# Package 'npsurvSS'

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Type Package

**Title** Sample Size and Power Calculation for Common Non-Parametric Tests in Survival Analysis

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Description A number of statistical tests have been proposed to compare two survival curves, including the difference in (or ratio of) t-year survival, difference in (or ratio of) p-th percentile survival, difference in (or ratio of) restricted mean survival time, and the weighted log-rank test.

Despite the multitude of options, the convention in survival studies is to assume proportional hazards and to use the unweighted log-rank test for design and analysis. This package provides sample size and power calculation for all of the above statistical tests with allowance for flexible accrual, censoring, and survival (eg. Weibull, piecewise-exponential, mixture cure). It is the companion R package to the paper by Yung and Liu (2019) <doi:10.1111/biom.13196>. Specific to the weighted log-rank test, users may specify which approximations they wish to use to estimate the large-sample mean and variance. The default option has been shown to provide substantial improvement over the conventional sample size and power equations based on Schoenfeld (1981) <doi:10.1093/biomet/68.1.316>.

**Depends** R (>= 3.4.0)

Imports stats, utils

License GPL-2

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Suggests knitr, rmarkdown, tidyverse, ggplot2

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URL http://github.com/godwinyung/npsurvSS

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BugReports http://github.com/godwinyung/npsurvSS/issues

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# **R** topics documented:

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create\_arm

Create an 'arm' object

# **Description**

Create an object of class 'arm' by specifying the trial parameters for a single arm, including the sample size, accrual distribution, survival distribution, loss to follow-up distribution, and study duration.

# Usage

```
create_arm(size, accr_time, accr_dist = "pieceuni",
   accr_interval = c(0, accr_time), accr_param = NA, surv_cure = 0,
   surv_interval = c(0, Inf), surv_shape = 1, surv_scale,
   loss_shape = 1, loss_scale, follow_time = Inf, total_time = Inf)
```

# Arguments

size sample size. If total sample size is unknown, provide the integer sample size relative to the opposing arm, e.g. 1 for 1:2 randomization ratio or 2 for 2:3.

accr\_time accrual duration.

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accr\_dist accrual distribution. Default is piecewise uniform. Alternatively, 'truncexp' allows for a truncated exponential distribution as proposed by Lachin and Foulkes (1986). Depending on the value of accr\_param, this distribution can be either convex or concave. accr\_interval accrual intervals. Defaults to the single interval spanning from 0 to accr\_time. If a piecewise uniform accrual with more than one interval is desired, specify accr\_interval as the vector of increasing changepoints (knots) starting from 0 and ending with accr\_time, e.g. c(0, 2, 4) defines a piecewise uniform distribution with two intervals, [0, 2) and [2, 4]. accr\_param additional accrual parameter(s). For a piecewise uniform accrual with more than one interval, specify accr\_param as the vector of probabilities a patient is enrolled in each interval. The probabilities should naturally sum to 1. For accr\_dist='truncexp', specify accr\_param as a single number. >0 results in a convex distribution and <0 results in a concave distribution. surv\_cure proportion of patients that are cured. survival intervals. Defaults to the single interval spanning from 0 to infinsurv\_interval ity. If a piecewise exponential survival is desired for uncured patients, specify surv\_interval as the vector of increasing changepoints (knots) starting from 0 and ending with infinity, e.g. c(0, 6, 10, Inf). surv\_shape Weibull shape parameter for the survival distribution of uncured patients. Weibull scale parameter for the survival distrubition of uncured patients. Piecesurv\_scale wise exponential survival may be defined by specifying surv\_shape=1 and surv\_scale as the vector of piecewise hazard rates. loss\_shape Weibull shape parameter for the loss to follow-up distribution. loss\_scale Weibull scale parameter for the loss to follow-up distribution. follow\_time follow-up duration. total study duration. Only 1 of the 2 parameters, follow\_time or total\_time, total\_time need to be defined. If neither is defined, total\_time is defaulted to max value

### Value

a list containing assumptions of size, accrual, censoring, survival, and follow-up for a single arm.

### References

Lachin, J. M. and Foulkes, M. A. (1986) Evaluation of sample size and power for analyses of survival with allowance for nonuniform patient entry, losses to follow-up, noncompliance, and stratification. *Biometrics*, **42**, 507-519.

#### See Also

create\_arm\_lachin for creating an object of subclass 'lachin'.

1e6.

4 create\_arm\_lachin

### **Examples**

```
# Example 1
example <- create_arm(size=120,
 accr_time=6,
                                 # uniform accrual
 surv scale=0.05.
                                 # exponential survival
 loss_scale=0.005,
                                 # exponential loss to follow-up
 follow_time=12)
                             # this example also satisfies properties of subclass 'lachin'
class(example)
# Example 2
create_arm(size=120,
                                 # truncated exponential accrual
 accr_time=6,
 accr_dist="truncexp",
 accr_param=0.1,
 surv_shape=2,
                                 # weibull survival
 surv_scale=0.05,
 loss_shape=1.5,
                                 # weilbull loss to follow-up
 loss_scale=0.005,
 total_time=18)
# Example 3
create_arm(size=120,
 accr_time=6,
                                 # piecewise uniform accrual
 accr_interval=c(0,2,4,6),
 accr_param=c(0.2,0.3,0.5),
                                 # 10% cure fraction
 surv_cure=0.1,
                                 # piecewise exponential survival for uncured patients
 surv_interval=c(0,6,10,Inf),
 surv_scale=c(0.05,0.04,0.03),
 loss_shape=0.7,
                                 # weibull loss to follow-up
 loss_scale=0.005,
 total_time=18)
```

create\_arm\_lachin

Create a 'lachin' object

### **Description**

Create an object of class 'lachin' by specifying the trial parameters for a single arm, including the sample size, accrual distribution, survival distribution, loss to follow-up distribution, and study duration. 'Lachin' objects are also 'arm' objects, but with accrual limited to the uniform and truncated exponential distributions, and survival and loss to follow-up limited to the exponential distribution. 'Lachin' objects have the advantage that expectations for certain counting processes have closed form equations and can therefore be calculated more efficiently (Lachin and Foulkes, 1986).

### Usage

```
create_arm_lachin(size, accr_time, accr_dist = "pieceuni",
    accr_param = NA, surv_median = NA, surv_exphazard = NA,
    surv_milestone = NA, loss_median = NA, loss_exphazard = NA,
    loss_milestone = NA, follow_time = Inf, total_time = Inf)
```

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### **Arguments**

size	sample size. If total sample size is unknown, provide the integer sample size relative to the opposing arm, e.g. 1 for 1:2 randomization ratio or 2 for 2:3.
accr_time	accrual duration.
accr_dist	accrual distribution. Default is uniform (piecewise uniform with one interval). Alternatively, 'truncexp' allows for a truncated exponential distribution as proposed by Lachin and Foulkes (1986). Depending on the value of accr_param, this distribution can be either convex or concave.
accr_param	additional accrual parameter for accr_dist='truncexp'. accr_param>0 specifies a convex distribution and accr_param<0 specifies a concave distribution.
surv_median	median survival.
surv_exphazard	exponential hazard rate for the survival distribution.
surv_milestone	a tuple c(milestone, probability) that uniquely defines the exponential survival distribution, e.g. $c(12,0.8)$ corresponds to the exponential distribution with $80\%$ survival rate at 12 months.
loss_median	median loss to follow-up.
loss_exphazard	exponential hazard rate for the loss to follow-up distribution.
loss_milestone	a tuple c(milestone, probability) that uniquely defines the exponential loss to follow-up distribution, e.g. $c(12, 0.99)$ corresponds to the exponential distribution with $1\%$ loss to follow-up at $12$ months.
follow_time	Follow-up duration. Either follow_time or total_time (below) should be specified.
total_time	Total study duration. Either follow_time (above) or total_time should be specified.

# Value

a list containing assumptions of size, accrual, censoring, survival, and follow-up for a single arm.

# References

Lachin, J. M. and Foulkes, M. A. (1986) Evaluation of sample size and power for analyses of survival with allowance for nonuniform patient entry, losses to follow-up, noncompliance, and stratification. *Biometrics*, **42**, 507-519.

# See Also

create\_arm for creating an object of class 'arm'.

# **Examples**

```
# 3 arms with similar survival and loss to follow-up
create_arm_lachin(size=120, accr_time=6,
    surv_median=14,
    loss_median=140,
    follow_time=12)
```

6 dacer

```
create_arm_lachin(size=120, accr_time=6,
   surv_exphazard=0.05,
   loss_exphazard=0.005,
   follow_time=12)
create_arm_lachin(size=120, accr_time=6,
   accr_dist="truncexp",
   accr_param=0.1,
   surv_milestone=c(14, 0.5),
   loss_milestone=c(140, 0.5),
   total_time=18)
```

daccr

Accrual

# **Description**

Density, distribution function, quantile function, and random generation for the accrual distribution.

# Usage

```
daccr(x, arm)
paccr(q, arm, lower.tail = T)
qaccr(p, arm)
raccr(n = 1, arm)
```

# **Arguments**

# Value

daccr gives the density, paccr gives the distribution function, qaccr gives the quantile function, and raccr generates random deviates.

### See Also

dloss 7

to follow-up

dloss	Loss

# Description

Density, distribution function, hazard function, quantile function, and random generation for the loss to follow-up distribution.

# Usage

```
dloss(x, arm)
ploss(q, arm, lower.tail = T)
hloss(x, arm)
qloss(p, arm, lower.tail = T)
rloss(n = 1, arm)
```

# **Arguments**

```
x, q vector of quantiles.  \text{arm} \qquad \text{object of class 'arm'}. \\  \text{lower.tail} \qquad \text{logical; if TRUE, probabilities are } P(X \leq x); \text{ otherwise, } P(X > x). \\  \text{p} \qquad \text{vector of probabilities.} \\  \text{n} \qquad \text{number of observations.}
```

# Value

dloss gives the density, ploss gives the distribution function, hloss gives the hazard function, qloss gives the quantile function, and rloss generates random deviates.

# See Also

```
create_arm and create_arm_lachin for creating an object of class 'arm'.
```

8 dmaxU

amaxu
-------

Maximum observed time

# **Description**

Density, distribution function, and expected value for the maximum observed time in a single arm of patients.

# Usage

```
dmaxU(x, arm, include_cens = T)
pmaxU(q, arm, include_cens = T, lower.tail = T)
emaxU(arm, include_cens = T)
```

### **Arguments**

x, q vector of quantiles.

arm object of class 'arm'.

include\_cens logical; if TRUE, include time-to-censoring as potential observed time; otherwise, observed time equals time-to-event.

logical; if TRUE, probabilities are  $P(X \le x)$ ; otherwise, P(X > x).

# **Details**

lower.tail

Given a patient's time-to-event  $T_i$  and time-to-censoring  $C_i$ ,  $U_i = \min(T_i, C_i)$  defines the patient's observed time. The maximum observed time over patients of a single arm is then  $\max_i U_i$ .

### Value

dmaxU gives the density, pmaxU gives the distribution function, and emaxU gives the expected value.

### See Also

dminimaxU 9

dminimaxU	Minimax observed time	
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### **Description**

Density, distribution function, quantile function, and expected value for the minimum of the maximum observed time over two treatment arms.

# Usage

```
dminimaxU(x, arm0, arm1, include_cens = T)
pminimaxU(q, arm0, arm1, include_cens = T, lower.tail = T)
qminimaxU(p, arm0, arm1, include_cens = T, margin = 0.01)
eminimaxU(arm0, arm1, include_cens = T)
```

### **Arguments**

x,q	vector of quantiles.
arm0	object of class 'arm'.
arm1	object of class 'arm'.
include_cens	logical; if TRUE, include time-to-censoring as potential observed time; otherwise, observed time equals time-to-event.
lower.tail	logical; if TRUE, probabilities are $P(X \le x)$ ; otherwise, $P(X > x)$ .
p	vector of probabilities.
margin	margin of accuracy.

### **Details**

Given a patient in arm  $X_i = j$  with time-to-event  $T_i$  and time-to-censoring  $C_i$ ,  $U_i = \min(T_i, C_i)$  defines the patient's observed time. The maximum observed time over patients of arm j is then  $\max_{i:X_i=j} U_i$ , and the minimax observed time over two arms is  $\min_j (\max_{i:X_i=j} U_i)$ .

### Value

dminimaxU gives the density, pminimaxU gives the distribution function, qminimaxU gives the quantile function, and eminimaxU gives the expected value.

### See Also

10 dsurv

Survival
----------

# Description

Density, distribution function, hazard function, quantile function, and random generation for the survival distribution.

# Usage

```
dsurv(x, arm, include_cured = T)
psurv(q, arm, include_cured = T, lower.tail = T)
hsurv(x, arm, include_cured = T)
qsurv(p, arm, include_cured = T, lower.tail = T)
rsurv(n = 1, arm, include_cured = T)
```

# **Arguments**

x, q	vector of quantiles.
arm	object of class arm.
include_cured	logical; if TRUE, mixture distribution of cured and uncured patients is considered; otherwise, only the distribution for uncured patients is considered.
lower.tail	logical; if TRUE, probabilities are $P(X \le x)$ ; otherwise, $P(X > x)$ .
p	vector of probabilities.
n	number of observations.

### Value

dsurv gives the density, psurv gives the distribution function, hsurv gives the hazard function, qsurv gives the quantile function, and rsurv generates random deviates.

## See Also

exp\_duration 11

# **Description**

Given one or two treatment arms, calculate the time  $\tau$  at which the expected number of events equals d.

# Usage

```
exp_duration(arm0 = NA, arm1 = NA, d, search_start = 10,
    search_prec = 0.01, max_duration = 1000)
```

# Arguments

arm0 object of class 'arm'.

arm1 object of class 'arm'.

d vector of number of events.

search\_start value at which the search for duration tau starts.

search\_prec value controlling the desired precision before terminating the search.

max\_duration maximum  $\tau$  for consideration.

### Value

expected trial duration.

### See Also

exp\_events for calculating expected events d at time tau, create\_arm and create\_arm\_lachin for creating an object of class 'arm'.

# Examples

```
arm0 <- create_arm(size=120, accr_time=6, surv_scale=0.05, loss_scale=0.005, follow_time=12)
arm1 <- create_arm(size=120, accr_time=6, surv_scale=0.03, loss_scale=0.005, follow_time=12)
exp_duration(arm0, d=61)
exp_duration(arm0, arm1, d=103)
exp_duration(arm0, d=c(35,61))
exp_duration(arm0, arm1, d=c(57,103))</pre>
```

12 per2haz

exp	0111	n+ 0
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Expected number of events

# Description

Given one or two treatment arms, calculate the expected number of events d at time  $\tau$ .

# Usage

```
exp_events(arm0 = NA, arm1 = NA, tau = NA)
```

# Arguments

arm0 object of class 'arm'.
arm1 object of class 'arm'.

tau vector of times. Defaults to total study duration.

#### Value

expected number of events.

### See Also

exp\_duration for calculating time to achieve expected events d, create\_arm and create\_arm\_lachin for creating an object of class 'arm'.

# **Examples**

```
arm0 <- create_arm(size=120, accr_time=6, surv_scale=0.05, loss_scale=0.005, follow_time=12)
arm1 <- create_arm(size=120, accr_time=6, surv_scale=0.03, loss_scale=0.005, follow_time=12)
exp_events(arm0)
exp_events(arm0, arm1)
exp_events(arm0, tau=c(10,NA))
exp_events(arm0, arm1, tau=c(10,NA))</pre>
```

per2haz

Convert exponential parameters

### **Description**

Convert exponential survival percentile or hazard rate to the other.

### Usage

```
per2haz(x, per = 0.5)
```

power\_two\_arm 13

### Arguments

x survival percentile or exponential hazard rate per (per)th percentile

### Details

$$y = -\log(1 - per)/x$$

#### Value

survival percentile or hazard rate.

# **Examples**

```
per2haz(14)  # hazard rate for exponential with 14-month median

per2haz(0.05)  # median survival for exponential with hazard rate 0.05

per2haz(14, 0.8)  # hazard rate for exponential with 80th percentile survival at 14 months

per2haz(0.27, 0.8)  # 80th percentile survival for exponential with hazard rate 0.27
```

power\_two\_arm

Power

# **Description**

Calculate power for a two-arm survival study.

### Usage

```
power_two_arm(arm0, arm1, test = list(test = "weighted logrank"),
  alpha = 0.025, sides = 1)
```

#### **Arguments**

arm0 object of class 'arm'.
arm1 object of class 'arm'.

test

list or list of lists. Each list must contain at minimum the key 'test' describing the type of statistical test. Default test is the "weighted logrank". Kaplan-Meier based tests ("survival difference", "survival ratio", "rmst difference", "rmst ratio", "percentile difference", and "percentile ratio") require the user to define an additional key, either the desired 'milestone' or 'percentile'. The weighted log-rank test does not require additional keys. However, user may choose which weight function ("1"=unweighted, "n"=Gehan-Breslow, "sqrtN"=Tarone-Ware, "FH\_p[a]\_q[b]"= Fleming-Harrington with p=a and q=b) and which approximation for the large-sample mean ("asymptotic", "generalized schoenfeld", "event driven") and variance ("1", "block[ randomization]", "simple[ randomization]") they wish to use. Default choice is 'weight'="1", 'mean.approx'="asymptotic", and 'var.approx'="1". For more details regarding the different mean and variance approximations for the weight log-rank test, please see Yung and Liu (2019).

14 simulate\_arm

```
alpha type 1 error rate
```

sides 1=1-sided test, 2=2-sided test

### Value

power.

### References

Yung, G and Liu, Y. (2019). Sample size and power for the weighted log-rank test and Kaplan-Meier based tests with allowance for non-proportional hazards. *Biometrics*. <doi:10.1111/biom.13196>

#### See Also

```
create_arm for creating an object of class 'arm'.
```

### **Examples**

```
arm0 <- create_arm(size=120, accr_time=6, surv_scale=0.05, loss_scale=0.005, follow_time=12)
arm1 <- create_arm(size=120, accr_time=6, surv_scale=0.03, loss_scale=0.005, follow_time=12)
power_two_arm(arm0, arm1)
power_two_arm(arm0, arm1, list(test="weighted logrank",
    weight="n",
    mean.approx="generalized schoenfeld",
    var.approx="block"))
power_two_arm(arm0, arm1, list(test="survival difference", milestone=12))
power_two_arm(arm0, arm1, list(test="rmst ratio", milestone=12))
power_two_arm(arm0, arm1, list(test="percentile difference", percentile=0.25))
power_two_arm(arm0, arm1, list(
    list(test="weighted logrank"),
    list(test="survival difference", milestone=12)))</pre>
```

simulate\_arm

Simulate complete data for a single arm

### **Description**

Simulate the complete data for a single arm, including time to accrual, event, and loss of followup. No cutoff (by number of events or time) is applied. Hence, no patients are administratively censored.

### Usage

```
simulate_arm(arm, label = 1)
```

# Arguments

```
arm object of class 'arm'.
```

label numeric label for the simulated arm, e.g. 0 for control, 1 for treatment

simulate\_trial 15

#### Value

arm label

time.accr time to accrual

time.obs time to observation from accrual

time.total time to observation from start of study

censor 0=censor, 1=event

reason event description ('[experience ]event', '[loss to ]followup', 'administration[

censoring]')

time.surv time to event

time.loss time to loss of follow-up

#### See Also

create\_arm for creating an object of class 'arm'.

# **Examples**

```
arm0 <- create_arm(size=120, accr_time=6, surv_scale=0.05, loss_scale=0.005, follow_time=12)
simulate_arm(arm0, label=0)</pre>
```

simulate\_trial

Simulate a clinical trial

### **Description**

Simulate a single- or two-arm clinical trial, where end of study (EOS) is triggered after a number of events has been observed or a certain time has elapsed. Whereas simulate\_arm provides complete data for patients, including time to event and loss of follow-up, simulate\_trial mimicks an actual survival study by providing only the observed time (minimum of time to event or censoring) and censoring indicator.

### Usage

```
simulate_trial(arm0 = NA, arm1 = NA, events = NA, duration = Inf)
```

### **Arguments**

arm0 object of class 'arm'.
arm1 object of class 'arm'.

events number of required events to trigger end of study; overrides study duration de-

fined within arm0 and arm1.

duration time from first-patient-in to trigger end of study; overrides study duration de-

fined within arm0 and arm1. If both events and duration are specified, end of

study is triggered by either criteria, whichever occurs first.

size\_two\_arm

### Value

arm 0=arm0, 1=arm1 time.accr time to accrual

time.obs time to observation from accrual time.total time to observation from start of study

censor 0=censor, 1=event

reason event description ('[experience ]event', '[loss to ]followup', 'administration[

censoring]')

#### See Also

simulate\_arm for simulating complete data for a single arm, create\_arm for creating an object of class 'arm'.

### **Examples**

```
arm0 <- create_arm(size=120, accr_time=6, surv_scale=0.05, loss_scale=0.005, follow_time=12)
arm1 <- create_arm(size=120, accr_time=6, surv_scale=0.03, loss_scale=0.005, follow_time=12)
simulate_trial(arm0, duration=10)
simulate_trial(arm0, arm1, events=50)</pre>
```

size\_two\_arm

Sample size

### **Description**

Calculate required sample size and expected number of events for a two-arm survival study.

# Usage

```
size_two_arm(arm0, arm1, test = list(test = "weighted logrank"),
power = 0.8, alpha = 0.025, sides = 1)
```

#### **Arguments**

arm0 object of class 'arm'.
arm1 object of class 'arm'.

test list or list of lists. Each list must contain at minimum the key 'test' describing

the type of statistical test. Default test is the "weighted logrank". Kaplan-Meier based tests ("survival difference", "survival ratio", "rmst difference", "rmst ratio", "percentile difference", and "percentile ratio") require the user to define an additional key, either the desired 'milestone' or 'percentile'. The weighted log-rank test does not require additional keys. However, user may choose which weight function ("1"=unweighted, "n"=Gehan-Breslow, "sqrtN"=Tarone-Ware,

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"FH\_[a]\_[b]"= Fleming-Harrington with p=a and q=b) and which approximation for the large-sample mean ("asymptotic", "generalized schoenfeld", "event driven") and variance ("1", "block[ randomization]", "simple[ randomization]") they wish to use. Default choice is 'weight'="1", 'mean.approx'="asymptotic", and 'var.approx'="1". For more details regarding the different mean and variance approximations for the weight log-rank test, please see Yung and Liu (in press).

power	1 - type 2 error rate
alpha	type 1 error rate
sides	1=1-sided test, 2=2-sided test

#### Value

n0	sample size for arm0
n1	sample size for arm1
n	total sample size
d0	expected number of events for arm0
d1	expected number of events for arm1
d	total expected number of events; can be used to convert a time-driven trial to an event-driven trial.

#### References

Yung, G and Liu, Y. (2019). Sample size and power for the weighted log-rank test and Kaplan-Meier based tests with allowance for non-proportional hazards. *Biometrics*. <doi:10.1111/biom.13196>

# See Also

create\_arm for creating an object of class 'arm'.

# **Examples**

```
arm0 <- create_arm(size=120, accr_time=6, surv_scale=0.05, loss_scale=0.005, follow_time=12)
arm1 <- create_arm(size=120, accr_time=6, surv_scale=0.03, loss_scale=0.005, follow_time=12)
size_two_arm(arm0, arm1)
size_two_arm(arm0, arm1, list(test="weighted logrank",
    weight="n",
    mean.approx="generalized schoenfeld",
    var.approx="block"))
size_two_arm(arm0, arm1, list(test="survival difference", milestone=12))
size_two_arm(arm0, arm1, list(test="rmst ratio", milestone=12))
size_two_arm(arm0, arm1, list(test="percentile difference", percentile=0.25))
size_two_arm(arm0, arm1, list(
    list(test="weighted logrank"),
    list(test="survival difference", milestone=12)))</pre>
```

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