

Package ‘NPCox’

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Type Package

Title Nonparametric and Semiparametric Proportional Hazards Model

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Description An estimation procedure for the analysis of nonparametric proportional hazards model (e.g. $h(t) = h_0(t)\exp(b(t)'Z)$), providing estimation of $b(t)$ and its pointwise standard errors, and semiparametric proportional hazards model (e.g. $h(t) = h_0(t)\exp(b(t)'Z_1 + c*Z_2)$), providing estimation of $b(t)$, c and their standard errors. More details can be found in Lu Tian et al. (2005) <[doi:10.1198/016214504000000845](https://doi.org/10.1198/016214504000000845)>.

License GPL-3

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| | |
|------------|--|
| bandselect | <i>Bandwidth selection function for 'npcox' and 'spcox'.</i> |
|------------|--|

Description

The embeded bandwidth selection function in 'npcox' and 'spcox' together with prediction error calculation.

Usage

```
bandselect(cva, delta, obstime, bandwidth = FALSE)
```

Arguments

| | |
|-----------|---|
| cva | Covariate Z in $h(t) = h_0(t)\exp(b(t)'Z)$ |
| delta | Right censoring indicator for the model |
| obstime | The observed time = min(censoring time, observed failure time) |
| bandwidth | Bandwidth for kernel function, which can be specified. The default value is FALSE and can be selected through least prediction error over all subjects. |

Details

This is some description of this function.

'bandwidth_selection' function can provide the prediction error calculation with given bandwidth, or produce the optimal bandwidth based on an arithmetic progression of bandwidths.

Value

Prediction error on given bandwidth or list that contains the optimal bandwidth, an arithmetic progression of bandwidths with corresponding prediction error.

Examples

```
data(pbc)
dta = na.omit(pbc[,c('time', 'status', 'age', "edema")])
dta = dta[dta$time>600 & dta$time<2000,]
dta[, 'status'] = sign(dta[, 'status'])
colnames(dta) = c('time', 'status', 'age', "edema")
res = bandselect(cva = dta[,3:4], delta = dta$status, obstime = dta$time, bandwidth = 700)
```

npcox

*Nonparametric and semiparametric Cox regression model.***Description**

Estimation of proportional hazards (PH) model with time-varying coefficients. Users should anticipate a significant increase in estimation time when using the 'SE = TRUE' option. Both the number of covariates and the sample size can lead to estimation time increasing quadratically.

Usage

```
npcox(cva, delta, obstime, SE = FALSE, bandwidth = FALSE, resamp = 100)
```

Arguments

| | |
|-----------|---|
| cva | Covariate Z in $h(t) = h_0(t)\exp(b(t)'Z)$ |
| delta | Right censoring indicator for the model |
| obstime | The observed time = $\min(\text{censoring time, observed failure time})$ |
| SE | Whether or not the estimation of standard error through resampling method will be done. The default value is FALSE. |
| bandwidth | Bandwidth for kernel function, which can be specified. The default value is FALSE and can be selected through least prediction error over all subjects. |
| resamp | Number of resampling for estimation of pointwise standard error. The default value is 100. |

Details

This is some description of this function.

'npcox' function is designed for PH model with time-varying coefficients, $h(t) = h_0(t)\exp(b(t)'Z)$, providing estimation of $b(t)$ and its pointwise standard errors on $[\text{bandwidth, max}(\text{obstime})-\text{bandwidth}]$.

Value

a list that contain the estimation result of temporal coefficients, standard error estimation, selected or predesigned bandwidth, dataset, unconverged time points.

Examples

```
data(pbc)
pbc = pbc[(pbc$time < 3000) & (pbc$time > 800), ]
Z = pbc[,c("age", "edema")]
colnames(Z) = c("age", "edema")
del = sign(pbc$status)
tim = pbc$time
res = npc Cox(cva = Z, delta = del, obstime = tim, bandwidth = 500)
```

npplot

Nonparametric and semiparametric Cox regression model.

Description

Plot of time-varying coefficient estimated through function 'npcox' or 'spcox'.

Usage

```
npplot(temp, xrange = NULL, CIlevel = 0.95)
```

Arguments

| | |
|---------|---|
| temp | Estimation result from function 'npcox' or 'spcox'. |
| xrange | Illustration range for x-axis. |
| CIlevel | The default confidence level stands at 0.95. |

Details

This is some description of this function.

This is the plot function for

Value

The plot of nonparametric coefficients function 'npcox' or 'spcox'.

Examples

```
data(pbc)
pbc = pbc[(pbc$time < 3000) & (pbc$time > 800), ]
Z = pbc[,c("age", "edema")]
colnames(Z) = c("age", "edema")
del = sign(pbc$status)
tim = pbc$time
res = npcoc(cva = Z, delta = del, obstime = tim, bandwidth = 500)
op = par(mfrow = c(1,2))
npplot(res)
par(op)
```

| | |
|--------|---|
| npsimu | <i>Simulation data generation of PH model with time-varying coefficients.</i> |
|--------|---|

Description

An example of data generation for nonparametric PH model.

Usage

```
npsimu(n, cenpara = NULL)
```

Arguments

| | |
|---------|--|
| n | The number of sample size, which can be self-determined. |
| cenpara | Censoring parameter, which is supposed to be positive, for adjustment of censoring rate. |

Details

This is some description of this function.

'npsimu' is designed for PH model with time-varying coefficients, $h(t) = h_0(t)\exp(b(t)'Z)$, generating the covariates, observed time and censoring indicator.

Value

a list that contain covariates, observed time and censoring indicator.

Examples

```
data = npsimu(200)
```

| | |
|------|---|
| pbcc | <i>Mayo Clinic Primary Biliary Cholangitis Data</i> |
|------|---|

Description

Primary biliary cholangitis is an autoimmune disease leading to destruction of the small bile ducts in the liver. Progression is slow but inexorable, eventually leading to cirrhosis and liver decompensation. The condition has been recognised since at least 1851 and was named "primary biliary cirrhosis" in 1949. Because cirrhosis is a feature only of advanced disease, a change of its name to "primary biliary cholangitis" was proposed by patient advocacy groups in 2014.

This data is from the Mayo Clinic trial in PBC conducted between 1974 and 1984. A total of 424 PBC patients, referred to Mayo Clinic during that ten-year interval, met eligibility criteria for the randomized placebo controlled trial of the drug D-penicillamine. The first 312 cases in the data set

participated in the randomized trial and contain largely complete data. The additional 112 cases did not participate in the clinical trial, but consented to have basic measurements recorded and to be followed for survival. Six of those cases were lost to follow-up shortly after diagnosis, so the data here are on an additional 106 cases as well as the 312 randomized participants.

A nearly identical data set found in appendix D of Fleming and Harrington; this version has fewer missing values.

Usage

```
pbc
data(pbc, package="NPCox")
```

Format

| | |
|-----------|---|
| age: | in years |
| albumin: | serum albumin (g/dl) |
| alk.phos: | alkaline phosphatase (U/liter) |
| ascites: | presence of ascites |
| ast: | aspartate aminotransferase, once called SGOT (U/ml) |
| bili: | serum bilirunbin (mg/dl) |
| chol: | serum cholesterol (mg/dl) |
| copper: | urine copper (ug/day) |
| edema: | 0 no edema, 0.5 untreated or successfully treated 1 edema despite diuretic therapy |
| hepato: | presence of hepatomegaly or enlarged liver |
| id: | case number |
| platelet: | platelet count |
| protime: | standardised blood clotting time |
| sex: | m/f |
| spiders: | blood vessel malformations in the skin |
| stage: | histologic stage of disease (needs biopsy) |
| status: | status at endpoint, 0/1/2 for censored, transplant, dead |
| time: | number of days between registration and the earlier of death, transplantation, or study analysis in July, 1986 |
| trt: | 1/2/NA for D-penicillmain, placebo, not randomised |
| trig: | triglycerides (mg/dl) |

Source

T Therneau and P Grambsch (2000), *Modeling Survival Data: Extending the Cox Model*, Springer-Verlag, New York. ISBN: 0-387-98784-3.

spcox

*Nonparametric and semiparametric Cox regression model.***Description**

Estimation of proportional hazards (PH) model with time-varying coefficients and constant coefficients. Users should anticipate a significant increase in estimation time when using the 'SE = TRUE' option. Both the number of covariates and the sample size can lead to estimation time increasing quadratically.

Usage

```
spcox(
  cva_cons,
  cva_time,
  delta,
  obstime,
  SE = FALSE,
  bandwidth = FALSE,
  resamp = 100
)
```

Arguments

| | |
|-----------|---|
| cva_cons | Covariate Z1 with constant coefficient c in $h(t) = h_0(t)\exp(c'Z1 + b(t)'Z2)$ |
| cva_time | Covariate Z2 with time-varying coefficient b(t) in $h(t) = h_0(t)\exp(c'Z1 + b(t)'Z2)$ |
| delta | Right censoring indicator for the model |
| obstime | The observed time = min(censoring time, observed failure time) |
| SE | Whether or not the estimation of standard error through resampling method will be done. The default value is FALSE. |
| bandwidth | Bandwidth for kernel function, which can be specified. The default value is FALSE and can be selected through least prediction error over all subjects. |
| resamp | Number of resampling for estimation of pointwise standard error. The default value is 100. |

Details

This is some description of this function.

'spcox' is designed for PH model with both time-varying and constant coefficients, $h(t) = h_0(t)\exp(b(t)'Z1 + c*Z2)$, providing estimation of b(t), c and their standard errors.

Value

a list that contain the estimation result of both temporal and constant coefficients, standard error estimation, selected or predesigned bandwidth, dataset, unconverged time points.

Examples

```
data(pbc)
pbc = pbc[(pbc$time < 3000) & (pbc$time > 800), ]
Z1 = as.matrix(pbc[,5])
Z2 = as.matrix(pbc[,c('albumin')])
colnames(Z1) = c('age')
colnames(Z2) = c('albumin')
del = sign(pbc$status)
tim = pbc$time
res1 = spcox(cva_cons = Z1, cva_time = Z2, delta = del, obstime = tim, bandwidth = 500)
```


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