Package 'tbd'

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Title Estimation of Causal Effects with Outcomes Truncated by Death		
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Description Estimation of the survivor average causal effect under outcomes truncated by death, which requires the existence of a substitution variable. It can be applied to both experimental and observational data.		
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2 boot.ci

boot.ci

Estimate the confidence interval of SACE using bootstrap.

Description

Give quantiles of bootstrap samples SACE.

Usage

```
boot.ci(object, nboot = 1000, seed = 100:(100 + nboot - 1), alpha = 0.05,
max.step = 1000, singular.ok = FALSE, print.progress = TRUE)
```

Arguments

object an object of class sace.

nboot a positive integer. The number of bootstrap samples desired.

seed an integer vector with length nboot. Seed to generate samples.

alpha confidence level.

max.step see documentation of sace.
singular.ok see documentation of sace.

print.progress logical. Need progress be printed?

Value

a list with 4 elements:

nskip number of failures during bootstrap.

sace.boot.record

a vector with length nboot-skip. SACE estimates of all bootstrap samples.

boot.sd scaler. Standard deviation of SACE estimates of all bootstrap samples.

ci a vector with length 2. Estimated confidence interval.

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print.sace

Print results of sace

Description

print.sace prints estimation of the SACE (survivor average causal effect).

Usage

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

Arguments

x an object of class sace.... additional arguments.

Value

the input object is returned silently.

sace

Estimation of causal effects with outcomes truncated by death

Description

sace estimates survivor average causal effects (SACE) with outcomes truncated by death.

Usage

```
sace(Z, S, Y, X, A, subset, optim.method = "BFGS", max.step = 1000,
    singular.ok = TRUE, need.variance = TRUE, hessian = TRUE)
```

Arguments

Z	a logical vector. Exposure indicator. Convetionally, 1 means treatment and 0 means control. Must not have missing values.
S	a logical vector. Survival indicator. 1 means survival and \emptyset means death. Must not have missing values.
Υ	a numeric vector. (Univariate) outcomes. May have NA where $S=0$ (since Y is not well-defined where $S=0$).
Χ	an optional numeric matrix or vector. Baseline covariates.
A	an optional numeric matrix or vector. Substitution variable(s) which satisfies the assumptions of "exclusion restriction" and "substitution relevance". See references. If A == NULL, then the naive method, namely OLS, will be used.

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subset an optional vector specifying a subset of obervations to be used.

optim.method The method to be used for maximum likelihood optimization. See optim.

max.step integer. Maximum iterating steps of maximum likelihood optimization.

singular.ok logical. Refers to the OLS estimation of the coefficients alpha_1 and alpha_2

using lm. If FALSE (default), a singular fit raises an error.

need.variance logical. Is variance of parameters and estimators needed? See details.

hessian logical. If TRUE, the hessian returned by optim will be used to compute the in-

formation matrix. If FALSE, the matrix will be calculated by an explicit formula.

Details

This function sace, gives estimation of average causal effects (ACE) with outcomes truncated by death. The identification of SACE relies on the existence of a substitution variable and requires the assumptions of monotonicity, ignorability, exclusion restriction, and relevance. While the naive estimates given by the coefficient of Z from $lm(Y \sim Z + X + A)$, subset = S == 1) are restricted among survivors and therefore may be subject to selection bias, this method gives consistent estimates of the SACE (survivor average causal effect), defined as the average causal effect among the subgroup consisting of subjects who would survive under either exposure, i.e. among the always-survivor group (G = LL). See references for details of the assumptions and the model parameterizations.

Parameters beta and gamma are estimated by MLE, using optim.

If need.variance == TRUE, the asymptotic variance estimators of both parameters and estimators will be given. This requires the **numDeriv** package.

Value

a list with following elements:

CALL function call.

data used (within the specified subset).

optim.method method used for optimization.

need.variance is variance of parameters and estimators needed?

n sample size.

mu_0_LL average potential outcomes among control group, E[Y(0)|G=LL]. mu_1_LL average potential outcomes among treatment group, E[Y(1)|G=LL].

sace survivor average causal effect, equals mu_1_LL-mu_0_LL.

beta $PrS(1) = 1|X, A = expit(\beta_0 + X'\beta_1 + A\beta_2)$, estimated by MLE.

gamma $PrS(0)=1|X,A/PrS(1)=1|X,A=expit(\gamma_0+X'\gamma_1+A\gamma_2), \text{ estimated }$

by MLE.

beta_gamma.convergence

indicator of convergence of MLE optimization of beta and gamma. 0 means

convergence. See optim.

alpha_1 $E[Y(0)|Z=0,G=LL,X,A] = \alpha_{10} + X'\alpha_{11} + A\alpha_{12}, \text{ coefficients of lm(Y } \sim 1.5 \text{ MeV})$

1 + X + A, subset = Z == 0).

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alpha_2
$$E[Y(1)|Z=1, G=LL, X, A] = \alpha_{20} + X'\alpha_{21} + G\alpha_{22}$$
, coefficients of lm(Y ~ 1 + X + W. expit, subset = (Z == 1 & S == 1)).

The following items will be given only if need.variance == TRUE:

beta.var estimated asymptotic covariance matrix of beta.

gamma.var estimated asymptotic covariance matrix of gamma.

relevance.Pvalue

P value of the asymptotic chi-squared test on the relevance assumption for the substitution variable. A large P value suggests that the relevance assumption may not hold, namely, the substitution variable(s) may have little impact on the latent survival type.

alpha_1.var estimated asymptotic covariance matrix of alpha_1. alpha_2.var estimated asymptotic covariance matrix of alpha 2.

mu_0_LL.var estimated asymptotic variance of mu_0_LL.
mu_1_LL.var estimated asymptotic variance of mu_1_LL.
sace.var estimated asymptotic variance of the SACE.

Note

The length of vectors Z, Y, S, as well as the row number of matrix X and A must equal the sample size n.

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References

Linbo Wang, Xiao-Hua Zhou, Thomas S. Richardson; Identification and estimation of causal effects with outcomes truncated by death, Biometrika, Volume 104, Issue 3, 1 September 2017, Pages 597-612, https://doi.org/10.1093/biomet/asx034

Examples

```
attach(simulated_data)
X <- cbind(X.X1, X.V2, X.V3)
sace.result <- sace(Z, S, Y, X, A)
sace</pre>
```

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simulated_data

Simulated data with known SACE

Description

This simulated dataset is to illustrate how to use sace to estimate the SACE, and compare it with other naive methods. In this simulated data, by design, there is confounding between Z and Y caused by X, and confounding between S and Y caused by X.

Format

A data frame with 5000 observations and 7 variables. Z, A, Y, S are 1-dimensional, and X is 3-dimensional. The variables are as follows:

- Z Binary treatment
- **X.X1** A factor covariate with 2 levels (1 and -1)
- X.V2 A continuous covariate
- X.V3 A contunuous covariate
- A The substitution variable which is continuous
- Y The continuous outcome. NA where S=0
- **S** The survival indicator. 1 means survival and 0 means death.

Source

The dataset is generated by the simulation design of Wang et al. 2017 with $\delta_1 = 1$ and $\delta_0 = 1$, which allows confounding between Z and Y caused by X, and confounding between S and Y caused by X.

References

Linbo Wang, Xiao-Hua Zhou, Thomas S. Richardson; Identification and estimation of causal effects with outcomes truncated by death, Biometrika, Volume 104, Issue 3, 1 September 2017, Pages 597-612, https://doi.org/10.1093/biomet/asx034

summary.sace

Summarize results of sace

Description

summary.sace summary estimation of the SACE (survivor average causal effect) and all other model parameters.

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Usage

```
## S3 method for class 'sace'
summary(object, ...)
```

Arguments

```
object an object of class sace.
... additional arguments.
```

Value

the input object is returned silently.

Note

If need.variance is TRUE, sace must have been called with need.variance == TRUE, so that the information needed was recorded.

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