

Package ‘crossurr’

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Type Package

Title Cross-Fitting for Doubly Robust Evaluation of High-Dimensional Surrogate Markers

Version 1.1.2

Description Doubly robust methods for evaluating surrogate markers as outlined in: Agniel D, Hejblum BP, Thiebaut R & Parast L (2022). “Doubly robust evaluation of high-dimensional surrogate markers”, *Biostatistics* <[doi:10.1093/biostatistics/kxac020](https://doi.org/10.1093/biostatistics/kxac020)>. You can use these methods to determine how much of the overall treatment effect is explained by a (possibly high-dimensional) set of surrogate markers.

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Depends R (>= 3.6.0)

Imports dplyr, gbm, glmnet, glue, parallel, pbapply, purrr, ranger, RCAL, rlang, SIS, stats, SuperLearner, tibble, tidy

Encoding UTF-8

RoxygenNote 7.3.2

NeedsCompilation no

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sim_data	<i>A simple function to simulate example data.</i>
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Description

A simple function to simulate example data.

Usage

```
sim_data(n, p)
```

Arguments

n	number of simulated observations
p	number of simulated variables

Value

toy dataset used for demonstrating the methods with outcome y , treatment a , covariates $x.1$, $x.2$, and surrogates $s.1$, $s.2$, ...

xfr_surrogate	<i>A function for estimating the proportion of treatment effect explained using repeated cross-fitting.</i>
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Description

A function for estimating the proportion of treatment effect explained using repeated cross-fitting.

Usage

```
xfr_surrogate(
  ds,
  x = NULL,
  s,
  y,
  a,
  splits = 50,
  K = 5,
  outcome_learners = NULL,
  ps_learners = NULL,
  interaction_model = TRUE,
  trim_at = 0.05,
  outcome_family = gaussian(),
  mthd = "superlearner",
```

```

    n_ptb = 0,
    ...
  )

```

Arguments

<code>ds</code>	a <code>data.frame</code> .
<code>x</code>	names of all covariates in <code>ds</code> that should be included to control for confounding (eg. age, sex, etc). Default is <code>NULL</code> .
<code>s</code>	names of surrogates in <code>ds</code> .
<code>y</code>	name of the outcome in <code>ds</code> .
<code>a</code>	treatment variable name (eg. groups). Expect a binary variable made of 1s and 0s.
<code>splits</code>	number of data splits to perform.
<code>K</code>	number of folds for cross-fitting. Default is 5.
<code>outcome_learners</code>	string vector indicating learners to be used for estimation of the outcome function (e.g., "SL.ridge"). See the SuperLearner package for details.
<code>ps_learners</code>	string vector indicating learners to be used for estimation of the propensity score function (e.g., "SL.ridge"). See the SuperLearner package for details.
<code>interaction_model</code>	logical indicating whether outcome functions for treated and control should be estimated separately. Default is <code>TRUE</code> .
<code>trim_at</code>	threshold at which to trim propensity scores. Default is <code>0.05</code> .
<code>outcome_family</code>	default is 'gaussian' for continuous outcomes. Other choice is 'binomial' for binary outcomes.
<code>mthd</code>	selected regression method. Default is 'superlearner', which uses the SuperLearner package for estimation. Other choices include 'lasso' (which uses <code>glmnet</code>), 'sis' (which uses SIS), 'cal' (which uses RCAL).
<code>n_ptb</code>	Number of perturbations. Default is 0 which means asymptotic standard errors are used.
<code>...</code>	additional parameters (in particular for <code>super_learner</code>)

Value

a tibble with columns:

- `Rm`: estimate of the proportion of treatment effect explained, computed as the median over the repeated splits.
- `R_se0` standard error for the PTE, accounting for the variability due to splitting.
- `R_cil0` lower confidence interval value for the PTE.
- `R_cih0` upper confidence interval value for the PTE.
- `Dm`: estimate of the overall treatment effect, computed as the median over the repeated splits.

- D_{se0} standard error for the overall treatment effect, accounting for the variability due to splitting.
- D_{ci10} lower confidence interval value for the overall treatment effect.
- D_{cih0} upper confidence interval value for the overall treatment effect.
- D_{sm} : estimate of the residual treatment effect, computed as the median over the repeated splits.
- $D_{s_{se0}}$ standard error for the residual treatment effect, accounting for the variability due to splitting.
- $D_{s_{ci10}}$ lower confidence interval value for the residual treatment effect.
- $D_{s_{cih0}}$ upper confidence interval value for the residual treatment effect.

Examples

```
n <- 100
p <- 20
q <- 2
wds <- sim_data(n = n, p = p)

if(interactive()){
  lasso_est <- xfr_surrogate(ds = wds,
    x = paste('x.', 1:q, sep = ''),
    s = paste('s.', 1:p, sep = ''),
    a = 'a',
    y = 'y',
    splits = 2,
    K = 2,
    trim_at = 0.01,
    mthd = 'lasso',
    ncores = 1)
}
```

xf_surrogate

A function for estimating the proportion of treatment effect explained using cross-fitting.

Description

A function for estimating the proportion of treatment effect explained using cross-fitting.

Usage

```
xf_surrogate(
  ds,
  x = NULL,
  s,
  y,
  a,
```

```

K = 5,
outcome_learners = NULL,
ps_learners = outcome_learners,
interaction_model = TRUE,
trim_at = 0.05,
outcome_family = gaussian(),
mthd = "superlearner",
n_ptb = 0,
ncores = parallel::detectCores() - 1,
...
)

```

Arguments

ds	a data.frame.
x	names of all covariates in ds that should be included to control for confounding (eg. age, sex, etc). Default is NULL.
s	names of surrogates in ds.
y	name of the outcome in ds.
a	treatment variable name (eg. groups). Expect a binary variable made of 1s and 0s.
K	number of folds for cross-fitting. Default is 5.
outcome_learners	string vector indicating learners to be used for estimation of the outcome function (e.g., "SL.ridge"). See the SuperLearner package for details.
ps_learners	string vector indicating learners to be used for estimation of the propensity score function (e.g., "SL.ridge"). See the SuperLearner package for details.
interaction_model	logical indicating whether outcome functions for treated and control should be estimated separately. Default is TRUE.
trim_at	threshold at which to trim propensity scores. Default is 0.05.
outcome_family	default is 'gaussian' for continuous outcomes. Other choice is 'binomial' for binary outcomes.
mthd	selected regression method. Default is 'superlearner', which uses the SuperLearner package for estimation. Other choices include 'lasso' (which uses glmnet), 'sis' (which uses SIS), 'cal' (which uses RCAL).
n_ptb	Number of perturbations. Default is 0 which means asymptotic standard errors are used.
ncores	number of CPUs used for parallel computations. Default is parallel::detectCores()-1
...	additional parameters (in particular for super_learner)

Value

a tibble with columns:

- R: estimate of the proportion of treatment effect explained, equal to $1 - \text{deltahat}_s/\text{deltahat}$.
- R_se standard error for the PTE.
- deltat_hat_s: residual treatment effect estimate.
- deltat_hat_s_se: standard error for the residual treatment effect.
- pi_o: estimate of the proportion of overlap.
- R_o: PTE only in the overlap region.
- R_o_se: the standard error for R_o.
- deltat_hat_s_o: residual treatment effect in overlap region,
- deltat_hat_s_se_o: standard error for deltat_hat_s_o.
- deltat_hat: overall treatment effect estimate.
- deltat_hat_se: standard error for overall treatment effect estimate.
- delta_diff: difference between the treatment effects, equal to the numerator of PTE.
- dd_se: standard error for delta_diff

Examples

```

n <- 300
p <- 50
q <- 2
wds <- sim_data(n = n, p = p)

if(interactive()){
  sl_est <- xf_surrogate(ds = wds,
    x = paste('x.', 1:q, sep = ''),
    s = paste('s.', 1:p, sep = ''),
    a = 'a',
    y = 'y',
    K = 4,
    trim_at = 0.01,
    mthd = 'superlearner',
    outcome_learners = c("SL.mean", "SL.lm", "SL.svm", "SL.ridge"),
    ps_learners = c("SL.mean", "SL.glm", "SL.svm", "SL.lda"),
    ncores = 1)

  lasso_est <- xf_surrogate(ds = wds,
    x = paste('x.', 1:q, sep = ''),
    s = paste('s.', 1:p, sep = ''),
    a = 'a',
    y = 'y',
    K = 4,
    trim_at = 0.01,
    mthd = 'lasso',
    ncores = 1)
}

```

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