.color = "wheat", lollipop.circle.width = 0.5,
  lollipop.label.ratio = 1.4, lollipop.label.min.font.size = 10,
  lollipop.color.scheme = "accent", highlight.text.angle = "90",
  title.text = "", title.font = "normal 16px Arial",
  title.color = "#424242", title.alignment = "middle",
  title.dy = "0.35em", anno.height = 30, anno.margin = list(top = 4,
  bottom = 0), anno.background = "transparent",
  anno.bar.fill = "#e5e3e1", anno.bar.margin = list(top = 2, bottom =
  2), domain.color.scheme = "category10", domain.margin = list(top = 0,
  bottom = 0), domain.text.font = "normal 11px Arial",
  domain.text.color = "#f2f2f2", brush = TRUE,
  brush.selection.background = "#666", brush.selection.opacity = 0.2,
  brush.border.color = "#969696", brush.handler.color = "#333",
  brush.border.width = 1, legend = TRUE, tooltip = TRUE,
  zoom = TRUE)
```

## Arguments

chart.width	chart width. Default 800.
chart.type	<i>pie</i> or <i>circle</i> . Default <i>circle</i> .
chart.margin	specify chart margin in <code>_list_</code> format. Default <code>list(left = 40, right = 20, top = 15, bottom = 25)</code> .

chart.background	chart background. Default <i>transparent</i> .
transition.time	animation transition time when clicking lollipop pops to show labels (in millisecond). Default 600.
y.axis.label	Y-axis label text. Default <i>"# of mutations"</i> .
axis.label.font	css font style shorthand ( <i>font-style font-variant font-weight font-size/line-height font-family</i> ). Default <i>"normal 12px Arial"</i> .
axis.label.color	axis label text color. Default <i>#4f4f4f</i> .
axis.label.alignment	axis label text alignment (start/end/middle). Default <i>middle</i> .
axis.label.dy	text adjustment of axis label text. Default <i>-2em</i> .
y.axis.line.color	color of y-axis in-chart lines (ticks). Default <i>#c4c8ca</i> .
y.axis.line.style	style of y-axis in-chart lines (ticks), "dash" or "line". Default <i>dash</i> .
y.axis.line.width	width of y-axis in-chart lines (ticks). Default 1.
y.max.range.ratio	ratio of y-axis range to data value range. Default 1.1.
legend.margin	legend margin in <i>list</i> . Default <code>list(left = 10, right = 0, top = 5, bottom = 5)</code> .
legend.interactive	legend interactive mode. Default TRUE.
legend.title	legend title. If NA, <i>factor.col</i> in <a href="#">g3Lollipop</a> is used. Default is NA.
lollipop.track.height	height of lollipop track. Default 420.
lollipop.track.background	background of lollipop track. Default <i>rgb(244,244,244)</i>
lollipop.pop.min.size	lollipop pop minimal size. Default 2.
lollipop.pop.max.size	lollipop pop maximal size. Default 12.
lollipop.pop.info.limit	threshold of lollipop pop size to show count information in middle of pop. Default 8.
lollipop.pop.info.color	lollipop pop information text color. Default <i>#EEE</i> .
lollipop.pop.info.dy	y-axis direction text adjustment of lollipop pop information. Default <i>-0.35em</i> .
lollipop.line.color	lollipop line color. Default <i>rgb(42,42,42)</i> .

`lollipop.line.width`  
lollipop line width. Default 0.5.

`lollipop.circle.color`  
lollipop circle border color. Default *wheat*.

`lollipop.circle.width`  
lollipop circle border width. Default 0.5.

`lollipop.label.ratio`  
lollipop click-out label font size to circle size ratio. Default 1.4.

`lollipop.label.min.font.size`  
lollipop click-out label minimal font size. Default 10.

`lollipop.color.scheme`  
color scheme to fill lollipop pops. Default *accent*.

`highlight.text.angle`  
pop-on-click highlight text angle. Default 90.

`title.text`  
title of chart. Default is empty.

`title.font`  
font of chart title. Default *normal 16px Arial*.

`title.color`  
color of chart title. Default *#424242*.

`title.alignment`  
text alignment of chart title (start/middle/end). Default *middle*.

`title.dy`  
text adjustment of chart title. Default *0.35em*.

`anno.height`  
height of protein structure annotation track. Default 30.

`anno.margin`  
margin of protein structure annotation track. Default `list(top = 4, bottom = 0)`.

`anno.background`  
background of protein structure annotation track. Default *transparent*.

`anno.bar.fill`  
background of protein bar in protein structure annotation track. Default *#e5e3e1*.

`anno.bar.margin`  
margin of protein bar in protein structure annotation track. Default `list(top = 2, bottom = 2)`.

`domain.color.scheme`  
color scheme of protein domains. Default *category10*.

`domain.margin`  
margin of protein domains. Default `list(top = 0, bottom = 0)`.

`domain.text.font`  
domain label text font in shorthand format. Default *normal 11px Arial*.

`domain.text.color`  
domain label text color. Default *#f2f2f2*.

`brush`  
if show brush. Default TRUE.

`brush.selection.background`  
background color of selection brush. Default *#666*.

`brush.selection.opacity`  
background opacity of selection brush. Default 0.2.

`brush.border.color`  
border color of selection brush. Default *#969696*.

`brush.handler.color`  
color of left and right handlers of selection brush. Default *#333*.

brush.border.width	border width of selection brush. Default 1.
legend	if show legend. Default TRUE.
tooltip	if show tooltip. Default TRUE.
zoom	if enable zoom feature. Default TRUE.

**Value**

a list with g3Lollipop plot options

---

g3Lollipop.theme	<i>G3Lollipop chart options of built-in themes.</i>
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---

**Description**

G3Lollipop chart options of built-in themes.

**Usage**

```
g3Lollipop.theme(theme.name = "default", title.text = "",
  y.axis.label = "# of mutations", legend.title = NA)
```

**Arguments**

theme.name	theme name, including <i>default</i> , <i>cbiportal</i> , <i>nature</i> , <i>nature2</i> , <i>dark</i> , <i>blue</i> , <i>ggplot2</i> , and <i>simple</i> . Default <i>default</i> .
title.text	title of chart. Default is empty.
y.axis.label	Y-axis label text. Default <i>"# of mutations"</i> .
legend.title	legend title. If NA, <i>factor.col</i> in <a href="#">g3Lollipop</a> is used. Default is NA.

**Value**

a list with g3Lollipop plot options

---

```
getMutationsFromCbioportal
```

*Query cancer genomic mutation data from cBioPortal*

---

### Description

Retrieve and parse mutation data from cBioPortal by the given cBioPortal cancer study ID and the gene symbol.

### Usage

```
getMutationsFromCbioportal(study.id, gene.symbol, output.file = NA,
  mutation.type.to.class.df = NA,
  cgds.url = "http://www.cbioportal.org/", test.cgds = FALSE)
```

### Arguments

<code>study.id</code>	cbioprotal study ID
<code>gene.symbol</code>	HGNC gene symbol.
<code>output.file</code>	if specified, output to a file in CSV format. Default is NA.
<code>mutation.type.to.class.df</code>	mapping table from mutation type to class. See <a href="#">mapMutationTypeToMutationClass</a> for details. Default NA, which indicates to use default mappings.
<code>cgds.url</code>	the URL for the public CGDS server (Cancer Genomic Data Server). Default is <a href="http://www.cbioportal.org/">http://www.cbioportal.org/</a> . Check <i>cgdsr</i> R-package for details.
<code>test.cgds</code>	if test CGDS connection. Default is FALSE

### Value

a data frame with columns

**Hugo\_Symbol** Hugo gene symbol

**Protein\_Change** Protein change information (cBioportal uses *HGVS<sub>p</sub>* format)

**Sample\_ID** Sample ID

**Mutation\_Type** mutation type, aka, variant classification.

**Chromosome** chromosome

**Start\_Position** start position

**End\_Position** end position

**Reference\_Allele** reference allele

**Variante\_Allele** variant allele

**Mutation\_Class** mutation class (e.g., Truncating/Missense/Inframe/Other)

**AA\_Position** amino-acid position of the variant; if the variant is not in protein-coding region, NA



## Examples

```
## Not run:
# Usage:
# Connection to CGDS (Cange Genomic Data Server). Internet access required.
# Note: this may need more than 10 seconds, and sometimes it may fail.
library(cgdsr)
cgds <- CGDS("http://www.cbioportal.org/")

# test if connection is OK (warning: sometimes it may fail)
test(cgds)

# list all studies of cBioPortal
all.studies <- getCancerStudies(cgds)

# First, select a cancer study that contains mutation data set ("caner_study_id")
# then, query genomic mutation data using a HGNC gene symbol,
# for example
mutation.dat <- getMutationsFromCbioportal("msk_impact_2017", "TP53")
mutation.dat <- getMutationsFromCbioportal("all_stjude_2016", "TP53")

## End(Not run)
```

---

guessMAFColumnName      *Guess column name for MAF file*

---

## Description

Guess column name for MAF file

## Usage

```
guessMAFColumnName(maf.df, alt.column.names)
```

## Arguments

maf.df                    MAF data frame  
alt.column.names            a vector of alternative column names

## Value

if hit one alternative column name, return the name; otherwise, return NA

hgnc2pfam

*Map from Hugo symbol to Pfam domains***Description**

Mapping from Hugo symbol to Pfam-A domain composition. If the given Hugo symbol has multiple UniProt ID mappings, and `guess == TRUE`, the longest UniProt protein is selected. Return is either a list of a JSON.

**Usage**

```
hgnc2pfam(hgnc.symbol, guess = TRUE, uniprot.id = NA,
          output.format = "json")
```

**Arguments**

<code>hgnc.symbol</code>	primary Hugo symbol
<code>guess</code>	if the given Hugo symbol links to multiple UniProt IDs, choose the longest one ( <code>guess == TRUE</code> ); otherwise NA ( <code>guess == FALSE</code> ). Default TRUE.
<code>uniprot.id</code>	UniProt ID, in case that gene symbol maps to multiple UniProt entries.
<code>output.format</code>	output format: JSON or list

**Value**

A list or a JSON with attributes: *symbol*, *uniprot*, *length*, and a list of *Pfam* entries, including *hmm.acc*, *hmm.name*, *start*, *end*, and *type*.

**Examples**

```
# general usage
hgnc2pfam("TP53")
hgnc2pfam("TP53", output.format = "json")
hgnc2pfam("TP53", output.format = "list")
hgnc2pfam("TP53", output.format = "json", uniprot.id = "P04637") # OK

# for gene mapping to multiple UniProt enties
hgnc2pfam("GNAS", guess = TRUE)
hgnc2pfam("GNAS", guess = FALSE)
hgnc2pfam("GNAS", output.format = "list")
hgnc2pfam("GNAS", output.format = "list", uniprot.id = "P84996")
## Not run:
hgnc2pfam("GNAS", output.format = "list", uniprot.id = "P84997") # not exists, returns FALSE

## End(Not run)

hgnc2pfam("PRAMEF9", output.format = "list") # no Pfam mappings
```

---

hgnc2pfam.df	<i>Mapping table between gene.symbol, uniprot.id, and pfam</i>
--------------	--

---

**Description**

A dataset containing the mapping table between Hugo symbol, UniProt ID, and Pfam ACC.

**Usage**

```
hgnc2pfam.df
```

**Format**

A data frame with columns:

**symbol** Gene symbol

**uniprot** UniProt ID

**length** protein length

**start** starting position of Pfam domain

**end** ending position of Pfam domain

**hmm.acc** Pfam accession number

**hmm.name** Pfam name

**type** Pfam type, i.e., domain/family/motif/repeat/disordered/coiled-coil

**Source**

Pfam (v31.0) and UniProt

**Examples**

```
hgnc2pfam.df
```

---

hgnc2uniprot	<i>Mapping from Hugo symbol to UniProt IDs</i>
--------------	--

---

**Description**

Mapping from Hugo Symbol to UniProt ID using internal mapping table. Return a data frame with columns *symbol* (Hugo symbol), *uniprot* (UniProt ID), and *length* (protein length).

**Usage**

```
hgnc2uniprot(hgnc.symbol)
```

**Arguments**

hgnc.symbol     primary HUGO symbol

**Value**

a data frame with columns *symbol* (Hugo symbol), *uniprot* (UniProt ID), and *length* (protein length).

**Examples**

```
# maps to single UniProt entry
hgnc2uniprot("TP53")

# maps to multiple UniProt entries
hgnc2uniprot("GNAS")
hgnc2uniprot("AKAP7")
```

---

mapMutationTypeToMutationClass

*Map from mutation type (aka, variant classification) to mutation class*

---

**Description**

Map from mutation type (aka, variant classification) to mutation class. Default mappings are as follows,

- Missense
  - *Missense\_Mutation* — a point mutation in which a single nucleotide change results in a codon that codes for a different amino acid See [https://en.wikipedia.org/wiki/Missense\\_mutation](https://en.wikipedia.org/wiki/Missense_mutation).
- Inframe
  - *In\_Frame\_Del* — a deletion that keeps the sequence in frame
  - *In\_Frame\_Ins* — an insertion that keeps the sequence in frame
  - *Silent* — variant is in coding region of the chosen transcript, but protein structure is identical (i.e., a synonymous mutation)
  - *Targeted\_Region* — targeted region
- Truncating
  - *Frame\_Shift* — a variant caused by indels of a number of nucleotides in a DNA sequence that is not divisible by three. See [https://en.wikipedia.org/wiki/Frameshift\\_mutation](https://en.wikipedia.org/wiki/Frameshift_mutation).
  - *Frame\_Shift\_Ins* — a variant caused by insertion that moves the coding sequence out of frame. See [https://en.wikipedia.org/wiki/Frameshift\\_mutation](https://en.wikipedia.org/wiki/Frameshift_mutation).
  - *Frame\_Shift\_Del* — a variant caused by deletion that moves the coding sequence out of frame. See [https://en.wikipedia.org/wiki/Frameshift\\_mutation](https://en.wikipedia.org/wiki/Frameshift_mutation).

- *Nonsense\_Mutation* — a premature stop codon that is created by the variant. See [https://en.wikipedia.org/wiki/Nonsense\\_mutation](https://en.wikipedia.org/wiki/Nonsense_mutation).
- *Nonstop\_Mutation* — a variant that removes stop codon.
- *Splice\_Site* — a variant that is within two bases of a splice site.
- *Splice\_Region* — a variant that is within splice region.
- Other
  - *5'UTR* — a variant that is on the 5'UTR for the chosen transcript.
  - *3'UTR* — a variant that is on the 3'UTR for the chosen transcript.
  - *5'Flank* — a variant that is upstream of the chosen transcript (generally within 3kb).
  - *3'Flank* — a variant that is downstream of the chosen transcript (generally within 3kb).
  - *Fusion* — a gene fusion.
  - *IGR* — an intergenic region. Does not overlap any transcript.
  - *Intron* — a variant that lies between exons within the bounds of the chosen transcript.
  - *Translation\_Start\_Site* — a variant that is in translation start site.
  - *De\_novo\_Start\_InFrame* — a novel start codon that is created by the given variant using the chosen transcript. However, it is in frame relative to the coded protein.
  - *De\_novo\_Start\_OutOfFrame* — a novel start codon that is created by the given variant using the chosen transcript. However, it is out of frame relative to the coded protein.
  - *Start\_Codon\_SNP* — a point mutation that overlaps the start codon.
  - *Start\_Codon\_Ins* — an insertion that overlaps the start codon.
  - *Start\_Codon\_Del* — a deletion that overlaps the start codon.
  - *RNA* — a variant that lies on one of the RNA transcripts.
  - *lincRNA* — a variant that lies on one of the lincRNAs.
  - *Unknown* — Unknown

## Usage

```
mapMutationTypeToMutationClass(mutation.type.vec,
  mutation.type.to.class.df = NA)
```

## Arguments

```
mutation.type.vec
  a vector of mutation type information

mutation.type.to.class.df
  A mapping table from mutation type (header Mutation_Type) to mutation class
  (header Mutation_Class). Default NA, which indicates to use default mappings.
```

## Value

```
a vector of mapped mutation class information
```

---

mutation.table.df	<i>Default mapping table between mutation type (aka, variant classification) to mutation class</i>
-------------------	--

---

### Description

A dataset containing the mapping table between genomic mutation type (aka, variant classification) to mutation class. See [mapMutationTypeToMutationClass](#) for details.

### Usage

```
mutation.table.df
```

### Format

A data frame with three columns:

**Mutation\_Type** Mutation type, aka, variant classification

**Mutation\_Class** mutation class

**Short\_Name** short name of mutation type

### Examples

```
mutation.table.df
```

---

parseProteinChange	<i>Extract amino_acid_position from Protein_Change</i>
--------------------	--

---

### Description

Parse *amino\_acid\_position* according to HGVS<sub>p</sub>\_short format.

For example, *p.Q16Rfs\*28*, amino-acid position is 16. See <http://varnomen.hgvs.org/recommendations/protein/> or <https://www.hgvs.org/mutnomen/recs-prot.html>.

### Usage

```
parseProteinChange(protein.change.vec, mutation.class.vec)
```

### Arguments

protein.change.vec

a vector of strings with protein change information, usually in HGVS<sub>p</sub>\_short format.

mutation.class.vec

a vector of strings with mutation class (or so-called variant classification) information.

**Value**

a vector of parsed amino-acid position

---

readMAF	<i>Read MAF file</i>
---------	----------------------

---

**Description**

Read mutation information from MAF file. For MAF format specification, see [https://docs.gdc.cancer.gov/Data/File\\_Formats/MAF\\_Format/](https://docs.gdc.cancer.gov/Data/File_Formats/MAF_Format/).

**Usage**

```
readMAF(maf.file, gene.symbol.col = "Hugo_Symbol",
        variant.class.col = c("Variant_Classification", "Mutation_Type"),
        protein.change.col = c("Protein_Change", "HGVS_Short"),
        if.parse.aa.pos = TRUE, if.parse.mutation.class = TRUE,
        mutation.class.col = "Mutation_Class", aa.pos.col = "AA_Position",
        mutation.type.to.class.df = NA, sep = "\t", quote = "", ...)
```

**Arguments**

maf.file	MAF file name. Gzipped input file allowed, with ".gz" file extension.
gene.symbol.col	Column name of Hugo gene symbols (e.g., TP53). Default <i>Hugo_Symbol</i> .
variant.class.col	Column name for variant class information (e.g., <i>Missense_Mutation</i> , <i>Nonsense_Mutation</i> ). Default is the first match of <i>Variant_Classification</i> or <i>Mutation_Type</i> .
protein.change.col	Column name for protein change information (e.g., p.K960R, G658S, L14Sfs*15). Default is the first match of <i>Protein_Change</i> or <i>HGVS_Short</i> .
if.parse.aa.pos	if parse amino-acid position of mutations. Default is TRUE.
if.parse.mutation.class	if parse mutation class from mutation type (variant classification) information. Default is TRUE.
mutation.class.col	Column name of the parsed mutation class. Default <i>Mutation_Class</i> .
aa.pos.col	Column name of the parsed amino-acid change position. Default <i>AA_Position</i> .
mutation.type.to.class.df	mapping table from mutation type to class. <a href="#">mapMutationTypeToMutationClass</a> for details. Default NA, which indicates to use default mappings.
sep	separator of columns. Default sep = "\t".
quote	the set of quoting characters. To disable quoting altogether, use quote = "". Default quote = "".
...	additional parameters pass to <a href="#">read.table</a> .

**Value**

a data frame containing MAF information, plus optional columns of the parsed *Mutation\_Class* and *Protein\_Position*.

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uniprot2pfam	<i>From UniProt ID to Pfam-A domain composition</i>
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**Description**

Map from UniProt ID to Pfam-A domain composition.

**Usage**

```
uniprot2pfam(uniprot.id)
```

**Arguments**

```
uniprot.id    UniProt ID
```

**Value**

a data frame with columns

- *uniprot* — UniProt ID
- *length* — protein length
- *hmm.acc* — accession number of Pfam HMM model, e.g., PF08563
- *hmm.name* — Pfam name, e.g., P53\_TAD
- *start* — Pfam domain start position
- *end* — Pfam domain end position
- *type* — Pfam type, including domain/motif/family

**Examples**

```
uniprot2pfam("Q5VWM5") # PRAMEF9; PRAMEF15
uniprot2pfam("P04637")
```



# Index

## \*Topic **datasets**

- hgnc2pfam.df, [11](#)
- mutation.table.df, [14](#)

- g3Lollipop, [2](#), [5](#), [7](#)
- g3Lollipop-shiny, [3](#)
- g3Lollipop.options, [2](#), [4](#)
- g3Lollipop.theme, [7](#)
- g3LollipopOutput (g3Lollipop-shiny), [3](#)
- getMutationsFromCbioportal, [8](#)
- guessMAFColumnName, [9](#)

- hgnc2pfam, [10](#)
- hgnc2pfam.df, [11](#)
- hgnc2uniprot, [11](#)

- mapMutationTypeToMutationClass, [8](#), [12](#),  
[14](#), [15](#)
- mutation.table.df, [14](#)

- parseProteinChange, [14](#)

- read.table, [15](#)
- readMAF, [15](#)
- renderG3Lollipop (g3Lollipop-shiny), [3](#)
- uniprot2pfam, [16](#)