

Package ‘evalITR’

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Title Evaluating Individualized Treatment Rules

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Description Provides various statistical methods for evaluating Individualized Treatment Rules under randomized data. The provided metrics include Population Average Value (PAV), Population Average Prescription Effect (PAPE), Area Under Prescription Effect Curve (AUPEC). It also provides the tools to analyze Individualized Treatment Rules under budget constraints. Detailed reference in Imai and Li (2023) <[doi:10.1080/01621459.2021.1923511](https://doi.org/10.1080/01621459.2021.1923511)>.

License GPL (>= 2)

URL <https://github.com/MichaelLLi/evalITR>,
<https://michaellli.github.io/evalITR/>,
<https://jialul.github.io/causal-ml/>

BugReports <https://github.com/MichaelLLi/evalITR/issues>

Depends dplyr (>= 1.0), MASS (>= 7.0), Matrix (>= 1.0), quadprog (>= 1.0), R (>= 3.5.0), stats

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AUPEC	<i>Estimation of the Area Under Prescription Evaluation Curve (AU-PEC) in Randomized Experiments</i>
-------	--

Description

This function estimates AUPEC. The details of the methods for this design are given in Imai and Li (2019).

Usage

```
AUPEC(T, tau, Y, centered = TRUE)
```

Arguments

T	A vector of the unit-level binary treatment receipt variable for each sample.
tau	A vector of the unit-level continuous score for treatment assignment. We assume those that have $\tau < 0$ should not have treatment. Conditional Average Treatment Effect is one possible measure.
Y	A vector of the outcome variable of interest for each sample.
centered	If TRUE, the outcome variables would be centered before processing. This minimizes the variance of the estimator. Default is TRUE.

Value

A list that contains the following items:

aupec	The estimated Area Under Prescription Evaluation Curve
sd	The estimated standard deviation of AUPEC.
vec	A vector of points outlining the AUPEC curve across each possible budget point for the dataset. Each step increases the budget by $1/n$ where n is the number of data points.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2019). “Experimental Evaluation of Individualized Treatment Rules”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
tau = c(0,0.1,0.2,0.3,0.4,0.5,0.6,0.7)
Y = c(4,5,0,2,4,1,-4,3)
aupeclist <- AUPEC(T,tau,Y)
aupeclist$aupec
aupeclist$sd
aupeclist$vec
```

AUPECcv

Estimation of the Area Under Prescription Evaluation Curve (AU-PEC) in Randomized Experiments Under Cross Validation

Description

This function estimates AUPEC. The details of the methods for this design are given in Imai and Li (2019).

Usage

```
AUPECcv(T, tau, Y, ind, centered = TRUE)
```

Arguments

T	A vector of the unit-level binary treatment receipt variable for each sample.
tau	A matrix where the <i>i</i> th column is the unit-level continuous score for treatment assignment generated in the <i>i</i> th fold.
Y	The outcome variable of interest.
ind	A vector of integers (between 1 and number of folds inclusive) indicating which testing set does each sample belong to.
centered	If TRUE, the outcome variables would be centered before processing. This minimizes the variance of the estimator. Default is TRUE.

Value

A list that contains the following items:

aupec	The estimated AUPEC.
sd	The estimated standard deviation of AUPEC.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2019). “Experimental Evaluation of Individualized Treatment Rules”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
tau = matrix(c(0,0.1,0.2,0.3,0.4,0.5,0.6,0.7,-0.5,-0.3,-0.1,0.1,0.3,0.5,0.7,0.9),nrow = 8, ncol = 2)
Y = c(4,5,0,2,4,1,-4,3)
ind = c(rep(1,4),rep(2,4))
aupeclist <- AUPECcv(T, tau, Y, ind)
aupeclist$aupec
aupeclist$sd
```

compute_qoi	<i>Compute Quantities of Interest (PAPE, PAPEp, PAPDp, AUPEC, GATE, GATEcv)</i>
-------------	---

Description

Compute Quantities of Interest (PAPE, PAPEp, PAPDp, AUPEC, GATE, GATEcv)

Usage

```
compute_qoi(fit_obj, algorithms)
```

Arguments

- fit_obj An output object from fit_itr function.
- algorithms Machine learning algorithms

compute_qoi_user	<i>Compute Quantities of Interest (PAPE, PAPEp, PAPDp, AUPEC, GATE, GATEcv) with user defined functions</i>
------------------	---

Description

Compute Quantities of Interest (PAPE, PAPEp, PAPDp, AUPEC, GATE, GATEcv) with user defined functions

Usage

```
compute_qoi_user(user_itr, Tcv, Ycv, data, ngates, budget, ...)
```

Arguments

user_itr	A user-defined function to create an ITR. The function should take the data as input and return an unit-level continuous score for treatment assignment. We assume those that have score less than 0 should not have treatment. The default is NULL, which means the ITR will be estimated from the estimate_itr.
Tcv	A vector of the unit-level binary treatment.
Ycv	A vector of the unit-level continuous outcome.
data	A data frame containing the variables of interest.
ngates	The number of gates to be used in the GATE function.
budget	The maximum percentage of population that can be treated under the budget constraint.
...	Additional arguments to be passed to the user-defined function.

consist.test	<i>The Consistency Test for Grouped Average Treatment Effects (GATEs) in Randomized Experiments</i>
--------------	---

Description

Tests whether treatment effects are nondecreasing across score groups.

Usage

```
consist.test(T, tau, Y, ngates = 5, nsim = 10000, centered = TRUE)
```

Arguments

T	Binary treatment indicator (0 or 1).
tau	Continuous score vector.
Y	Outcome vector.
ngates	Number of groups (at least 2).
nsim	Number of Monte Carlo draws (at least 2).
centered	Whether to center outcomes before estimation.

Details

See [GATE](#) for inference details.

Value

A list with test statistic `stat` and p-value `pval`.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2022). “Statistical Inference for Heterogeneous Treatment Effects Discovered by Generic Machine Learning in Randomized Experiments”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
tau = c(0,0.1,0.2,0.3,0.4,0.5,0.6,0.7)
Y = c(4,5,0,2,4,1,-4,3)
consisttestlist <- consist.test(T,tau,Y,ngates=2)
consisttestlist$stat
consisttestlist$pval
```

consistcv.test	<i>The Consistency Test for Grouped Average Treatment Effects (GATEs) under Cross Validation in Randomized Experiments</i>
----------------	--

Description

Tests whether treatment effects are nondecreasing across cross-validated score groups.

Usage

```
consistcv.test(T, tau, Y, ind, ngates = 5, nsim = 10000, centered = TRUE)
```

Arguments

T	Binary treatment indicator (0 or 1).
tau	A matrix of scores with one column per fold. Column i contains predictions for all observations from a model trained without fold i.
Y	Outcome vector.
ind	Integer validation-fold labels starting at 1.
ngates	Number of groups (at least 2).
nsim	Number of Monte Carlo draws (at least 2).
centered	Whether to center outcomes before estimation.

Details

See [GATEcv](#) for inference details.

Value

A list with test statistic `stat` and p-value `pval`.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2022). “Statistical Inference for Heterogeneous Treatment Effects Discovered by Generic Machine Learning in Randomized Experiments”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
tau = matrix(c(0,0.1,0.2,0.3,0.4,0.5,0.6,0.7,-0.5,-0.3,-0.1,0.1,0.3,0.5,0.7,0.9),nrow = 8, ncol = 2)
Y = c(4,5,0,2,4,1,-4,3)
ind = c(rep(1,4),rep(2,4))
consisttestlist <- consistcv.test(T,tau,Y,ind,ngates=2)
consisttestlist$stat
consisttestlist$pval
```

create_ml_args

Create general arguments

Description

Create general arguments

Usage

```
create_ml_args(data)
```

Arguments

`data` A dataset

create_ml_args_bart	Create arguments for bartMachine
---------------------	----------------------------------

Description

Create arguments for bartMachine

Usage

```
create_ml_args_bart(data)
```

Arguments

data	A dataset
------	-----------

create_ml_args_bartc	Create arguments for bartCause
----------------------	--------------------------------

Description

Create arguments for bartCause

Usage

```
create_ml_args_bartc(data)
```

Arguments

data	A dataset
------	-----------

create_ml_args_causalforest	Create arguments for causal forest
-----------------------------	------------------------------------

Description

Create arguments for causal forest

Usage

```
create_ml_args_causalforest(data)
```

Arguments

data	A dataset
------	-----------

create_ml_args_lasso	Create arguments for LASSO
----------------------	----------------------------

Description

Create arguments for LASSO

Usage

```
create_ml_args_lasso(data)
```

Arguments

data	A dataset
------	-----------

create_ml_args_superLearner	Create arguments for super learner
-----------------------------	------------------------------------

Description

Create arguments for super learner

Usage

```
create_ml_args_superLearner(data)
```

Arguments

data	A dataset
------	-----------

create_ml_args_svm	Create arguments for SVM
--------------------	--------------------------

Description

Create arguments for SVM

Usage

```
create_ml_args_svm(data)
```

Arguments

data	A dataset
------	-----------

create_ml_args_svm_cls	<i>Create arguments for SVM classification</i>
------------------------	--

Description

Create arguments for SVM classification

Usage

create_ml_args_svm_cls(data)

Arguments

data	A dataset
------	-----------

create_ml_arguments	<i>Create arguments for ML algorithms</i>
---------------------	---

Description

Create arguments for ML algorithms

Usage

create_ml_arguments(outcome, treatment, data)

Arguments

outcome	Outcome of interests
treatment	Treatment variable
data	A dataset

estimate_itr	<i>Estimate individual treatment rules (ITR)</i>
--------------	--

Description

Estimate individual treatment rules (ITR)

Usage

```
estimate_itr(
  treatment,
  form,
  data,
  algorithms,
  budget,
  n_folds = 5,
  split_ratio = 0,
  ngates = 5,
  preProcess = NULL,
  weights = NULL,
  trControl = caret::trainControl(method = "none"),
  tuneGrid = NULL,
  tuneLength = ifelse(trControl$method == "none", 1, 3),
  user_model = NULL,
  SL_library = NULL,
  ...
)
```

Arguments

treatment	Treatment variable
form	a formula object that takes the form $y \sim T + x_1 + x_2 + \dots$
data	A data frame that contains the outcome y and the treatment T .
algorithms	List of machine learning algorithms to be used.
budget	The maximum percentage of population that can be treated under the budget constraint.
n_folds	Number of cross-validation folds. Default is 5.
split_ratio	Split ratio between train and test set under sample splitting. Default is 0.
ngates	The number of groups to separate the data into. The groups are determined by tau. Default is 5.
preProcess	caret parameter
weights	caret parameter
trControl	caret parameter
tuneGrid	caret parameter

tuneLength	caret parameter
user_model	A user-defined function to create an ITR. The function should take the data as input and return a model to estimate the ITR.
SL_library	A list of machine learning algorithms to be used in the super learner.
...	Additional arguments passed to <code>caret::train</code>

Value

An object of `itr` class

evaluate_itr	<i>Evaluate ITR</i>
--------------	---------------------

Description

Evaluate ITR

Usage

```
evaluate_itr(
  fit = NULL,
  user_itr = NULL,
  outcome = c(),
  treatment = c(),
  data = list(),
  budget = 1,
  ngates = 5,
  ...
)
```

Arguments

fit	Fitted model. Usually an output from <code>estimate_itr</code>
user_itr	A user-defined function to create an ITR. The function should take the data as input and return an unit-level continuous score for treatment assignment. We assume those that have score less than 0 should not have treatment. The default is <code>NULL</code> , which means the ITR will be estimated from the <code>estimate_itr</code> .
outcome	A character string of the outcome variable name.
treatment	A character string of the treatment variable name.
data	A data frame containing the variables specified in <code>outcome</code> , <code>treatment</code> , and <code>tau</code> .
budget	The maximum percentage of population that can be treated under the budget constraint.

ngates	The number of gates to use for the ITR. The default is 5. A user-defined function to create an ITR. The function should take the data as input and return an ITR. The output is a vector of the unit-level binary treatment that would have been assigned by the individualized treatment rule. The default is NULL, which means the ITR will be estimated from the estimate_itr. See ?evaluate_itr for an example.
...	Further arguments passed to the function.

Value

An object of itr class

fit_itr	<i>Estimate ITR for Single Outcome</i>
---------	--

Description

Estimate ITR for Single Outcome

Usage

```
fit_itr(data, algorithms, params, folds, budget, user_model, ...)
```

Arguments

data	A dataset.
algorithms	Machine learning algorithms.
params	A list of parameters.
folds	Number of folds.
budget	The maximum percentage of population that can be treated under the budget constraint.
user_model	User's own function to estimated the ITR.
...	Additional arguments passed to caret::train

Value

A list of estimates.

GATE	<i>Estimation of the Grouped Average Treatment Effects (GATEs) in Randomized Experiments</i>
------	--

Description

Estimates grouped average treatment effects from a continuous score.

Usage

```
GATE(T, tau, Y, ngates = 5, centered = FALSE)
```

Arguments

T	Binary treatment indicator (0 or 1).
tau	Continuous score vector.
Y	Outcome vector.
ngates	Number of groups (at least 2).
centered	Whether to center outcomes before estimation.

Details

Standard errors account for estimated group cutoffs and assume continuous treatment effects at the boundaries. Ties are broken randomly; use `set.seed()` for reproducibility. Both arms need at least two observations. Empty group arms return NA standard errors.

Value

A list with group estimates `gate` and standard errors `sd`, ordered by increasing score.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2022). “Statistical Inference for Heterogeneous Treatment Effects Discovered by Generic Machine Learning in Randomized Experiments”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
tau = c(0,0.1,0.2,0.3,0.4,0.5,0.6,0.7)
Y = c(4,5,0,2,4,1,-4,3)
gatelist <- GATE(T,tau,Y,ngates=2)
gatelist$gate
gatelist$sd
```

GATEcv	<i>Estimation of the Grouped Average Treatment Effects (GATEs) in Randomized Experiments Under Cross Validation</i>
--------	---

Description

Estimates grouped average treatment effects from cross-validated scores.

Usage

```
GATEcv(T, tau, Y, ind, ngates = 5, centered = FALSE)
```

Arguments

T	Binary treatment indicator (0 or 1).
tau	A matrix of scores with one column per fold. Column <i>i</i> contains predictions for all observations from a model trained without fold <i>i</i> .
Y	Outcome vector.
ind	Integer validation-fold labels starting at 1.
ngates	Number of groups (at least 2).
centered	Whether to center outcomes before estimation.

Details

Inference assumes a fixed number of folds and stable scoring rules. Each fold needs at least two observations per treatment arm. See [GATE](#) for grouping and covariance details.

Value

A list with group estimates `gate` and standard errors `sd`, ordered by increasing score.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2022). “Statistical Inference for Heterogeneous Treatment Effects Discovered by Generic Machine Learning in Randomized Experiments”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
tau = matrix(c(0,0.1,0.2,0.3,0.4,0.5,0.6,0.7,-0.5,-0.3,-0.1,0.1,0.3,0.5,0.7,0.9),nrow = 8, ncol = 2)
Y = c(4,5,0,2,4,1,-4,3)
ind = c(rep(1,4),rep(2,4))
gatelist <- GATEcv(T, tau, Y, ind, ngates = 2)
gatelist$gate
gatelist$sd
```

het.test	<i>The Heterogeneity Test for Grouped Average Treatment Effects (GATEs) in Randomized Experiments</i>
----------	---

Description

Tests whether treatment effects are equal across score groups.

Usage

```
het.test(T, tau, Y, ngates = 5, centered = TRUE)
```

Arguments

T	Binary treatment indicator (0 or 1).
tau	Continuous score vector.
Y	Outcome vector.
ngates	Number of groups (at least 2).
centered	Whether to center outcomes before estimation.

Details

See [GATE](#) for inference details.

Value

A list with test statistic stat and p-value pval.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2022). “Statistical Inference for Heterogeneous Treatment Effects Discovered by Generic Machine Learning in Randomized Experiments”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
tau = c(0,0.1,0.2,0.3,0.4,0.5,0.6,0.7)
Y = c(4,5,0,2,4,1,-4,3)
hettestlist <- het.test(T,tau,Y,ngates=2)
hettestlist$stat
hettestlist$pval
```

hetcv.test	<i>The Heterogeneity Test for Grouped Average Treatment Effects (GATEs) under Cross Validation in Randomized Experiments</i>
------------	--

Description

Tests whether treatment effects are equal across cross-validated score groups.

Usage

```
hetcv.test(T, tau, Y, ind, ngates = 5, centered = TRUE)
```

Arguments

T	Binary treatment indicator (0 or 1).
tau	A matrix of scores with one column per fold. Column i contains predictions for all observations from a model trained without fold i.
Y	Outcome vector.
ind	Integer validation-fold labels starting at 1.
ngates	Number of groups (at least 2).
centered	Whether to center outcomes before estimation.

Details

See [GATEcv](#) for inference details.

Value

A list with test statistic stat and p-value pval.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2022). “Statistical Inference for Heterogeneous Treatment Effects Discovered by Generic Machine Learning in Randomized Experiments”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
tau = matrix(c(0,0.1,0.2,0.3,0.4,0.5,0.6,0.7,-0.5,-0.3,-0.1,0.1,0.3,0.5,0.7,0.9),nrow = 8, ncol = 2)
Y = c(4,5,0,2,4,1,-4,3)
ind = c(rep(1,4),rep(2,4))
hettestlist <- hetcv.test(T,tau,Y,ind,ngates=2)
hettestlist$stat
hettestlist$pval
```

PAPD	<i>Estimation of the Population Average Prescription Difference in Randomized Experiments</i>
------	---

Description

This function estimates the Population Average Prescription Difference with a budget constraint. The details of the methods for this design are given in Imai and Li (2019).

Usage

```
PAPD(T, Thatfp, Thatgp, Y, budget, centered = TRUE)
```

Arguments

T	A vector of the unit-level binary treatment receipt variable for each sample.
Thatfp	A vector of the unit-level binary treatment that would have been assigned by the first individualized treatment rule. Please ensure that the percentage of treatment units of That is lower than the budget constraint.
Thatgp	A vector of the unit-level binary treatment that would have been assigned by the second individualized treatment rule. Please ensure that the percentage of treatment units of That is lower than the budget constraint.
Y	A vector of the outcome variable of interest for each sample.
budget	The maximum percentage of population that can be treated under the budget constraint. Should be a decimal between 0 and 1.
centered	If TRUE, the outcome variables would be centered before processing. This minimizes the variance of the estimator. Default is TRUE.

Value

A list that contains the following items:

papd	The estimated Population Average Prescription Difference
sd	The estimated standard deviation of PAPD.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2019). “Experimental Evaluation of Individualized Treatment Rules”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
That = c(0,1,1,0,0,1,1,0)
That2 = c(1,0,0,1,1,0,0,1)
Y = c(4,5,0,2,4,1,-4,3)
papdlist <- PAPD(T,That,That2,Y,budget = 0.5)
papdlist$papd
papdlist$sd
```

PAPDcv

Estimation of the Population Average Prescription Difference in Randomized Experiments Under Cross Validation

Description

This function estimates the Population Average Prescription Difference with a budget constraint under cross validation. The details of the methods for this design are given in Imai and Li (2019).

Usage

```
PAPDcv(T, Thatfp, Thatgp, Y, ind, budget, centered = TRUE)
```

Arguments

T	A vector of the unit-level binary treatment receipt variable for each sample.
Thatfp	A matrix where the <i>i</i> th column is the unit-level binary treatment that would have been assigned by the first individualized treatment rule generated in the <i>i</i> th fold. Please ensure that the percentage of treatment units of That is lower than the budget constraint.
Thatgp	A matrix where the <i>i</i> th column is the unit-level binary treatment that would have been assigned by the second individualized treatment rule generated in the <i>i</i> th fold. Please ensure that the percentage of treatment units of That is lower than the budget constraint.
Y	The outcome variable of interest.
ind	A vector of integers (between 1 and number of folds inclusive) indicating which testing set does each sample belong to.

budget	The maximum percentage of population that can be treated under the budget constraint. Should be a decimal between 0 and 1.
centered	If TRUE, the outcome variables would be centered before processing. This minimizes the variance of the estimator. Default is TRUE.

Value

A list that contains the following items:

papd	The estimated Population Average Prescription Difference.
sd	The estimated standard deviation of PAPD.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2019). “Experimental Evaluation of Individualized Treatment Rules”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
That = matrix(c(0,1,1,0,0,1,1,0,1,0,0,1,1,0,0,1), nrow = 8, ncol = 2)
That2 = matrix(c(0,0,1,1,0,0,1,1,1,1,0,0,1,1,0,0), nrow = 8, ncol = 2)
Y = c(4,5,0,2,4,1,-4,3)
ind = c(rep(1,4),rep(2,4))
papdlist <- PAPDcv(T, That, That2, Y, ind, budget = 0.5)
papdlist$papd
papdlist$sd
```

PAPE	<i>Estimation of the Population Average Prescription Effect in Randomized Experiments</i>
------	---

Description

This function estimates the Population Average Prescription Effect with and without a budget constraint. The details of the methods for this design are given in Imai and Li (2019).

Usage

```
PAPE(T, That, Y, budget = NA, centered = TRUE)
```

Arguments

T	A vector of the unit-level binary treatment receipt variable for each sample.
That	A vector of the unit-level binary treatment that would have been assigned by the individualized treatment rule. If <code>budget</code> is specified, please ensure that the percentage of treatment units of <code>That</code> is lower than the budget constraint.
Y	A vector of the outcome variable of interest for each sample.
budget	The maximum percentage of population that can be treated under the budget constraint. Should be a decimal between 0 and 1. Default is NA which assumes no budget constraint.
centered	If TRUE, the outcome variables would be centered before processing. This minimizes the variance of the estimator. Default is TRUE.

Value

A list that contains the following items:

pape	The estimated Population Average Prescription Effect.
sd	The estimated standard deviation of PAPE.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2019). “Experimental Evaluation of Individualized Treatment Rules”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
That = c(0,1,1,0,0,1,1,0)
Y = c(4,5,0,2,4,1,-4,3)
papelist <- PAPE(T,That,Y)
papelist$pape
papelist$sd
```

PAPEcv

Estimation of the Population Average Prescription Effect in Randomized Experiments Under Cross Validation

Description

This function estimates the Population Average Prescription Effect with and without a budget constraint. The details of the methods for this design are given in Imai and Li (2019).

Usage

```
PAPEcv(T, That, Y, ind, budget = NA, centered = TRUE)
```

Arguments

T	A vector of the unit-level binary treatment receipt variable for each sample.
That	A matrix where the <i>i</i> th column is the unit-level binary treatment that would have been assigned by the individualized treatment rule generated in the <i>i</i> th fold. If budget is specified, please ensure that the percentage of treatment units of That is lower than the budget constraint.
Y	The outcome variable of interest.
ind	A vector of integers (between 1 and number of folds inclusive) indicating which testing set does each sample belong to.
budget	The maximum percentage of population that can be treated under the budget constraint. Should be a decimal between 0 and 1. Default is NA which assumes no budget constraint.
centered	If TRUE, the outcome variables would be centered before processing. This minimizes the variance of the estimator. Default is TRUE.

Value

A list that contains the following items:

pape	The estimated Population Average Prescription Effect.
sd	The estimated standard deviation of PAPE.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2019). “Experimental Evaluation of Individualized Treatment Rules”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
That = matrix(c(0,1,1,0,0,1,1,0,1,0,0,1,1,0,0,1), nrow = 8, ncol = 2)
Y = c(4,5,0,2,4,1,-4,3)
ind = c(rep(1,4),rep(2,4))
papelist <- PAPEcv(T, That, Y, ind)
papelist$pape
papelist$sd
```

PAV	<i>Estimation of the Population Average Value in Randomized Experiments</i>
-----	---

Description

This function estimates the Population Average Value. The details of the methods for this design are given in Imai and Li (2019).

Usage

```
PAV(T, That, Y, centered = TRUE)
```

Arguments

T	A vector of the unit-level binary treatment receipt variable for each sample.
That	A vector of the unit-level binary treatment that would have been assigned by the individualized treatment rule. If budget is specified, please ensure that the percentage of treatment units of That is lower than the budget constraint.
Y	A vector of the outcome variable of interest for each sample.
centered	If TRUE, the outcome variables would be centered before processing. This minimizes the variance of the estimator. Default is TRUE.

Value

A list that contains the following items:

pav	The estimated Population Average Value.
sd	The estimated standard deviation of PAV.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2019). “Experimental Evaluation of Individualized Treatment Rules”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
That = c(0,1,1,0,0,1,1,0)
Y = c(4,5,0,2,4,1,-4,3)
pavlist <- PAV(T,That,Y)
pavlist$pav
pavlist$sd
```


PAVcv

*Estimation of the Population Average Value in Randomized Experiments Under Cross Validation***Description**

This function estimates the Population Average Value. The details of the methods for this design are given in Imai and Li (2019).

Usage

```
PAVcv(T, That, Y, ind, centered = TRUE)
```

Arguments

T	A vector of the unit-level binary treatment receipt variable for each sample.
That	A matrix where the i th column is the unit-level binary treatment that would have been assigned by the individualized treatment rule generated in the i th fold. If budget is specified, please ensure that the percentage of treatment units of That is lower than the budget constraint.
Y	The outcome variable of interest.
ind	A vector of integers (between 1 and number of folds inclusive) indicating which testing set does each sample belong to.
centered	If TRUE, the outcome variables would be centered before processing. This minimizes the variance of the estimator. Default is TRUE.

Value

A list that contains the following items:

pav	The estimated Population Average Value.
sd	The estimated standard deviation of PAV.

Author(s)

Michael Lingzhi Li, Technology and Operations Management, Harvard Business School <mili@hbs.edu>, <https://www.michaellz.com/>;

References

Imai and Li (2019). “Experimental Evaluation of Individualized Treatment Rules”,

Examples

```
T = c(1,0,1,0,1,0,1,0)
That = matrix(c(0,1,1,0,0,1,1,0,1,0,0,1,1,0,0,1), nrow = 8, ncol = 2)
Y = c(4,5,0,2,4,1,-4,3)
ind = c(rep(1,4),rep(2,4))
pavlist <- PAVcv(T, That, Y, ind)
pavlist$pav
pavlist$sd
```

plot.itr	<i>Plot the AUPEC curve</i>
----------	-----------------------------

Description

Plot the AUPEC curve

Usage

```
## S3 method for class 'itr'
plot(x, ...)
```

Arguments

- x An object of evaluate_itr() class. This is typically an output of evaluate_itr() function.
- ... Further arguments passed to the function.

Value

A plot of ggplot2 object.

plot_estimate	<i>Plot the GATE estimate</i>
---------------	-------------------------------

Description

Plot the GATE estimate

Usage

```
plot_estimate(x, type, ...)
```

Arguments

- x An table object. This is typically an output of evaluate_itr() function.
- type The metric that you wish to plot. One of GATE, PAPE, PAPEp, or PAPDp.
- ... Further arguments passed to the function.

Value

A plot of ggplot2 object.

```
print.summary.itr
```

Print

Description

Print

Usage

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

Arguments

x	An object of summary.itr class. This is typically an output of summary.itr() function.
...	Other parameters. Currently not supported.

```
print.summary.test_itr
```

Print

Description

Print

Usage

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

Arguments

x	An object of summary.test_itr class. This is typically an output of summary.test_itr() function.
...	Other parameters.

star

Tennessee's Student/Teacher Achievement Ratio (STAR) project

Description

A longitudinal study experimentally evaluating the impacts of class size in early education on various outcomes (Mosteller, 1995)

Usage

```
star
```

Format

A data frame with 1911 observations and 14 variables:

treatment A binary treatment indicating whether a student is assigned to small class and regular class without an aid

g3tlangss A continuous variable measuring student's writing scores

g3treadss A continuous variable measuring student's reading scores

g3tmathss A continuous variable measuring student's math scores

gender Students' gender

race Students' race

birthmonth Students' birth month

birthyear Students' birth year

SCHLURBN Urban or rural

GKENRMNT Enrollment size

GRDRANGE Grade range

GKFRLNCH Number of students on free lunch

GKBUSED Number of students on school buses

GKWHITE Percentage of white students

summary.itr	<i>Summarize estimate_itr output</i>
-------------	--------------------------------------

Description

Summarize estimate_itr output

Usage

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

Arguments

object	An object of estimate_itr class (typically an output of estimate_itr() function).
...	Other parameters.

summary.test_itr	<i>Summarize test_itr output</i>
------------------	----------------------------------

Description

Summarize test_itr output

Usage

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

Arguments

object	An object of test_itr class (typically an output of test_itr() function).
...	Other parameters.

test_itr	<i>Conduct hypothesis tests</i>
----------	---------------------------------

Description

Conduct hypothesis tests

Usage

```
test_itr(fit, nsim = 1000, centered = TRUE, ...)
```

Arguments

fit	Fitted model. Usually an output from <code>estimate_itr</code>
nsim	Number of Monte Carlo simulations used to simulate the null distributions. Default is 1000.
centered	Whether to center outcomes before testing.
...	Further arguments passed to the function.

Value

An object of `test_itr` class

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