

# Package ‘addhazard’

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**Title** Fit Additive Hazards Models for Survival Analysis

**Version** 1.1.0

**Date** 2017-03-20

**Description** Contains tools to fit the additive hazards model to data from a cohort, random sampling, two-phase Bernoulli sampling and two-phase finite population sampling, as well as calibration tool to incorporate phase I auxiliary information into the two-phase data model fitting. This package provides regression parameter estimates and their model-based and robust standard errors. It also offers tools to make prediction of individual specific hazards.

**LazyData** true

**Depends** R (>= 3.3.1)

**Imports** ahaz, survival, rootSolve

**RoxygenNote** 5.0.1

**License** GPL-2

**NeedsCompilation** no

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**ah***Fit Additive Hazards Regression Models*

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**Description**

Fit a semiparametric additive hazard model 'ah'

$$\lambda(t|Z = z) = \lambda_0(t) + \beta'z.$$

The estimating procedures follow Lin & Ying (1994).

**Usage**

```
ah(formula, data, robust, weights, ties, ...)
```

**Arguments**

formula	a formula object for the regression model of the form response ~ predictors. The outcome is a survival object created by <a href="#">Surv</a> .
data	a data frame. Input dataset.
robust	a logical variable. Robust standard errors are provided if robust == TRUE.
weights	a numeric vector. The weight of each observation.
ties	a logical variable. FALSE if there are no ties in the censored failure times.
...	additional arguments to be passed to the low level regression fitting functions.

**Value**

An object of class 'ah' representing the fit.

**Note**

The response variable is a survival object. If there are ties in the survival time, in the current version we recommend users to break ties by adding a small random number to the survival time. An example is provided. The regression model can be univariate or multivariate. This package is built upon the function [ahaz](#) by Anders Gorst-Rasmussen.

**References**

Lin, D.Y. & Ying, Z. (1994). Semiparametric analysis of the additive risk model. *Biometrika*; 81:61-71.

**See Also**

[predict.ah](#) for prediction based on fitted [ah](#) model, [nwtsc](#) for the description of nwtsc dataset

**Examples**

```

library(survival)
### using the first 100 rows in nwtsco to build an additive hazards model
nwts<- nwtsco[1:100,]

### fit the additive hazards model to the data
### the model-based standard errors are reported when setting robust = FALSE
fit1 <- ah(Surv(trel,relaps) ~ age + instit, ties = FALSE, data = nwts, robust = FALSE)
summary(fit1)

### fit the additive hazards model to the data with robust standard errors
fit2 <- ah(Surv(trel,relaps) ~ age + instit, ties = FALSE, data = nwts, robust = TRUE)
summary(fit2)

### when there are ties, break the ties first
nwts_all <- nwtsco
nwts_all$trel <- nwtsco$trel + runif(dim(nwts_all)[1],0,1)*1e-8
fit3 <- ah(Surv(trel,relaps) ~ age + instit, ties = FALSE, data = nwts_all, robust = TRUE)
summary(fit3)

```

ah.2ph

*Fit Additive Hazards Regression Models to Two-phase Sampling***Description**

The function fits a semiparametric additive hazards model

$$\lambda(t|Z = z) = \lambda_0(t) + \beta'z.$$

to two-phase sampling data. The estimating procedures follow Hu (2014).

**Usage**

```
ah.2ph(formula, data, R, Pi, ties, robust = FALSE,
       calibration.variables = NULL, ...)
```

**Arguments**

formula	a formula object for the regression model of the form response ~ predictors. The outcome is a survival object created by <a href="#">Surv</a> .
data	a data frame. Input dataset.
R	a phase II membership indicator. A vector of values of 0 and 1. The subject is selected to phase II if R = 1.
Pi	the probability of a subject to be selected to the phase II subsample.
ties	a logical variable. FALSE if there are no ties in the censored failure times.
robust	a logical variable. Robust standard errors are provided if robust = TRUE.

calibration.variables

a vector of some column names of the data. These are the variables available for every observation. They are used to calibrate the weight assigned to each subject in order to improve estimation efficiency.

... additional arguments to be passed to the low level regression fitting functions.

### Value

An object of class 'ah.2h' representing the fit.

### Note

This function estimates both model-based and robust standard errors. It can be used to analyze case-cohort studies. It allows subsampling among cases. It can incorporate the calibration procedure and analyze the combined dataset of phase I and phase II samples.

### References

Jie Hu (2014) A Z-estimation System for Two-phase Sampling with Applications to Additive Hazards Models and Epidemiologic Studies. Dissertation, University of Washington.

### See Also

[predict.ah.2ph](#) for prediction based on fitted additive hazards model with two-phase sampling and [nwtsco](#) for the description of nwtsco dataset.

### Examples

```
library(survival)
### fit an additive hazards model to two-phase sampling data without calibration
nwts2ph$trel <- nwts2ph$trel + runif(dim(nwts2ph)[1],0,1)*1e-8
fit1 <- ah.2ph(Surv(trel,relaps) ~ age + histol, ties = FALSE, data = nwts2ph, R = in.ph2, Pi = Pi,
  robust = FALSE, calibration.variables = NULL)
summary(fit1)

### fit an additive hazards model with calibration on age
fit2 <- ah.2ph(Surv(trel,relaps) ~ age + histol, ties = FALSE, data = nwts2ph, R = in.ph2, Pi = Pi,
  robust = FALSE, calibration.variables = "age")
summary(fit2)

### calibrate on age square
### note if users create a calibration variable, then
### the new variable should be added to the original data frame
nwts2ph$age2 <- nwts2ph$age^2
fit3 <- ah.2ph(Surv(trel,relaps) ~ age + histol, ties = FALSE, data = nwts2ph, R = in.ph2, Pi = Pi,
  robust = FALSE, calibration.variables = "age2")
summary(fit3)

#####
### When phase II samples are obtained by finite Sampling #####
#####
```

```

### calculating the sample size for each stratum
### calculate the strata size
strt.size <- table(nwts2ph$strt)
ph2.strt.size <- table(subset(nwts2ph, in.ph2 == 1)$strt)
### fit an additive hazards model with finite stratified sampling
### calculate the sampling fractions
frac <- ph2.strt.size/strt.size
### treating the problem as bernoulli sampling coupled with calibration on strata sizes
### using frac as the sampling probabilities
nwts2ph_by_FPSS <- nwts2ph
nwts2ph_by_FPSS$Pi <- NULL
for (i in 1:length(strt.size)){
  nwts2ph_by_FPSS$Pi[nwts2ph_by_FPSS$strt ==i] <- frac[i]
}

### create strt indicators, which become our calibration variables
for (i in 1:length(strt.size)){
  nwts2ph_by_FPSS$strt_ind <- as.numeric(nwts2ph_by_FPSS$strt ==i)
  names(nwts2ph_by_FPSS)[ncol(nwts2ph_by_FPSS)] = paste0("strt", i)
}
### fit an additive hazards model with finite sampling
fit4 <- ah.2ph(Surv(trel,relaps) ~ age + histol,
              data = nwts2ph_by_FPSS, ties = FALSE,
              R = in.ph2, Pi = Pi,
              robust = FALSE,
              calibration.variables = c("strt1","strt2","strt3"))

summary(fit4)

```

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nwts2ph

*An hypothetical two-phase sampling dataset based on [nwtsco](#) dataset from the National Wilms Tumor Study (NWTS)*

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## Description

An hypothetical two-phase sampling dataset based on [nwtsco](#) dataset from the National Wilms Tumor Study (NWTS)

## Usage

```
nwts2ph
```

## Format

A data frame with 3915 rows and 15 variables:

We create a hypothetical two-phase sampling (stratified sampling) dataset. In this dataset, only the subjects who have relapses and some of the controls have their samples sent to the central laboratory for more accurate histology examination.

Institutional histology is examined in the local hospital. It is usually available for all the samples. Central histology is more expensive to obtain since the samples have to be sent to the central laboratory and the work requires experienced lab scientists. It is reasonable to assume only some samples were tested for central histology. Following the two-phase sampling design, we selected subjects for central histology measurements based on their outcomes and institutional histology results.

- trel** Time to relapse or last date seen (yr), continuous
- tsur** Time to death or last date seen (yr), continuous
- relaps** Indicator of relapse, 0 = Alive no prior relapse when last seen, 1 = Relapsed after trel years
- dead** Indicator of death, 0 = Alive when last seen, 1 = Died after tsur years
- study** NWTS study, 3 = NWTS-3, 4 = NWTS-4
- stage** Stage of disease, 1=I, 2=II, 3=III, 4=IV
- histol** Central Path histology, 0 = Favorable (FH) and the subject is selected into the phase II subsample (in.ph2 = 1), 1 = Unfavorable (UH) and the subject is selected into the phase II subsample (in.ph2 = 1), NA = subject is NOT selected into the phase II subsample (in.ph2 = 1)
- instit** Institutional histology, 0 = Favorable (FH), 1 = Unfavorable (UH)
- age** Age at diagnosis (yr), continuous
- yr** Year of diagnosis, calendar year
- specwgt** Weight of tumor bearing specimen, in grams (continuous)
- tumdiam** Diameter of tumor, in centimeters (continuous)
- strat** Strata, 1 = Unfavorable Institutional histology and no relapse, 2 = favorable Institutional histology and no relapse, 3 = relapse
- Pi** Sampling probability, 0.5 for strata =1, 0.9 for strata = 2, 0.9 for strata = 3
- in.ph2** Phase II membership, 1 = selected into the phase II subsample, 0 = not selected into the phase II subsample

## Source

This dataset was created based on [nwtsco](#) dataset from the National Wilms Tumor Study (NWTS)

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nwts2ph.generate	<i>This file generate the example dataset nwts2ph importFrom("stats", "rbinom")</i>
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## Description

This file generate the example dataset nwts2ph importFrom("stats", "rbinom")

## Usage

```
nwts2ph.generate(data, seed = 20)
```

**Arguments**

<code>data</code>	the full cohort data
<code>seed</code>	the random seed we use for generating this dataset

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<code>nwtsco</code>	<i>Dataset from the National Wilms Tumor Study (NWTS)</i>
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**Description**

Dataset from the National Wilms Tumor Study (NWTS)

**Usage**

`nwtsco`

**Format**

A data frame with 3915 rows and 12 variables:

**trel** Time to relapse or last date seen (yr), continuous

**tsur** Time to death or last date seen (yr), continuous

**relaps** Indicator of relapse, 0 = Alive no prior relapse when last seen, 1 = Relapsed after trel years

**dead** Indicator of death, 0 = Alive when last seen, 1 = Died after tsur years

**study** NWTS study, 3 = NWTS-3, 4 = NWTS-4

**stage** Stage of disease, 1=I, 2=II, 3=III, 4=IV

**histol** Central Path histology, 0 = Favorable (FH), 1 = Unfavorable (UH)

**instit** Institutional histology, 0 = Favorable (FH), 1 = Unfavorable (UH)

**age** Age at diagnosis (yr), continuous

**yr** Year of diagnosis, calendar year

**specwgt** Weight of tumor bearing specimen, in grams (continuous)

**tumdiam** Diameter of tumor, in centimeters (continuous)

**Source**

Originally used by M. Kulich and D.Y. Lin: Improving the efficiency of relative-risk estimation in case-cohort studies. *J Amer Statis Assoc* 99:832-844, 2004.

predict.ah

*Prediction Based on the Fitted Additive Hazards Model***Description**

This function predicts a subject's overall hazard rates at given time points based on this subject's covariate values. The prediction function is an additive hazards model fitted using [ah](#).

**Usage**

```
## S3 method for class 'ah'
predict(object, newdata, newtime, ...)
```

**Arguments**

object	an object of class inhering from <a href="#">ah</a> .
newdata	a dataframe of an individual's predictors.
newtime	a given sequence of time points at which the prediction is performed. The time should be on the same scale as the survival time in <a href="#">Surv</a> .
...	further arguments passed to or from other methods.

**Value**

A dataframe including the time points for prediction, predicted values and their standard errors.

**See Also**

[ah](#) for fitting the additive hazards model, [nwtsc0](#) for the description of nwtsc0 dataset

**Examples**

```
library(survival)
### fit the additive hazards model to the data
nwtsc0 <- nwtsc0[1:100,]
fit <- ah(Surv(trel,relaps) ~ age + instit, data = nwtsc0, ties = FALSE, robust = FALSE)

### see the covariate names in the prediction function
fit$call
### the newdata should be a dataframe
### the variable names are the same as the covariate names in
### the prediction function
newdata <- data.frame(age=60, instit =1)

### an alternative way to give the newdata
newdata <- nwtsc0[101,]

### based on this subject's covariate values, the function predicts individual specific
### hazard rates at time points 3 and 5
predict(fit, newdata, newtime = c(3,5))
```



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predict.ah.2ph	<i>Prediction Based on the Additive Hazards Model Fitted from Two-phase Sampling</i>
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### Description

This function predicts a subject's overall hazard rates at given time points based on this subject's covariate values. The prediction function is an object from [ah.2ph](#). The estimating procedures follow Hu (2014).

### Usage

```
## S3 method for class 'ah.2ph'  
predict(object, newdata, newtime, ...)
```

### Arguments

object	an object of class inhering from 'ah.2ph'.
newdata	a dataframe of an individual's predictors.
newtime	a given sequence of time points at which the prediction is performed.
...	further arguments passed to or from other methods.

### Value

A dataframe including the given time points, predicted hazards, their standard errors, their variances, the phase I component of the variance for predicted hazards and the phase II component of the variance.

### References

Jie Hu (2014) A Z-estimation System for Two-phase Sampling with Applications to Additive Hazards Models and Epidemiologic Studies. Dissertation, University of Washington.

### See Also

[ah.2ph](#) for fitting the additive hazards model with two-phase

### Examples

```
library(survival)  
### load data  
nwtsc <- nwtsc[1:100,]  
  
### create strata based on institutional histology and disease status  
nwtsc$strat <- 1+nwtsc$instit  
### add a stratum containing all (relapsed) cases  
nwtsc$strat[nwtsc$relaps==1] <- 3
```

```

### assign phase II subsampling probabilities
### oversample unfavorable histology (instit =1) and cases
### Pi = 0.5 for instit =0, Pi =1 for instit =1 and relaps =1
nwts$Pi<- 0.5 * (nwts$strt == 1) + 1 * (nwts$strt == 2) + 1 * (nwts$strt == 3)

### generate phase II sampling indicators
N <- dim(nwts)[1]
nwts$in.ph2 <- rbinom(N, 1, nwts$Pi)

### fit an additive hazards model to two-phase sampling data without calibration
fit1 <- ah.2ph(Surv(trel,relaps) ~ age + histol,
              data = nwts,
              ties = FALSE,
              R = in.ph2, Pi = Pi,
              robust = FALSE)

### input the new data for prediction
newdata <- nwtsco[101,]
### based on the fitted model fit1, perform prediction at time points t =3 and t= 5
predict(fit1, newdata, newtime = c(3,5))

### fit an additive hazards model to two-phase sampling data with calibration
### The calibration variable is stage
fit2 <- ah.2ph(Surv(trel,relaps) ~ age + histol, data = nwts, R = in.ph2, Pi = Pi,
              ties = FALSE, robust = FALSE, calibration.variables = "stage")

### based on the fitted model fit2, perform prediction at time points t =3 and t= 5
predict(fit2, newdata, newtime = c(3,5))

## Not run:
### The calibration variable is stage, when set robust = TRUE
fit3 <- ah.2ph(Surv(trel,relaps) ~ age + histol, data = nwts, R = in.ph2, Pi = Pi,
              ties = FALSE, robust = TRUE, calibration.variables = "stage")

### based on the fitted model fit2, perform prediction at time points t =3 and t= 5
predict(fit3, newdata, newtime = c(3,5))

## End(Not run)

```

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