Package 'ctmle'

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|--|
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| Description Implements the general template for collaborative targeted maximum likelihood estimation. It also provides several commonly used C-TMLE instantiation, like the vanilla/scalable var able-selection C-TMLE (Ju et al. (2017) <doi:10.1177 0962280217729845="">) and the glmnet-C TMLE algorithm (Ju et al. (2017) <arxiv:1706.10029>).</arxiv:1706.10029></doi:10.1177> |
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bound

set outliers to min/max allowable values. It assumes x contains only numerical data

Description

set outliers to min/max allowable values. It assumes x contains only numerical data

Usage

```
bound(x, bounds)
```

Arguments

x input data

bounds a vector with length 2, contains the min and max of the bound

Value

x truncated input x by min/max in bounds

Examples

```
x <- rnorm(1000)
x <- bound(x, c(-1, 1))</pre>
```

build_gn_seq

Help function to build the sequence of gn candidates in ctmleGeneral

Description

This function helps building gn candidates for ctmleGeneral. It returns gn_candidates_cv, gn_candidates, and folds, which could be directly applied to ctmleGeneral.

Usage

```
build_gn_seq(A, W, SL.library, folds, verbose = TRUE)
```

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Arguments

| Α | binary treatment indicator, 1 - treatment, 0 - control |
|------------|---|
| W | vector, matrix, or dataframe containing baseline covariates for Q bar |
| SL.library | a vector of the names of the estimators for ctmle (need to be prepared in the format for SL, see more details in SuperLearner package), The theory of ctmle requires the estimators are ordered by the model complexity, with the last one be a consistent estimator. |
| folds | The list of indices for the ctmle cross-validation step |
| verbose | A boolean. If print out the training log for Super Learne |

Value

gn_candidates_cv matrix or dataframe, each column stand for a estimate of propensity score. Estimate in the column with larger index should have smaller empirical loss

gn_candidates matrix or dataframe, each column stand for a the cross-validated estimate. For example, the (i,j)-th element is the predicted propensity score by j-th estimator, for i-th observation, when it is in the validation set

folds The list of indices for the ctmle cross-validation step

details The SuperLearner object used to generate gn_candidates_cv

```
N <- 1000
p = 100
V = 5
Wmat <- matrix(rnorm(N * p), ncol = p)</pre>
gcoef \leftarrow matrix(c(-1,-1,rep(-(3/((p)-2)),(p)-2)),ncol=1)
W <- as.data.frame(Wmat)</pre>
g <- 1/(1+exp(Wmat%*%gcoef / 3))</pre>
A \leftarrow rbinom(N, 1, prob = g)
folds <-by(sample(1:N,N), rep(1:V, length=N), list)</pre>
lasso_fit <- cv.glmnet(x = as.matrix(W), y = A, alpha = 1, nlambda = 100, nfolds = 10)</pre>
lasso_lambdas <- lasso_fit$lambda[lasso_fit$lambda <= lasso_fit$lambda.min][1:5]</pre>
# Build template for glmnet
SL.glmnet_new <- function (Y, X, newX, family, obsWeights, id, alpha = 1,
                             nlambda = 100, lambda = 0,...)
    # browser()
    if (!is.matrix(X)) {
           X \leftarrow model.matrix(\sim -1 + ., X)
          newX <- model.matrix(~-1 + ., newX)</pre>
   fit <- glmnet::glmnet(x = X, y = Y,</pre>
                            lambda = lambda,
                            family = family$family, alpha = alpha)
```

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```
pred <- predict(fit, newx = newX, type = "response")</pre>
     fit <- list(object = fit)</pre>
   class(fit) <- "SL.glmnet"</pre>
   out <- list(pred = pred, fit = fit)</pre>
   return(out)
}
# Use a sequence of estimator to build gn sequence:
SL.cv1lasso <- function (..., alpha = 1, lambda = lasso_lambdas[1]){</pre>
   SL.glmnet_new(... , alpha = alpha, lambda = lambda)
}
SL.cv2lasso <- function (..., alpha = 1, lambda = lasso_lambdas[2]){</pre>
    SL.glmnet_new(... , alpha = alpha, lambda = lambda)
}
SL.cv3lasso <- function (..., alpha = 1, lambda = lasso_lambdas[3]){</pre>
    SL.glmnet_new(... , alpha = alpha, lambda = lambda)
}
SL.cv4lasso <- function (..., alpha = 1, lambda = lasso_lambdas[4]){</pre>
     SL.glmnet_new(... , alpha = alpha, lambda = lambda)
SL.library = c('SL.cv1lasso', 'SL.cv2lasso', 'SL.cv3lasso', 'SL.cv4lasso', 'SL.glm')
gn_seq <- build_gn_seq(A = A, W = W, SL.library = SL.library, folds = folds)</pre>
gn_seq$gn_candidates_cv
gn_seq$gn_candidates
```

ctmleDiscrete

Discrete Collaborative Targeted Minimum-loss based Estimation

Description

This function computes the discrete Collaborative Targeted Minimum-loss based Estimator for variable selection. It includes the greedy C-TMLE algorithm (Gruber and van der Laan 2010), and scalable C-TMLE algorithm (Ju, Gruber, and Lendle et al. 2016) with a user-specified order.

Usage

```
ctmleDiscrete(Y, A, W, Wg = W, Q = NULL, preOrder = FALSE, order = NULL,
  patience = FALSE, Qbounds = NULL, cvQinit = FALSE, Qform = NULL,
  SL.library = NULL, alpha = 0.995, family = "gaussian", gbound = 0.025,
  like_type = "RSS", fluctuation = "logistic", verbose = FALSE,
  detailed = FALSE, PEN = FALSE, V = 5, folds = NULL,
  stopFactor = 10^6)
```

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Arguments

| Υ | continuous or binary outcome variable |
|-------------|--|
| Α | binary treatment indicator, 1 for treatment, 0 for control |
| W | vector, matrix, or dataframe containing baseline covariates for Q bar |
| Wg | vector, matrix, or dataframe containing baseline covariates for propensity score model (defaults to W if not supplied by user) |
| Q | n by 2 matrix of initial values for Q0W, Q1W in columns 1 and 2, respectively. Current version does not support SL for automatic initial estimation of Q bar |
| preOrder | boolean indicator for using scalable C-TMLE algorithm or not |
| order | the use-specified order of covariables. Only used when (preOrder = TRUE). If not supplied by user, it would automatically order covariates from W_1 to W_p |
| patience | a number to stop early when the score in the CV function does not improve after so many covariates. Used only when (preOrder = TRUE) |
| Qbounds | bound on initial Y and predicted values for Q. |
| cvQinit | if TRUE, cross-validate initial values for Q to avoid overfits |
| Qform | optional regression formula for estimating initial Q |
| SL.library | optional vector of prediction algorithms for data adaptive estimation of Q, defaults to glm, and glmnet |
| alpha | used to keep predicted initial values bounded away from $(0,1)$ for logistic fluctuation, 0.995 (default) |
| family | family specification for working regression models, generally 'gaussian' for continuous outcomes (default), 'binomial' for binary outcomes |
| gbound | bound on P(A=1 W), defaults to 0.025 |
| like_type | 'RSS' or 'loglike'. The metric to use for forward selection and cross-validation |
| fluctuation | 'logistic' (default) or 'linear', for targeting step |
| verbose | print status messages if TRUE |
| detailed | boolean number. If it is TRUE, return more detailed results |
| PEN | boolean. If true, penalized loss is used in cross-validation step |
| V | Number of folds. Only used if folds is not specified |
| folds | The list of indices for cross-validation step. We recommend the cv-splits in C-TMLE matchs that in gn_candidate_cv |
| stopFactor | Numerical value with default 1e6. If the current empirical likelihood is stopFactor times larger than the best previous one, the construction would stop |
| | |

Value

best_k the index of estimate that selected by cross-validation est estimate of psi_0 $CI\ IC$ -based 95 PV pvalue pvalue for the null hypothesis that PSi = 0

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likelihood sum of squared residuals, based on selected estimator evaluated on all obs or, logistic loglikelihood if like_type != 'RSS'

varIC empirical variance of the influence curve adjusted for estimation of g

varDstar empirical variance of the influence curve

var.psi variance of the estimate

varIC.cv cross-validated variance of the influence curve

penlikelihood.cv penalized cross-validated likelihood

cv.res all cross-validation results for each fold

```
## Not run:
N <- 1000
p = 10
Wmat <- matrix(rnorm(N * p), ncol = p)</pre>
beta1 <- 4+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
beta0 <- 2+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
tauW <- 2
tau <- 2
gcoef \leftarrow matrix(c(-1,-1,rep(-(3/((p)-2)),(p)-2)),ncol=1)
Wm <- as.matrix(Wmat)</pre>
g <- 1/(1+exp(Wm%*%gcoef))</pre>
A \leftarrow rbinom(N, 1, prob = g)
sigma <- 1
epsilon <-rnorm(N,0,sigma)</pre>
Y <- beta0 + tauW*A + epsilon
# Initial estimate of Q
Q \leftarrow cbind(rep(mean(Y[A == 0]), N), rep(mean(Y[A == 1]), N))
# User-suplied initial estimate
time_greedy <- system.time(</pre>
ctmle_discrete_fit1 <- ctmleDiscrete(Y = Y, A = A, W = data.frame(Wmat), Q = Q,</pre>
                                       preOrder = FALSE)
)
# If there is no input Q, then intial Q would be estimated by SL with Sl.library
ctmle_discrete_fit2 <- ctmleDiscrete(Y = Y, A = A, W = data.frame(Wmat),</pre>
                                       preOrder = FALSE, detailed = TRUE)
# scalable C-TMLE with pre-order option; order is user-specified,
# If 'order' is not specified takes order from W1 to Wp.
time_preorder <- system.time(</pre>
ctmle_discrete_fit3 <- ctmleDiscrete(Y = Y, A = A, W = data.frame(Wmat), Q = Q,</pre>
                                       preOrder = TRUE,
                                       order = rev(1:p), detailed = TRUE)
)
# Compare the running time
time_greedy
```

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```
time_preorder
## End(Not run)
```

ctmleGeneral General Template for Collaborative Targeted Maximum Likelihood
Estimation

Description

This function computes the Collaborative Targeted Maximum Likelihood Estimator.

Usage

```
ctmleGeneral(Y, A, W, Wg = W, Q, ctmletype, gn_candidates,
  gn_candidates_cv = NULL, alpha = 0.995, family = "gaussian",
  gbound = 0.025, like_type = "RSS", fluctuation = "logistic",
  verbose = FALSE, detailed = FALSE, PEN = FALSE, g1W = NULL,
  g1WPrev = NULL, V = 5, folds = NULL, stopFactor = 10^6)
```

Arguments

| Υ | continuous or binary outcome variable | |
|------------------|--|--|
| Α | binary treatment indicator, 1 for treatment, 0 for control | |
| W | vector, matrix, or dataframe containing baseline covariates for Q bar | |
| Wg | vector, matrix, or dataframe containing baseline covariates for propensity score model (defaults to W if not supplied by user) | |
| Q | n by 2 matrix of initial values for Q0W, Q1W in columns 1 and 2, respectively. Current version does not support SL for automatic initial estimation of Q bar $^{\circ}$ | |
| ctmletype | 1 or 3. Type of general C-TMLE. Type 1 uses cross-validation to select best gn, while Type 3 directly solves extra clever covariates. | |
| gn_candidates | matrix or dataframe, each column stand for a estimate of propensity score. Estimate in the column with larger index should have smaller empirical loss | |
| gn_candidates_cv | | |
| | matrix or dataframe, each column stand for a the cross-validated estimate. For example, the (i,j)-th element is the predicted propensity score by j-th estimator, for i-th observation, when it is in the validation set | |
| alpha | used to keep predicted initial values bounded away from $(0,1)$ for logistic fluctuation, 0.995 (default) | |
| family | family specification for working regression models, generally 'gaussian' for continuous outcomes (default), 'binomial' for binary outcomes | |
| gbound | bound on P(A=1 W), defaults to 0.025 | |
| like_type | 'RSS' or 'loglike'. The metric to use for forward selection and cross-validation | |
| fluctuation | 'logistic' (default) or 'linear', for targeting step | |
| | | |

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| verbose | print status messages if TRUE |
|------------|--|
| detailed | boolean number. If it is TRUE, return more detailed results |
| PEN | boolean. If true, penalized loss is used in cross-validation step |
| g1W | Only used when type is 3. a user-supplied propensity score estimate. |
| g1WPrev | Only used when type is 3. a user-supplied propensity score estimate, with small fluctuation compared to g1W. |
| V | Number of folds. Only used if folds is not specified |
| folds | The list of indices for cross-validation step. We recommend the cv-splits in C-TMLE matchs that in gn_candidate_cv |
| stopFactor | Numerical value with default 1e6. If the current empirical likelihood is stopFactor times larger than the best previous one, the construction would stop |

Value

best_k the index of estimate that selected by cross-validation est estimate of psi $_0$

CI IC-based 95

pvalue pvalue for the null hypothesis that Psi = 0

likelihood sum of squared residuals, based on selected estimator evaluated on all obs or, logistic loglikelihood if like_type != "RSS"

varIC empirical variance of the influence curve adjusted for estimation of g

varDstar empirical variance of the influence curve

var.psi variance of the estimate

varIC.cv cross-validated variance of the influence curve

penlikelihood.cv penalized cross-validatedlikelihood

cv.res all cross-validation results for each fold

```
N <- 1000
p = 100
V = 5
Wmat <- matrix(rnorm(N * p), ncol = p)
gcoef <- matrix(c(-1,-1,rep(-(3/((p)-2)),(p)-2)),ncol=1)
W <- as.data.frame(Wmat)
g <- 1/(1+exp(Wmat%*%gcoef / 3))
A <- rbinom(N, 1, prob = g)

# Build potential outcome pairs, and the observed outcome Y
beta1 <- 4+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]
beta0 <- 2+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]

tau = 2
sigma <- 1</pre>
```

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```
epsilon <-rnorm(N,0,sigma)</pre>
Y <- beta0 + tau * A + epsilon
# Initial estimate of Q
Q \leftarrow cbind(rep(mean(Y[A == 1]), N), rep(mean(Y[A == 0]), N))
folds <-by(sample(1:N,N), rep(1:V, length=N), list)</pre>
lasso_fit <- cv.glmnet(x = as.matrix(W), y = A, alpha = 1, nlambda = 100, nfolds = 10)
lasso_lambdas <- lasso_fit$lambda[lasso_fit$lambda <= lasso_fit$lambda.min][1:5]
# Build template for glmnet
SL.glmnet_new <- function (Y, X, newX, family, obsWeights, id, alpha = 1,
                           nlambda = 100, lambda = 0,...)
{
    # browser()
    if (!is.matrix(X)) {
          X \leftarrow model.matrix(\sim-1 + ., X)
         newX <- model.matrix(~-1 + ., newX)</pre>
   fit \leftarrow glmnet::glmnet(x = X, y = Y,
                          lambda = lambda,
                          family = family$family, alpha = alpha)
   pred <- predict(fit, newx = newX, type = "response")</pre>
     fit <- list(object = fit)</pre>
   class(fit) <- "SL.glmnet"</pre>
   out <- list(pred = pred, fit = fit)</pre>
   return(out)
}
# Use a sequence of LASSO estimators to build gn sequence:
SL.cv1lasso <- function (... , alpha = 1, lambda = lasso_lambdas[1]){</pre>
   SL.glmnet_new(..., alpha = alpha, lambda = lambda)
}
SL.cv2lasso <- function (..., alpha = 1, lambda = lasso_lambdas[2]){</pre>
    SL.glmnet_new(... , alpha = alpha, lambda = lambda)
}
SL.cv3lasso <- function (..., alpha = 1, lambda = lasso_lambdas[3]){</pre>
    SL.glmnet_new(..., alpha = alpha, lambda = lambda)
SL.cv4lasso \leftarrow function (..., alpha = 1, lambda = lasso_lambdas[4]){}
     SL.glmnet_new(..., alpha = alpha, lambda = lambda)
}
SL.library = c('SL.cv1lasso', 'SL.cv2lasso', 'SL.cv3lasso', 'SL.cv4lasso', 'SL.glm')
# Build the sequence. See more details in the function build_gn_seq, and package SuperLearner
gn_seq <- build_gn_seq(A = A, W = W, SL.library = SL.library, folds = folds)</pre>
# Use the output of build_gn_seq for ctmle general templates
ctmle_fit <- ctmleGeneral(Y = Y, A = A, W = W, Q = Q, ctmletype = 1,</pre>
```

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```
gn_candidates = gn_seq$gn_candidates,
gn_candidates_cv = gn_seq$gn_candidates_cv,
folds = gn_seq$folds, V = length(folds))
```

ctmleGlmnet

Collaborative Targeted Maximum Likelihood Estimation for hyperparameter tuning of LASSO

Description

This function computes the Collaborative Maximum Likelihood Estimation for hyper-parameter tuning of LASSO.

Usage

```
ctmleGlmnet(Y, A, W, Wg = W, Q, lambdas = NULL, ctmletype, V = 5,
  folds = NULL, alpha = 0.995, family = "gaussian", gbound = 0.025,
  like_type = "RSS", fluctuation = "logistic", verbose = FALSE,
  detailed = FALSE, PEN = FALSE, g1W = NULL, g1WPrev = NULL,
  stopFactor = 10^6)
```

Arguments

| Υ | continuous or binary outcome variable |
|-----------|---|
| Α | binary treatment indicator, 1 for treatment, 0 for control |
| W | vector, matrix, or dataframe containing baseline covariates for Q bar |
| Wg | vector, matrix, or dataframe containing baseline covariates for propensity score model (defaults to W if not supplied by user) |
| Q | n by 2 matrix of initial values for Q0W, Q1W in columns 1 and 2, respectively. Current version does not support SL for automatic initial estimation of Q bar |
| lambdas | numeric vector of lambdas (regularization parameter) for glmnet estimation of propensity score, with decreasing order. We recommend the first lambda is selected by external cross-validation. |
| ctmletype | 1, 2 or 3. Type of general C-TMLE. Type 1 uses cross-validation to select best gn, Type 3 directly solves extra clever covariates, and Type 2 uses both cross-validation and extra covariate. See more details in !!! |
| V | Number of folds. Only used if folds is not specified |
| folds | The list of indices for cross-validation step. We recommend the cv-splits in C-TMLE matchs that in gn_candidate_cv |
| alpha | used to keep predicted initial values bounded away from $(0,1)$ for logistic fluctuation, 0.995 (default) |
| family | family specification for working regression models, generally 'gaussian' for continuous outcomes (default), 'binomial' for binary outcomes |
| gbound | bound on P(A=1 W), defaults to 0.025 |
| | |

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'RSS' or 'loglike'. The metric to use for forward selection and cross-validation like_type 'logistic' (default) or 'linear', for targeting step fluctuation print status messages if TRUE verbose detailed boolean number. If it is TRUE, return more detailed results PEN boolean. If true, penalized loss is used in cross-validation step g1W Only used when type is 3. a user-supplied propensity score estimate. g1WPrev Only used when type is 3. a user-supplied propensity score estimate, with small fluctuation compared to g1W. stopFactor Numerical value with default 1e6. If the current empirical likelihood is stopFactor times larger than the best previous one, the construction would stop

Value

best_k the index of estimate that selected by cross-validation

est estimate of psi_0

CI IC-based 95

pvalue pvalue for the null hypothesis that Psi = 0

likelihood sum of squared residuals, based on selected estimator evaluated on all obs or, logistic loglikelihood if like_type != 'RSS'

varIC empirical variance of the influence curve adjusted for estimation of g

varDstar empirical variance of the influence curve

var.psi variance of the estimate

varIC.cv cross-validated variance of the influence curve

penlikelihood.cv penalized cross-validatedlikelihood

cv.res all cross-validation results for each fold

```
## Not run:
set.seed(123)
N <- 1000
p = 10
Wmat <- matrix(rnorm(N * p), ncol = p)</pre>
beta1 <- 4+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
beta0 <- 2+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
tau <- 2
gcoef <- matrix(c(-1,-1,rep(0,(p)-2)),ncol=1)
Wm <- as.matrix(Wmat)</pre>
g <- 1/(1+exp(Wm%*%gcoef / 3))
A \leftarrow rbinom(N, 1, prob = g)
sigma <- 1
epsilon <-rnorm(N,0,sigma)</pre>
Y <- beta0 + tau * A + epsilon
# ctmleGlmnet must provide user-specified Q
W_tmp <- data.frame(Wm[,1:3])</pre>
```

print.ctmle

```
treated<- W_tmp[which(A==1),]</pre>
untreated<-W_tmp[which(A==0),]</pre>
Y1<-Y[which(A==1)]
Y0<-Y[which(A==0)]
# Initial Q-estimate
beta1hat <- predict(lm(Y1~.,data=treated),newdata=W_tmp)</pre>
beta0hat <- predict(lm(Y0~., data=untreated),newdata=W_tmp)</pre>
Q <- matrix(c(beta0hat,beta1hat),ncol=2)</pre>
W = Wm
glmnet_fit <- cv.glmnet(y = A, x = Wm,</pre>
                         family = 'binomial', nlambda = 40)
start = which(glmnet_fit$lambda==glmnet_fit$lambda.min))
end = length(glmnet_fit$lambda)
lambdas <-glmnet_fit$lambda[start:end]</pre>
ctmle_fit1 <- ctmleGlmnet(Y=Y, A=A,</pre>
                           W=data.frame(W=W),
                           Q = Q, lambdas = lambdas,
                           ctmletype=1, alpha=.995,
                           family="gaussian",
                           gbound=0.025,like_type="loglik" ,
                           fluctuation="logistic",
                           verbose=FALSE,
                           detailed=FALSE, PEN=FALSE,
                           V=5, stopFactor=10<sup>6</sup>)
## End(Not run)
```

print.ctmle

print a ctmle object

Description

print a ctmle object

Usage

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

Arguments

```
x a ctmle object ... other parameter
```

```
## Not run:
N <- 1000
p = 10
```

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```
Wmat <- matrix(rnorm(N * p), ncol = p)</pre>
beta1 <- 4+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
beta0 <- 2+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
tauW <- 2
tau <- 2
gcoef \leftarrow matrix(c(-1,-1,rep(-(3/((p)-2)),(p)-2)),ncol=1)
Wm <- as.matrix(Wmat)</pre>
g <- 1/(1+exp(Wm%*%gcoef))</pre>
A \leftarrow rbinom(N, 1, prob = g)
sigma <- 1
epsilon <-rnorm(N,0,sigma)</pre>
Y <- beta0 + tauW*A + epsilon
# Initial estimate of Q
Q \leftarrow cbind(rep(mean(Y[A == 0]), N), rep(mean(Y[A == 1]), N))
# User-suplied initial estimate
time_greedy <- system.time(</pre>
ctmle_discrete_fit1 <- ctmleDiscrete(Y = Y, A = A, W = data.frame(Wmat), Q = Q,</pre>
                                       preOrder = FALSE)
)
ctmle_summary = summary(ctmle_discrete_fit1)
ctmle_summary
ctmle_discrete_fit1
## End(Not run)
```

Description

print the summary of a ctmle object

Usage

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

Arguments

```
x a summary.ctmle object ... other parameter
```

```
## Not run:
N <- 1000
p = 10
```

summary.ctmle

```
Wmat <- matrix(rnorm(N * p), ncol = p)</pre>
beta1 <- 4+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
beta0 <- 2+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
tauW <- 2
tau <- 2
gcoef \leftarrow matrix(c(-1,-1,rep(-(3/((p)-2)),(p)-2)),ncol=1)
Wm <- as.matrix(Wmat)</pre>
g <- 1/(1+exp(Wm%*%gcoef))</pre>
A \leftarrow rbinom(N, 1, prob = g)
sigma <- 1
epsilon <-rnorm(N,0,sigma)
Y <- beta0 + tauW*A + epsilon
# Initial estimate of Q
Q \leftarrow cbind(rep(mean(Y[A == 0]), N), rep(mean(Y[A == 1]), N))
# User-suplied initial estimate
time_greedy <- system.time(</pre>
ctmle_discrete_fit1 <- ctmleDiscrete(Y = Y, A = A, W = data.frame(Wmat), Q = Q,</pre>
                                       preOrder = FALSE)
)
ctmle_summary = summary(ctmle_discrete_fit1)
ctmle_summary
ctmle_discrete_fit1
## End(Not run)
```

summary.ctmle

Summarise a ctmle object

Description

Summarise a ctmle object

Usage

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

Arguments

```
object a ctmle object ... other parameter
```

```
## Not run:
N <- 1000
p = 10
```

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```
Wmat <- matrix(rnorm(N * p), ncol = p)</pre>
beta1 <- 4+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
beta0 <- 2+2*Wmat[,1]+2*Wmat[,2]+2*Wmat[,5]+2*Wmat[,6]+2*Wmat[,8]</pre>
tauW <- 2
tau <- 2
gcoef \leftarrow matrix(c(-1,-1,rep(-(3/((p)-2)),(p)-2)),ncol=1)
Wm <- as.matrix(Wmat)</pre>
g <- 1/(1+exp(Wm%*%gcoef))</pre>
A \leftarrow rbinom(N, 1, prob = g)
sigma <- 1
epsilon <-rnorm(N,0,sigma)</pre>
Y <- beta0 + tauW*A + epsilon
# Initial estimate of Q
Q \leftarrow cbind(rep(mean(Y[A == 0]), N), rep(mean(Y[A == 1]), N))
# User-suplied initial estimate
time_greedy <- system.time(</pre>
ctmle_discrete_fit1 <- ctmleDiscrete(Y = Y, A = A, W = data.frame(Wmat), Q = Q,</pre>
                                       preOrder = FALSE)
ctmle_summary = summary(ctmle_discrete_fit1)
ctmle_summary
ctmle_discrete_fit1
## End(Not run)
```

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