

# Package ‘reportRmd’

January 13, 2023

**Title** Tidy Presentation of Clinical Reporting

**Version** 0.0.2

**Description** Streamlined statistical reporting in 'Rmarkdown' environments. Facilitates the automated reporting of descriptive statistics, multiple univariate models, multivariable models and tables combining these outputs. Plotting functions include customisable survival curves, forest plots from logistic and ordinal regression and bivariate comparison plots.

**License** MIT + file LICENSE

**Suggests** rmarkdown, testthat (>= 3.0.0)

**Config/testthat/edition** 3

**Encoding** UTF-8

**RoxygenNote** 7.2.3

**Imports** aod, cmprsk, geopack, ggplot2, ggpubr, gridExtra, kableExtra, knitr, MASS, pander, plyr, rlang, rstatix, scales, survival

**Collate** 'helper.R' 'main.R' 'globals.R' 'data.R'

**Depends** R (>= 2.10)

**LazyData** true

**VignetteBuilder** knitr, rmarkdown

**NeedsCompilation** no

**Author** Lisa Avery [cre, aut],  
Ryan Del Bel [aut],  
Osvaldo Espin-Garcia [aut],  
Tyler Pittman [aut] (0000-0002-5013-6980),  
Yanning Wang [ctr],  
Jessica Weiss [aut],  
Wei Xu [aut]

**Maintainer** Lisa Avery <lisa.avery@uhn.ca>

**Repository** CRAN

**Date/Publication** 2023-01-13 18:50:02 UTC

**R topics documented:**

addspace . . . . .	3
boxcoxfitRx . . . . .	3
cap . . . . .	4
covsum . . . . .	4
crrRx . . . . .	6
ctDNA . . . . .	7
excelCol . . . . .	7
forestplot2 . . . . .	8
formatp . . . . .	9
geoR_boxcoxfit . . . . .	9
ggkmcif . . . . .	10
ggkmcif_paste . . . . .	14
hbld . . . . .	15
lbld . . . . .	15
lpvalue . . . . .	16
mvsum . . . . .	16
nestTable . . . . .	17
nicename . . . . .	19
niceNum . . . . .	19
outTable . . . . .	20
pembrolizumab . . . . .	21
plotuv . . . . .	22
psthr . . . . .	24
pstprn . . . . .	24
pvalue . . . . .	24
rmds . . . . .	25
rm_covsum . . . . .	25
rm_mvsum . . . . .	28
rm_survdiff . . . . .	30
rm_survsum . . . . .	31
rm_survtime . . . . .	33
rm_uvsum . . . . .	35
rm_uv_mv . . . . .	38
sanitizestr . . . . .	39
testData . . . . .	40
uvsum . . . . .	40
<b>Index</b>	<b>43</b>

---

addspace	<i>Add spaces to strings in LaTeX</i>
----------	---------------------------------------

---

**Description**

Add spaces to strings in LaTeX. Returns appends ~~~ before the string

**Usage**

```
addspace(x)
```

**Arguments**

x	string
---	--------

---

boxcoxfitRx	<i>fit box cox transformed linear model</i>
-------------	---

---

**Description**

Wrapper function to fit fine and gray competing risk model using function crr from package cmprsk

**Usage**

```
boxcoxfitRx(f, data, lambda = FALSE)
```

**Arguments**

f	formula for the model. Currently the formula only works by using the name of the column in a dataframe. It does not work by using \$ or [] notation.
data	dataframe containing data
lambda	boolean indicating if you want to output the lamda used in the boxcox transformation. If so the function will return a list of length 2 with the model as the first element and a vector of length 2 as the second.

**Value**

a list containing the linear model (lm) object and, if requested, lambda

---

cap	<i>Capitalize a string</i>
-----	----------------------------

---

**Description**

Capitalize a string

**Usage**

```
cap(x)
```

**Arguments**

x	string
---	--------

---

covsum	<i>Get covariate summary dataframe</i>
--------	--

---

**Description**

Returns a dataframe corresponding to a descriptive table.

**Usage**

```
covsum(
  data,
  covs,
  maincov = NULL,
  digits = 1,
  numobs = NULL,
  markup = TRUE,
  sanitize = TRUE,
  nicenames = TRUE,
  IQR = FALSE,
  all.stats = FALSE,
  pvalue = TRUE,
  effSize = FALSE,
  show.tests = FALSE,
  dropLevels = TRUE,
  excludeLevels = NULL,
  full = TRUE,
  digits.cat = 0,
  testcont = c("rank-sum test", "ANOVA"),
  testcat = c("Chi-squared", "Fisher"),
  include_missing = FALSE,
  percentage = c("column", "row")
)
```

**Arguments**

<code>data</code>	dataframe containing data
<code>covs</code>	character vector with the names of columns to include in table
<code>maincov</code>	covariate to stratify table by
<code>digits</code>	number of digits for summarizing mean data, does not affect p-values
<code>numobs</code>	named list overriding the number of people you expect to have the covariate
<code>markup</code>	boolean indicating if you want latex markup
<code>sanitize</code>	boolean indicating if you want to sanitize all strings to not break LaTeX
<code>nicenames</code>	boolean indicating if you want to replace . and _ in strings with a space
<code>IQR</code>	boolean indicating if you want to display the inter quantile range (Q1,Q3) as opposed to (min,max) in the summary for continuous variables
<code>all.stats</code>	boolean indicating if all summary statistics (Q1,Q3 + min,max on a separate line) should be displayed. Overrides IQR.
<code>pvalue</code>	boolean indicating if you want p-values included in the table
<code>effSize</code>	boolean indicating if you want effect sizes included in the table. Can only be obtained if pvalue is also requested. Effect sizes are calculated with the rstatix package using Cramer V for categorical and Eta Squared for continuous covariates.
<code>show.tests</code>	boolean indicating if the type of statistical used should be shown in a column beside the pvalues. Ignored if pvalue=FALSE.
<code>dropLevels</code>	logical, indicating if empty factor levels be dropped from the output, default is TRUE.
<code>excludeLevels</code>	a named list of covariate levels to exclude from statistical tests in the form list(varname =c('level1','level2')). These levels will be excluded from association tests, but not the table. This can be useful for levels where there is a logical skip (ie not missing, but not presented). Ignored if pvalue=FALSE.
<code>full</code>	boolean indicating if you want the full sample included in the table, ignored if maincov is NULL
<code>digits.cat</code>	number of digits for the proportions when summarizing categorical data (default: 0)
<code>testcont</code>	test of choice for continuous variables,one of <i>rank-sum</i> (default) or <i>ANOVA</i>
<code>testcat</code>	test of choice for categorical variables,one of <i>Chi-squared</i> (default) or <i>Fisher</i>
<code>include_missing</code>	Option to include NA values of maincov. NAs will not be included in statistical tests
<code>percentage</code>	choice of how percentages are presented ,one of <i>column</i> (default) or <i>row</i>

**Details**

Comparisons for categorical variables default to chi-square tests, but if there are counts of <5 then the Fisher Exact test will be used and if this is unsuccessful then a second attempt will be made

computing p-values using MC simulation. If `testcont='ANOVA'` then the t-test with unequal variance will be used for two groups and an ANOVA will be used for three or more. The statistical test used can be displayed by specifying `show.tests=TRUE`.

The number of decimal places to display the statistics can be changed with `digits`, but this will not change the display of p-values. If more significant digits are required for p-values then use `tableOnly=TRUE` and `format` as desired.

### See Also

[fisher.test](#), [chisq.test](#), [wilcox.test](#), [kruskal.test](#), and [anova](#)

---

crrRx

*fit crr model*

---

### Description

Wrapper function to fit fine and gray competing risk model using function `crr` from package `cmprsk`

### Usage

```
crrRx(f, data)
```

### Arguments

<code>f</code>	formula for the model. Currently the formula only works by using the name of the column in a dataframe. It does not work by using <code>\$</code> or <code>[]</code> notation.
<code>data</code>	dataframe containing data

### Value

a competing risk model with the call appended to the list

### See Also

[crr](#)

### Examples

```
# From the crr help file:
set.seed(10)
ftime <- rexp(200)
fstatus <- sample(0:2,200,replace=TRUE)
cov <- matrix(runif(600),nrow=200)
dimnames(cov)[[2]] <- c('x1','x2','x3')
df <- data.frame(ftime,fstatus,cov)
m1 <- crrRx(as.formula('ftime+fstatus~x1+x2+x3'),df)
# Nicely output to report:
rm_mvsum(m1,data=df,showN = TRUE,vif=TRUE)
```

---

ctDNA	<i>Tumour size change over time Longitudinal changes in tumour size since baseline for patients by changes in ctDNA status (clearance, decrease or increase) since baseline.</i>
-------	--

---

**Description**

Tumour size change over time

Longitudinal changes in tumour size since baseline for patients by changes in ctDNA status (clearance, decrease or increase) since baseline.

**Usage**

ctDNA

**Format**

A data frame with 270 rows and 5 variables:

**id** Patient ID

**cohort** Study Cohort: A = Squamous cell carcinoma of soft pallate, B = Triple negative breast cancer, C = Ovarian, high grade serous, D = Melanoma, E = Other Solid Tumor

**ctdna\_status** Change in ctDNA since baseline

**time** Number of weeks on treatment

**size\_change** Percentage change in tumour measurement

**Source**

<https://www.nature.com/articles/s43018-020-0096-5>

---

excelCol	<i>Retrieve columns number from spreadsheet columns specified as unquoted letters</i>
----------	---

---

**Description**

Retrieve columns number from spreadsheet columns specified as unquoted letters

**Usage**

excelCol(...)

**Arguments**

... unquoted excel column headers (i.e. excelCol(A,CG,AA)) separated by commas

**Value**

a numeric vector corresponding to columns in a spreadsheet

**Examples**

```
## Find the column numbers for excel columns AB, CE and BB
excelCol(AB,CE,bb)
```

---

 forestplot2

---

*Create a forest plot using ggplot2*


---

**Description**

This function will accept a log or logistic regression fit from glm or geeglm, and display the OR or RR for each variable on the appropriate log scale.

**Usage**

```
forestplot2(
  model,
  conf.level = 0.95,
  orderByRisk = TRUE,
  colours = "default",
  showEst = TRUE,
  rmRef = FALSE,
  logScale = TRUE,
  nxBreaks = 5
)
```

**Arguments**

model	an object output from the glm or geeglm function, must be from a logistic regression
conf.level	controls the width of the confidence interval
orderByRisk	logical, should the plot be ordered by risk
colours	can specify colours for risks less than, 1 and greater than 1.0. Default is red, black, green
showEst	logical, should the risks be displayed on the plot in text
rmRef	logical, should the reference levels be removed for the plot?
logScale	logical, should OR/RR be shown on log scale, defaults to TRUE. See <a href="https://doi.org/10.1093/aje/kwr156">https://doi.org/10.1093/aje/kwr156</a> for why you may prefer a linear scale.
nxBreaks	Number of tick marks supplied to the log_breaks function to produce

**Value**

a plot object



**Examples**

```
glm_fit = glm(orr~change_ctdna_group+sex+age+l_size,
data=pembrolizumab,family = 'binomial')
forestplot2(glm_fit)
```

---

formatp	<i>Specific p-value formatting</i>
---------	------------------------------------

---

**Description**

If  $p < 0.001$  returns "<0.001", if  $p < 0.01$  returns p to 3 decimal places otherwise returns p to 2 decimal places

**Usage**

```
formatp(pvalues)
```

**Arguments**

pvalues            a vector of p values

---

geoR_boxcoffit	<i>Parameter Estimation for the Box-Cox Transformation</i>
----------------	--

---

**Description**

This function is copied from the geoR package which has been removed from the CRAN repository.

**Usage**

```
geoR_boxcoffit(object, xmat, lambda, lambda2 = NULL, add.to.data = 0)
```

**Arguments**

object            a vector with the data

xmat             a matrix with covariates values. Defaults to `rep(1, length(y))`.

lambda            numerical value(s) for the transformation parameter lambda. Used as the initial value in the function for parameter estimation. If not provided default values are assumed. If multiple values are passed the one with highest likelihood is used as initial value.

lambda2           logical or numerical value(s) of the additional transformation (see DETAILS below). Defaults to NULL. If TRUE this parameter is also estimated and the initial value is set to the absolute value of the minimum data. A numerical value is provided it is used as the initial value. Multiple values are allowed as for lambda.

add.to.data      a constant value to be added to the data.

## Details

For more information see: <https://cran.r-project.org/web/packages/geoR/index.html>

---

ggkmcif

*Plot KM and CIF curves with ggplot*

---

## Description

This function will plot a KM or CIF curve with option to add the number at risk. You can specify if you want confidence bands, the hazard ratio, and pvalues, as well as the units of time used.

## Usage

```
ggkmcif(  
  response,  
  cov = NULL,  
  data,  
  type = NULL,  
  pval = TRUE,  
  HR = FALSE,  
  HR_pval = FALSE,  
  conf.curves = FALSE,  
  conf.type = "log",  
  table = TRUE,  
  times = NULL,  
  xlab = "Time",  
  ylab = NULL,  
  main = NULL,  
  stratalabs = NULL,  
  strataname = niceName(cov),  
  stratalabs.table = NULL,  
  strataname.table = strataname,  
  median.text = FALSE,  
  median.lines = FALSE,  
  median.CI = FALSE,  
  set.time.text = NULL,  
  set.time.line = FALSE,  
  set.time = 5,  
  set.time.CI = FALSE,  
  censor.marks = TRUE,  
  censor.size = 3,  
  censor.stroke = 1.5,  
  fsize = 10,  
  nsize = 3,  
  lsize = 1,  
  psize = 3.5,
```

```

median.size = 3,
median.pos = NULL,
median.lsize = 1,
set.size = 3,
set.pos = NULL,
set.lsize = 1,
ylim = c(0, 1),
col = NULL,
linetype = NULL,
xlim = NULL,
legend.pos = NULL,
pval.pos = NULL,
plot.event = 1,
event = c("col", "linetype"),
flip.CIF = FALSE,
cut = NULL,
eventlabs = NULL,
event.name = NULL,
Numbers_at_risk_text = "Numbers at risk",
HR.digits = 2,
HR.pval.digits = 3,
pval.digits = 3,
median.digits = 3,
set.time.digits = 3,
returns = FALSE,
print.n.missing = TRUE
)

```

### Arguments

response	character vector with names of columns to use for response
cov	String specifying the column name of stratification variable
data	dataframe containing your data
type	string indicating the type of univariate model to fit. The function will try and guess what type you want based on your response. If you want to override this you can manually specify the type. Options include "KM", and "CIF"
pval	boolean to specify if you want p-values in the plot (Log Rank test for KM and Gray's test for CIF)
HR	boolean to specify if you want hazard ratios included in the plot
HR_pval	boolean to specify if you want HR p-values in the plot
conf.curves	boolean to specify if you want confidence interval bands
conf.type	One of "none"(the default), "plain", "log", "log-log" or "logit". Only enough of the string to uniquely identify it is necessary. The first option causes confidence intervals not to be generated. The second causes the standard intervals curve $\pm k * se(\text{curve})$ , where k is determined from conf.int. The log option calculates intervals based on the cumulative hazard or log(survival). The log-log option

	bases the intervals on the log hazard or $\log(-\log(\text{survival}))$ , and the logit option on $\log(\text{survival}/(1-\text{survival}))$ .
table	Logical value. If TRUE, includes the number at risk table
times	Numeric vector of times for the x-axis
xlab	String corresponding to xlabel. By default is "Time"
ylab	String corresponding to ylabel. When NULL uses "Survival probability" for KM curves, and "Probability of an event" for CIF
main	String corresponding to main title. When NULL uses Kaplan-Meier Plot s, and "Cumulative Incidence Plot for CIF"
stratalabs	string corresponding to the labels of the covariate, when NULL will use the levels of the covariate
strataname	String of the covariate name default is <code>nicename(cov)</code>
stratalabs.table	String corresponding to the levels of the covariate for the number at risk table, when NULL will use the levels of the covariate. Can use a string of "-" when the labels are long
strataname.table	String of the covariate name for the number at risk table default is <code>nicename(cov)</code>
median.text	boolean to specify if you want the median values added to the legend (or as added text if there are no covariates), for KM only
median.lines	boolean to specify if you want the median values added as lines to the plot, for KM only
median.CI	boolean to specify if you want the 95% with the median text (Only for KM)
set.time.text	string for the text to add survival at a specified time (eg. year OS)
set.time.line	boolean to specify if you want the survival added as lines to the plot at a specified point
set.time	Numeric values of the specific time of interest, default is 5 (Multiple values can be entered)
set.time.CI	boolean to specify if you want the 95% interval with the set time text
cursor.marks	logical value. If TRUE, includes cursor marks (only for KM curves)
cursor.size	size of cursor marks, default is 3
cursor.stroke	stroke of cursor marks, default is 1.5
fsize	font size
nsize	font size for numbers in the numbers at risk table
lsize	line size
psize	size of the pvalue
median.size	size of the median text (Only when there are no covariates)
median.pos	vector of length 2 corresponding to the median position (Only when there are no covariates)
median.lsize	line size of the median lines

<code>set.size</code>	size of the survival at a set time text (Only when there are no covariates)
<code>set.pos</code>	vector of length 2 corresponding to the survival at a set point position (Only when there are no covariates)
<code>set.lsize</code>	line size of the survival at set points
<code>ylim</code>	vector of length 2 corresponding to limits of y-axis. Default to NULL
<code>col</code>	vector of colours
<code>linetype</code>	vector of line types
<code>xlim</code>	vector of length 2 corresponding to limits of x-axis. Default to NULL
<code>legend.pos</code>	Can be either a string corresponding to the legend position ("left", "top", "right", "bottom", "none") or a vector of length 2 corresponding to the legend position (uses normalized units (ie the c(0.5,0.5) is the middle of the plot))
<code>pval.pos</code>	vector of length 2 corresponding to the p-value position
<code>plot.event</code>	Which event(s) to plot (1,2, or c(1,2))
<code>event</code>	String specifying if the event should be mapped to the colour, or linetype when plotting both events to colour = "col", line type
<code>flip.CIF</code>	boolean to flip the CIF curve to start at 1
<code>cut</code>	numeric value indicating where to divide a continuous covariate (default is the median)
<code>eventlabs</code>	String corresponding to the event type names
<code>event.name</code>	String corresponding to the label of the event types
<code>Numbers_at_risk_text</code>	String for the label of the number at risk
<code>HR.digits</code>	Number of digits printed of the hazard ratio
<code>HR.pval.digits</code>	Number of digits printed of the hazard ratio pvalue
<code>pval.digits</code>	Number of digits printed of the Gray's/log rank pvalue
<code>median.digits</code>	Number of digits printed of the median pvalue
<code>set.time.digits</code>	Number of digits printed of the probability at a specified time
<code>returns</code>	Logical value returns a list with all ggplot objects in a list
<code>print.n.missing</code>	Logical, should the number of missing be shown !Needs to be checked

## Details

Note that for proper pdf output of special characters the following code needs to be included in the first chunk of the rmd knitr::opts\_chunk\$set(dev="cairo\_pdf")

## Value

Nothing is returned unless `returns = TRUE` is used. With `returns = TRUE`, if `table=TRUE` (the default) a table style graphic with survival plot and number at risk table is returned. Otherwise a plot with the survival curves is returned.

**Examples**

```
# Simple plot without confidence intervals
ggkmcif(response = c('os_time','os_status'),
cov='cohort',
data=pembrolizumab)

# Plot with median survival time
ggkmcif(response = c('os_time','os_status'),
cov='sex',
data=pembrolizumab,
median.text = TRUE,median.lines=TRUE,conf.curves=TRUE)

# Plot with specified survival times and log-log CI
ggkmcif(response = c('os_time','os_status'),
cov='sex',
data=pembrolizumab,
median.text = FALSE,set.time.text = 'mo OS',
set.time = c(12,24),conf.type = 'log-log',conf.curves=TRUE)

# KM plot with 95% CI and censor marks
ggkmcif(c('os_time','os_status'),'sex',data = pembrolizumab, type = 'KM',
HR=TRUE, HR_pval = TRUE, conf.curves = TRUE,conf.type='log-log',
set.time.CI = TRUE, censor.marks=TRUE)
```

ggkmcif\_paste

*Plot KM and CIF curves with ggplot***Description**

This function puts together a survival curve, and a number at risk table

**Usage**

```
ggkmcif_paste(list_gg)
```

**Arguments**

`list_gg` list containing the results of ggkmcif

**Value**

a gtable with three elements, the survival curve, a spacer and the number at risk table

**Examples**

```
plot <- ggkmcif(response=c('pfs_time','pfs_status'),
data=pembrolizumab,returns = TRUE)

# Highlighting a section of the curve
```

```
plot[[1]] <- plot[[1]] +  
ggplot2::geom_rect(xmin=4,xmax=8,ymin=0.15,ymax=0.4,alpha=0.01,fill='yellow')  
  
# Putting the curve back together  
ggkmcif_paste(plot)
```

---

**hbld***Bold strings in HTML*

---

**Description**

Bold strings in HTML

**Usage**

```
hbld(strings)
```

**Arguments**

strings      A vector of strings to bold.

---

**lbld***Bold strings in LaTeX*

---

**Description**

Bold strings in LaTeX

**Usage**

```
lbld(strings)
```

**Arguments**

strings      A vector of strings to bold.

---

lpvalue	<i>Formats p-values for LaTeX</i>
---------	-----------------------------------

---

**Description**

Returns <0.001 if pvalue is <0.001. Else rounds the pvalue to specified significant digits. Will bold the p-value if it is <= 0.05

**Usage**

```
lpvalue(x, sigdigits = 2)
```

**Arguments**

x	an integer
sigdigits	number of significant digit to report

---

mvsum	<i>Get multivariate summary dataframe</i>
-------	---

---

**Description**

Returns a dataframe with the model summary and global p-value for multi-level variables.

**Usage**

```
mvsum(
  model,
  data,
  digits = 2,
  showN = FALSE,
  markup = TRUE,
  sanitize = TRUE,
  nicenames = TRUE,
  CIwidth = 0.95,
  vif = TRUE
)
```

**Arguments**

model	fitted model object
data	dataframe containing data
digits	number of digits to round to
showN	boolean indicating sample sizes should be shown for each comparison, can be useful for interactions



markup	boolean indicating if you want latex markup
sanitize	boolean indicating if you want to sanitize all strings to not break LaTeX
nicenames	boolean indicating if you want to replace . and _ in strings with a space.
CIwidth	width for confidence intervals, defaults to 0.95
vif	boolean indicating if the variance inflation factor should be included. See details

## Details

Global p-values are likelihood ratio tests for lm, glm and polr models. For lme models an attempt is made to re-fit the model using ML and if successful LRT is used to obtain a global p-value. For coxph models the model is re-run without robust variances with and without each variable and a LRT is presented. If unsuccessful a Wald p-value is returned. For GEE and CRR models Wald global p-values are returned.

If the variance inflation factor is requested (VIF=T) then a generalised VIF will be calculated in the same manner as the car package.

VIF for competing risk models is computed by fitting a linear model with a dependent variable comprised of the sum of the model independent variables and then calculating VIF from this linear model.

## References

John Fox & Georges Monette (1992) Generalized Collinearity Diagnostics, Journal of the American Statistical Association, 87:417, 178-183, DOI: 10.1080/01621459.1992.10475190

John Fox and Sanford Weisberg (2019). An R Companion to Applied Regression, Third Edition. Thousand Oaks CA: Sage. URL: <https://socialsciences.mcmaster.ca/jfox/Books/Companion>

---

nestTable	<i>Combine two table columns into a single column with levels of one nested within levels of the other.</i>
-----------	---

---

## Description

This function accepts a data frame (via the data argument) and combines two columns into a single column with values from the head\_col serving as headers and values of the to\_col displayed underneath each header. The resulting table is then passed to outTable for printing and output, to use the grouped table as a data frame specify tableOnly=TRUE. By default the headers will be bolded and the remaining values indented.

## Usage

```
nestTable(
  data,
  head_col,
  to_col,
  colHeader = "",
```

```

caption = NULL,
indent = TRUE,
boldheaders = TRUE,
hdr_prefix = "",
hdr_suffix = "",
digits = 2,
tableOnly = FALSE
)

```

### Arguments

data	dataframe
head_col	character value specifying the column name with the headers
to_col	character value specifying the column name to add the headers into
colHeader	character with the desired name of the first column. The default is to leave this empty for output or, for table only output to use the column name 'col1'.
caption	table caption
indent	Boolean should the original values in the to_col be indented
boldheaders	Boolean should the header column values be bolded
hdr_prefix	character value that will prefix headers
hdr_suffix	character value that will suffix headers
digits	number of digits to round numeric columns to, wither a single number or a vector corresponding to the number of numeric columns
tableOnly	boolean indicating if the table should be formatted for printing or returned as a data frame

### Details

Note that it is possible to combine multiple tables (more than two) with this function.

### Value

A character vector of the table source code, unless tableOnly=TRUE in which case a data frame is returned

### Examples

```

## Investigate models to predict baseline ctDNA and tumour size and display together
## (not clinically useful!)
fit1 <- lm(baseline_ctdna~age+l_size+pd11,data=pembrolizumab)
m1 <- rm_mvsum(fit1,tableOnly=TRUE)
m1$Response = 'ctDNA'
fit2 <- lm(l_size~age+baseline_ctdna+pd11,data=pembrolizumab)
m2 <- rm_mvsum(fit2,tableOnly=TRUE)
m2$Response = 'Tumour Size'
rbind(m1,m2)
nestTable(rbind(m1,m2),head_col='Response',to_col='Covariate')

```

---

nicename	<i>Lean strings for printing</i>
----------	----------------------------------

---

**Description**

Returns strings with . and \_ replaced by a space. This is nice when printing column names of your dataframe in a report

**Usage**

```
nicename(strings, check_numbers = TRUE)
```

**Arguments**

strings            vector of strings to give a nice name  
check\_numbers    boolean indicating if numbers with decimals should be checked for and retained.

---

niceNum	<i>Round retaining digits</i>
---------	-------------------------------

---

**Description**

Round retaining digits

**Usage**

```
niceNum(x, digits = 2)
```

**Arguments**

x                    a vector  
digits                numeric

---

 outTable

*Print tables to PDF/Latex HTML or Word*


---

**Description**

Output the table nicely to whatever format is appropriate. This is the output function used by the `rm_*` printing functions.

**Usage**

```
outTable(
  tab,
  row.names = NULL,
  to_indent = numeric(0),
  bold_headers = TRUE,
  rows_bold = numeric(0),
  bold_cells = NULL,
  caption = NULL,
  digits,
  align,
  applyAttributes = TRUE,
  keep.rownames = FALSE,
  fontsize,
  chunk_label
)
```

**Arguments**

<code>tab</code>	a table to format
<code>row.names</code>	a string specifying the column name to assign to the rownames. If NULL (the default) then rownames are removed.
<code>to_indent</code>	numeric vector indicating which rows to indent in the first column.
<code>bold_headers</code>	boolean indicating if the column headers should be bolded
<code>rows_bold</code>	numeric vector indicating which rows to bold
<code>bold_cells</code>	array indices indicating which cells to bold. These will be in addition to rows bolded by <code>rows_bold</code> .
<code>caption</code>	table caption
<code>digits</code>	number of digits to round numeric columns to, wither a single number or a vector corresponding to the number of numeric columns in tab
<code>align</code>	string specifying column alignment, defaults to left alignment of the first column and right alignment of all other columns. The <code>align</code> argument accepts a single string with 'l' for left, 'c' for centre and 'r' for right, with no separations. For example, to set the left column to be centred, the middle column right-aligned and the right column left aligned use: <code>align='crl'</code>

`applyAttributes` boolean indicating if the function should use `to_indent` and `bold_cells` formatting attributes. This will only work properly if the dimensions of the table output from `rm_covsum`, `rm_uvsum` etc haven't changed.

`keep.rownames` should the row names be included in the output

`fontsize` PDF/HTML output only, manually set the table fontsize

`chunk_label` only used knitting to Word docs to allow cross-referencing

### Details

Entire rows can be bolded, or specific cells. Currently indentation refers to the first column only. By default, underscores in column names are converted to spaces. To disable this set `rm_` to FALSE

### Value

A character vector of the table source code, unless `tableOnly=TRUE` in which case a data frame is returned

### Examples

```
# To make custom changes or change the fontsize in PDF/HTML
tab <- rm_covsum(data=pembrolizumab,maincov = 'change_ctdna_group',
covs=c('age','sex','pd11','tmb','l_size'),show.tests=TRUE,tableOnly = TRUE)
outTable(tab, fontsize=7)

# To bold columns with the variable names
rows_bold <- c(1,4,7,10,13)
outTable(tab,rows_bold = rows_bold)

# To bold the estimates for male/female
bold_cells <- as.matrix(expand.grid(5:6,1:ncol(tab)))
outTable(tab,bold_cells= bold_cells)
```

---

pembrolizumab	<i>Survival data Survival status and ctDNA levels for patients receiving pembrolizumab</i>
---------------	--

---

### Description

Survival data

Survival status and ctDNA levels for patients receiving pembrolizumab

### Usage

```
pembrolizumab
```

**Format**

A data frame with 94 rows and 15 variables:

**id** Patient ID

**age** Age at study entry

**sex** Patient Sex

**cohort** Study Cohort: A = Squamous cell carcinoma of soft pallate, B = Triple negative breast cancer, C = Ovarian, high grade serous, D = Melanoma, E = Other Solid Tumor

**l\_size** Target lesion size at baseline

**pdl1** PD L1 percent

**tmb** log of TMB

**baseline\_ctdna** Baseline ctDNA

**change\_ctdna\_group** Did ctDNA increase or decrease from baseline to cycle 3

**orr** Objective Response

**cbr** Clinical Beneficial Response

**os\_status** Overall survival status, 0 = alive, 1 = deceased

**os\_time** Overall survival time in months

**pfs\_status** Progression free survival status, 0 = progression free, 1 = progressed

**pfs\_time** Progression free survival time in months

**Source**

<https://www.nature.com/articles/s43018-020-0096-5>

---

plotuv

*Plot multiple bivariate relationships in a single plot*

---

**Description**

This function is designed to accompany `uvsum` as a means of visualising the results, and uses similar syntax.

**Usage**

```
plotuv(
  response,
  covs,
  data,
  showN = FALSE,
  showPoints = TRUE,
  na.rm = TRUE,
  response_title = NULL,
  return_plotlist = FALSE,
  ncol = 2,
  p_margins = c(0, 0.2, 1, 0.2)
)
```

**Arguments**

response	character vector with names of columns to use for response
covs	character vector with names of columns to use for covariates
data	dataframe containing your data
showN	boolean indicating whether sample sizes should be shown on the plots
showPoints	boolean indicating whether individual data points should be shown when $n > 20$ in a category
na.rm	boolean indicating whether na values should be shown or removed
response_title	character value with title of the plot
return_plotlist	boolean indicating that the list of plots should be returned instead of a plot, useful for applying changes to the plot, see details
ncol	the number of columns of plots to be display in the ggarrange call, defaults to 2
p_margins	sets the TRBL margins of the individual plots, defaults to <code>c(0,0.2,1,.2)</code>

**Details**

Plots are displayed as follows: If response is continuous For a numeric predictor scatterplot For a categorical predictor: If 20+ observations available boxplot, otherwise dotplot with median line If response is a factor For a numeric predictor: If 20+ observations available boxplot, otherwise dotplot with median line For a categorical predictor barplot Response variables are shown on the ordinate (y-axis) and covariates on the abscissa (x-axis)

**Value**

a list containing plots for each variable in covs  
a plot object

**See Also**

[ggplot](#) and [ggarrange](#)

**Examples**

```
## Run multiple univariate analyses on the pembrolizumab dataset to predict cbr and
## then visualise the relationships.
rm_uvsum(data=pembrolizumab,
response='cbr',covs=c('age','sex','l_size','baseline_ctdna'))
plotuv(data=pembrolizumab, response='cbr',
covs=c('age','sex','l_size','baseline_ctdna'),showN=TRUE)
```

---

psthr *Round and paste with parentheses*

---

**Description**

Round and paste with parentheses

**Usage**

```
psthr(x, y = 2)
```

**Arguments**

x                    a numeric vector  
y                    integer corresponding to the number of digits to round by

---

pstprn *Paste with parentheses*

---

**Description**

Paste with parentheses

**Usage**

```
pstprn(x)
```

**Arguments**

x                    a vector

---

pvalue *Formats p-values*

---

**Description**

Returns <0.001 if pvalue is <0.001. Else rounds the pvalue to specified significant digits

**Usage**

```
pvalue(x, digits)
```

**Arguments**

x                    an integer  
digits                the number of significant digits to return



---

rmds	<i>Replace dollar signs with html for proper HTML output</i>
------	--

---

**Description**

Replace dollar signs with html for proper HTML output

**Usage**

```
rmds(s)
```

**Arguments**

s                    a character vector

---

rm_covsum	<i>Outputs a descriptive covariate table</i>
-----------	--

---

**Description**

Returns a data frame corresponding to a descriptive table.

**Usage**

```
rm_covsum(  
  data,  
  covs,  
  maincov = NULL,  
  caption = NULL,  
  tableOnly = FALSE,  
  covTitle = "",  
  digits = 1,  
  digits.cat = 0,  
  nicenames = TRUE,  
  IQR = FALSE,  
  all.stats = FALSE,  
  pvalue = TRUE,  
  effSize = FALSE,  
  unformattedp = FALSE,  
  show.tests = FALSE,  
  testcont = c("rank-sum test", "ANOVA"),  
  testcat = c("Chi-squared", "Fisher"),  
  full = TRUE,  
  include_missing = FALSE,  
  percentage = c("column", "row"),
```

```

    dropLevels = TRUE,
    excludeLevels = NULL,
    numobs = NULL,
    chunk_label
  )

```

## Arguments

<code>data</code>	dataframe containing data
<code>covs</code>	character vector with the names of columns to include in table
<code>maincov</code>	covariate to stratify table by
<code>caption</code>	character containing table caption (default is no caption)
<code>tableOnly</code>	Logical, if TRUE then a dataframe is returned, otherwise a formatted printed object is returned (default).
<code>covTitle</code>	character with the names of the covariate (predictor) column. The default is to leave this empty for output or, for table only output to use the column name 'Covariate'.
<code>digits</code>	number of digits for summarizing mean data
<code>digits.cat</code>	number of digits for the proportions when summarizing categorical data (default: 0)
<code>nicenames</code>	boolean indicating if you want to replace . and _ in strings with a space
<code>IQR</code>	boolean indicating if you want to display the inter quantile range (Q1,Q3) as opposed to (min,max) in the summary for continuous variables
<code>all.stats</code>	boolean indicating if all summary statistics (Q1,Q3 + min,max on a separate line) should be displayed. Overrides IQR.
<code>pvalue</code>	boolean indicating if you want p-values included in the table
<code>effSize</code>	boolean indicating if you want effect sizes included in the table. Can only be obtained if pvalue is also requested. Effect sizes are calculated with the rstatix package using Cramer V for categorical and Eta Squared for continuous covariates.
<code>unformattedp</code>	boolean indicating if you would like the p-value to be returned unformatted (ie not rounded or prefixed with '<'). Best used with tableOnly = T and outTable function. See examples.
<code>show.tests</code>	boolean indicating if the type of statistical used should be shown in a column beside the pvalues. Ignored if pvalue=FALSE.
<code>testcont</code>	test of choice for continuous variables, one of <i>rank-sum</i> (default) or <i>ANOVA</i>
<code>testcat</code>	test of choice for categorical variables, one of <i>Chi-squared</i> (default) or <i>Fisher</i>
<code>full</code>	boolean indicating if you want the full sample included in the table, ignored if maincov is NULL
<code>include_missing</code>	Option to include NA values of maincov. NAs will not be included in statistical tests
<code>percentage</code>	choice of how percentages are presented, one of <i>column</i> (default) or <i>row</i>

dropLevels	logical, indicating if empty factor levels be dropped from the output, default is TRUE.
excludeLevels	a named list of covariate levels to exclude from statistical tests in the form list(varname =c('level1','level2')). These levels will be excluded from association tests, but not the table. This can be useful for levels where there is a logical skip (ie not missing, but not presented). Ignored if pvalue=FALSE.
numobs	named list overriding the number of people you expect to have the covariate
chunk_label	only used if output is to Word to allow cross-referencing

### Details

Comparisons for categorical variables default to chi-square tests, but if there are counts of <5 then the Fisher Exact test will be used and if this is unsuccessful then a second attempt will be made computing p-values using MC simulation. If testcont='ANOVA' then the t-test with unequal variance will be used for two groups and an ANOVA will be used for three or more. The statistical test used can be displayed by specifying show.tests=TRUE.

Effect size can be obtained when p-value is requested.

Further formatting options are available using tableOnly=TRUE and outputting the table with a call to outTable.

### Value

A character vector of the table source code, unless tableOnly=TRUE in which case a data frame is returned

### See Also

[covsum](#), [fisher.test](#), [chisq.test](#), [wilcox.test](#), [kruskal.test](#), [anova](#), [cramer\\_v](#), [eta\\_squared](#), and [outTable](#)

### Examples

```
rm_covsum(data=pembrolizumab, maincov = 'orr',
covs=c('age', 'sex', 'pd11', 'tmb', 'l_size', 'change_ctdna_group'),
show.tests=TRUE)

# To Show Effect Sizes
rm_covsum(data=pembrolizumab, maincov = 'orr',
covs=c('age', 'sex'),
effSize=TRUE)

# To make custom changes or change the fontsize in PDF/HTML
tab <- rm_covsum(data=pembrolizumab,maincov = 'change_ctdna_group',
covs=c('age', 'sex', 'pd11', 'tmb', 'l_size'),show.tests=TRUE,tableOnly = TRUE)
outTable(tab, fontsize=7)

# To return unformatted p-values
tab <- rm_covsum(data=pembrolizumab, maincov = 'orr',
covs=c('age', 'sex', 'pd11', 'tmb', 'l_size', 'change_ctdna_group'),
```

```
show.tests=TRUE,unformattedp=TRUE,tableOnly=TRUE)
outTable(tab,digits=5)
outTable(tab,digits=5, applyAttributes=FALSE) # remove bold/indent
```

---

 rm\_mvsum

*Format a regression model nicely for 'Rmarkdown'*


---

## Description

Multivariable (or univariate) regression models are re-formatted for reporting and a global p-value is added for the evaluation of factor variables.

## Usage

```
rm_mvsum(
  model,
  data,
  digits = 2,
  covTitle = "",
  showN = FALSE,
  CIwidth = 0.95,
  vif = TRUE,
  caption = NULL,
  tableOnly = FALSE,
  p.adjust = "none",
  unformattedp = FALSE,
  chunk_label,
  nicenames = TRUE
)
```

## Arguments

model	model fit
data	data that model was fit on (an attempt will be made to extract this from the model)
digits	number of digits to round estimates to, does not affect p-values
covTitle	character with the names of the covariate (predictor) column. The default is to leave this empty for output or, for table only output to use the column name 'Covariate'.
showN	boolean indicating sample sizes should be shown for each comparison, can be useful for interactions
CIwidth	width for confidence intervals, defaults to 0.95
vif	boolean indicating if the variance inflation factor should be included. See details
caption	table caption
tableOnly	boolean indicating if unformatted table should be returned

p.adjust	p-adjustments to be performed (Global p-values only)
unformattedp	boolean indicating if you would like the p-value to be returned unformatted (ie not rounded or prefixed with '<'). Should be used in conjunction with the digits argument.
chunk_label	only used if output is to Word to allow cross-referencing
nicenames	boolean indicating if you want to replace . and _ in strings with a space

## Details

Global p-values are likelihood ratio tests for lm, glm and polr models. For lme models an attempt is made to re-fit the model using ML and if successful LRT is used to obtain a global p-value. For coxph models the model is re-run without robust variances with and without each variable and a LRT is presented. If unsuccessful a Wald p-value is returned. For GEE and CRR models Wald global p-values are returned.

If the variance inflation factor is requested (VIF=T) then a generalised VIF will be calculated in the same manner as the car package.

The number of decimals places to display the statistics can be changed with digits, but this will not change the display of p-values. If more significant digits are required for p-values then use tableOnly=TRUE and format as desired.

## Value

A character vector of the table source code, unless tableOnly=TRUE in which case a data frame is returned

## References

John Fox & Georges Monette (1992) Generalized Collinearity Diagnostics, Journal of the American Statistical Association, 87:417, 178-183, DOI: 10.1080/01621459.1992.10475190

John Fox and Sanford Weisberg (2019). An R Companion to Applied Regression, Third Edition. Thousand Oaks CA: Sage. URL: <https://socialsciences.mcmaster.ca/jfox/Books/Companion>

## Examples

```
glm_fit = glm(change_ctdna_group~sex:age+baseline_ctdna+l_size,
data=pembrolizumab,family = 'binomial')
rm_mvsum(glm_fit)

#linear model with p-value adjustment
lm_fit=lm(baseline_ctdna~age+sex+l_size+tmb,data=pembrolizumab)
rm_mvsum(lm_fit,p.adjust = "bonferroni")
#Coxph
require(survival)
res.cox <- coxph(Surv(os_time, os_status) ~ sex+age+l_size+tmb, data = pembrolizumab)
rm_mvsum(res.cox, vif=TRUE)
```

---

rm_survdiff	<i>Display event counts, expected event counts and logrank test of differences</i>
-------------	--

---

### Description

This is a wrapper function around the `survdiff` function to display overall event rates and group-specific rates along with the log-rank test of a difference in survival between groups in a single table suitable for markdown output. Median survival times are included by default but can be removed setting `median=FALSE`

### Usage

```
rm_survdiff(
  data,
  time,
  status,
  covs,
  strata,
  includeVarNames = FALSE,
  digits = 1,
  showCols = c("N", "Observed", "Expected"),
  CIwidth = 0.95,
  conf.type = "log",
  caption = NULL,
  tableOnly = FALSE
)
```

### Arguments

<code>data</code>	data frame containing survival data
<code>time</code>	string indicating survival time variable
<code>status</code>	string indicating event status variable
<code>covs</code>	character vector indicating variables to group observations by
<code>strata</code>	string indicating the variable to stratify observations by
<code>includeVarNames</code>	boolean indicating if the variable names should be included in the output table, default is <code>FALSE</code>
<code>digits</code>	the number of digits in the survival rate
<code>showCols</code>	character vector indicating which of the optional columns to display, defaults to <code>c('N','Observed','Expected')</code>
<code>CIwidth</code>	width of the median survival estimates, default is <code>95%</code>
<code>conf.type</code>	type of confidence interval see <a href="#">survfit</a> for details. Default is <code>'log'</code> .
<code>caption</code>	table caption
<code>tableOnly</code>	should a dataframe or a formatted object be returned

**Value**

A character vector of the survival table source code, unless tableOnly=TRUE in which case a data frame is returned

**See Also**

[survdiff](#)

**Examples**

```
#' # Differences between sex
rm_survdiff(data=pembrolizumab,time='os_time',status='os_status',
covs='sex',digits=1)

# Differences between sex, stratified by cohort
rm_survdiff(data=pembrolizumab,time='os_time',status='os_status',
covs='sex',strata='cohort',digits=1)
# Differences between sex/cohort groups
rm_survdiff(data=pembrolizumab,time='os_time',status='os_status',
covs=c('sex','cohort'),digits=1)
```

---

rm\_survsum

*Summarise survival data by group*

---

**Description**

Displays event counts, median survival time and survival rates at specified times points for the entire cohort and by group. The logrank test of differences in survival curves is displayed.

**Usage**

```
rm_survsum(
  data,
  time,
  status,
  group = NULL,
  survtimes = NULL,
  survtimeunit,
  survtimesLbls = NULL,
  CIwidth = 0.95,
  unformattedp = FALSE,
  conf.type = "log",
  na.action = "na.omit",
  showCounts = TRUE,
  digits = 2,
  caption = NULL,
  tableOnly = FALSE
)
```

**Arguments**

<code>data</code>	data frame containing survival data
<code>time</code>	string indicating survival time variable
<code>status</code>	string indicating event status variable
<code>group</code>	string or character vector indicating the variable(s) to group observations by. If this is left as NULL (the default) then summaries are provided for the entire cohort.
<code>survtimes</code>	numeric vector specifying when survival probabilities should be calculated.
<code>survtimeunit</code>	unit of time to suffix to the time column label if survival probabilities are requested, should be plural
<code>survtimesLbls</code>	if supplied, a vector the same length as <code>survtimes</code> with descriptions (useful for displaying years with data provided in months)
<code>CIwidth</code>	width of the survival probabilities, default is 95%
<code>unformattedp</code>	boolean indicating if you would like the p-value to be returned unformatted (ie not rounded or prefixed with '<'). Should be used in conjunction with the <code>digits</code> argument.
<code>conf.type</code>	type of confidence interval see <a href="#">survfit</a> for details. Default is 'log'.
<code>na.action</code>	default is to omit missing values, but can be set to throw an error using <code>na.action='na.fail'</code>
<code>showCounts</code>	boolean indicating if the at risk, events and censored columns should be output, default is TRUE
<code>digits</code>	the number of digits in the survival rate, default is 2.
<code>caption</code>	table caption for markdown output
<code>tableOnly</code>	should a dataframe or a formatted object be returned

**Details**

This summary table is supplied for simple group comparisons only. To examine differences in groups with stratification see [rm\\_survdiff](#). To summarise differences in survival rates controlling for covariates see [rm\\_survtime](#).

**Value**

A character vector of the survival table source code, unless `tableOnly=TRUE` in which case a data frame is returned

**See Also**

[survfit](#)

**Examples**

```
# Simple median survival table
rm_survsum(data=pembrolizumab,time='os_time',status='os_status')

# Survival table with yearly survival rates
```



```

rm_survsum(data=pembrolizumab,time='os_time',status='os_status',
survtimes=c(12,24),survtimesLbbs=1:2, survtimeunit='yr')

#Median survival by group
rm_survsum(data=pembrolizumab,time='os_time',status='os_status',group='sex')

# Survival Summary by cohort, displayed in years
rm_survsum(data=pembrolizumab,time='os_time',status='os_status',
group="cohort",survtimes=seq(12,72,12),
survtimesLbbs=seq(1,6,1),
survtimeunit='years')

# Survival Summary by Sex and ctDNA group
rm_survsum(data=pembrolizumab,time='os_time',status='os_status',
group=c('sex','change_ctdna_group'),survtimes=c(12,24),survtimeunit='mo')

```

---

rm\_survtime

---

*Display survival rates and events for specified times*


---

## Description

This is a wrapper for the survfit function to output a tidy display for reporting. Either Kaplan Meier or Cox Proportional Hazards models may be used to estimate the survival probabilities.

## Usage

```

rm_survtime(
  data,
  time,
  status,
  covs = NULL,
  strata = NULL,
  type = "KM",
  survtimes,
  survtimeunit,
  strata.prefix = NULL,
  survtimesLbbs = NULL,
  showCols = c("At Risk", "Events", "Censored"),
  CIwidth = 0.95,
  conf.type = "log",
  na.action = "na.omit",
  showCounts = TRUE,
  digits = 2,
  caption = NULL,
  tableOnly = FALSE
)

```

**Arguments**

data	data frame containing survival data
time	string indicating survival time variable
status	string indicating event status variable
covs	character vector with the names of variables to adjust for in coxph fit
strata	string indicating the variable to group observations by. If this is left as NULL (the default) then event counts and survival rates are provided for the entire cohort.
type	survival function, if no covs are specified defaults to Kaplan-Meier, otherwise the Cox PH model is fit. Use type='PH' to fit a Cox PH model with no covariates.
survtimes	numeric vector specifying when survival probabilities should be calculated.
survtimeunit	unit of time to suffix to the time column label if survival probabilities are requested, should be plural
strata.prefix	character value describing the grouping variable
survtimesLbls	if supplied, a vector the same length as survtimes with descriptions (useful for displaying years with data provided in months)
showCols	character vector specifying which of the optional columns to display, defaults to c('At Risk','Events','Censored')
CIwidth	width of the survival probabilities, default is 95%
conf.type	type of confidence interval see <a href="#">survfit</a> for details. Default is 'log'.
na.action	default is to omit missing values, but can be set to throw an error using na.action='na.fail'
showCounts	boolean indicating if the at risk, events and censored columns should be output, default is TRUE
digits	the number of digits in the survival rate, default is 2.
caption	table caption for markdown output
tableOnly	should a dataframe or a formatted object be returned

**Details**

If covariates are supplied then a Cox proportional hazards model is fit for the entire cohort and each strata. Otherwise the default is for Kaplan-Meier estimates. Setting type = 'PH' will force a proportional hazards model.

**Value**

A character vector of the survival table source code, unless tableOnly=TRUE in which case a data frame is returned

**See Also**

[survfit](#)

**Examples**

```
# Kaplan-Meier survival probabilities with time displayed in years
rm_survtime(data=pembrolizumab,time='os_time',status='os_status',
strata="cohort",type='KM',survtimes=seq(12,72,12),
survtimesLbbs=seq(1,6,1),
survtimeunit='years')

# Cox Proportional Hazards survival probabilities
rm_survtime(data=pembrolizumab,time='os_time',status='os_status',
strata="cohort",type='PH',survtimes=seq(12,72,12),survtimeunit='months')

# Cox Proportional Hazards survival probabilities controlling for age
rm_survtime(data=pembrolizumab,time='os_time',status='os_status',
covs='age',strata="cohort",survtimes=seq(12,72,12),survtimeunit='months')
```

---

rm\_uvsum

*Output several univariate models nicely in a single table*


---

**Description**

A table with the model parameters from running separate univariate models on each covariate. For factors with more than two levels a Global p-value is returned.

**Usage**

```
rm_uvsum(
  response,
  covs,
  data,
  digits = 2,
  covTitle = "",
  caption = NULL,
  tableOnly = FALSE,
  removeInf = FALSE,
  p.adjust = "none",
  unformattedp = FALSE,
  chunk_label,
  gee = FALSE,
  id = NULL,
  corstr = NULL,
  family = NULL,
  type = NULL,
  strata = 1,
  nicenames = TRUE,
  showN = TRUE,
  CIwidth = 0.95,
  refllevel = NULL,
```

```

    returnModels = FALSE
  )

```

### Arguments

response	string vector with name of response
covs	character vector with the names of columns to fit univariate models to
data	dataframe containing data
digits	number of digits to round estimates and CI to. Does not affect p-values.
covTitle	character with the names of the covariate (predictor) column. The default is to leave this empty for output or, for table only output to use the column name 'Covariate'.
caption	character containing table caption (default is no caption)
tableOnly	boolean indicating if unformatted table should be returned
removeInf	boolean indicating if infinite estimates should be removed from the table
p.adjust	p-adjustments to be performed (Global p-values only)
unformattedp	boolean indicating if you would like the p-value to be returned unformatted (ie not rounded or prefixed with '<'). Should be used in conjunction with the digits argument.
chunk_label	only used if output is to Word to allow cross-referencing
gee	boolean indicating if gee models should be fit to account for correlated observations. If TRUE then the id argument must specify the column in the data which indicates the correlated clusters.
id	character vector which identifies clusters. Only used for geeglm
corstr	character string specifying the correlation structure. Only used for geeglm. The following are permitted: "independence", "exchangeable", "ar1", "unstructured" and "userdefined"
family	description of the error distribution and link function to be used in the model. Only used for geeglm
type	string indicating the type of univariate model to fit. The function will try and guess what type you want based on your response. If you want to override this you can manually specify the type. Options include "linear", "logistic", "poisson", "coxph", "crr", "boxcox", "ordinal", "geeglm"
strata	character vector of covariates to stratify by. Only used for coxph and crr
nicenames	boolean indicating if you want to replace . and _ in strings with a space
showN	boolean indicating if you want to show sample sizes
CIwidth	width of confidence interval, default is 0.95
reflevel	manual specification of the reference level. Only used for ordinal regression. This will allow you to see which model is not fitting if the function throws an error
returnModels	boolean indicating if a list of fitted models should be returned. If this is TRUE then the models will be returned, but the output will be suppressed. In addition to the model elements a data element will be appended to each model so that the fitted data can be examined, if necessary. See Details

## Details

Global p-values are likelihood ratio tests for lm, glm and polr models. For lme models an attempt is made to re-fit the model using ML and if successful LRT is used to obtain a global p-value. For coxph models the model is re-run without robust variances with and without each variable and a LRT is presented. If unsuccessful a Wald p-value is returned. For GEE and CRR models Wald global p-values are returned.

The number of decimal places to display the statistics can be changed with digits, but this will not change the display of p-values. If more significant digits are required for p-values then use tableOnly=TRUE and format as desired.

## Value

A character vector of the table source code, unless tableOnly=TRUE in which case a data frame is returned

## See Also

[uvsum](#), [lm](#), [glm](#), [crr](#), [coxph](#), [lme](#), [geeglm](#), [polr](#)

## Examples

```
# Examples are for demonstration and are not meaningful
# Coxph model with 90% CI
rm_uvsum(response = c('os_time', 'os_status'),
covs=c('age', 'sex', 'baseline_ctdna', 'l_size', 'change_ctdna_group'),
data=pembrolizumab, CIwidth=.9)

# Linear model with default 95% CI
rm_uvsum(response = 'baseline_ctdna',
covs=c('age', 'sex', 'l_size', 'pd11', 'tmb'),
data=pembrolizumab)

# Logistic model with default 95% CI
rm_uvsum(response = 'os_status',
covs=c('age', 'sex', 'l_size', 'pd11', 'tmb'),
data=pembrolizumab, family = binomial)
# Poisson models returned as model list
mList <- rm_uvsum(response = 'baseline_ctdna',
covs=c('age', 'sex', 'l_size', 'pd11', 'tmb'),
data=pembrolizumab, returnModels=TRUE)
# mList$sex$data # will expose the modelled data

# GEE on correlated outcomes
rm_uvsum(response = 'size_change',
covs=c('time', 'ctdna_status'),
gee=TRUE,
id='id', corstr="exchangeable",
family=gaussian("identity"),
data=ctDNA, showN=TRUE)
```

---

 rm\_uv\_mv
 

---



---

*Combine univariate and multivariable regression tables*


---

## Description

This function will combine `rm_uvsum` and `rm_mvsum` outputs into a single table. The `tableOnly` argument must be set to `TRUE` when tables to be combined are created. The resulting table will be in the same order as the `uvsum` table and will contain the same columns as the `uvsum` and `mvsum` tables, but the p-values will be combined into a single column. There must be a variable overlapping between the `uvsum` and `mvsum` tables and all variables in the `mvsum` table must also appear in the `uvsum` table.

## Usage

```
rm_uv_mv(
  uvsumTable,
  mvsumTable,
  covTitle = "",
  vif = FALSE,
  caption = NULL,
  tableOnly = FALSE,
  chunk_label
)
```

## Arguments

<code>uvsumTable</code>	Output from <code>rm_uvsum</code> , with <code>tableOnly=TRUE</code>
<code>mvsumTable</code>	Output from <code>rm_mvsum</code> , with <code>tableOnly=TRUE</code>
<code>covTitle</code>	character with the names of the covariate (predictor) column. The default is to leave this empty for output or, for table only output to use the column name 'Covariate'.
<code>vif</code>	boolean indicating if the variance inflation factor should be shown if present in the <code>mvsumTable</code> . Default is <code>FALSE</code> .
<code>caption</code>	table caption
<code>tableOnly</code>	boolean indicating if unformatted table should be returned
<code>chunk_label</code>	only used if output is to Word to allow cross-referencing

## Value

A character vector of the table source code, unless `tableOnly=TRUE` in which case a data frame is returned

## See Also

[rm\\_uvsum](#), [rm\\_mvsum](#)

**Examples**

```

require(survival)

uvTab <- rm_uvsum(response = c('os_time', 'os_status'),
  covs=c('age', 'sex', 'baseline_ctdna', 'l_size', 'change_ctdna_group'),
  data=pembrolizumab, tableOnly=TRUE)
mv_surv_fit <- coxph(Surv(os_time, os_status)~age+sex+
  baseline_ctdna+l_size+change_ctdna_group, data=pembrolizumab)
uvTab <- rm_mvsum(mv_surv_fit)

#linear model
uvtab<-rm_uvsum(response = 'baseline_ctdna',
  covs=c('age', 'sex', 'l_size', 'pd11', 'tmb'),
  data=pembrolizumab, tableOnly=TRUE)
lm_fit=lm(baseline_ctdna~age+sex+l_size+tmb, data=pembrolizumab)
mvtab<-rm_mvsum(lm_fit, tableOnly = TRUE)
rm_uv_mv(uvtab, mvtab, tableOnly=TRUE)

#logistic model
uvtab<-rm_uvsum(response = 'os_status',
  covs=c('age', 'sex', 'l_size', 'pd11', 'tmb'),
  data=pembrolizumab, family = binomial, tableOnly=TRUE)
logis_fit<-glm(os_status~age+sex+l_size+pd11+tmb, data = pembrolizumab, family = 'binomial')
mvtab<-rm_mvsum(logis_fit, tableOnly = TRUE)
rm_uv_mv(uvtab, mvtab, tableOnly=TRUE)

```

---

 sanitizestr

*Sanitizes strings to not break LaTeX*


---

**Description**

Strings with special characters will break LaTeX if returned 'asis' by knitr. This happens every time we use one of the main reportRx functions. We first sanitize our strings with this function to stop LaTeX from breaking.

**Usage**

```
sanitizestr(str)
```

**Arguments**

str                    a vector of strings to sanitize

---

testData	<i>Funky ctDNA data There is a weird factor with all one level, the cohort variable contains a cohort level (and Cohort A) and for one cohort all the size changes are missing</i>
----------	--

---

### Description

Funky ctDNA data

There is a weird factor with all one level, the cohort variable contains a cohort level (and Cohort A) and for one cohort all the size changes are missing

### Usage

```
testData
```

### Format

A data frame with 270 rows and 6 variables:

**id** Patient ID

**cohort** Study Cohort: A = Squamous cell carcinoma of soft pallate, B = Triple negative breast cancer, C = Ovarian, high grade serous, D = Melanoma, E = Other Solid Tumor, cohort -for testing only

**badfactor** for testing

**ctdna\_status** Change in ctDNA since baseline

**time** Number of weeks on treatment

**size\_change** Percentage change in tumour measurement

### Source

<https://www.nature.com/articles/s43018-020-0096-5>

---

uvsum	<i>Get univariate summary dataframe</i>
-------	---

---

### Description

Returns a dataframe corresponding to a univariate regression table



**Usage**

```
uvsum(
  response,
  covs,
  data,
  digits = 2,
  id = NULL,
  corstr = NULL,
  family = NULL,
  type = NULL,
  gee = FALSE,
  strata = 1,
  markup = TRUE,
  sanitize = TRUE,
  nicenames = TRUE,
  showN = TRUE,
  CIwidth = 0.95,
  reflevel = NULL,
  returnModels = FALSE
)
```

**Arguments**

response	string vector with name of response
covs	character vector with the names of columns to fit univariate models to
data	dataframe containing data
digits	number of digits to round to
id	character vector which identifies clusters. Used for GEE and coxph models.
corstr	character string specifying the correlation structure. Only used for geeglm. The following are permitted: "independence", "exchangeable", "ar1", "unstructured" and "userdefined"
family	specify details of the model used. This argument does not need to be specified and should be used with caution. By default, gaussian errors are used for linear models, the binomial family with logit link is used for logistic regression and poisson with log link is used for poisson regression. This can be specified with the type argument, or will be inferred from the data type. See <a href="#">family</a> . Ignored for ordinal and survival regression and if the type argument is not explicitly specified.
type	string indicating the type of univariate model to fit. The function will try and guess what type you want based on your response. If you want to override this you can manually specify the type. Options include "linear", "logistic", "poisson", "coxph", "crr", "boxcox" and "ordinal"
gee	boolean indicating if gee models should be fit to account for correlated observations. If TRUE then the id argument must specify the column in the data which indicates the correlated clusters.
strata	character vector of covariates to stratify by. Only used for coxph and crr

markup	boolean indicating if you want latex markup
sanitize	boolean indicating if you want to sanitize all strings to not break LaTeX
nicenames	boolean indicating if you want to replace . and _ in strings with a space
showN	boolean indicating if you want to show sample sizes
CIwidth	width of confidence interval, default is 0.95
reflevel	manual specification of the reference level. Only used for ordinal. This may allow you to debug if the function throws an error.
returnModels	boolean indicating if a list of fitted models should be returned.

### Details

Univariate summaries for a number of covariates, the type of model can be specified. If unspecified the function will guess the appropriate model based on the response variable.

Confidence intervals are extracted using `confint` where possible. Otherwise Student t distribution is used for linear models and the Normal distribution is used for proportions.

`returnModels` can be used to return a list of the univariate models, which will be the same length as `covs`. The data used to run each model will include all cases with observations on the response and covariate. For gee models the data are re-ordered so that the ids appear sequentially and proper estimates are given.

### See Also

[lm](#), [glm](#), [crr](#), [coxph](#), [lme](#), [geeglm](#), [polr](#)

# Index

- \* **dataframe**
  - covsum, 4
  - mvsum, 16
  - rm\_covsum, 25
  - uvsum, 40
- \* **datasets**
  - ctDNA, 7
  - pembrolizumab, 21
  - testData, 40
- \* **helper**
  - addspace, 3
  - cap, 4
  - formatp, 9
  - hbld, 15
  - lbld, 15
  - lpvalue, 16
  - nicename, 19
  - niceNum, 19
  - psthr, 24
  - pstprn, 24
  - pvalue, 24
  - rmds, 25
  - sanitizestr, 39
- \* **model**
  - boxcoxfitRx, 3
  - crrRx, 6
- \* **plot**
  - forestplot2, 8
  - plotuv, 22
- addspace, 3
- anova, 6, 27
- boxcoxfitRx, 3
- cap, 4
- chisq.test, 6, 27
- covsum, 4, 27
- coxph, 37, 42
- cramer\_v, 27
- crr, 6, 37, 42
- crrRx, 6
- ctDNA, 7
- eta\_squared, 27
- excelCol, 7
- family, 41
- fisher.test, 6, 27
- forestplot2, 8
- formatp, 9
- geeglm, 37, 42
- geoR\_boxcoxfit, 9
- ggarrange, 23
- ggkmcif, 10
- ggkmcif\_paste, 14
- ggplot, 23
- glm, 37, 42
- hbld, 15
- kruskal.test, 6, 27
- lbld, 15
- lm, 37, 42
- lme, 37, 42
- lpvalue, 16
- mvsum, 16
- nestTable, 17
- nicename, 19
- niceNum, 19
- outTable, 20, 27
- pembrolizumab, 21
- plotuv, 22
- polr, 37, 42
- psthr, 24

pstprn, [24](#)  
pvalue, [24](#)

rm\_covsum, [25](#)  
rm\_mvsum, [28](#), [38](#)  
rm\_survdif, [30](#), [32](#)  
rm\_survsum, [31](#)  
rm\_survtime, [32](#), [33](#)  
rm\_uv\_mv, [38](#)  
rm\_uvsum, [35](#), [38](#)  
rmds, [25](#)

sanitizestr, [39](#)  
survdif, [31](#)  
survfit, [30](#), [32](#), [34](#)

testData, [40](#)

uvsum, [22](#), [37](#), [40](#)

wilcox.test, [6](#), [27](#)