

# Package ‘Publish’

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**Type** Package

**Title** Format Output of Various Routines in a Suitable Way for Reports and Publication

**Description**

A bunch of convenience functions that transform the results of some basic statistical analyses into table format nearly ready for publication. This includes descriptive tables, tables of logistic regression and Cox regression results as well as forest plots.

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**Maintainer** Thomas A. Gerds <tag@biostat.ku.dk>

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**License** GPL (>= 2)

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**Author** Thomas A. Gerds [aut, cre],  
Christian Torp-Pedersen [ctb],  
Klaus K Holst [ctb],  
Brice Ozenne [aut]

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Publish-package	<i>Publish package</i>
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### Description

This package processes results of descriptive statistics and regression analysis into final tables and figures of a manuscript

acut	<i>Automatic selection and formatting of breaks in cut</i>
------	--

### Description

A version of cut that easily formats the labels and places breaks by default.

### Usage

```
acut(
  x,
  n = 5,
  type = "default",
  format = NULL,
  format.low = NULL,
  format.high = NULL,
  dig.lab = 3,
  right = TRUE,
  breaks,
  labels = TRUE,
  ...
)
```

### Arguments

x	a numeric vector which is to be converted to a factor by cutting (passed directly to cut).
n	number of bins to create based on the empirical quantiles of x. This will be overruled if breaks is supplied.
type	a high-level formatting option. For now, the only other option than the default setting is "age". See details and examples.
format	string used to make labels. %l and %u identifies the lower and upper value of the breaks respectively. See examples.

<code>format.low</code>	string used specifically on the lowest label.
<code>format.high</code>	string used specifically on the highest label.
<code>dig.lab</code>	integer which is used when labels are not given. It determines the number of digits used in formatting the break numbers. (Passed directly to <code>cut</code> .)
<code>right</code>	logical, indicating if the intervals should be closed on the right (and open on the left) or vice versa (passed directly to <code>cut</code> ).
<code>breaks</code>	specify breaks manually as in <code>cut</code> .
<code>labels</code>	logical, indicating whether or not to make labels or simply use ordered numbers. If <code>TRUE</code> , the labels are constructed as discribed above.
<code>...</code>	further arguments passed to <code>cut</code> .

### Details

The formats are supplied by specifying the text around the lower (`%l`) and upper (`%u`) value (see examples). If user specified breaks are supplied, the default labels from `cut` are used. If automatic breaks are used, the default labels are a slight modification at the end point of the default from `cut`. All this can of course be adjusted manually through the format functionality (see below).

By default, 5 breaks are constructed according to the quantiles with of the input `x`. The number of breaks can be adjusted, and default specifying breaks (as in `cut`) can be supplied instead.

If `type` is changed from "default" to another option, a different formatting template is used. For now the only other option is "age", which is designed to be well suited to easily group age variables. When `type="age"` only the `breaks` argument is used, and it behaves different from otherwise. If a single number is supplied, intervals of length `breaks` will automatically be constructed (starting from 0). If a vector is supplied, the intervals are used as in `cut` but formatted differently, see examples.

### Value

same as for `cut`. A vector of 'factors' is created, unless 'labels=FALSE'.

### Author(s)

Anders Munch

### Examples

```
data(Diabetes) # load dataset

## The default uses format similar to cut
chol.groups <- acut(Diabetes$chol)
table(chol.groups)

## The formatting can easily be changed
chol.groups <- acut(Diabetes$chol,format="%l-%u",n=5)
table(chol.groups)

## The default is to automatic place the breaks, so the number of this can easily be changed.
chol.groups <- acut(Diabetes$chol,n=7)
```

```

table(chol.groups)

## Manually setting format and breaks
age.groups <- acut(Diabetes$age,format="%l-%u",breaks=seq(0,100,by=10))
table(age.groups)

## Other variations
age.groups <- acut(Diabetes$age,
                  format="%l-%u",
                  format.low="below %u",
                  format.high="above %l",
                  breaks=c(0, seq(20,80,by=10), Inf))
table(age.groups)

BMI.groups <- acut(Diabetes$BMI,
                  format="BMI between %l and %u",
                  format.low="BMI below %u",
                  format.high="BMI above %l")
table(BMI.groups)
org(as.data.frame(table(BMI=BMI.groups)))

## Instead of using the quantiles, we can specify equally spaced breaks,
## but still get the same formatting
BMI.grouping <-
  seq(min(Diabetes$BMI,na.rm=TRUE), max(Diabetes$BMI,na.rm=TRUE), length.out=6)
BMI.grouping[1] <- -Inf # To get all included
BMI.groups <- acut(Diabetes$BMI,
                  breaks=BMI.grouping,
                  format="BMI between %l and %u",
                  format.low="BMI below %u",
                  format.high="BMI above %l")
table(BMI.groups)
org(as.data.frame(table(BMI=BMI.groups)))

## Using type="age"
## When using type="age", categories of 10 years are constructed by default.
## The are formatted to be easier to read when the values are ages.
table(acut(Diabetes$age, type="age"))

## This can be changes with the breaks argument.
## Note that this is diffent from cut when breaks is a single number.
table(acut(Diabetes$age, type="age", breaks=20))

## Of course We can also supply the breaks manually.
## The formatting depends on whether or not all the values fall within the breaks:
## All values within the breaks
table(acut(Diabetes$age, type="age", breaks=c(0, 30, 50, 80, 100)))
## Some values below and above the breaks
table(acut(Diabetes$age, type="age", breaks=c(30, 50, 80)))

```

---

ci.mean	<i>Compute mean values with confidence intervals</i>
---------	--

---

**Description**

Compute mean values with confidence intervals

**Usage**

```
ci.mean(x, ...)
```

**Arguments**

x	object passed to methods
...	passed to methods

**Details**

Normal approximation

**Value**

a list with mean values and confidence limits

---

ci.mean.default	<i>Compute mean values with confidence intervals</i>
-----------------	--

---

**Description**

Compute mean values with confidence intervals

**Usage**

```
## Default S3 method:  
ci.mean(  
  x,  
  alpha = 0.05,  
  normal = TRUE,  
  na.rm = TRUE,  
  statistic = "arithmetic",  
  ...  
)
```

**Arguments**

x	numeric vector
alpha	level of significance
normal	If TRUE use quantile of t-distribution else use normal approximation and quantile of normal approximation. Do you think this is confusing?
na.rm	If TRUE remove missing values from x.
statistic	Decide which mean to compute: either "arithmetic" or "geometric"
...	not used

**Details**

Normal approximation

**Value**

a list with mean values and confidence limits

**Author(s)**

Thomas Gerds

---

CiTable

*CiTable data*

---

**Description**

These data are used for testing Publish package functionality.

**Format**

A data frame with 27 observations on the following 9 variables.

**Drug**

**Time**

**Drug.Time**

**Dose**

**Mean**

**SD**

**n**

**HazardRatio**

**lower**

**upper**

**p**

**Examples**

```

data(CiTable)
labellist <- split(CiTable[,c("Dose", "Mean", "SD", "n")], CiTable[, "Drug"])
labellist
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")], labels=labellist)

```

---

coxphSeries

*Run a series of Cox regression models*


---

**Description**

Run a series of Cox regression analyses for a list of predictor variables and summarize the results in a table. The Cox models can be adjusted for a fixed set of covariates

This function runs on coxph from the survival package.

**Usage**

```
coxphSeries(formula, data, vars, ...)
```

**Arguments**

formula	The fixed part of the regression formula. For univariate analyses this is simply <code>Surv(time,status)~1</code> where <code>Surv(time,status)</code> is the outcome variable. When the aim is to control the effect of vars in each element of the series by a fixed set of variables it is <code>Surv(time,status)~x1+x2</code> where again <code>Surv(time,status)</code> is the outcome and x1 and x2 are confounders.
data	A data.frame in which the formula gets evaluated.
vars	A list of variable names, the changing part of the regression formula.
...	passed to <code>publish.coxph</code>

**Value**

matrix with results

**Author(s)**

Thomas Alexander Gerds



**Examples**

```

library(survival)
data(pbc)
## collect hazard ratios from three univariate Cox regression analyses
pbc$edema <- factor(pbc$edema,levels=c("0","0.5","1"),labels=c("0","0.5","1"))
uni.hr <- coxphSeries(Surv(time,status==2)~1,vars=c("edema","bili","protime"),data=pbc)
uni.hr

## control the logistic regression analyses for age and gender
## but collect only information on the variables in `vars`.
controlled.hr <- coxphSeries(Surv(time,status==2)~age+sex,vars=c("edema","bili","protime"),data=pbc)
controlled.hr

```

---

Diabetes

*Diabetes data of Dr John Schorling*


---

**Description**

These data are courtesy of Dr John Schorling, Department of Medicine, University of Virginia School of Medicine. The data consist of 19 variables on 403 subjects from 1046 subjects who were interviewed in a study to understand the prevalence of obesity, diabetes, and other cardiovascular risk factors in central Virginia for African Americans. According to Dr John Hong, Diabetes Mellitus Type II (adult onset diabetes) is associated most strongly with obesity. The waist/hip ratio may be a predictor in diabetes and heart disease. DM II is also associated with hypertension - they may both be part of "Syndrome X". The 403 subjects were the ones who were actually screened for diabetes. Glycosolated hemoglobin > 7.0 is usually taken as a positive diagnosis of diabetes.

**Format**

A data frame with 205 observations on the following 12 variables.

**id** subject id  
**chol** Total Cholesterol  
**stab.glu** Stabilized Glucose  
**hdl** High Density Lipoprotein  
**ratio** Cholesterol/HDL Ratio  
**glyhb** Glycosolated Hemoglobin  
**location** a factor with levels (Buckingham,Louisa)  
**age** age (years)  
**gender** male or female  
**height** height (inches)  
**height.europe** height (cm)  
**weight** weight (pounds)

**weight.europe** weight (kg)  
**frame** a factor with levels (small,medium,large)  
**bp.1s** First Systolic Blood Pressure  
**bp.1d** First Diastolic Blood Pressure  
**bp.2s** Second Diastolic Blood Pressure  
**bp.2d** Second Diastolic Blood Pressure  
**waist** waist in inches  
**hip** hip in inches  
**time.ppn** Postprandial Time when Labs were Drawn in minutes  
**AgeGroups** Categorized age  
**BMI** Categorized BMI

## References

Willems JP, Saunders JT, DE Hunt, JB Schorling: Prevalence of coronary heart disease risk factors among rural blacks: A community-based study. Southern Medical Journal 90:814-820; 1997  
 Schorling JB, Roach J, Siegel M, Baturka N, Hunt DE, Guterbock TM, Stewart HL: A trial of church-based smoking cessation interventions for rural African Americans. Preventive Medicine 26:92-101; 1997.

## Examples

```
data(Diabetes)
```

---

```
fixRegressionTable    Expand regression coefficient table
```

---

## Description

Expand regression coefficient table

## Usage

```

fixRegressionTable(
  x,
  varnames,
  reference.value,
  reference.style = NULL,
  factorlevels,
  scale = NULL,
  nmiss,
  intercept
)

```

**Arguments**

x	object resulting from lm, glm or coxph.
varnames	Names of variables
reference.value	Reference value for reference categories
reference.style	Style for showing results for categorical variables. If "extraline" show an additional line for the reference category.
factorlevels	Levels of the categorical variables.
scale	Scale for some or all of the variables
nmiss	Number of missing values
intercept	Intercept

**Details**

This function expands results from "regressionTable" with extralines and columns

For factor variables the reference group is shown. For continuous variables the units are shown and for transformed continuous variables also the scale. For all variables the numbers of missing values are added.

**Value**

a table with regression coefficients

**Author(s)**

Thomas Alexander Gerds <tag@biostat.ku.dk>

---

followupTable	<i>Summary tables for a given followup time point.</i>
---------------	--

---

**Description**

Summarize baseline variables in groups defined by outcome at a given followup time point

**Usage**

```
followupTable(formula, data, followup.time, compare.groups, ...)
```

**Arguments**

formula	Formula A formula whose left hand side is a Hist object. In some special cases it can also be a Surv response object. The right hand side is as in <a href="#">utable</a> .
data	A data.frame in which all the variables of formula can be interpreted.
followup.time	Time point at which to evaluate outcome status.
compare.groups	Method for comparing groups.
...	Passed to utable. All arguments of utable can be controlled in this way except for compare.groups which is set to "Cox". See details.

**Details**

If compare.groups!=FALSE, p-values are obtained from stopped Cox regression, i.e., all events are censored at follow-up time. A univariate Cox regression model is fitted to assess the effect of each variable on the right hand side of the formula on the event hazard and shown is the p-value of anova(fit), see [anova.coxph](#).

**Value**

Summary table.

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

univariateTable

**Examples**

```
library(survival)
data(pbc)
pbc$edema <- factor(pbc$edema, levels=c("0", "0.5", "1"), labels=c("0", "0.5", "1"))
pbc$sex <- factor(pbc$sex, levels=c("m", "f"), labels=c("m", "f"))
followupTable(Hist(time, status)~age+edema+sex, data=pbc, followup.time=1000)
```

---

formatCI

*Formatting confidence intervals*


---

**Description**

Format confidence intervals

**Usage**

```
formatCI(
  x,
  lower,
  upper,
  show.x = FALSE,
  handler = "sprintf",
  format = "[l;u]",
  degenerated = "asis",
  digits = 2,
  nsmall = digits,
  sep = "",
  reference.pos,
  reference.label = "",
  ...
)
```

**Arguments**

x	not used (for compatibility with format)
lower	Numeric vector of lower limits
upper	Numeric vector of upper limits
show.x	Logical. If TRUE show value of x in front of confidence interval.
handler	Function to format numeric values. Default is <code>sprintf</code> , also supported are <code>format</code> and <code>prettyNum</code>
format	Character string in which l will be replaced by the value of the lower limit (argument lower) and u by the value of the upper upper limit. For example, (l,u) yields confidence intervals in round parenthesis in which the upper and lower limits are comma separated. Default is [l;u].
degenerated	String to show when lower==upper. Default is '-'
digits	If handler <code>format</code> or <code>prettyNum</code> used format numeric vectors.
nsmall	If handler <code>format</code> or <code>prettyNum</code> used format numeric vectors.
sep	Field separator
reference.pos	Position of factor reference
reference.label	Label for factor reference
...	passed to handler

**Details**

The default format for confidence intervals is [lower; upper].

**Value**

String vector with confidence intervals

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

plot.ci ci.mean

**Examples**

```
x=ci.mean(rnorm(10))
formatCI(lower=x[3],upper=x[4])
formatCI(lower=c(0.001,-2.8413),upper=c(1,3.0008884))
# change format
formatCI(lower=c(0.001,-2.8413),upper=c(1,3.0008884),format="(l, u)")
# show x
formatCI(x=x$mean,lower=x$lower,upper=x$upper,format="(l, u)",show.x=TRUE)

# change of handler function
l <- c(-0.0890139,0.0084736,144.898333,0.000000001)
u <- c(0.03911392,0.3784706,3338944.8821221,0.00001)
cbind(format=formatCI(lower=l,upper=u,format="[l;u]",digits=2,nsmall=2,handler="format"),
      prettyNum=formatCI(lower=l,upper=u,format="[l;u]",digits=2,nsmall=2,handler="prettyNum"),
      sprintf=formatCI(lower=l,upper=u,format="[l;u]",digits=2,nsmall=2,handler="sprintf"))
```

---

glmSeries

*Run a series of generalized linear regression analyses*

---

**Description**

Run a series of generalized linear regression analyses for a list of predictor variables and summarize the results in a table. The regression models can be adjusted for a fixed set of covariates.

**Usage**

```
glmSeries(formula, data, vars, ...)
```

**Arguments**

formula	The fixed part of the regression formula. For univariate analyses this is simply $y \sim 1$ where $y$ is the outcome variable. When the aim is to control the effect of vars in each element of the series by a fixed set of variables it is $y \sim x_1 + x_2$ where again $y$ is the outcome and $x_1$ and $x_2$ are confounders.
data	A data.frame in which we evaluate the formula.
vars	A list of variable names, the changing part of the regression formula.
...	passed to glm

**Value**

Matrix with regression coefficients, one for each element of vars.

**Author(s)**

Thomas Alexander Gerds

**Examples**

```
data(Diabetes)
Diabetes$hyper1 <- factor(1*(Diabetes$bp.1s>140))
## collect odds ratios from three univariate logistic regression analyses
uni.odds <- glmSeries(hyper1~1,vars=c("chol","hdl","location"),data=Diabetes,family=binomial)
uni.odds
## control the logistic regression analyses for age and gender
## but collect only information on the variables in `vars`.
controlled.odds <- glmSeries(hyper1~age+gender,
                             vars=c("chol","hdl","location"),
                             data=Diabetes, family=binomial)
controlled.odds
```

---

labelUnits

*labelUnits*

---

**Description**

Label output tables

**Usage**

```
labelUnits(x, ...)
```

**Arguments**

x	A matrix obtained with univariateTable.
...	not used

**Details**

Modify labels and values of variables in summary tables

**Value**

The re-labeled matrix

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

univariateTable

**Examples**

```
data(Diabetes)
tab <- summary(univariateTable(gender~AgeGroups+chol+waist,data=Diabetes))
publish(tab)
ltab <- labelUnits(tab,"chol"="Cholesterol (mg/dL)","<40"="younger than 40")
publish(ltab)

## pass labels immediately to utable
utable(gender~AgeGroups+chol+waist,data=Diabetes,
       "chol"="Cholesterol (mg/dL)","<40"="younger than 40")

## sometimes useful to state explicitly which variables value
## should be re-labelled
utable(gender~AgeGroups+chol+waist,data=Diabetes,
       "chol"="Cholesterol (mg/dL)","AgeGroups.<40"="younger than 40")
```

---

lazyDateCoding

*Efficient coding of date variables*

---

**Description**

This function eases the process of generating date variables. All variables in a data.frame which match a regular expression are included

**Usage**

```
lazyDateCoding(data, format, pattern, varnames, testlength = 10)
```

**Arguments**

data	Data frame in which to search for date variables.
format	passed to as.Date
pattern	match date variables
varnames	variable names
testlength	how many rows of data should be evaluated to guess the format.

**Details**

The code needs to be copy-and-pasted from the R-output buffer into the R-code buffer. This can be customized for the really efficiently working people, e.g., in emacs.



**Value**

R-code one line for each variable.

**Author(s)**

Thomas Alexander Gerds

**Examples**

```
d <- data.frame(x0="190101", x1=c("12/8/2019"), x2="12-8-2019", x3="20190812", stringsAsFactors=FALSE)
lazyDateCoding(d, pattern="x")
lazyDateCoding(d, pattern="3")
```

---

lazyFactorCoding      *Efficient coding of factor levels*

---

**Description**

This function eases the process of generating factor variables with relevant labels. All variables in a data.frame with less than a user set number of levels result in a line which suggests levels and labels. The result can then be modified for use.

**Usage**

```
lazyFactorCoding(data, max.levels = 10)
```

**Arguments**

data	Data frame in which to search for categorical variables.
max.levels	Treat non-factor variables only if the number of unique values less than max.levels. Defaults to 10.

**Details**

The code needs to be copy-and-pasted from the R-output buffer into the R-code buffer. This can be customized for the really efficiently working people e.g. in emacs.

**Value**

R-code one line for each variable.

**Author(s)**

Thomas Alexander Gerds

**Examples**

```
data(Diabetes)
lazyFactorCoding(Diabetes)
```

---

org

*Wrapper function for publish with output format org*

---

**Description**

Wrapper for `publish(..., org=TRUE)`

**Usage**

```
org(x, ...)
```

**Arguments**

x	object to format as org
...	passed to publish

**Value**

See `publish`

**Author(s)**

Thomas Alexander Gerds

---

parseInteractionTerms *Parse interaction terms*

---

**Description**

Parse interaction terms for regression tables

**Usage**

```

parseInteractionTerms(
  terms,
  xlevels,
  units,
  format.factor,
  format.contrast,
  format.scale,
  format.scale.unit,
  sep = ": ",
  ...
)

```

**Arguments**

terms	Terms of a formula
xlevels	Factor levels corresponding to the variables in terms
units	named list with unit labels. names should match variable names in formula.
format.factor	For categorical variables. A string which specifies the print format for factor labels. The string has to contain the keywords "var" and "level" which will be replaced by the name of the variable and the current level, respectively. Default is "var(level)".
format.contrast	For categorical variables. A string which specifies the print format for contrast statements. The string has to contain the keywords "var", "level" and "ref" which will be replaced by the name of the variable, the current level and the reference level, respectively.
format.scale	A string which specifies the print format for continuous variables without units. The string has to contain the keyword "var" which will be replaced by the name of the variable and the unit, respectively. Default is "var".
format.scale.unit	A string which specifies the print format for continuous variables with units. The string has to contain the keywords "var" and "unit" which will be replaced by the name of the variable and the unit, respectively. Default is "var(unit)".
sep	a character string to separate the terms. Default is ": ".
...	Not yet used

**Details**

Prepare a list of contrasts which combines regression coefficients to describe statistical interactions.

**Value**

List of contrasts which can be passed to `lava::estimate`.

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

lava::estimate

**Examples**

```
tt <- terms(formula(SBP~age+sex*BMI))
xlev <- list(sex=c("male", "female"), BMI=c("normal", "overweight", "obese"))
parseInteractionTerms(terms=tt, xlevels=xlev)
parseInteractionTerms(terms=tt, xlevels=xlev, format.factor="var level")
parseInteractionTerms(terms=tt, xlevels=xlev, format.contrast="var(level:ref)")

tt2 <- terms(formula(SBP~age*factor(sex)+BMI))
xlev2 <- list("factor(sex)"=c("male", "female"))
parseInteractionTerms(terms=tt2, xlevels=xlev2)
parseInteractionTerms(terms=tt2, xlevels=xlev2, units=list(age="yrs"))

data(Diabetes)
fit <- glm(bp.2s~age*factor(gender)+BMI, data=Diabetes)
parseInteractionTerms(terms=terms(fit$formula), xlevels=fit$xlevels,
                      format.scale="var -- level:ref", units=list("age"='years'))
parseInteractionTerms(terms=terms(fit$formula), xlevels=fit$xlevels,
                      format.scale.unit="var [unit]", units=list("age"='years'))
it <- parseInteractionTerms(terms=terms(fit$formula), xlevels=fit$xlevels)
ivars <- unlist(lapply(it, function(x) attr(x, "variables")))
lava::estimate(fit, function(p) lapply(unlist(it), eval, envir=sys.parent(-1)))
```

---

plot.ci

*Plot confidence intervals*

---

**Description**

Function to plot confidence intervals

**Usage**

```
## S3 method for class 'ci'
plot(x, xlim, xlab = "", labels, ...)
```

**Arguments**

x	List, data.frame or other object of this form containing point estimates (first element) and the corresponding confidence intervals as elements lower and upper.
xlim	Limit of the x-axis
xlab	Label for the y-axis
labels	labels
...	Used to transport arguments to plotConfidence.

**Details**

Function to plot means and other point estimates with confidence intervals

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**Examples**

```

data(Diabetes)
x=ci.mean(bp.2s~AgeGroups,data=Diabetes)
plot(x,title.labels="Age groups",xratio=c(0.4,0.3))
x=ci.mean(bp.2s/500~AgeGroups+gender,data=Diabetes)
plot(x,xratio=c(0.4,0.2))
plot(x,xratio=c(0.4,0.2),
      labels=split(x$labels["AgeGroups"],x$labels["gender"]),
      title.labels="Age groups")
## Not run:
plot(x, leftmargin=0, rightmargin=0)
plotConfidence(x, leftmargin=0, rightmargin=0)

data(CiTable)
with(CiTable,plotConfidence(x=list(HazardRatio),
                             lower=lower,
                             upper=upper,
                             labels=CiTable[,2:6],
                             factor.reference.pos=c(1,10,19),
                             format="(u-1)",
                             points.col="blue",
                             digits=2))

with(CiTable,Publish::plot.ci(x=list(HazardRatio),
                               lower=lower,
                               upper=upper,
                               labels=CiTable[,2:6],
                               factor.reference.pos=c(1,10,19),
                               format="(u-1)",
                               points.col="blue",
                               digits=2,
                               leftmargin=-2,

```

```

title.labels.cex=1.1,
labels.cex=0.8,values.cex=0.8))

## End(Not run)

```

---

plot.regressionTable *Plotting regression coefficients with confidence limits*

---

## Description

Plotting regression coefficients with confidence limits

## Usage

```

## S3 method for class 'regressionTable'
plot(x, xlim, xlab, style = 1, ...)

```

## Arguments

x	regression table obtained with regressionTable
xlim	Limits for x-axis
xlab	Label for x-axis
style	Determines how to arrange variable names and their corresponding units
...	passed to plotConfidence

## Author(s)

Thomas A. Gerds <tag@biostat.ku.dk>

## See Also

regressionTable

## Examples

```

## linear regression
data(Diabetes)
f <- glm(bp.1s~AgeGroups+chol+gender+location,data=Diabetes)
rtf <- regressionTable(f,factor.reference = "inline")
plot(rtf,cex=1.3)

## logistic regression
data(Diabetes)
f <- glm(I(BMI>25)~bp.1s+AgeGroups+chol+gender+location,data=Diabetes,family=binomial)
rtf <- regressionTable(f,factor.reference = "inline")
plot(rtf,cex=1.3)

## Poisson regression

```

```
data(trace)
fit <- glm(dead ~ smoking+ sex+ age+Time+offset(log(ObsTime)), family = poisson,data=trace)
rtab <- regressionTable(fit,factor.reference = "inline")
plot(rtab,xlim=c(0.85,1.15),cex=1.8,xaxis.cex=1.5)

## Cox regression
library(survival)
data(pbc)
coxfit <- coxph(Surv(time,status!=0)~age+log(bili)+log(albumin)+factor(edema)+sex,data=pbc)
pubcox <- publish(coxfit)
plot(pubcox,cex=1.5,xratio=c(0.4,0.2))
```

---

plot.subgroupAnalysis *plot.subgroupAnalysis*

---

## Description

This function operates on a "subgroupAnalysis" object to produce a formatted table and a forest plot

## Usage

```
## S3 method for class 'subgroupAnalysis'
plot(x, ...)
```

## Arguments

x                   - a subgroupAnalysis object  
...                   - passed on to plotConfidence

## Details

This function produces a formatted table of a subgroupAnalysis object and adds a forest plot. If further details needs attention before plotting is is advisable use adjust the table produced by the summary function and then plotting with the plotConfidence function

## Author(s)

Christian Torp-Pedersen

## See Also

subgroupAnalysis, plotConfidence

**Examples**

```

#load libraries
library(Publish)
library(survival)
library(data.table)
data(traceR) #get dataframe traceR
setDT(traceR)
traceR[, ':='(wmi2=factor(wallMotionIndex<0.9, levels=c(TRUE, FALSE),
                      labels=c("bad", "good")),
            abd2=factor(abdominalCircumference<95, levels=c(TRUE, FALSE),
                      labels=c("slim", "fat")),
            sex=factor(sex))]
fit_cox <- coxph(Surv(observationTime, dead)~treatment, data=traceR)
# Selected subgroups - univariable analysis
sub_cox <- subgroupAnalysis(fit_cox, traceR, treatment="treatment",
                           subgroup=c("smoking", "sex", "wmi2", "abd2")) # subgroups as character string
plot(sub_cox)

```

---

plotConfidence

*Plot confidence intervals*


---

**Description**

Function to plot confidence intervals with their values and additional labels. One anticipated use of this function involves first the generation of a regression object, then arrangement of a result table with "regressionTable", further arrangement of table with with e.g. "fixRegressionTable" and various user defined changes - and then finally table along with forest plot using the current function.

**Usage**

```

plotConfidence(
  x,
  y.at,
  lower,
  upper,
  pch = 16,
  cex = 1,
  lwd = 1,
  col = 4,
  xlim,
  xlab,
  labels,
  title.labels,
  values,
  title.values,
  section.pos,
  section.sep,
  section.title = NULL,

```



```

    section.title.x,
    section.title.offset,
    order,
    leftmargin = 0.025,
    rightmargin = 0.025,
    stripes,
    factor.reference.pos,
    factor.reference.label = "Reference",
    factor.reference.pch = 16,
    refline = 1,
    title.line = TRUE,
    xratio,
    y.offset = 0,
    y.title.offset,
    digits = 2,
    format,
    extremearrows.length = 0.05,
    extremearrows.angle = 30,
    add = FALSE,
    layout = TRUE,
    xaxis = TRUE,
    ...
)

```

### Arguments

x	Either a vector containing the point estimates or a list whose first element contains the point estimates. Further list elements can contain the confidence intervals and labels. In this case the list needs to have names 'lower' and 'upper' to indicate the values of the lower and the upper limits of the confidence intervals, respectively, and may have an element 'labels' which is a vector or matrix or list with labels.
y.at	Optional vector of y-position for the confidence intervals and corresponding values and labels.
lower	Lower confidence limits. Used if object x is a vector and if x is a list lower overwrites element x\$lower.
upper	Upper confidence limits. Used if object x is a vector and if x is a list upper overwrites element x\$upper.
pch	Symbol for points.
cex	Defaults size of all figures and plotting symbol. Single elements are controlled separately. See . . . .
lwd	Default width of all lines Single elements are controlled separately. See . . . .
col	Default colour of confidence intervals.
xlim	Plotting limits for the confidence intervals. See also xratio on how to control the layout.
xlab	Label for the x-axis.

<code>labels</code>	Vector or matrix or list with labels. Used if object <code>x</code> is a vector and if <code>x</code> is a list it overwrites element <code>x\$labels</code> . To avoid drawing of labels, set <code>labels=FALSE</code> .
<code>title.labels</code>	Main title for the column which shows the labels. If <code>labels</code> is a matrix or list <code>title.labels</code> should be a vector with as many elements as <code>labels</code> has columns or elements.
<code>values</code>	Either logical or vector, matrix or list with values. If <code>values=TRUE</code> values are constructed according to <code>format</code> from lower and upper overwrites constructed values. If <code>values=FALSE</code> do not draw values.
<code>title.values</code>	Main title for the column values. If <code>values</code> is a matrix or list <code>title.labels</code> should be a vector with as many elements as <code>values</code> has columns or elements.
<code>section.pos</code>	Vector with y-axis positions for <code>section.titles</code> .
<code>section.sep</code>	Amount of space between paragraphs (applies only if <code>labels</code> is a named list)
<code>section.title</code>	Intermediate section headings.
<code>section.title.x</code>	x-position for <code>section.titles</code>
<code>section.title.offset</code>	Y-offset for <code>section.titles</code>
<code>order</code>	Order of the three columns: labels, confidence limits, values. See examples.
<code>leftmargin</code>	Percentage of plotting region used for leftmargin. Default is 0.025. See also Details.
<code>rightmargin</code>	Percentage of plotting region used for rightmargin. Default is 0.025. See also Details.
<code>stripes</code>	Vector of up to three Logicals. If <code>TRUE</code> draw stripes into the background. The first applies to the labels, the second to the graphical presentation of the confidence intervals and the third to the values. Thus, stripes
<code>factor.reference.pos</code>	Position at which factors attain reference values.
<code>factor.reference.label</code>	Label to use at <code>factor.reference.pos</code> instead of values.
<code>factor.reference.pch</code>	Plotting symbol to use at <code>factor.reference.pos</code>
<code>refline</code>	Position of a vertical line to indicate the null hypothesis. Default is 1 which would work for odds ratios and hazard ratios.
<code>title.line</code>	Position of a horizontal line to separate the title line from the plot
<code>xratio</code>	One or two values between 0 and 1 which determine how to split the plot window in horizontal x-direction. If there are two columns (labels, CI) or (CI, values) only one value is used and the default is 0.618 (goldener schnitt) which gives the graphical presentation of the confidence intervals 38.2 graph. The remaining 61.8 If there are three columns (labels, CI, values), <code>xratio</code> has two values which default to fractions of 0.7 according to the relative widths of labels and values, thus by default only 0.3 are used for the graphical presentation of the confidence intervals. The remaining 30 confidence intervals. See examles.

<code>y.offset</code>	Either a single value or a vector determining the vertical offset of all rows. If it is a single value all rows are shifted up (or down if negative) by this value. This can be used to add a second set of confidence intervals to an existing graph or to achieve a visual grouping of rows that belong together. See examples.
<code>y.title.offset</code>	Numeric value by which to vertically shift the titles of the labels and values.
<code>digits</code>	Number of digits, passed to <code>pubformat</code> and <code>formatCI</code> .
<code>format</code>	Format for constructing values of confidence intervals. Defaults to <code>'(u;l)'</code> if there are negative lower or upper values and to <code>'(u-l)'</code> otherwise.
<code>extremearrows.length</code>	Length of the arrows in case of confidence intervals that stretch beyond <code>xlim</code> .
<code>extremearrows.angle</code>	Angle of the arrows in case of confidence intervals that stretch beyond <code>xlim</code> .
<code>add</code>	Logical. If TRUE do not draw labels or values and add confidence intervals to existing plot.
<code>layout</code>	Logical. If FALSE do not call layout. This is useful when several <code>plotConfidence</code> results should be combined in one graph and hence layout is called externally.
<code>xaxis</code>	Logical. If FALSE do not draw x-axis.
<code>...</code>	Used to control arguments of the following subroutines: <code>plot</code> : Applies to plotting frame of the graphical presentation of confidence intervals. Use arguments of <code>plot</code> , e.g., <code>plot.main="Odds ratio"</code> . <code>points</code> , <code>arrows</code> : Use arguments of points and arrows, respectively. E.g., <code>points.pch=8</code> and <code>arrows.lwd=2</code> . <code>refline</code> : Use arguments of segments, e.g., <code>refline.lwd=2</code> . See <a href="#">segments</a> . <code>labels</code> , <code>values</code> , <code>title.labels</code> , <code>title.values</code> : Use arguments of text, e.g., <code>labels.col="red"</code> or <code>title.values.cex=1.8</code> . <code>xaxis</code> : Use arguments of axis, e.g., <code>xaxis.at=c(-0.3,0,0.3)</code> <code>xlab</code> : Use arguments of <code>mtext</code> , e.g., <code>xlab.line=2</code> . <code>stripes</code> : Use arguments of <code>stripes</code> . See examples. See examples for usage.

## Details

Function to plot means and other point estimates with confidence intervals, their values and additional labels. Horizontal margins as determined by `par()$mar` are ignored. Instead `layout` is used to divide the plotting region horizontally into two or three parts plus `leftmargin` and `rightmargin`.

When `values` is FALSE there are only two parts. The default order is labels on the left confidence intervals on the right. When no labels are given or `labels` is FALSE there are only two parts. The default order is confidence intervals on the left values on the right.

The default order of three parts from left to right is labels, confidence intervals, values. The order can be changed as shown by the examples below. The relative widths of the two or three parts need to be adapted to the actual size of the text of the labels. This depends on the plotting device and the size of the font and figures and thus has to be adjusted manually.

`Oma` can be used to further control horizontal margins, e.g., `par(oma=c(0,4,0,4))`.

If confidence limits extend beyond the range determined by `xlim`, then arrows are drawn at the `x-lim` borders to indicate that the confidence limits continue.

## Value

List of dimensions and coordinates

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**Examples**

```

library(Publish)
data(CiTable)

## A first draft version of the plot is obtained as follows
plotConfidence(x=CiTable[,c("HazardRatio","lower","upper","p")],
               labels=CiTable[,c("Drug.Time","Dose","Mean","SD","n")])

## if argument labels is a named list the table is subdivided:
labellist <- split(CiTable[,c("Dose","Time","Mean","SD","n")],CiTable[, "Drug"])
labellist
## the data need to be ordered accordingly
CC= data.table::rbindlist(split(CiTable[,c("HazardRatio","lower","upper")],CiTable[, "Drug"]))
plotConfidence(x=CC, labels=labellist)

## The graph consist of at most three columns:
##
## column 1: labels
## column 2: printed values of the confidence intervals
## column 3: graphical presentation of the confidence intervals
##
## NOTE: column 3 appears always, the user decides if also
##       column 1, 2 should appear
##
## The columns are arranged with the function layout
## and the default order is 1,3,2 such that the graphical
## display of the confidence intervals appears in the middle
##
## the order of appearance of the three columns can be changed as follows
plotConfidence(x=CiTable[,c("HazardRatio","lower","upper")],
               labels=CiTable[,c("Drug.Time","Dose","Mean","SD","n")],
               order=c(1,3,2))
plotConfidence(x=CiTable[,c("HazardRatio","lower","upper")],
               labels=CiTable[,c("Drug.Time","Dose","Mean","SD","n")],
               order=c(2,3,1))
## if there are only two columns the order is 1, 2
plotConfidence(x=CiTable[,c("HazardRatio","lower","upper")],
               labels=CiTable[,c("Drug.Time","Dose","Mean","SD","n")],
               values=FALSE,
               order=c(2,1))
plotConfidence(x=CiTable[,c("HazardRatio","lower","upper")],
               labels=CiTable[,c("Drug.Time","Dose","Mean","SD","n")],
               values=FALSE,
               order=c(1,2))

```

```

## The relative size of the columns needs to be controlled manually
## by using the argument xratio. If there are only two columns
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
               xratio=c(0.4,0.15))

## The amount of space on the left and right margin can be controlled
## as follows:
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
               xratio=c(0.4,0.15),
               leftmargin=0.1, rightmargin=0.00)

## The actual size of the current graphics device determines
## the size of the figures and the space between them.
## The sizes and line widths are increased as follows:
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               xlab="Hazard ratio",
               labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
               points.cex=3,
               cex=2,
               lwd=3,
               xaxis.lwd=1.3,
               xaxis.cex=1.3)

## Note that 'cex' of axis ticks is controlled via 'par' but
## cex of the label via argument 'cex' of 'mtext'.
## The sizes and line widths are decreased as follows:
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
               cex=0.8,
               lwd=0.8,
               xaxis.lwd=0.8,
               xaxis.cex=0.8)

## Another good news is that all figures can be controlled separately

## The size of the graphic device can be controlled in the usual way, e.g.:
## Not run:
pdf("~/tmp/testCI.pdf", width=8, height=8)
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")])
dev.off()

## End(Not run)

## More control of the x-axis and confidence intervals that
## stretch outside the x-range end in an arrow.
## the argument xlab.line adjusts the distance of the x-axis
## label from the graph
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               xlab="Hazard ratio",
               xlab.line=1.8,

```

```

    xaxis.at=c(0.8,1,1.3),
    labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
    xlim=c(0.8,1.3))

## log-scale
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
  xlab="Hazard ratio",
  xlab.line=1.8,
  xaxis.at=c(0.8,1,1.3),
  labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
  xlim=c(0.8,1.3),plot.log="x")

## More pronounced arrows
## Coloured xlab expression
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
  xlab=expression(HR[1](s)),
  xlab.line=1.8,
  xlab.col="darkred",
  extremearrows.angle=50,
  extremearrows.length=0.1,
  labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
  xlim=c(0.8,1.3))

## Controlling the labels and their titles
## and the values and their titles
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
  labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
  xlab="Hazard ratio",
  title.values=expression(bold(HR (CI[95]))),
  title.labels=c("Drug/Time", "Dose", "Mean", "St.dev.", "N"),
  factor.reference.pos=c(1,10,19),
  factor.reference.pch=16,
  cex=1.3,
  xaxis.at=c(0.75,1,1.25,1.5,2))

## For factor reference groups, one may want to replace the
## confidence intervals by the word Reference, as in the previous example.
## To change the word 'Reference' we use the argument factor.reference.label:
## To change the plot symbol for the reference lines factor.reference.pch
## To remove the plot symbol in the reference lines use 'NA' as follows:
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
  labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
  xlab="Hazard ratio",
  factor.reference.label="Ref",
  title.values=expression(bold(HR (CI[95]))),
  title.labels=c("Drug/Time", "Dose", "Mean", "St.dev.", "N"),
  factor.reference.pos=c(1,10,19),
  factor.reference.pch=NA,
  cex=1.3,
  xaxis.at=c(0.75,1,1.25,1.5,2))

## changing the style of the graphical confidence intervals
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],

```

```

        labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
        xlab="Hazard ratio",
        factor.reference.pos=c(1,10,19),
        points.pch=15,
        points.col=rainbow(27),
        points.cex=2,
        arrows.col="darkblue",
        cex=1.3,
        order=c(1,3,2),
        xaxis.at=c(0.75,1,1.25,1.5))

## the values column of the graph can have multiple columns as well
## to illustrate this we create the confidence intervals
## before calling the function and then cbind them
## to the pvalues
HR <- pubformat(CiTable[,6])
CI95 <- formatCI(lower=CiTable[,7], upper=CiTable[,8], format="(l-u)")
pval <- format.pval(CiTable[,9], digits=3, eps=10^{-3})
pval[pval=="NA"] <- ""
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
               values=list("HR"=HR, "CI-95"=CI95, "P-value"=pval),
               cex=1.2,
               xratio=c(0.5,0.3))

## Finally, vertical columns can be delimited with background color
## NOTE: this may slow things down and potentially create
##       large figures (many bytes)
col1 <- rep(c(prodlim::dimColor("green", density=22),
              prodlim::dimColor("green")), length.out=9)
col2 <- rep(c(prodlim::dimColor("orange", density=22),
              prodlim::dimColor("orange")), length.out=9)
col3 <- rep(c(prodlim::dimColor("blue", density=22),
              prodlim::dimColor("blue")), length.out=9)
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
               stripes=c(1,0,1),
               stripes.col=c(col1, col2, col3))
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
               stripes=c(1,1,1),
               stripes.col=c(col1, col2, col3))

threegreens <- c(prodlim::dimColor("green", density=55),
                 prodlim::dimColor("green", density=33),
                 prodlim::dimColor("green", density=22))
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               labels=CiTable[,c("Drug.Time", "Dose", "Mean", "SD", "n")],
               values=FALSE,
               xlim=c(0.75,1.5),
               stripes=c(1,1,1),
               xratio=c(0.5,0.15),
               stripes.horizontal=c(0,9,18,27)+0.5,

```

```

        stripes.col=threegreens)

# combining multiple plots into one
layout(t(matrix(1:5)))
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               labels=CiTable[,c("Mean", "n")],
               layout=FALSE)
plotConfidence(x=CiTable[,c("HazardRatio", "lower", "upper")],
               layout=FALSE)

```

---

print.ci

*Print confidence intervals*


---

### Description

Print confidence intervals

### Usage

```

## S3 method for class 'ci'
print(x, se = FALSE, print = TRUE, ...)

```

### Arguments

x	Object containing point estimates and the corresponding confidence intervals
se	If TRUE add the standard error.
print	Logical: if FALSE do not actually print confidence intervals but just return them invisibly.
...	passed to summary.ci

### Details

This format of the confidence intervals is user-manipulable.

### Value

A string: the formatted confidence intervals

### Author(s)

Thomas A. Gerds <tag@biostat.ku.dk>

### See Also

ci plot.ci formatCI summary.ci



**Examples**

```
library(lava)
m <- lvm(Y~X)
m <- categorical(m,Y~X,K=4)
set.seed(4)
d <- sim(m,24)
ci.mean(Y~X,data=d)
x <- ci.mean(Y~X,data=d)
print(x,format="(l,u)")
```

---

```
print.subgroupAnalysis
```

*Printing univariate tables*

---

**Description**

Print function for subgroupAnalysis

**Usage**

```
## S3 method for class 'subgroupAnalysis'
print(x, ...)
```

**Arguments**

x                   - An object obtained with subgroupAnalysis  
...                 Passed to summary.subgroupAnalysis

**Details**

This function is simply calling summary.subgroupAnalysis

**Value**

The result of summary.subgroupAnalysis(x)

**Author(s)**

Christian Torp-Pedersen (ctp@heart.dk)

**See Also**

subgroupAnalysis

---

print.table2x2      *print results of 2x2 contingency table analysis*

---

### Description

print results of 2x2 contingency table analysis

### Usage

```
## S3 method for class 'table2x2'  
print(x, digits = 1, ...)
```

### Arguments

x	object obtained with table2x2
digits	rounding digits
...	not used

### Value

invisible x

### Author(s)

Thomas A. Gerds <tag@biostat.ku.dk>

### See Also

table2x2

### Examples

```
table2x2(table("marker"=rbinom(100,1,0.4),"response"=rbinom(100,1,0.1)))  
table2x2(matrix(c(71,18,38,8),ncol=2),stats="table")  
table2x2(matrix(c(71,18,38,8),ncol=2),stats=c("rr","fisher"))
```

---

print.univariateTable *Printing univariate tables*

---

**Description**

Print function for univariate tables

**Usage**

```
## S3 method for class 'univariateTable'  
print(x, ...)
```

**Arguments**

x                    An object obtained with univariateTable  
...                   Passed to summary.univariateTable

**Details**

This function is simply calling summary.univariateTable

**Value**

The result of summary.univariateTable(x)

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

univariateTable

---

pubformat                    *Format numbers for publication*

---

**Description**

Format numbers according to a specified handler function. Currently supported are sprintf, format and prettyNum.

**Usage**

```
pubformat(x, digits = 2, nsmall = digits, handler = "sprintf", ...)
```

**Arguments**

x	numeric vector
digits	number of digits
nsmall	see handler
handler	String specifying the name of the function which should perform the formatting. See <code>sprintf</code> , <code>format</code> and <code>prettyNum</code> .
...	Passed to handler function if applicable, i.e., not to <code>sprintf</code> .

**Value**

Formatted number

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

`sprintf`, `format`, `prettyNum`

**Examples**

```
pubformat(c(0.000143,12.8,1))
pubformat(c(0.000143,12.8,1),handler="format")
pubformat(c(0.000143,12.8,1),handler="format",trim=TRUE)
pubformat(c(0.000143,12.8,1),handler="prettyNum")
```

---

publish

*Publishing tables and figures*

---

**Description**

Publish provides summary functions for data and results of statistical analysis in ready-for-publication design

**Usage**

```
publish(object, ...)
```

**Arguments**

object	object to be published
...	Passed to method.

**Details**

Some warnings are currently suppressed.

**Value**

Tables and figures

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

publish.CauseSpecificCox publish.ci publish.coxph publish.glm publish.riskRegression publish.survdiff

---

publish.CauseSpecificCox

*Tabulizing cause-specific hazard ratio from all causes with confidence limits and Wald test p-values.*

---

**Description**

Publish cause-specific Cox models

**Usage**

```
## S3 method for class 'CauseSpecificCox'
publish(
  object,
  cause,
  confint.method,
  pvalue.method,
  factor.reference = "extraline",
  units = NULL,
  print = TRUE,
  ...
)
```

**Arguments**

object	Cause-specific hazard model obtained with CSC.
cause	Show a table for this cause. If omitted, list all causes.
confint.method	See regressionTable
pvalue.method	See regressionTable
factor.reference	See regressionTable

units	See regressionTable
print	If TRUE print the table(s).
...	passed on to control formatting of parameters, confidence intervals and p-values. See summary.regressionTable.

**Details**

The cause-specific hazard ratio's are combined into one table.

**Value**

Table with cause-specific hazard ratios, confidence limits and p-values.

**Author(s)**

Thomas Alexander Gerds <tab@biostat.ku.dk>

**Examples**

```
if (requireNamespace("riskRegression", quietly=TRUE)){
  library(riskRegression)
  library(prodlim)
  library(survival)
  data(Melanoma, package="riskRegression")
  fit1 <- CSC(list(Hist(time, status)~sex, Hist(time, status)~invasion+epicel+age),
             data=Melanoma)
  publish(fit1)
  publish(fit1, pvalue.stars=TRUE)
  publish(fit1, factor.reference="inline", units=list("age"="years"))

  # wide format (same variables in both Cox regression formula)
  fit2 <- CSC(Hist(time, status)~invasion+epicel+age, data=Melanoma)
  publish(fit2)

  # with p-values
  x <- publish(fit2, print=FALSE)
  table <- cbind(x[[1]]$regressionTable,
                x[[2]]$regressionTable[, -c(1,2)])
}
```

---

publish.ci

*Publish tables with confidence intervals*

---

**Description**

Publish tables with confidence intervals

**Usage**

```
## S3 method for class 'ci'  
publish(object, format = "[u;l]", se = FALSE, ...)
```

**Arguments**

object	Object of class ci containing point estimates and the corresponding confidence intervals
format	A string which indicates the format used for confidence intervals. The string is passed to <code>formatCI</code> with two arguments: the lower and the upper limit. For example '(l;u)' yields confidence intervals with round parenthesis in which the upper and the lower limits are separated by semicolon.
se	If TRUE add standard error.
...	passed to <code>publish</code>

**Details**

This function calls `summary.ci` with `print=FALSE` and then `publish`

**Value**

table with confidence intervals

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

`summary.ci`

**Examples**

```
data(Diabetes)  
publish(ci.mean(chol~location+gender, data=Diabetes), org=TRUE)
```

---

publish.coxph

*Tabulize hazard ratios with confidence intervals and p-values.*

---

**Description**

Tabulize the part of the result of a Cox regression analysis which is commonly shown in publications.

**Usage**

```
## S3 method for class 'coxph'
publish(
  object,
  confint.method,
  pvalue.method,
  print = TRUE,
  factor.reference = "extraline",
  units = NULL,
  probindex = FALSE,
  ...
)
```

**Arguments**

object	A coxph object.
confint.method	See regressionTable
pvalue.method	See regressionTable
print	If FALSE do not print results.
factor.reference	See regressionTable
units	See regressionTable
probindex	Logical. If TRUE show coefficients on probabilistic index scale instead of hazard ratio scale.
...	passed to summary.regressionTable and also to labelUnits.

**Details**

Transforms the log hazard ratios to hazard ratios and returns them with confidence limits and p-values. If explanatory variables are log transformed or log2 transformed, a scaling factor is multiplied to both the log-hazard ratio and its standard-error.

**Value**

Table with hazard ratios, confidence intervals and p-values.

**Author(s)**

Thomas Alexander Gerds

**Examples**

```
library(survival)
data(pbc)
pbc$edema <- factor(pbc$edema,
  levels=c("0", "0.5", "1"), labels=c("0", "0.5", "1"))
fit = coxph(Surv(time, status!=0)~age+sex+edema+log(bili)+log(albumin),
```



```

        data=na.omit(pbc))
publish(fit)
## forest plot
plot(publish(fit),cex=1.3)

publish(fit,ci.digits=2,pvalue.eps=0.01,pvalue.digits=2,pvalue.stars=TRUE)
publish(fit,ci.digits=2,ci.handler="prettyNum",pvalue.eps=0.01,
        pvalue.digits=2,pvalue.stars=TRUE)
publish(fit, ci.digits=2, ci.handler="sprintf", pvalue.eps=0.01,
        pvalue.digits=2,pvalue.stars=TRUE, ci.trim=FALSE)

fit2 = coxph(Surv(time,status!=0)~age+sex+edema+log(bili,base=2)+log(albumin)+log(protime),
            data=na.omit(pbc))
publish(fit2)

# with cluster variable
fit3 = coxph(Surv(time,status!=0)~age+cluster(sex)+edema+log(bili,base=2)
            +log(albumin)+log(protime),
            data=na.omit(pbc))
publish(fit3)

# with strata and cluster variable
fit4 = coxph(Surv(time,status!=0)~age+cluster(sex)+strata(edema)+log(bili,base=2)
            +log(albumin)+log(protime),
            data=pbc)
publish(fit4)

```

---

publish.glm

*Tabulize regression coefficients with confidence intervals and p-values.*


---

## Description

Tabulate the results of a generalized linear regression analysis.

## Usage

```

## S3 method for class 'glm'
publish(
  object,
  confint.method,
  pvalue.method,
  digits = c(2, 4),
  print = TRUE,
  factor.reference = "extraline",
  intercept = ifelse((is.null(object$family) || object$family$family == "gaussian"), 1L,
                    0L),
  units = NULL,
  ...
)

```

**Arguments**

object	A glm object.
confint.method	See regressionTable.
pvalue.method	See regressionTable.
digits	A vector of two integer values. These determine how to round numbers (first value) and p-values (second value). E.g., c(1,3) would mean 1 digit for all numbers and 3 digits for p-values. The actual rounding is done by summary.regressionTable.
print	If FALSE do not print results.
factor.reference	Style for showing results for categorical. See regressionTable.
intercept	See regressionTable.
units	See regressionTable.
...	passed to summary.regressionTable and also to labelUnits.
reference	Style for showing results for categorical variables. If "extraline" show an additional line for the reference category.

**Details**

The table shows changes in mean for linear regression and odds ratios for logistic regression (family = binomial).

**Value**

Table with regression coefficients, confidence intervals and p-values.

**Author(s)**

Thomas Alexander Gerds <tag@biostat.ku.dk>

**Examples**

```
data(Diabetes)
## Linear regression
f = glm(bp.2s~frame+gender+age, data=Diabetes)
publish(f)
publish(f, factor.reference="inline")
publish(f, pvalue.stars=TRUE)
publish(f, ci.format="(1,u)")

### interaction
fit = glm(bp.2s~frame+gender*age, data=Diabetes)
summary(fit)
publish(fit)

Fit = glm(bp.2s~frame*gender+age, data=Diabetes)
publish(Fit)
```

```

## Logistic regression
Diabetes$hyper1 <- factor(1*(Diabetes$bp.1s>140))
lrfit <- glm(hyper1~frame+gender+age,data=Diabetes,family=binomial)
publish(lrfit)

### interaction
lrfit1 <- glm(hyper1~frame+gender*age,data=Diabetes,family=binomial)
publish(lrfit1)

lrfit2 <- glm(hyper1~frame*gender+age,data=Diabetes,family=binomial)
publish(lrfit2)

## Poisson regression
data(trace)
trace <- Units(trace,list("age"="years"))
fit <- glm(dead ~ smoking+sex+age+Time+offset(log(ObsTime)), family="poisson",data=trace)
rtf <- regressionTable(fit,factor.reference = "inline")
summary(rtf)
publish(fit)

## gls regression
if (requireNamespace("nlme",quietly=TRUE)){
  requireNamespace("lava",quietly=TRUE)
  library(lava)
  library(nlme)
  m <- lvm(Y ~ X1 + gender + group + Interaction)
  distribution(m, ~gender) <- binomial.lvm()
  distribution(m, ~group) <- binomial.lvm(size = 2)
  constrain(m, Interaction ~ gender + group) <- function(x){x[,1]*x[,2]}
  d <- sim(m, 1e2)
  d$gender <- factor(d$gender, labels = letters[1:2])
  d$group <- factor(d$group)

  e.gls <- gls(Y ~ X1 + gender*group, data = d,
              weights = varIdent(form = ~1|group))
  publish(e.gls)

## lme
fm1 <- lme(distance ~ age*Sex,
           random = ~1|Subject,
           data = Orthodont)
res <- publish(fm1)
}

```

---

publish.htest

*Pretty printing of test results.*


---

## Description

Pretty printing of test results.

**Usage**

```
## S3 method for class 'htest'
publish(object, title, ...)
```

**Arguments**

object	Result of <code>t.test</code> or <code>wilcox.test</code>
title	Decoration also used to name output
...	Used to transport arguments <code>ci.arg</code> and <code>pvalue.arg</code> to subroutines <code>format.pval</code> and <code>formatCI</code> . See also <code>prodlm::SmartControl</code> .

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**Examples**

```
data(Diabetes)
publish(t.test(bp.2s~gender,data=Diabetes))
publish(wilcox.test(bp.2s~gender,data=Diabetes))
publish(with(Diabetes,t.test(bp.2s,bp.1s,paired=TRUE)))
publish(with(Diabetes,wilcox.test(bp.2s,bp.1s,paired=TRUE)))
```

---

publish.matrix

*Publishing a matrix in raw, org, latex, or muse format*

---

**Description**

This is the heart of the Publish package

**Usage**

```
## S3 method for class 'matrix'
publish(
  object,
  title,
  colnames = TRUE,
  rownames = TRUE,
  col1name = "",
  digits = 4,
  try.convert = TRUE,
  sep = " ",
  endhead,
  endrow,
  style,
  inter.lines,
```

```

    latex = FALSE,
    wiki = FALSE,
    org = FALSE,
    markdown = FALSE,
    tabular = TRUE,
    latex.table.format = NA,
    latex.hline = 1,
    latex.nodollar = FALSE,
    ...
)

```

### Arguments

object	Matrix to be published
title	Title for table, only in wiki and muse format
colnames	If TRUE show column names
rownames	If TRUE show row names
col1name	Name for first column
digits	Numbers are rounded according to digits
try.convert	Logical. If TRUE try to convert also non-numeric formats such as character to numeric before rounding. Default is TRUE.
sep	Field separator when style is "none"
endhead	String to be pasted at the end of the first row (header)
endrow	String to be pasted at the end of each row
style	Table style for export to "latex", "org", "markdown", "wiki", "none". Overwritten by arguments below.
inter.lines	A named list which contains strings to be placed between the rows of the table. An element with name "0" is used to place a line before the first column, elements with name "r" are placed between line r and r+1.
latex	If TRUE use latex table format
wiki	If TRUE use mediawiki table format
org	If TRUE use emacs orgmode table format
markdown	If TRUE use markdown table format
tabular	For style latex only: if TRUE enclose the table in begin/end tabular environment.
latex.table.format	For style latex only: format of the tabular environment.
latex.hline	For style latex only: if TRUE add hline statements add the end of each line.
latex.nodollar	For style latex only: if TRUE do not enclose numbers in dollars.
...	Used to transport arguments. Currently supports wiki.class.

**Examples**

```
x <- matrix(1:12,ncol=3)
publish(x)

# rounding the numeric part of data mixtures
y <- cbind(matrix(letters[1:12],ncol=3),x,matrix(rnorm(12),ncol=3))
publish(y,digits=1)

publish(x,latex=TRUE,
inter.lines=list("1"="text between line 1 and line 2",
                 "3"="text between line 3 and line 4"))
```

---

publish.MIresult	<i>Present logistic regression and Cox regression obtained with mi-tools::MIcombine based on smcfcs::smcfcs multiple imputation analysis</i>
------------------	--

---

**Description**

Regression tables after multiple imputations

**Usage**

```
## S3 method for class 'MIresult'
publish(
  object,
  confint.method,
  pvalue.method,
  digits = c(2, 4),
  print = TRUE,
  factor.reference = "extraline",
  intercept,
  units = NULL,
  fit,
  data,
  ...
)
```

**Arguments**

object	Object obtained with mitools::MIcombine based on smcfcs::smcfcs multiple imputation analysis
confint.method	No options here. Only Wald type confidence intervals.
pvalue.method	No options here. Only Wald type tests.
digits	Rounding digits for all numbers but the p-values.

print	If FALSE suppress printing of the results
factor.reference	Style for showing results for categorical. See regressionTable.
intercept	See regressionTable.
units	See regressionTable.
fit	One fitted model using the same formula as object. This can be the fit to the complete case data or the fit to one of the completed data. It is used to get xlevels, formula and terms. For usage see examples. is used to fit
data	Original data set which includes the missing values
...	passed to summary.regressionTable, labelUnits and publish.default.

### Details

Show results of smcfcs based multiple imputations of missing covariates in publishable format

### Author(s)

Thomas A. Gerds <tag@biostat.ku.dk>

### Examples

```
## Not run:
if (requireNamespace("riskRegression", quietly=TRUE)
    & requireNamespace("mitools", quietly=TRUE)
    & requireNamespace("smcfcs", quietly=TRUE)){
library(riskRegression)
library(mitools)
library(smcfcs)
## continuous outcome: linear regression
# lava some data with missing values
set.seed(7)
d=sampleData(78)
## generate missing values
d[X1==1,X6:=NA]
d[X2==1,X3:=NA]
d=d[,.(X8,X4,X3,X6,X7)]
sapply(d,function(x)sum(is.na(x)))

# multiple imputation (should set m to a large value)

set.seed(17)
f= smcfcs(d,smtpe="lm",
          smformula=X8~X4+X3+X6+X7,
          method=c("","","logreg","norm",""),m=3)
ccfit=lm(X8~X4+X3+X6+X7,data=d)
mifit=MIcombine(with(imputationList(f$impDatasets),
                      lm(X8~X4+X3+X6+X7)))
publish(mifit,fit=ccfit,data=d)
publish(ccfit)
```

```

## binary outcome
# lava some data with missing values
set.seed(7)
db=sampleData(78,outcome="binary")
## generate missing values
db[X1==1,X6:=NA]
db[X2==1,X3:=NA]
db=db[,.(Y,X4,X3,X6,X7)]
sapply(db,function(x)sum(is.na(x)))

# multiple imputation (should set m to a large value)
set.seed(17)
fb= smcfcs(db,smtpe="logistic",
           smformula=Y~X4+X3+X6+X7,
           method=c("","","logreg","norm",""),m=2)
ccfit=glm(Y~X4+X3+X6+X7,family="binomial",data=db)
mifit=MIcombine(with(imputationList(fb$impDatasets),
                    glm(Y~X4+X3+X6+X7,family="binomial")))
publish(mifit,fit=ccfit)
publish(ccfit)

## survival: Cox regression
library(survival)
# lava some data with missing values
set.seed(7)
ds=sampleData(78,outcome="survival")
## generate missing values
ds[X5==1,X6:=NA]
ds[X2==1,X3:=NA]
ds=ds[,.(time,event,X4,X3,X6,X7)]
sapply(ds,function(x)sum(is.na(x)))

set.seed(17)
fs= smcfcs(ds,smtpe="coxph",
           smformula="Surv(time,event)~X4+X3+X6+X7",
           method=c("","","logreg","norm",""),m=2)
ccfit=coxph(Surv(time,event)~X4+X3+X6+X7,data=ds)
mifit=MIcombine(with(imputationList(fs$impDatasets),
                    coxph(Surv(time,event)~X4+X3+X6+X7)))
publish(mifit,fit=ccfit,data=ds)
publish(ccfit)

## competing risks: Cause-specific Cox regression
library(survival)
# lava some data with missing values
set.seed(7)
dcr=sampleData(78,outcome="competing.risks")
## generate missing values
dcr[X5==1,X6:=NA]
dcr[X2==1,X3:=NA]
dcr=dcr[,.(time,event,X4,X3,X6,X7)]
sapply(dcr,function(x)sum(is.na(x)))

```



```

set.seed(17)
fcr= smcfcs(dcr,smtpe="compet",
            smformula=c("Surv(time,event==1)~X4+X3+X6+X7",
                        "Surv(time,event==2)~X4+X3+X6+X7"),
            method=c("","","","logreg","norm",""),m=2)
## cause 2
ccfit2=coxph(Surv(time,event==2)~X4+X3+X6+X7,data=dcr)
mifit2=MIcombine(with(imputationList(fcr$impDatasets),
                      coxph(Surv(time,event==2)~X4+X3+X6+X7)))
publish(mifit2,fit=ccfit2,data=dcr)
publish(ccfit2)
}

## End(Not run)

```

---

publish.riskRegression

*Publishing results of riskRegression*

---

## Description

Preparing a publishable table from riskRegression results

## Usage

```

## S3 method for class 'riskRegression'
publish(object, digits = c(2, 4), print = TRUE, ...)

```

## Arguments

object	object of class riskRegression as obtained with functions ARR and LRR.
digits	Number of digits for regression coefficients
print	If FALSE do not print the results
...	passed to <a href="#">publish.matrix</a>

## Value

Table with regression coefficients, confidence intervals and p-values

## Author(s)

Thomas A. Gerds <tag@biostat.ku.dk>

## See Also

ARR LRR

**Examples**

```

if (requireNamespace("riskRegression",quietly=TRUE)){
  library(riskRegression)
  library(prodlim)
  library(lava)
  library(survival)
  set.seed(20)
  d <- SimCompRisk(20)
  f <- ARR(Hist(time,event)~X1+X2,data=d,cause=1)
  publish(f)
  publish(f,digits=c(1,3))
}

```

---

publish.Score

*Publish predictive accuracy results*


---

**Description**

Write output of riskRegression::Score in tables

**Usage**

```

## S3 method for class 'Score'
publish(object, metrics, score = TRUE, contrasts = TRUE, level = 3, ...)

```

**Arguments**

object	Object obtained with riskRegression::Score
metrics	Which metrics to put into tables. Defaults to object\$metrics.
score	Logical. If TRUE print the score elements, i.e., metric applied to the risk prediction models.
contrasts	Logical. If TRUE print the contrast elements (if any). These compare risk prediction models according to metrics.
level	Level of subsection headers, i.e., ** for level 2 and *** for level 3 (useful for emacs org-users). Default is plain subsection headers no stars. A negative value will suppress subsection headers.
...	Passed to publish

**Details**

Collect prediction accuracy results in tables

**Value**

Results of Score in tabular form

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**Examples**

```
if (requireNamespace("riskRegression", quietly=TRUE)){
  library(riskRegression)
  library(survival)
  learn = sampleData(100)
  val= sampleData(100)
  f1=CSC(Hist(time,event)~X1+X8,data=learn)
  f2=CSC(Hist(time,event)~X1+X5+X6+X8,learn)
  xs=Score(list(f1,f2),data=val,formula=Hist(time,event)~1)
  publish(xs)
}
```

---

publish.summary.aov     *Format summary table of aov results*

---

**Description**

Format summary table of aov results

**Usage**

```
## S3 method for class 'summary.aov'
publish(
  object,
  print = TRUE,
  handler = "sprintf",
  digits = c(2, 4),
  nsmall = digits,
  ...
)
```

**Arguments**

object	glm object
print	Logical. Decide about whether or not to print the results.
handler	see pubformat
digits	see pubformat
nsmall	see pubformat
...	used to transport further arguments

## Examples

```
data(Diabetes)
f <- glm(bp.l~age+chol+gender+location,data=Diabetes)
publish(summary(aov(f)),digits=c(1,2))
```

---

publish.survdiff	<i>Alternative summary of survdiff results</i>
------------------	--

---

## Description

Alternative summary of survdiff results

## Usage

```
## S3 method for class 'survdiff'
publish(object, digits = c(2, 4), print = TRUE, ...)
```

## Arguments

object	Object obtained with <code>survival::survdiff</code> .
digits	Vector with digits for rounding numbers: the second for pvalues, the first for all other numbers.
print	If FALSE do not print results.
...	Not (yet) used.

## Author(s)

Thomas A. Gerds <tag@biostat.ku.dk>

## Examples

```
library(survival)
data(pbc)
sd <- survdiff(Surv(time,status!=0)~sex,data=pbc)
publish(sd)
publish(sd,digits=c(3,2))
```

---

regressionTable	<i>Regression table</i>
-----------------	-------------------------

---

**Description**

Tabulate the results of a regression analysis.

**Usage**

```
regressionTable(
  object,
  param.method = "coef",
  confint.method = c("default", "profile", "robust", "simultaneous"),
  pvalue.method = c("default", "robust", "simultaneous"),
  factor.reference = "extraline",
  intercept = 0L,
  units = NULL,
  noterms = NULL,
  probindex = 0L,
  ...
)
```

**Arguments**

object	Fitted regression model obtained with <code>lm</code> , <code>glm</code> or <code>coxph</code> .
param.method	Method to obtain model coefficients.
confint.method	Method to obtain confidence intervals. Default is 'default' which leads to Wald type intervals using the model based estimate of standard error. 'profile' yields profile likelihood confidence intervals, available from library MASS for <code>lm</code> and <code>glm</code> objects. 'robust' uses the sandwich form standard error to construct Wald type intervals (see <code>lava::estimate.default</code> ). 'simultaneous' calls <code>multcomp::glht</code> to obtain simultaneous confidence intervals.
pvalue.method	Method to obtain p-values. If 'default' show raw p-values. If 'robust' use p-value corresponding to robust standard error as provided by <code>lava::estimate.default</code> . If 'simultaneous' call <code>multcomp::glht</code> to obtain p-values.
factor.reference	Style for showing results for categorical variables. If 'extraline' show an additional line for the reference category. If 'inline' display as level vs. reference.
intercept	Logical. If FALSE suppress intercept.
units	List of units for continuous variables. See examples.
noterms	Position of terms that should be ignored. E.g., for a Cox model with a <code>cluster(id)</code> term, there will be no hazard ratio for variable <code>id</code> .
probindex	Logical. If TRUE show coefficients on probabilistic index scale instead of hazard ratio scale.
...	Not yet used

**Details**

The basic use of this function is to generate a near publication worthy table from a regression object. As with `summary(object)` reference levels of factor variables are not included. Expansion of the table with such values can be performed using the `"fixRegressionTable"` function. Forest plot can be added to the output with `"plotRegressionTable"`.

`regressionTable` produces an object (list) with the parameters deriveds. The summary function creates a data frame which can be used as a (near) publication ready table.

The table shows changes in mean for linear regression, odds ratios for logistic regression (family = binomial) and hazard ratios for Cox regression.

**Value**

List of regression blocks

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**Examples**

```
# linear regression
data(Diabetes)
f1 <- glm(bp.1s~age+gender+frame+chol,data=Diabetes)
summary(regressionTable(f1))
summary(regressionTable(f1,units=list("chol"="mmol/L", "age"="years")))
## with interaction
f2 <- glm(bp.1s~age*gender+frame+chol,data=Diabetes)
summary(regressionTable(f2))
#Add reference values
summary(regressionTable(f2))
f3 <- glm(bp.1s~age+gender*frame+chol,data=Diabetes)
publish(f3)
regressionTable(f3)

# logistic regression
Diabetes$hyp1 <- factor(1*(Diabetes$bp.1s>140))
l1 <- glm(hyp1~age+gender+frame+chol,data=Diabetes,family="binomial")
regressionTable(l1)
publish(l1)
plot(regressionTable(l1))

## with interaction
l2 <- glm(hyp1~age+gender+frame*chol,data=Diabetes,family="binomial")
regressionTable(l2)
l3 <- glm(hyp1~age*gender+frame*chol,data=Diabetes,family="binomial")
regressionTable(l3)

# Cox regression
library(survival)
data(pbc)
```

```

pbc$edema <- factor(pbc$edema, levels=c("0", "0.5", "1"), labels=c("0", "0.5", "1"))
c1 <- coxph(Surv(time, status!=0)~log(bili)+age+protime+sex+edema, data=pbc)
regressionTable(c1)
# with interaction
c2 <- coxph(Surv(time, status!=0)~log(bili)+age+protime*sex+edema, data=pbc)
regressionTable(c2)
c3 <- coxph(Surv(time, status!=0)~edema*log(bili)+age+protime+sex+edema+edema:sex, data=pbc)
regressionTable(c3)

if (requireNamespace("nlme", quietly=TRUE)){
## gls regression
library(lava)
library(nlme)
m <- lvm(Y ~ X1 + gender + group + Interaction)
distribution(m, ~gender) <- binomial.lvm()
distribution(m, ~group) <- binomial.lvm(size = 2)
constrain(m, Interaction ~ gender + group) <- function(x){x[,1]*x[,2]}
d <- sim(m, 1e2)
d$gender <- factor(d$gender, labels = letters[1:2])
d$group <- factor(d$group)

e.gls <- gls(Y ~ X1 + gender*group, data = d,
             weights = varIdent(form = ~1|group))
regressionTable(e.gls)
summary(regressionTable(e.gls))
}

```

---

SpaceT

*A study was made of all 26 astronauts on the first eight space shuttle flights (Bungo et.al., 1985). On a voluntary basis 17 astronauts consumed large quantities of salt and fluid prior to landing as a countermeasure to space deconditioning, while nine did not.*

---

## Description

A study was made of all 26 astronauts on the first eight space shuttle flights (Bungo et.al., 1985). On a voluntary basis 17 astronauts consumed large quantities of salt and fluid prior to landing as a countermeasure to space deconditioning, while nine did not.

## Format

A data frame with 52 observations on the following 4 variables:

**Status** Factor with levels Post (after flight) and Pre (before flight)

**HR** Supine heart rate (beats per minute)

**Treatment** Countermeasure salt/fluid (1= yes, 0=no)

**ID** Person id

**References**

Altman, Practical statistics for medical research, Page 223, Ex. 9.1. Bungo et.al., 1985

**Examples**

```
data(SpaceT)
```

---

<code>spaghettiogram</code>	<i>Spaghettiogram</i>
-----------------------------	-----------------------

---

**Description**

A spaghettiogram is showing repeated measures (longitudinal data)

**Usage**

```
spaghettiogram(
  formula,
  data,
  xlim,
  ylim,
  xlab = "",
  ylab = "",
  axes = TRUE,
  col,
  lwd,
  lty,
  pch,
  legend = FALSE,
  add = FALSE,
  background = TRUE,
  ...
)
```

**Arguments**

<code>formula</code>	A formula which specifies the variables for the spaghettiograms. If $Y \sim X + id(Z)$ then for each value of $Z$ the spaghettiogram is the graph $(X,Y)$ in the subset defined by the value of $Z$ . Data are expected to be in the "long" format. $Y$ is a numeric vector and $X$ is a factor whose levels define the X-axis. Each level of the id-vector corresponds to one line (spaghetti) in the plot.
<code>data</code>	data set in which variables $X$ , $Y$ and $Z$ are defined.
<code>xlim</code>	Limits for x-axis
<code>ylim</code>	Limits for y-axis
<code>xlab</code>	Label for x-axis



ylab	Label for x-axis
axes	Logical indicating if axes should be drawn.
col	Colors for the spaghettiograms
lwd	Widths for the spaghettiograms
lty	Type for the spaghettiograms
pch	Point-type for the spaghettiograms
legend	If TRUE add a legend. Argument A of legend is controlled as legend.A. E.g., when legend.cex=2 legend will be called with argument cex=2.
add	If TRUE add to existing plot device.
background	Control the background color of the graph.
...	used to transport arguments which are passed to the following subroutines: "plot", "lines", "legend", "background", "axis1", "axis2".

**Value**

List with data of each subject

**Examples**

```
data(SpaceT)
Spaghettiogram(HR~Status+id(ID),
               data=SpaceT)
```

---

specialFrame	<i>Special frame</i>
--------------	----------------------

---

**Description**

Extract data and design matrix including specials from call

**Usage**

```
specialFrame(
  formula,
  data,
  unspecial.design = TRUE,
  specials,
  specials.factor = TRUE,
  specials.design = FALSE,
  strip.specials = TRUE,
  strip.arguments = NULL,
  strip.alias = NULL,
  strip.unspecials = NULL,
  drop.intercept = TRUE,
  response = TRUE,
  na.action = options()$na.action
)
```

**Arguments**

formula	Formula whose left hand side specifies the event history, i.e., either via <code>Surv()</code> or <code>Hist()</code> .
data	Data frame in which the formula is interpreted
unspecials.design	Passed as is to <code>model.design</code> .
specials	Character vector of special function names. Usually the body of the special functions is <code>function(x)x</code> but e.g., <code>strata</code> from the survival package does treat the values
specials.factor	Passed as is to <code>model.design</code> .
specials.design	Passed as is to <code>model.design</code>
strip.specials	Passed as specials to <code>strip.terms</code>
strip.arguments	Passed as arguments to <code>strip.terms</code>
strip.alias	Passed as alias.names to <code>strip.terms</code>
strip.unspecials	Passed as unspecials to <code>strip.terms</code>
drop.intercept	Passed as is to <code>model.design</code>
response	If FALSE do not get response data.
na.action	Decide what to do with missing values.

**Details**

Obtain a list with the data used for event history regression analysis. This function cannot be used directly on the user level but inside a function to prepare data for survival analysis.

**Value**

A list which contains - the response - the design matrix (see `model.design`) - one entry for each special (see `model.design`)

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

`model.frame` `model.design` `Hist`

**Examples**

```

## Here are some data with an event time and no competing risks
## and two covariates X1 and X2.
## Suppose we want to declare that variable X1 is treated differently
## than variable X2. For example, X1 could be a cluster variable, or
## X1 should have a proportional effect on the outcome.
d <- data.frame(y=1:7,
                X2=c(2.24,3.22,9.59,4.4,3.54,6.81,5.05),
                X3=c(1,1,1,1,0,0,1),
                X4=c(44.69,37.41,68.54,38.85,35.9,27.02,41.84),
                X1=factor(c("a","b","a","c","c","a","b"),
                          levels=c("c","a","b")))
## define special functions prop and cluster
prop <- function(x)x
cluster <- function(x)x
## We pass a formula and the data
e <- specialFrame(y~prop(X1)+X2+cluster(X3)+X4,
                 data=d,
                 specials=c("prop","cluster"))
## The first element is the response
e$response
## The other elements are the design, i.e., model.matrix for the non-special covariates
e$design
## and a data.frame for the special covariates
e$prop
## The special covariates can be returned as a model.matrix
e2 <- specialFrame(y~prop(X1)+X2+cluster(X3)+X4,
                 data=d,
                 specials=c("prop","cluster"),
                 specials.design=TRUE)
e2$prop
## and the non-special covariates can be returned as a data.frame
e3 <- specialFrame(y~prop(X1)+X2+cluster(X3)+X4,
                 data=d,
                 specials=c("prop","cluster"),
                 specials.design=TRUE,
                 unspecial.design=FALSE)
e3$design

```

**Description**

Plotting the prediction of a logistic regression model with confidence bands against one continuous variable.

**Usage**

```
splinePlot.lrm(
  object,
  xvar,
  xvalues,
  xlim = range(xvalues),
  ylim,
  xlab = xvar,
  ylab = scale[[1]],
  col = 1,
  lty = 1,
  lwd = 3,
  confint = TRUE,
  newdata = NULL,
  scale = c("risk", "odds"),
  add = FALSE,
  ...
)
```

**Arguments**

<code>object</code>	Logistic regression model fitted with <code>rms::lrm</code>
<code>xvar</code>	Name of the variable to show on x-axis
<code>xvalues</code>	Sequence of <code>xvar</code> values
<code>xlim</code>	x-axis limits
<code>ylim</code>	y-axis limits
<code>xlab</code>	x-axis labels
<code>ylab</code>	y-axis labels
<code>col</code>	color of the line
<code>lty</code>	line style
<code>lwd</code>	line width
<code>confint</code>	Logical. If TRUE show confidence shadows
<code>newdata</code>	How to adjust
<code>scale</code>	Character string that determines the outcome scale (y-axis). Choose between "risk" and "odds".
<code>add</code>	Logical. If TRUE add lines to an existing graph
<code>...</code>	Further arguments passed to <code>plot</code> . Only if <code>add</code> is FALSE.

**Details**

Function which extracts from a logistic regression model fitted with `rms::lrm` the predicted risks or odds.

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**Examples**

```
data(Diabetes)
Diabetes$hypertension= 1*(Diabetes$bp.1s>140)
library(rms)
uu <- datadist(Diabetes)
options(datadist="uu")
fit=lrn(hypertension~rcs(age)+gender+hdl,data=Diabetes)
splinePlot.lrn(fit,xvar="age",xvalues=seq(30,50,1))
```

---

stripes

*Background and grid color control.*

---

**Description**

Some users like background colors, and it may be helpful to have grid lines to read off e.g. probabilities from a Kaplan-Meier graph. Both things can be controlled with this function. However, it mainly serves [plot.prodlm](#).

**Usage**

```
stripes(
  xlim,
  ylim,
  col = "white",
  lwd = 1,
  gridcol = "gray77",
  fill = "white",
  horizontal = NULL,
  vertical = NULL,
  border = "black",
  xpd = FALSE
)
```

**Arguments**

xlim	Limits for the horizontal x-dimension. Defaults to par("usr")[1:2].
ylim	Limits for the vertical y-dimension.
col	Colors use for the stripes. Can be a vector of colors which are then repeated appropriately.
lwd	Line width
gridcol	Color of grid lines
fill	Color to fill the background rectangle given by par("usr").

horizontal	Numerical values at which to show horizontal grid lines, and at which to change the color of the stripes.
vertical	Numerical values at which to show vertical grid lines.
border	If a fill color is provided, the color of the border around the background.
xpd	From help(par): A logical value or NA. If FALSE, all plotting is clipped to the plot region, if TRUE, all plotting is clipped to the figure region, and if NA, all plotting is clipped to the device region. See also clip.

**Author(s)**

Thomas Alexander Gerds <tag@biostat.ku.dk>

**Examples**

```
plot(0,0)
backGround(bg="beige",fg="red",vertical=0,horizontal=0)

plot(0,0)
stripes(col=c("yellow","green"),gridcol="red",xlim=c(-1,1),horizontal=seq(0,1,.1))
stripes(col=c("yellow","green"),gridcol="red",horizontal=seq(0,1,.1))
```

---

subgroupAnalysis      *Subgroup Analysis - Interactions and estimates*

---

**Description**

The function can examine Cox regression, logistic regression and Poisson regression (Poisson regression for survival analysis) where the effect of one variable is of particular interest. This function systematically checks for effect modification with a list of other variables.

In randomised studies the main regression analysis is often univariate and includes only the exposure of interest. In observational studies the main regression analysis can readily be adjusted for other variables including those which may modify the effect of the variable of interest.

**Usage**

```
subgroupAnalysis(
  object,
  data,
  treatment,
  subgroups,
  confint.method = "default",
  factor.reference = "extraline",
  ...
)
```

**Arguments**

object - glm, coxph or cph object for which subgroups should be analyzed.  
 data - Dataset including all relevant variables  
 treatment - Must be numeric - 0/1  
 subgroups - A vector of variable names presenting the factor variables where subgroups should be formed. These variables should all be "factors"  
 confint.method "default" creates Wald type confidence interval, "robust", creates creates robust standard errors - see regressionTable function.  
 factor.reference "extraline" creates an extraline for the reference, "inline" avoids this line.  
 ... additional arguments such as case weights, which are passed on to glm and coxph.

**Details**

The function can only handle a bivariate treatment, which MUST coded as zero or one. The p-value for interaction is obtained with a likelihood ratio test comparing the main regression analysis with the interaction model.

There are plot and print functions available for the function see helppages for plot.subgroupAnalysis and print.subgroupAnalysis

**Value**

A data.frame with subgroup specifications, number in each subgroup, parameter estimates and p-value for interaction. A forest plot can be obtained with "plotConfidence".

**Author(s)**

Christian Torp-Pedersen

**See Also**

coxph, glm, plotConfidence

**Examples**

```

#load libraries
library(data.table)
library(Publish)
library(survival)
data(traceR) #get dataframe traceR
data.table::setDT(traceR)
traceR[, ':='(wmi2=factor(wallMotionIndex<0.9, levels=c(TRUE, FALSE),
                        labels=c("bad", "good")),
            abd2=factor(abdominalCircumference<95, levels=c(TRUE, FALSE),
                        labels=c("slim", "fat")))]
traceR[, sex:=as.factor(sex)] # all subgroup variables needs to be factor
traceR[observationTime==0, observationTime:=1]

```

```

# remove missing covariate values
traceR=na.omit(traceR)
# univariate analysis of smoking in subgroups of age and sex
# Main regression analysis is a simple/univariate Cox regression
fit_cox <- coxph(Surv(observationTime,dead)~treatment,data=traceR)
sub_cox <- subgroupAnalysis(fit_cox,traceR,treatment="treatment",
  subgroups=c("smoking","sex","wmi2","abd2"))
sub_cox

# to see how the results are obtained consider the variable: smoking
fit_cox_smoke <- coxph(Surv(observationTime,dead)~treatment*smoking,data=traceR)
# the last three rows of the following output:
publish(fit_cox_smoke)
# are included in the first 3 rows of the result of the sub group analysis:
sub_cox[1:3,]
# the p-value is obtained as:
fit_cox_smoke_add <- coxph(Surv(observationTime,dead)~treatment+smoking,data=traceR)
anova(fit_cox_smoke_add,fit_cox_smoke,test="Chisq")

# Note that a real subgroup analysis would be to subset the data
fit_cox1a <- coxph(Surv(observationTime,dead)~treatment,data=traceR[smoking=="never"])
fit_cox1b <- coxph(Surv(observationTime,dead)~treatment,data=traceR[smoking=="current"])
fit_cox1c <- coxph(Surv(observationTime,dead)~treatment,data=traceR[smoking=="prior"])

## when the main analysis is already adjusted
fit_cox_adj <- coxph(Surv(observationTime,dead)~treatment+smoking+sex+wmi2+abd2,
  data=traceR)
sub_cox_adj <- subgroupAnalysis(fit_cox_adj,traceR,treatment="treatment",
  subgroups=c("smoking","sex","wmi2","abd2")) # subgroups as character string
sub_cox_adj

# When both start and end are in the Surv statement:
traceR[,null:=0]
fit_cox2 <- coxph(Surv(null,observationTime,dead)~treatment+smoking+sex+wmi2+abd2,data=traceR)
summary(regressionTable(fit_cox2))
sub_cox2 <- subgroupAnalysis(fit_cox2,traceR,treatment="treatment",
  subgroups=c("smoking","sex","wmi2","abd2"))
# Analysis with Poisson - and the unrealistic assumption of constant hazard
# and adjusted for age in all subgroups
fit_p <- glm(dead~treatment+age+offset(log(observationTime)),family="poisson",
  data=traceR)
sub_pois <- subgroupAnalysis(fit_p,traceR,treatment="treatment",
  subgroups=~smoking+sex+wmi2+abd2)
# Analysis with logistic regression - and very wrongly ignoring censoring
fit_log <- glm(dead~treatment+age,family="binomial",data=traceR)
sub_log <- subgroupAnalysis(fit_log,traceR,treatment="treatment",
  subgroups=~smoking+sex+wmi2+abd2, factor.reference="inline")

```



**Description**

Summarize confidence intervals

**Usage**

```
## S3 method for class 'ci'
summary(object, format = "[u;l]", se = FALSE, print = TRUE, ...)
```

**Arguments**

object	Object of class ci containing point estimates and the corresponding confidence intervals
format	A string which indicates the format used for confidence intervals. The string is passed to <code>formatCI</code> with two arguments: the lower and the upper limit. For example <code>'(l;u)'</code> yields confidence intervals with round parenthesis in which the upper and the lower limits are separated by semicolon.
se	If TRUE add standard error.
print	Logical: if FALSE do not actually print confidence intervals but just return them invisibly.
...	used to control formatting of numbers

**Details**

This format of the confidence intervals is user-manipulable.

**Value**

Formatted confidence intervals

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

ci plot.ci format.ci

**Examples**

```
library(lava)
m <- lvm(Y~X)
m <- categorical(m,Y~X,K=4)
set.seed(4)
d <- sim(m,24)
ci.mean(Y~X,data=d)
x <- summary(ci.mean(Y~X,data=d),digits=2)
x
x <- summary(ci.mean(Y~X,data=d),format="(u,l)",digits=2)
x <- summary(ci.mean(Y~X,data=d),format="(u,l)",digits=1,se=TRUE)
```

```
x <- summary(ci.mean(Y~X,data=d),format="(u,l)",digits=1,handler="format")
x <- summary(ci.mean(Y~X,data=d),format="(u,l)",digits=1,handler="prettyNum")
```

---

summary.regressionTable

*Formatting regression tables*

---

### Description

Preparing regression results for publication

### Usage

```
## S3 method for class 'regressionTable'
summary(object, show.missing = "ifany", print = TRUE, ...)
```

### Arguments

object	object obtained with regressionTable or summary.regressionTable.
show.missing	Decide if number of missing values are shown. Either logical or character. If 'ifany' then number missing values are shown if there are some.
print	If TRUE print results.
...	Used to control formatting of parameter estimates, confidence intervals and p-values. See examples.

### Value

List with two elements:

- regressionTable: the formatted regression table (a data.frame)
- rawTable: table with the unformatted values (a data.frame)

### Author(s)

Thomas A. Gerds <tag@biostat.ku.dk>

### See Also

publish.glm publish.coxph

**Examples**

```

library(survival)
data(pbc)
pbc$edema <- factor(pbc$edema, levels=c("0", "0.5", "1"), labels=c("0", "0.5", "1"))
fit = coxph(Surv(time, status!=0)~age+sex+edema+log(bili)+log(albumin)+log(protime),
            data=pbc)
u=summary(regressionTable(fit))
u$regressionTable
u$rawTable
summary(regressionTable(fit), handler="prettyNum")
summary(regressionTable(fit), handler="format")
summary(regressionTable(fit), handler="sprintf", digits=c(2,2), pValue.stars=TRUE)
summary(regressionTable(fit), handler="sprintf", digits=c(2,2), pValue.stars=TRUE, ci.format="(1,u)")

```

---

```

summary.subgroupAnalysis
      summary.subgroupAnalysis

```

---

**Description**

This function operates on a "subgroupAnalysis" object to produce a formatted table.

**Usage**

```

## S3 method for class 'subgroupAnalysis'
summary(
  object,
  digits = 3,
  eps = 0.001,
  subgroup.p = FALSE,
  keep.digital = FALSE,
  ...
)

```

**Arguments**

object	- a subgroupAnalysis object
digits	- number of digits for risk ratios
eps	- lowest value of p to be shown exactly, others will be "<eps"
subgroup.p	- present p-values for analyses in subgroups
keep.digital	- prevents formatting risk ratio and confidence limits. Useful for cases when further manipulations of rows and columns prior to adding a forest plot is relevant.
...	- not currently used

**Details**

This function produces a formatted or unformatted table of a subgroupAnalysis object. A forest plot can be added with the plot function.

**Value**

A data.frame with formatted values for subgroups

**Author(s)**

Christian Torp-Pedersen

**See Also**

subgroupAnalysis

**Examples**

```
#load libraries
library(Publish)
library(survival)
library(data.table)
data(traceR) #get dataframe traceR
setDT(traceR)
traceR[, ':='(wmi2=factor(wallMotionIndex<0.9, levels=c(TRUE,FALSE),
  labels=c("bad", "good")),
  abd2=factor(abdominalCircumference<95, levels=c(TRUE,FALSE),
  labels=c("slim", "fat")))]
traceR[,sex:=as.factor(sex)] # all subgroup variables needs to be factor
traceR[observationTime==0,observationTime:=1]
# univariate analysis of smoking in subgroups of age and sex
# Basic model from randomised study - but observed for 12 years
fit_cox <- coxph(Surv(observationTime,dead)~treatment,data=traceR)
sub_cox <- subgroupAnalysis(fit_cox,traceR,treatment="treatment",
  subgroup=c("smoking","sex","wmi2","abd2")) # subgroups as character string
summary(sub_cox)
```

---

summary.univariateTable

*Preparing univariate tables for publication*

---

**Description**

Summary function for univariate table

**Usage**

```
## S3 method for class 'univariateTable'
summary(
  object,
  n = "inNames",
  drop.reference = FALSE,
  pvalue.stars = FALSE,
  pvalue.digits = 4,
  show.missing = c("ifany", "always", "never"),
  show.pvalues,
  show.totals,
  ...
)
```

**Arguments**

object	univariateTable object as obtained with function univariateTable.
n	If not missing, show the number of subjects in each column. If equal to "inNames", show the numbers in parentheses in the column names. If missing the value object\$n is used.
drop.reference	Logical or character (vector). Decide if line with reference level should be suppressed for factors. If TRUE or "all" suppress for all categorical factors. If 'binary' suppress only for binary variables. Can be character vector in which case reference lines are suppressed for variables that are included in the vector.
pvalue.stars	If TRUE use symnum to parse p-values otherwise use format.pval.
pvalue.digits	Passed to format.pval.
show.missing	Decides if number of missing values are shown in table. Defaults to "ifany", and can also be set to "always" or "never".
show.pvalues	Logical. If set to FALSE the column p-values is removed. If missing the value object\$compare.groups[[1]]==TRUE is used.
show.totals	Logical. If set to FALSE the column Totals is removed. If missing the value object\$show.totals is used.
...	passed on to labelUnits. This overwrites labels stored in object\$labels

**Details**

Collects results of univariate table in a matrix.

**Value**

Summary table

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**Examples**

```

data(Diabetes)
u <- univariateTable(gender~age+location+Q(BMI)+height+weight,
                    data=Diabetes)

summary(u)
summary(u,n=NULL)
summary(u,pvalue.digits=2,"age"="Age (years)","height"="Body height (cm)")

u2 <- univariateTable(location~age+AgeGroups+gender+height+weight,
                    data=Diabetes)

summary(u2)
summary(u2,drop.reference=TRUE)
## same but more flexible
summary(u2,drop.reference=c("binary"))
## same but even more flexible
summary(u2,drop.reference=c("gender"))

```

---

sutable

*Fast summary of a univariate table*


---

**Description**

First apply univariateTable then call summary.

**Usage**

```
sutable(...)
```

**Arguments**

... Unnamed arguments and are passed to univariateTable as well as named arguments that match univariateTable's arguments, other arguments are passed to summary.univariateTable

**Value**

Summary table

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**See Also**

summary.univariateTable univariateTable

**Examples**

```
data(Diabetes)
sutable(gender~age+location+Q(BMI)+height+weight,data=Diabetes,BMI="Body mass index (kg/m^2)")
```

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table2x2	<i>2x2 table calculus for teaching</i>
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**Description**

2x2 table calculus for teaching

**Usage**

```
table2x2(
  x,
  digits = 1,
  conf.level = 0.95,
  stats = c("table", "rd", "rr", "or", "chisq", "fisher")
)
```

**Arguments**

x	2x2 table
digits	rounding digits
conf.level	Confidence level used for constructing confidence intervals. Default is 0.95.
stats	subset or all of c("table", "rd", "or", "rr", "chisq", "fisher") where rd= risk difference, rr = risk ratio, or = odds ratio, chisq = chi-square test, fisher= fisher's exact test and table = the 2x2 table

**Details**

2x2 table calculus for teaching

**Value**

see example

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**Examples**

```
table2x2(table("marker"=rbinom(100,1,0.4),"response"=rbinom(100,1,0.1)))
table2x2(matrix(c(71,18,38,8),ncol=2),stats="table")
table2x2(matrix(c(71,18,38,8),ncol=2),stats=c("rr","fisher"))
```

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trace	<i>trace data</i>
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## Description

These data are from screening to the TRACE study, a comparison between the angiotensin converting enzyme inhibitor trandolapril and placebo for large myocardial infarctions. A total of 6676 patients were screened for the study. Survival has been followed for the screened population for 16 years. The current data has been prepared for a poisson regression to examine survival. The data has been "split" in 0.5 year intervals (plitLexus function from Epi package) and then collapsed on all variables (aggregate function).

## Format

A data frame with 1832 observations on the following 6 variables.

**Time** Time after myocardial infarction, in 6 months intervals

**smoking** Smoking status. A factor with levels (Never, Current, Previous)

**sex** A factor with levels (Female, Male)

**age** Age in years at the time of myocardial infarction

**ObsTime** Cumulative risk time in each split

**dead** Count of deaths

## References

Kober et al 1995 Am. J. Cardiol 76,1-5

## Examples

```
data(trace)
Units(trace,list("age"="years"))
fit <- glm(dead ~ smoking+sex+age+Time+offset(log(ObsTime)), family="poisson",data=trace)
rtf <- regressionTable(fit,factor.reference = "inline")
summary(rtf)
publish(fit)
```



---

`traceR`*traceR data*

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## Description

These data are from the TRACE randomised trial, a comparison between the angiotensin converting enzyme inhibitor trandolapril and placebo for large myocardial infarctions. In all, 1749 patients were randomised. The current data are from a 15 year follow-up.

## Format

A data frame with 1749 observations on the following variables.

**weight** Weight in kilo

**height** Height in meters

**abdominalCircumference** in centimeters

**seCreatinine** in mmol per liter

**wallMotionIndex** left ventricular function 0-2, 0 worst, 2 normal

**observationTime** time to death or censor

**age** age in years

**sex** 0=female,1=male

**smoking** 0=never,1=prior,2=current

**dead** 0=censor,1=dead

**treatment** placebo or trandolapril

## References

Kober et al 1995 NEJM 333,1670

## Examples

```
data(trace)
Units(trace,list("age"="years"))
fit <- glm(dead ~ smoking+sex+age+Time+offset(log(ObsTime)), family="poisson",data=trace)
rtf <- regressionTable(fit,factor.reference = "inline")
summary(rtf)
publish(fit)
```

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Units	<i>Add units to data set</i>
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**Description**

Add variable units to data.frame (or data.table).

**Usage**

```
Units(object, units)
```

**Arguments**

object	A data.frame or data.table
units	Named list of units. Names are variable names. If omitted, show existing units.

**Details**

If the object has units existing units are replaced by given units.

**Value**

The object augmented with attribute "units"

**Author(s)**

Thomas A. Gerds <tag@biostat.ku.dk>

**Examples**

```
data(Diabetes)
Diabetes <- Units(Diabetes, list(BMI="kg/m^2"))
Units(Diabetes)
Diabetes <- Units(Diabetes, list(bp.1s="mm Hg", bp.2s="mm Hg"))
Units(Diabetes)
```

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univariateTable	<i>Univariate table</i>
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---

**Description**

Categorical variables are summarized using counts and frequencies and compared .

**Usage**

```

univariateTable(
  formula,
  data = parent.frame(),
  summary.format = "mean(x) (sd(x))",
  Q.format = "median(x) [iqr(x)]",
  freq.format = "count(x) (percent(x))",
  column.percent = TRUE,
  digits = c(1, 1, 3),
  big.mark = ",",
  short.groupnames,
  compare.groups = TRUE,
  show.totals = TRUE,
  n = "inNames",
  outcome = NULL,
  ...
)

```

**Arguments**

formula	Formula specifying the grouping variable (strata) on the left hand side (can be omitted) and on the right hand side the variables for which to obtain (descriptive) statistics.
data	Data set in which formula is evaluated
summary.format	Format for the numeric (non-factor) variables. Default is mean (SD). If different formats are desired, either special Q can be used or the function is called multiple times and the results are rbinded. See examples.
Q.format	Format for quantile summary of numerical variables: Default is median (inter quartile range).
freq.format	Format for categorical variables. Default is count (percentage).
column.percent	Logical, if TRUE and the default freq.format is used then column percentages are given instead of row percentages for categorical variables (factors).
digits	Number of digits
big.mark	For formatting large numbers (i.e., greater than 1,000). "" turn this off.
short.groupnames	If TRUE group names are abbreviated.
compare.groups	Method used to compare groups. If "logistic" and there are exactly two groups logistic regression is used instead of t-tests and Wilcoxon rank tests to compare numeric variables across groups.
show.totals	If TRUE show a column with totals.
n	If TRUE show the number of subjects as a separate row. If equal to "inNames", show the numbers in parentheses in the column names. If FALSE do not show number of subjects.
outcome	Outcome data used to calculate p-values when compare groups method is 'logistic' or 'cox'.
...	saved as part of the result to be passed on to labelUnits

## Details

This function can generate the baseline demographic characteristics that forms table 1 in many publications. It is also useful for generating other tables of univariate statistics.

The result of the function is an object (list) which contains the various data generated. In most applications the summary function should be applied which generates a data.frame with a (nearly) publication ready table. Standard manipulation can be used to modify, add or remove columns/rows and for users not accustomed to R the table generated can be exported to a text file which can be read by other software, e.g., via `write.csv(table,file="path/to/results/table.csv")`

By default, continuous variables are summarized by means and standard deviations and compared with t-tests. When continuous variables are summarized by medians and interquartile ranges the Deviations from the above defaults are obtained when the arguments `summary.format` and `freq.format` are combined with suitable summary functions.

## Value

List with one summary table element for each variable on the right hand side of formula. The summary tables can be combined with `rbind`. The function `summary.univariateTable` combines the tables, and shows p-values in custom format.

## Author(s)

Thomas A. Gerds

## See Also

`summary.univariateTable`, `publish.univariateTable`

## Examples

```
data(Diabetes)
library(data.table)
univariateTable(~age,data=Diabetes)
univariateTable(~gender,data=Diabetes)
univariateTable(~age+gender+ height+weight,data=Diabetes)
## same thing but less typing
utable(~age+gender+ height+weight,data=Diabetes)

## summary by location:
univariateTable(location~Q(age)+gender+height+weight,data=Diabetes)
## continuous variables marked with Q() are (by default) summarized
## with median (IQR) and kruskal.test (with two groups equivalent to wilcox.test)
## variables not marked with Q() are (by default) summarized
## with mean (sd) and anova.glm(...,test="Chisq")
## the p-value of anova(glm()) with only two groups is similar
## but not exactly equal to that of a t.test
## categorical variables are (by default) summarized by count
## (percent) and chi-square tests (\code{chisq.test}). When \code{compare.groups='logistic'}
## anova(glm(...,family=binomial,test="Chisq")) is used to calculate p-values.

## export result to csv
```

```

table1 = summary(univariateTable(location~age+gender+height+weight,data=Diabetes),
show.pvalues=FALSE)
# write.csv(table1,file=~"/table1.csv",rownames=FALSE)

## change labels and values
utable(location~age+gender+height+weight,data=Diabetes,
age="Age (years)",gender="Sex",
gender.female="Female",
gender.male="Male",
height="Body height (inches)",
weight="Body weight (pounds)")

## Use quantiles and rank tests for some variables and mean and standard deviation for others
univariateTable(gender~Q(age)+location+Q(BMI)+height+weight,
data=Diabetes)

## Factor with more than 2 levels
Diabetes$AgeGroups <- cut(Diabetes$age,
c(19,29,39,49,59,69,92),
include.lowest=TRUE)
univariateTable(location~AgeGroups+gender+height+weight,
data=Diabetes)

## Row percent
univariateTable(location~gender+age+AgeGroups,
data=Diabetes,
column.percent=FALSE)

## change of frequency format
univariateTable(location~gender+age+AgeGroups,
data=Diabetes,
column.percent=FALSE,
freq.format="percent(x) (n=count(x))")

## changing Labels
u <- univariateTable(location~gender+AgeGroups+ height + weight,
data=Diabetes,
column.percent=TRUE,
freq.format="count(x) (percent(x))")
summary(u,"AgeGroups"="Age (years)","height"="Height (inches)")

## more than two groups
Diabetes$frame=factor(Diabetes$frame,levels=c("small","medium","large"))
univariateTable(frame~gender+BMI+age,data=Diabetes)

Diabetes$sex=as.numeric(Diabetes$gender)
univariateTable(frame~sex+gender+BMI+age,
data=Diabetes,freq.format="count(x) (percent(x))")

## multiple summary formats
## suppose we want for some reason mean (range) for age
## and median (range) for BMI.
## method 1:

```



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