

# Package ‘causaldrf’

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**Type** Package

**Title** Estimating Causal Dose Response Functions

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**Description** Functions and data to estimate causal dose response functions given continuous, ordinal, or binary treatments. A description of the methods is given in Galagate (2016) <<https://drum.lib.umd.edu/handle/1903/18170>>.

**License** MIT + file LICENSE

**LazyData** TRUE

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add_spl_est	<i>The additive spline estimator</i>
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## Description

This function estimates the ADRF with an additive spline estimator described in Bia et al. (2014).

## Usage

```
add_spl_est(Y,
            treat,
            treat_formula,
            data,
            grid_val,
            knot_num,
            treat_mod,
            link_function,
            ...)
```

## Arguments

Y	is the the name of the outcome variable contained in data.
treat	is the name of the treatment variable contained in data.
treat_formula	an object of class "formula" (or one that can be coerced to that class) that regresses treat on a linear combination of X: a symbolic description of the model to be fitted.
data	is a dataframe containing Y, treat, and X.
grid_val	contains the treatment values to be evaluated.
knot_num	is the number of knots used in outcome model

treat_mod	a description of the error distribution to be used in the model for treatment. Options include: "Normal" for normal model, "LogNormal" for lognormal model, "Sqrt" for square-root transformation to a normal treatment, "Poisson" for Poisson model, "NegBinom" for negative binomial model, "Gamma" for gamma model.
link_function	is either "log", "inverse", or "identity" for the "Gamma" treat_mod.
...	additional arguments to be passed to the outcome regression fitting function.

### Details

This function estimates the ADRF using additive splines in the outcome model described in Bia et al. (2014).

### Value

add\_spl\_est returns an object of class "causaldrf", a list that contains the following components:

param	parameter estimates for a add_spl fit.
t_mod	the result of the treatment model fit.
out_mod	the result of the outcome model fit.
call	the matched call.

### References

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

Bia, Michela, et al. (2014). A Stata package for the application of semiparametric estimators of dose response functions. *Stata Journal* **14.3**, 580-604.

### See Also

[nw\\_est](#), [iw\\_est](#), [hi\\_est](#), [gam\\_est](#), [bart\\_est](#), etc. for other estimates.

[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

### Examples

```
## Example from Schafer (2015).
example_data <- sim_data
add_spl_list <- add_spl_est(Y = Y,
  treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data = example_data,
  grid_val = seq(8, 16, by = 1),
  knot_num = 3,
  treat_mod = "Normal")

sample_index <- sample(1:1000, 100)
plot(example_data$T[sample_index],
```

```

    example_data$Y[sample_index],
    xlab = "T",
    ylab = "Y",
    main = "additive spline estimate")

lines(seq(8, 16, by = 1),
      add_spl_list$param,
      lty = 2,
      lwd = 2,
      col = "blue")
legend('bottomright',
      "additive spline estimate",
      lty=2,
      lwd = 2,
      col = "blue",
      bty='Y', cex=1)

rm(example_data, add_spl_list, sample_index)

## See Vignette for more examples.

```

---

aipwee\_est

*Prediction with a residual bias correction estimator*


---

## Description

This method combines the regression estimator with a residual bias correction for estimating a parametric ADRF.

## Usage

```

aipwee_est(Y,
  treat,
  covar_formula = ~ 1,
  covar_lin_formula = ~ 1,
  covar_sq_formula = ~ 1,
  data,
  e_treat_1 = NULL,
  e_treat_2 = NULL,
  e_treat_3 = NULL,
  e_treat_4 = NULL,
  degree = 1,
  wt = NULL,
  method = "same",
  spline_df = NULL,
  spline_const = 1,
  spline_linear = 1,
  spline_quad = 1)

```

## Arguments

<code>Y</code>	is the the name of the outcome variable contained in data.
<code>treat</code>	is the name of the treatment variable contained in data.
<code>covar_formula</code>	is the formula to describe the covariates needed to estimate the constant term: $\sim X.1 + \dots$ . Can include higher order terms or interactions. i.e. $\sim X.1 + I(X.1^2) + X.1 * X.2 + \dots$ . Don't forget the tilde before listing the covariates.
<code>covar_lin_formula</code>	is the formula to describe the covariates needed to estimate the linear term, $t$ : $\sim X.1 + \dots$ . Can include higher order terms or interactions. i.e. $\sim X.1 + I(X.1^2) + X.1 * X.2 + \dots$ . Don't forget the tilde before listing the covariates.
<code>covar_sq_formula</code>	is the formula to describe the covariates needed to estimate the quadratic term, $t^2$ : $\sim X.1 + \dots$ . Can include higher order terms or interactions. i.e. $\sim X.1 + I(X.1^2) + X.1 * X.2 + \dots$ . Don't forget the tilde before listing the covariates.
<code>data</code>	is a dataframe containing <code>Y</code> , <code>treat</code> , and <code>X</code> .
<code>e_treat_1</code>	a vector, representing the conditional expectation of <code>treat</code> from <code>T_mod</code> .
<code>e_treat_2</code>	a vector, representing the conditional expectation of <code>treat^2</code> from <code>T_mod</code> .
<code>e_treat_3</code>	a vector, representing the conditional expectation of <code>treat^3</code> from <code>T_mod</code> .
<code>e_treat_4</code>	a vector, representing the conditional expectation of <code>treat^4</code> from <code>T_mod</code> .
<code>degree</code>	is 1 for linear and 2 for quadratic outcome model.
<code>wt</code>	is weight used in <code>lsfit</code> for outcome regression. Default is <code>wt = NULL</code> .
<code>method</code>	is "same" if the same set of covariates are used to estimate the constant, linear, and/or quadratic term. If <code>method = "different"</code> , then different sets of covariates can be used to estimate the constant, linear, and/or quadratic term. <code>covar_lin_formula</code> and <code>covar_sq_formula</code> must be specified if <code>method = "different"</code> .
<code>spline_df</code>	degrees of freedom. The default, <code>spline_df = NULL</code> , corresponds to no knots.
<code>spline_const</code>	is the number of spline terms needed to estimate the constant term.
<code>spline_linear</code>	is the number of spline terms needed to estimate the linear term.
<code>spline_quad</code>	is the number of spline terms needed to estimate the quadratic term.

## Details

This estimator bears a strong resemblance to general regression estimators in the survey literature, part of a more general class of calibration estimators (Deville and Sarndal, 1992). It is doubly robust, which means that it is consistent if either of the models is true (Scharfstein, Rotnitzky and Robins 1999). If the Y-model is correct, then the first term in the previous equation is unbiased for  $\xi$  and the second term has mean zero even if the T-model is wrong. If the Y-model is incorrect, the first term is biased, but the second term gives a consistent estimate of (minus one times) the bias from the Y-model if the T-model is correct.

This function is a doubly-robust estimator that fits an outcome regression model with a bias correction term. For details see Schafer and Galagate (2015).

**Value**

aipwee\_est returns an object of class "causaldrf\_lsfit", a list that contains the following components:

param	parameter estimates for a add_spl fit.
t_mod	the result of the treatment model fit.
out_mod	the result of the outcome model fit.
call	the matched call.

**References**

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

Schafer, Joseph L, Kang, Joseph (2008). Average causal effects from nonrandomized studies: a practical guide and simulated example. *Psychological methods*, **13.4**, 279.

Robins, James M and Rotnitzky, Andrea (1995). Semiparametric efficiency in multivariate regression models with missing data *Journal of the American Statistical Association*, **90.429**, 122–129.

Scharfstein, Daniel O and Rotnitzky, Andrea and Robins, James M (1999). Adjusting for non-ignorable drop-out using semiparametric nonresponse models *Journal of the American Statistical Association*, **94.448**, 1096–1120.

Deville, Jean-Claude and Sarndal, Carl-Erik (1992). Calibration estimators in survey sampling *Journal of the American Statistical Association*, **87.418**, 376–380.

**See Also**

[iptw\\_est](#), [ismw\\_est](#), [reg\\_est](#), [wtrg\\_est](#), `##'` etc. for other estimates.

[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

**Examples**

```
## Example from Schafer (2015).

example_data <- sim_data

t_mod_list <- t_mod(treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data = example_data,
  treat_mod = "Normal")

cond_exp_data <- t_mod_list$T_data
full_data <- cbind(example_data, cond_exp_data)

aipwee_list <- aipwee_est(Y = Y,
  treat = T,
  covar_formula = ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  covar_lin_formula = ~ 1,
  covar_sq_formula = ~ 1,
```

```
data = example_data,
e_treat_1 = full_data$est_treat,
e_treat_2 = full_data$est_treat_sq,
e_treat_3 = full_data$est_treat_cube,
e_treat_4 = full_data$est_treat_quartic,
degree = 1,
wt = NULL,
method = "same",
spline_df = NULL,
spline_const = 1,
spline_linear = 1,
spline_quad = 1)

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
      example_data$Y[sample_index],
      xlab = "T",
      ylab = "Y",
      main = "aipwee estimate")

abline(aipwee_list$param[1],
       aipwee_list$param[2],
       lty = 2,
       lwd = 2,
       col = "blue")

legend('bottomright',
       "aipwee estimate",
       lty = 2,
       lwd = 2,
       col = "blue",
       bty='Y',
       cex=1)

rm(example_data, t_mod_list, cond_exp_data, full_data, aipwee_list, sample_index)
```

---

bart\_est

*The BART estimator*

---

## Description

This function estimates the ADRF using Bayesian additive regression trees (BART).

## Usage

```
bart_est(Y,
        treat,
        outcome_formula,
        data,
```

```
grid_val,
...)
```

### Arguments

Y	is the the name of the outcome variable contained in data.
treat	is the name of the treatment variable contained in data.
outcome_formula	is the formula used for fitting the outcome surface. <code>gps</code> is one of the independent variables to use in the <code>outcome_formula</code> . ie. $Y \sim \text{treat} + X.1 + X.2 + \dots$ or a variation of this.
data	is a dataframe containing Y, treat, and X.
grid_val	contains the treatment values to be evaluated.
...	additional arguments to be passed to the <code>bart()</code> outcome function.

### Details

BART is a prediction model that is applicable to many settings, one of which is causal inference problems. It is a sum of trees fit, but the influence of each tree is held back by a regularization prior so that each tree only contributes a small amount to the overall fit. Priors are put on the parameters to avoid overfitting the data and so that no single tree has a significant influence on the model fit. For more details see Chipman (2010).

BART does not require fitting a treatment model. Instead, it fits a response surface to the whole dataset and if the response surface is correctly specified, then the causal effect estimate is unbiased. Although most of the focus on BART is for the binary treatment setting, Hill (2011) also mentions an extension to the continuous or multidose treatment setting. When using BART in this continuous treatment setting, Hill (2011) compares the outcomes of units with treatment level  $T_i = t$  to their outcomes had  $T_i = 0$ . This method infers the treatment effect of units had they not received treatment compared to their actual observed treatment. The comparison is between  $Y_i(0)|(I = 1, T_i = t)$  and  $Y_i(t)|(I = 1, T_i = t)$  where  $I = 1$  means that the unit is part of the treatment group. The causal effect is comparing the predicted outcome of units that received treatment with what their predicted outcome would have been had they received zero treatment.

This method performs well in simulation studies. One drawback from BART is the amount of computing time needed.

### Value

`bart_est` returns an object of class "causaldrf\_simple", a list that contains the following components:

param	parameter estimates for a bart fit.
out_mod	the result of the bart fit.
call	the matched call.

## References

- Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.
- Hill, Jennifer L. (2011). Bayesian nonparametric modeling for causal inference. *Journal of Computational and Graphical Statistics* **20.1** (2011).
- Chipman, Hugh A and George, Edward I and McCulloch, Robert E and others (2010). BART: Bayesian additive regression trees. *The Annals of Applied Statistics* **4.1**, 266–298.

## See Also

[nw\\_est](#), [iw\\_est](#), [hi\\_est](#), [gam\\_est](#), [add\\_spl\\_est](#), etc. for other estimates.  
[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

## Examples

```
## Example from Schafer (2015). bart takes a few minutes to run (depending on computer).

example_data <- sim_data

# This estimate takes a long time to run...
bart_list <- bart_est(Y = Y,
  treat = T,
  outcome_formula = Y ~ T + B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data = example_data,
  grid_val = seq(8, 16, by = 1))

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
  example_data$Y[sample_index],
  xlab = "T",
  ylab = "Y",
  main = "bart estimate")

lines(seq(8, 16, by = 1),
  bart_list$param,
  lty = 2,
  lwd = 2,
  col = "blue")

legend('bottomright',
  "bart estimate",
  lty=2,
  lwd = 2,
  col = "blue",
  bty='Y',
  cex=1)

rm(example_data, bart_list, sample_index)
```

gam\_est

*The GAM estimator***Description**

This estimates the ADRF using a method similar to that described in Hirano and Imbens (2004), but with spline basis terms in the outcome model.

**Usage**

```
gam_est(Y,
        treat,
        treat_formula,
        data,
        grid_val,
        treat_mod,
        link_function,
        ...)
```

**Arguments**

Y	is the the name of the outcome variable contained in data.
treat	is the name of the treatment variable contained in data.
treat_formula	an object of class "formula" (or one that can be coerced to that class) that regresses treat on a linear combination of X: a symbolic description of the model to be fitted.
data	is a dataframe containing Y and treat and X.
grid_val	contains the treatment values to be evaluated.
treat_mod	a description of the error distribution to be used in the model for treatment. Options include: "Normal" for normal model, "LogNormal" for lognormal model, "Sqrt" for square-root transformation to a normal treatment, "Poisson" for Poisson model, "NegBinom" for negative binomial model, "Gamma" for gamma model.
link_function	is either "log", "inverse", or "identity" for the "Gamma" treat_mod.
...	additional arguments to be passed to the gam() outcome function.

**Details**

This function estimates the ADRF similarly to the method described by Hirano and Imbens (2004), but with a generalized additive model in the outcome model.

**Value**

gam\_est returns an object of class "causaldrf", a list that contains the following components:

param	parameter estimates for a gam fit.
t_mod	the result of the treatment model fit.
out_mod	the result of the outcome model fit.
call	the matched call.

**References**

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

Hirano, Keisuke, Imbens, Guido W (2004). The propensity score with continuous treatments. *Applied Bayesian modeling and causal inference from incomplete-data perspectives*.

Flores, Carlos A and Flores-Lagunes, Alfonso and Gonzalez, Arturo and Neumann, Todd C (2012). Estimating the effects of length of exposure to instruction in a training program: the case of job corps. *Review of Economics and Statistics*. **94.1**, 153-171

**See Also**

[nw\\_est](#), [iw\\_est](#), [hi\\_est](#), [gam\\_est](#), [add\\_spl\\_est](#), [bart\\_est](#), etc. for other estimates.

[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

**Examples**

```
## Example from Schafer (2015).

example_data <- sim_data

gam_list <- gam_est(Y = Y,
  treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data = example_data,
  grid_val = seq(8, 16, by = 1),
  treat_mod = "Normal")

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
  example_data$Y[sample_index],
  xlab = "T",
  ylab = "Y",
  main = "gam estimate")

lines(seq(8, 16, by = 1),
  gam_list$param,
  lty = 2,
  lwd = 2,
  col = "blue")
```

```

legend('bottomright',
      "gam estimate",
      lty=2,
      lwd = 2,
      col = "blue",
      bty='Y',
      cex=1)

rm(example_data, gam_list, sample_index)

```

---

get\_ci *This calculates an upper and lower bound from bootstrap matrix*

---

### Description

This function takes a matrix containing the bootstrapped coefficients from a parametric ADRF estimator and returns upper and lower 95 percent confidence lines.

### Usage

```

get_ci(grid_val,
      coef_mat,
      degree)

```

### Arguments

grid\_val is the vector of grid values on treat axis  
coef\_mat contains the bootstrapped parameter estimates.  
degree is 1 for linear and 2 for quadratic outcome model

### Value

get\_ci returns upper and lower 95 percent confidence lines.

---

hi\_est *The Hirano and Imbens estimator*

---

### Description

This function estimates the GPS function and estimates the ADRF. The GPS score is based on different treatment models. The treatment is linearly related to Xs.

**Usage**

```
hi_est(Y,
      treat,
      treat_formula,
      outcome_formula,
      data,
      grid_val,
      treat_mod,
      link_function,
      ...)
```

**Arguments**

Y	is the the name of the outcome variable contained in data.
treat	is the name of the treatment variable contained in data.
treat_formula	an object of class "formula" (or one that can be coerced to that class) that regresses treat on a linear combination of X: a symbolic description of the model to be fitted.
outcome_formula	is the formula used for fitting the outcome surface. gps is one of the independent variables to use in the outcome_formula. ie. $Y \sim \text{treat} + I(\text{treat}^2) + \text{gps} + I(\text{gps}^2) + \text{treat} * \text{gps}$ or a variation of this. Use gps as the name of the variable representing the gps in outcome_formula.
data	is a dataframe containing Y, treat, and X.
grid_val	contains the treatment values to be evaluated.
treat_mod	a description of the error distribution to be used in the model for treatment. Options include: "Normal" for normal model, "LogNormal" for lognormal model, "Sqrt" for square-root transformation to a normal treatment, "Poisson" for Poisson model, "NegBinom" for negative binomial model, "Gamma" for gamma model, "Binomial" for binomial model.
link_function	For treat_mod = "Gamma" (fitted using glm) alternatives are "log" or "inverse". For treat_mod = "Binomial" (fitted using glm) alternatives are "logit", "probit", "cauchit", "log" and "cloglog".
...	additional arguments to be passed to the outcome lm() function.

**Details**

Hirano (2004) (HI) introduced this imputation-type method that includes a GPS component. The idea is to fit a parametric observable (outcome) model, which includes the estimated GPS as a covariate, to impute missing potential outcomes.

The method requires several steps. First, a model is used to relate treatment to the recorded covariates. For example,  $T_i | \mathbf{X}_i \sim \mathcal{N}(\mathbf{X}_i^T \boldsymbol{\beta}, \sigma^2)$  and then estimate the  $\boldsymbol{\beta}$  parameters. Next, the GPS for each unit is estimated

$$\hat{R}_i(t) = \frac{1}{\sqrt{2\pi\hat{\sigma}^2}} e^{-\frac{(t-x_i^T\hat{\beta})^2}{2\hat{\sigma}^2}}$$

These GPS estimates are used in the outcome or observable model. The outcome is modeled as a function of  $T_i$  and  $\hat{R}_i$  parametrically. For example,

$$E[Y_i|T_i, R_i] = \alpha_0 + \alpha_1 T_i + \alpha_2 T_i^2 + \alpha_3 \hat{R}_i + \alpha_4 \hat{R}_i^2 + \alpha_5 \hat{R}_i \cdot T_i$$

After collecting the estimated parameters in the outcome and treatment models, plug-in the treatment values into the model to estimate the missing potential outcomes of each individual at that treatment level. For example, if we plug in  $T_i = t$  into the estimated models, then each unit will have a potential outcome estimated at treatment level  $T_i = t$ .

$$\hat{Y}_i(t) = \hat{\alpha}_0 + \hat{\alpha}_1 t + \hat{\alpha}_2 t^2 + \hat{\alpha}_3 \hat{R}_i(t) + \hat{\alpha}_4 \hat{R}_i^2(t) + \hat{\alpha}_5 \hat{R}_i(t) \cdot t$$

The next step is to aggregate these estimated potential outcomes to get an average treatment effect at dose level  $T_i = t$ . The mean outcome at dose-level  $T_i = t$  is given by:

$$\hat{\mu}(t) = \frac{1}{N} \sum_i^N \hat{\alpha}_0 + \hat{\alpha}_1 t + \hat{\alpha}_2 t^2 + \hat{\alpha}_3 \hat{R}_i(t) + \hat{\alpha}_4 \hat{R}_i^2(t) + \hat{\alpha}_5 \hat{R}_i(t) \cdot t$$

Different treatment levels are plugged into the previous equation to estimate the missing potential outcomes. If many  $t$  values are evaluated, then it is possible to trace out an ADRF.

## Value

hi\_est returns an object of class "causaldrf", a list that contains the following components:

param	parameter estimates for a hi fit.
t_mod	the result of the treatment model fit.
out_mod	the result of the outcome model fit.
call	the matched call.

## References

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

Hirano, Keisuke, Imbens, Guido W (2004). The propensity score with continuous treatments. *Applied Bayesian modeling and causal inference from incomplete-data perspectives*.

## See Also

[nw\\_est](#), [iw\\_est](#), [hi\\_est](#), [gam\\_est](#), [add\\_spl\\_est](#), [bart\\_est](#), etc. for other estimates.

[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

**Examples**

```

## Example from Schafer (2015).

example_data <- sim_data

hi_list <- hi_est(Y = Y,
  treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  outcome_formula = Y ~ T + I(T^2) + gps + I(gps^2) + T * gps,
  data = example_data,
  grid_val = seq(8, 16, by = 1),
  treat_mod = "Normal")

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
  example_data$Y[sample_index],
  xlab = "T",
  ylab = "Y",
  main = "hi estimate")

lines(seq(8, 16, by = 1),
  hi_list$param,
  lty = 2,
  lwd = 2,
  col = "blue")

legend('bottomright',
  "hi estimate",
  lty=2,
  lwd = 2,
  col = "blue",
  bty='Y',
  cex=1)

rm(example_data, hi_list, sample_index)

## Example from van der Wal, Willem M., and Ronald B. Geskus. (2011)
#Simulate data with continuous confounder and outcome, binomial exposure.
#Marginal causal effect of exposure on outcome: 10.
n <- 1000
simdat <- data.frame(l = rnorm(n, 10, 5))
a.lin <- simdat$l - 10
pa <- exp(a.lin)/(1 + exp(a.lin))
simdat$a <- rbinom(n, 1, prob = pa)
simdat$y <- 10*simdat$a + 0.5*simdat$l + rnorm(n, -10, 5)
simdat[1:5,]
temp_hi <- hi_est(Y = y,
  treat = a,
  treat_formula = a ~ l,
  outcome_formula = y ~ gps,

```

```
data = simdat,  
grid_val = c(0, 1),  
treat_mod = "Binomial",  
link_function = "logit")  
  
temp_hi[[1]] # estimated coefficients
```

---

hi\_sim\_data

*Simulated data from Hirano and Imbens (2004)*

---

### Description

Simulated data used in the paper "The propensity score with continuous treatments."

### Usage

```
data(hi_sim_data)
```

### Format

A data frame with 1000 rows and 6 variables:

### Details

A dataset containing hi\_sim\_data.

### Source

use the hi\_sample function

### References

Hirano, Keisuke, and Guido W. Imbens. "The propensity score with continuous treatments." *Applied Bayesian modeling and causal inference from incomplete-data perspectives* (2004): 73-84.

Moodie, Erica EM, and David A. Stephens. "Estimation of dose-response functions for longitudinal data using the generalised propensity score." *Statistical methods in medical research* **21.2** (2012): 149-166.

### Examples

```
## Example from Hirano and Imbens (2004).  
data(hi_sim_data)  
head(hi_sim_data)
```

---

iptw\_est                      *The inverse probability of treatment weighting (iptw) estimator*

---

## Description

The iptw method or importance weighting method estimates the ADRF by weighting the data with stabilized or non-stabilized weights.

## Usage

```
iptw_est(Y,
        treat,
        treat_formula,
        numerator_formula,
        data,
        degree,
        treat_mod,
        link_function,
        ...)
```

## Arguments

Y	is the the name of the outcome variable contained in data.
treat	is the name of the treatment variable contained in data.
treat_formula	an object of class "formula" (or one that can be coerced to that class) that regresses treat on a linear combination of X: a symbolic description of the model to be fitted.
numerator_formula	an object of class "formula" (or one that can be coerced to that class) that regresses treat on a linear combination of X: a symbolic description of the model to be fitted. i.e. $treat \sim 1$ .
data	is a dataframe containing Y, treat, and X.
degree	is 1 for linear and 2 for quadratic outcome model.
treat_mod	a description of the error distribution to be used in the model for treatment. Options include: "Normal" for normal model, "LogNormal" for lognormal model, "Sqrt" for square-root transformation to a normal treatment, "Poisson" for Poisson model, "NegBinom" for negative binomial model, "Gamma" for gamma model, "Binomial" for binomial model, "Ordinal" for ordinal model, "Multinomial" for multinomial model.
link_function	specifies the link function between the variables in numerator or denominator and exposure, respectively. For <code>treat_mod = "Gamma"</code> (fitted using glm) alternatives are "log" or "inverse". For <code>treat_mod = "Binomial"</code> (fitted using glm) alternatives are "logit", "probit", "cauchit", "log" and "cloglog". For <code>treat_mod = "Multinomial"</code> this argument is ignored, and multinomial logistic regression models are always used (fitted using multinom). For <code>treat_mod = "Ordinal"</code> (fitted using polr) alternatives are "logit", "probit", "cauchit", and "cloglog".

... additional arguments to be passed to the low level treatment regression fitting functions.

### Details

This method uses inverse probability of treatment weighting to adjust for possible biases. For more details see Schafer and Galagate (2015) and Robins, Hernan, and Brumback (2000).

### Value

iptw\_est returns an object of class "causaldrf", a list that contains the following components:

param	parameter estimates for a iptw fit.
t_mod	the result of the treatment model fit.
num_mod	the result of the numerator model fit.
weights	the estimated weights.
weight_data	the weights.
out_mod	the outcome model.
call	the matched call.

### References

- Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.
- van der Wal, Willem M., and Ronald B. Geskus. "IPW: an R package for inverse probability weighting." *Journal of Statistical Software* **43.13** (2011): 1-23.
- Robins, James M and Hernan, Miguel Angel and Brumback, Babette. Marginal structural models and causal inference in epidemiology. *Epidemiology* **11.5** (2000): 550–560.
- Zhu, Yeying and Coffman, Donna L and Ghosh, Debashis. A Boosting Algorithm for Estimating Generalized Propensity Scores with Continuous Treatments. *Journal of Causal Inference* **3.1** (2015): 25–40.

### See Also

[iptw\\_est](#), [ismw\\_est](#), [reg\\_est](#), [aipwee\\_est](#), [wtrg\\_est](#), etc. for other estimates.  
[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

### Examples

```
## Example from Schafer (2015).

example_data <- sim_data

iptw_list <- iptw_est(Y = Y,
  treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  numerator_formula = T ~ 1,
  data = example_data,
```

```

        degree = 1,
        treat_mod = "Normal")

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
      example_data$Y[sample_index],
      xlab = "T",
      ylab = "Y",
      main = "iptw estimate")

abline(iptw_list$param[1],
       iptw_list$param[2],
       lty=2,
       lwd = 2,
       col = "blue")

legend('bottomright',
      "iptw estimate",
      lty=2,
      lwd = 2,
      col = "blue",
      bty='Y',
      cex=1)

rm(example_data, iptw_list, sample_index)

## Example from van der Wal, Willem M., and Ronald B. Geskus. (2011)
## Simulate data with continuous confounder and outcome, binomial exposure.
## Marginal causal effect of exposure on outcome: 10.
n <- 1000
simdat <- data.frame(l = rnorm(n, 10, 5))
a.lin <- simdat$l - 10
pa <- exp(a.lin)/(1 + exp(a.lin))
simdat$a <- rbinom(n, 1, prob = pa)
simdat$y <- 10*simdat$a + 0.5*simdat$l + rnorm(n, -10, 5)
simdat[1:5,]
temp_iptw <- iptw_est(Y = y,
                    treat = a,
                    treat_formula = a ~ 1,
                    numerator_formula = a ~ 1,
                    data = simdat,
                    degree = 1,
                    treat_mod = "Binomial",
                    link_function = "logit")

temp_iptw[[1]] # estimated coefficients

```

**Description**

This method estimates the ADRF by using weighting matrices instead of scalars. The weight matrices require conditional expectations of the treatment and higher order conditional expectations. It uses outputs from the `t_mod` function.

**Usage**

```
ismw_est(Y,
         treat,
         data,
         e_treat_1,
         e_treat_2,
         e_treat_3,
         e_treat_4,
         degree )
```

**Arguments**

<code>Y</code>	is the the name of the outcome variable contained in <code>data</code> .
<code>treat</code>	is the name of the treatment variable contained in <code>data</code> .
<code>data</code>	is a dataframe containing <code>Y</code> , <code>treat</code> , and <code>X</code> .
<code>e_treat_1</code>	a vector, representing the conditional expectation of <code>treat</code> from <code>t_mod</code> .
<code>e_treat_2</code>	a vector, representing the conditional expectation of <code>treat^2</code> from <code>t_mod</code> .
<code>e_treat_3</code>	a vector, representing the conditional expectation of <code>treat^3</code> from <code>t_mod</code> .
<code>e_treat_4</code>	a vector, representing the conditional expectation of <code>treat^4</code> from <code>t_mod</code> .
<code>degree</code>	is 1 for linear and 2 for quadratic outcome model.

**Details**

This function estimates the ADRF requires estimated moments and uses the outputs of the `t_mod` function as inputs. For more details, see Schafer and Galagate (2015).

**Value**

`ismw_est` returns an object of class "causaldrf\_simple", a list that contains the following components:

<code>param</code>	the estimated parameters.
<code>call</code>	the matched call.

**References**

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

**See Also**

[iptw\\_est](#), [ismw\\_est](#), [reg\\_est](#), [aipwee\\_est](#), [wtrg\\_est](#), etc. for other estimates.  
[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

**Examples**

```
## Example from Schafer (2015).

example_data <- sim_data

t_mod_list <- t_mod(treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data = example_data,
  treat_mod = "Normal")

cond_exp_data <- t_mod_list$T_data

full_data <- cbind(example_data, cond_exp_data)

ismw_list <- ismw_est(Y = Y,
  treat = T,
  data = full_data,
  e_treat_1 = full_data$est_treat,
  e_treat_2 = full_data$est_treat_sq,
  e_treat_3 = full_data$est_treat_cube,
  e_treat_4 = full_data$est_treat_quartic,
  degree = 1)

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
  example_data$Y[sample_index],
  xlab = "T",
  ylab = "Y",
  main = "ismw estimate")

abline(ismw_list$param[1],
  ismw_list$param[2],
  lty=2,
  lwd = 2,
  col = "blue")

legend('bottomright',
  "ismw estimate",
  lty=2,
  lwd = 2,
  col = "blue",
  bty='Y',
  cex=1)

rm(example_data, t_mod_list, cond_exp_data, full_data, ismw_list, sample_index)
```

---

iw_est	<i>The inverse weighting estimator (nonparametric method)</i>
--------	---

---

### Description

This is a nonparametric method that estimates the ADRF by using a local linear regression of  $Y$  on  $treat$  with weighted kernel function. For details, see Flores et. al. (2012).

### Usage

```
iw_est(Y,
      treat,
      treat_formula,
      data,
      grid_val,
      bandw,
      treat_mod,
      link_function,
      ...)
```

### Arguments

$Y$	is the the name of the outcome variable contained in data.
$treat$	is the name of the treatment variable contained in data.
$treat\_formula$	an object of class "formula" (or one that can be coerced to that class) that regresses $treat$ on a linear combination of $X$ : a symbolic description of the model to be fitted.
$data$	is a dataframe containing $Y$ , $treat$ , and $X$ .
$grid\_val$	contains the treatment values to be evaluated.
$bandw$	is the bandwidth. Default is 1.
$treat\_mod$	a description of the error distribution to be used in the model for treatment. Options include: "Normal" for normal model, "LogNormal" for lognormal model, "Sqrt" for square-root transformation to a normal treatment, "Poisson" for Poisson model, "NegBinom" for negative binomial model, "Gamma" for gamma model.
$link\_function$	is either "log", "inverse", or "identity" for the "Gamma" $treat\_mod$ .
...	additional arguments to be passed to the treatment regression function.

### Details

The ADRF is estimated by

$$(D_0(t)S_2(t) - D_1(t)S_1(t))/(S_0(t)S_2(t) - S_1^2(t))$$

where

$$D_j(t) = \sum_{i=1}^N \tilde{K}_{h,X}(T_i - t)(T_i - t)^j Y_i$$

and  $S_j(t) = \sum_{i=1}^N \tilde{K}_{h,X}(T_i - t)(T_i - t)^j$   $\tilde{K}_{h,X}(t) = K_h(t)/\hat{R}_i(t)$  which is a local linear regression. More details are given in Flores (2012).

## Value

`iw_est` returns an object of class "causaldrf", a list that contains the following components:

<code>param</code>	parameter estimates for a iw fit.
<code>t_mod</code>	the result of the treatment model fit.
<code>call</code>	the matched call.

## References

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

Flores, Carlos A., et al. "Estimating the effects of length of exposure to instruction in a training program: the case of job corps." *Review of Economics and Statistics* **94.1** (2012): 153-171.

## See Also

[nw\\_est](#), [iw\\_est](#), [hi\\_est](#), [gam\\_est](#), [add\\_spl\\_est](#), [bart\\_est](#), etc. for other estimates.

## Examples

```
## Example from Schafer (2015).

example_data <- sim_data

iw_list <- iw_est(Y = Y,
                 treat = T,
                 treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
                 data = example_data,
                 grid_val = seq(8, 16, by = 1),
                 bandwidth = bw.SJ(example_data$T),
                 treat_mod = "Normal")

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
     example_data$Y[sample_index],
     xlab = "T",
     ylab = "Y",
     main = "iw estimate")

lines(seq(8, 16, by = 1),
      iw_list$param,
      lty = 2,
```

```

      lwd = 2,
      col = "blue")

legend('bottomright',
      "iw estimate",
      lty=2,
      lwd = 2,
      col = "blue",
      bty='Y',
      cex=1)

rm(example_data, iw_list, sample_index)

## Example from Imai & van Dyk (2004).

data("nmes_data")
head(nmes_data)
# look at only people with medical expenditures greater than 0
nmes_nonzero <- nmes_data[which(nmes_data$TOTALEXP > 0), ]

iw_list <- iw_est(Y = TOTALEXP,
  treat = packyears,
  treat_formula = packyears ~ LASTAGE + I(LASTAGE^2) +
    AGESMOKE + I(AGESMOKE^2) + MALE + RACE3 + beltuse +
    educate + marital + SREGION + POVSTALB,
  data = nmes_nonzero,
  grid_val = seq(5, 100, by = 5),
  bandw = bw.SJ(nmes_nonzero$packyears),
  treat_mod = "LogNormal")

set.seed(307)
sample_index <- sample(1:nrow(nmes_nonzero), 1000)

plot(nmes_nonzero$packyears[sample_index],
  nmes_nonzero$TOTALEXP[sample_index],
  xlab = "packyears",
  ylab = "TOTALEXP",
  main = "iw estimate",
  ylim = c(0, 10000),
  xlim = c(0, 100))

lines(seq(5, 100, by = 5),
  iw_list$param,
  lty = 2,
  lwd = 2,
  col = "blue")

legend('topright',
  "iw estimate",
  lty=2,
  lwd = 2,
  col = "blue",

```

```
      bty='Y',  
      cex = 1)  
abline(0, 0)
```

---

nmes_data	<i>Data set containing data from the National Medical Expenditure Survey (NMES)</i>
-----------	---

---

### Description

Data set from the NMES. with 9708 observations and 12 variables.

### Usage

```
data(nmes_data)
```

### Format

A dataset containing 9708 observations and 12 variables.

### References

Imai, K., & van Dyk, D.A. (2004). Causal Inference With General Treatment Regimes: Generalizing the Propensity Score. *Journal of the American Statistical Association*, **99**(467).

National Center for Health Services Research and Health Care Technology Assessment. NATIONAL MEDICAL EXPENDITURE SURVEY, 1987: INSTITUTIONAL POPULATION COMPONENT. Rockville, MD: Westat, Inc. [producer], 1987. Ann Arbor, MI: Inter-university Consortium for Political and Social Research [distributor], 1990. doi:10.3886/ICPSR09280.v1

Bryer, Jason M. "TriMatch: An R Package for Propensity Score Matching of Non-binary Treatments." The R User Conference, useR! 2013 July 10-12 2013 University of Castilla-La Mancha, Albacete, Spain. Vol. 10. No. 30. 2013.

### Examples

```
data(nmes_data)  
head(nmes_data)
```

---

 nw\_est

*The Nadaraya-Watson modified estimator*


---

### Description

This is a kernel based regression method that uses a kernel as a weighting function to estimate the ADRF. The normal kernel is weighted by the inverse of the estimated GPS. See Flores et al. (2012) for more details.

### Usage

```
nw_est(Y,
       treat,
       treat_formula,
       data,
       grid_val,
       bandw,
       treat_mod,
       link_function,
       ...)
```

### Arguments

Y	is the the name of the outcome variable contained in data.
treat	is the name of the treatment variable contained in data.
treat_formula	an object of class "formula" (or one that can be coerced to that class) that regresses treat on a linear combination of X: a symbolic description of the model to be fitted.
data	is a dataframe containing Y and treat and X.
grid_val	contains the treatment values to be evaluated.
bandw	is the bandwidth. Default is 1.
treat_mod	a description of the error distribution to be used in the model for treatment. Options include: "Normal" for normal model, "LogNormal" for lognormal model, "Sqrt" for square-root transformation to a normal treatment, "Poisson" for Poisson model, "NegBinom" for negative binomial model, "Gamma" for gamma model.
link_function	is either "log", "inverse", or "identity" for the "Gamma" treat_mod.
...	additional arguments to be passed to the treatment regression function.

### Details

This method is a version of the Nadarya-Watson estimator Nadaraya (1964) which is a local constant regression but weighted by the inverse of the estimated GPS.

**Value**

nw\_est returns an object of class "causaldrf", a list that contains the following components:

param	parameter estimates for a nw fit.
t_mod	the result of the treatment model fit.
call	the matched call.

**References**

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

Flores, Carlos A., et al. "Estimating the effects of length of exposure to instruction in a training program: the case of job corps." *Review of Economics and Statistics* **94.1** (2012): 153-171.

Nadaraya, Elizbar A. "On estimating regression." *Theory of Probability and Its Applications* **9.1** (1964): 141–142.

**See Also**

[nw\\_est](#), [iw\\_est](#), [hi\\_est](#), [gam\\_est](#), [add\\_spl\\_est](#), [bart\\_est](#), etc. for other estimates.

[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

**Examples**

```
## Example from Schafer (2015).

example_data <- sim_data

nw_list <- nw_est(Y = Y,
  treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data = example_data,
  grid_val = seq(8, 16, by = 1),
  bandw = bw.SJ(example_data$T),
  treat_mod = "Normal")

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
  example_data$Y[sample_index],
  xlab = "T",
  ylab = "Y",
  main = "nw estimate")

lines(seq(8, 16, by = 1),
  nw_list$param,
  lty = 2,
  lwd = 2,
  col = "blue")

legend('bottomright',
```

```

      "nw estimate",
      lty=2,
      lwd = 2,
      col = "blue",
      bty='Y',
      cex=1)

rm(example_data, nw_list, sample_index)

```

---

 overlap\_fun

*This function creates an overlapping dataset*


---

### Description

This function ensures that the units overlap according to the estimated gps values. The overlapping dataset depends on the number of classes `n_class` to subclassify on.

### Usage

```

overlap_fun(Y,
            treat,
            treat_formula,
            data_set,
            n_class,
            treat_mod,
            link_function,
            ...)

```

### Arguments

<code>Y</code>	is the the name of the outcome variable contained in data.
<code>treat</code>	is the name of the treatment variable contained in data.
<code>treat_formula</code>	an object of class "formula" (or one that can be coerced to that class) that regresses <code>treat</code> on a linear combination of <code>X</code> : a symbolic description of the model to be fitted.
<code>data_set</code>	is a dataframe containing <code>Y</code> , <code>treat</code> , and <code>X</code> .
<code>n_class</code>	is the number of classes to split gps into.
<code>treat_mod</code>	a description of the error distribution to be used in the model for treatment. Options include: "Normal" for normal model, "LogNormal" for lognormal model, "Sqrt" for square-root transformation to a normal treatment, "Poisson" for Poisson model, "NegBinom" for negative binomial model, "Gamma" for gamma model.
<code>link_function</code>	is either "log", "inverse", or "identity" for the "Gamma" <code>treat_mod</code> .
<code>...</code>	additional arguments to be passed to the treatment regression function

**Value**

overlap\_fun returns a list containing the following elements:

overlap\_dataset            dataframe containing overlapping data.  
 median\_vec                a vector containing median values.  
 overlap\_treat\_result      the resulting treatment fit.

**References**

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

Bia, Michela, et al. "A Stata package for the application of semiparametric estimators of dose response functions." *Stata Journal* **14.3** (2014): 580-604.

**See Also**

[iptw\\_est](#), [ismw\\_est](#), [reg\\_est](#), [aipwee\\_est](#), [wtrg\\_est](#), etc. for other estimates.  
[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

**Examples**

```
## Example from Schafer (2015).

example_data <- sim_data

overlap_list <- overlap_fun(Y = Y,
  treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data_set = example_data,
  n_class = 3,
  treat_mod = "Normal")

overlapped_data <- overlap_list$overlap_dataset
summary(overlapped_data)

rm(example_data, overlap_list, overlapped_data)
```

---

 prop\_spline\_est

*The propensity-spline prediction estimator*


---

**Description**

This method estimates the linear or quadratic parameters of the ADRF by estimating a least-squares fit on the basis functions which are composed of combinations of the covariates, propensity spline basis, and treatment values.

**Usage**

```
prop_spline_est(Y,
               treat,
               covar_formula = ~ 1,
               covar_lin_formula = ~ 1,
               covar_sq_formula = ~ 1,
               data,
               e_treat_1 = NULL,
               degree = 1,
               wt = NULL,
               method = "same",
               spline_df = NULL,
               spline_const = 1,
               spline_linear = 1,
               spline_quad = 1)
```

**Arguments**

**Y** is the the name of the outcome variable contained in data.

**treat** is the name of the treatment variable contained in data.

**covar\_formula** is the formula to describe the covariates needed to estimate the constant term:  $\sim X.1 + \dots$ . Can include higher order terms or interactions. i.e.  $\sim X.1 + I(X.1^2) + X.1 * X.2 + \dots$ . Don't forget the tilde before listing the covariates.

**covar\_lin\_formula** is the formula to describe the covariates needed to estimate the linear term,  $t$ :  $\sim X.1 + \dots$ . Can include higher order terms or interactions. i.e.  $\sim X.1 + I(X.1^2) + X.1 * X.2 + \dots$ . Don't forget the tilde before listing the covariates.

**covar\_sq\_formula** is the formula to describe the covariates needed to estimate the quadratic term,  $t^2$ :  $\sim X.1 + \dots$ . Can include higher order terms or interactions. i.e.  $\sim X.1 + I(X.1^2) + X.1 * X.2 + \dots$ . Don't forget the tilde before listing the covariates.

**data** is a dataframe containing Y, treat, and X.

**e\_treat\_1** a vector, representing the conditional expectation of treat from T\_mod. Or, plug in gps estimates here to create splines from the gps values.

**degree** is 1 for linear and 2 for quadratic outcome model.

**wt** is weight used in lsfit for outcome regression. Default is wt = NULL.

**method** is "same" if the same set of covariates are used to estimate the constant, linear, and/or quadratic term with no spline terms. If method = "different", then different sets of covariates can be used to estimate the constant, linear, and/or quadratic term. To use spline terms, it is necessary to set method = "different". covar\_lin\_formula and covar\_sq\_formula must be specified if method = "different".

**spline\_df** degrees of freedom. The default, spline\_df = NULL, corresponds to no knots.

spline\_const is the number of spline terms to include when estimating the constant term.  
 spline\_linear is the number of spline terms to include when estimating the linear term.  
 spline\_quad is the number of spline terms to include when estimating the quadratic term.

### Details

This function estimates the ADRF by the method described in Schafer and Galagate (2015), that fits an outcome model using a function of the covariates and spline basis functions derived from the propensity function component.

### Value

prop\_spline\_est returns an object of class "causaldrf\_lsfit", a list that contains the following components:

param the estimated parameters.  
 out\_mod the result of the outcome model fit using lsfit.  
 call the matched call.

### References

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.  
 Little, Roderick and An, Hyonggin (2004). ROBUST LIKELIHOOD-BASED ANALYSIS OF MULTIVARIATE DATA WITH MISSING VALUES. *Statistica Sinica*. **14**: 949–968.  
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### See Also

[iptw\\_est](#), [ismw\\_est](#), [reg\\_est](#), [aipwee\\_est](#), [wtrg\\_est](#), etc. for other estimates.  
[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

### Examples

```
## Example from Schafer (2015).

example_data <- sim_data

t_mod_list <- t_mod(treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data = example_data,
  treat_mod = "Normal")

cond_exp_data <- t_mod_list$T_data
full_data <- cbind(example_data, cond_exp_data)

prop_spline_list <- prop_spline_est(Y = Y,
  treat = T,
```

```

covar_formula = ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
covar_lin_formula = ~ 1,
covar_sq_formula = ~ 1,
data = example_data,
e_treat_1 = full_data$est_treat,
degree = 1,
wt = NULL,
method = "different",
spline_df = 5,
spline_const = 4,
spline_linear = 4,
spline_quad = 4)

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
      example_data$Y[sample_index],
      xlab = "T",
      ylab = "Y",
      main = "propensity spline estimate")

abline(prop_spline_list$param[1],
       prop_spline_list$param[2],
       lty = 2,
       col = "blue",
       lwd = 2)

legend('bottomright',
       "propensity spline estimate",
       lty = 2,
       bty = 'Y',
       cex = 1,
       col = "blue",
       lwd = 2)

rm(example_data, prop_spline_list, sample_index)

```

---

reg\_est

*The regression prediction estimator*


---

### Description

This method estimates the linear or quadratic parameters of the ADRF by estimating a least-squares fit on the basis functions which are composed of combinations of the covariates and treatment values.

### Usage

```

reg_est(Y,
       treat,

```

```

covar_formula,
covar_lin_formula = NULL,
covar_sq_formula = NULL,
data,
degree,
wt = NULL,
method = "same")

```

## Arguments

Y	is the the name of the outcome variable contained in data.
treat	is the name of the treatment variable contained in data.
covar_formula	is the formula to describe the covariates needed to estimate the constant term: $\sim X.1 + \dots$ . Can include higher order terms or interactions. i.e. $\sim X.1 + I(X.1^2) + X.1 * X.2 + \dots$ . Don't forget the tilde before listing the covariates.
covar_lin_formula	is the formula to describe the covariates needed to estimate the linear term, t: $\sim X.1 + \dots$ . Can include higher order terms or interactions. i.e. $\sim X.1 + I(X.1^2) + X.1 * X.2 + \dots$ . Don't forget the tilde before listing the covariates.
covar_sq_formula	is the formula to describe the covariates needed to estimate the quadratic term, t <sup>2</sup> : $\sim X.1 + \dots$ . Can include higher order terms or interactions. i.e. $\sim X.1 + I(X.1^2) + X.1 * X.2 + \dots$ . Don't forget the tilde before listing the covariates.
data	is a dataframe containing Y, treat, and X.
degree	is 1 for linear and 2 for quadratic outcome model.
wt	is weight used in lsfit for outcome regression. Default is wt = NULL.
method	is "same" if the same set of covariates are used to estimate the constant, linear, and/or quadratic term. If method = "different", then different sets of covariates can be used to estimate the constant, linear, and/or quadratic term. covar_lin_formula and covar_sq_formula must be specified if method = "different".

## Details

This function estimates the ADRF by the method described in Schafer and Galagate (2015) that fits an outcome model using a function of the covariates.

## Value

reg\_est returns an object of class "causaldrf\_lsfit", a list that contains the following components:

param	the estimated parameters.
out_mod	the result of the outcome model fit using lsfit.
call	the matched call.

## References

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

Schafer, Joseph L, Kang, Joseph (2008). Average causal effects from nonrandomized studies: a practical guide and simulated example. *Psychological methods*, **13.4**, 279.

## See Also

[iptw\\_est](#), [ismw\\_est](#), [aipwee\\_est](#), [wtrg\\_est](#), etc. for other estimates.

[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

## Examples

```
## Example from Schafer (2015).

example_data <- sim_data

reg_list <- reg_est(Y = Y,
                  treat = T,
                  covar_formula = ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
                  covar_lin_formula = ~ 1,
                  covar_sq_formula = ~ 1,
                  data = example_data,
                  degree = 1,
                  wt = NULL,
                  method = "same")

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
     example_data$Y[sample_index],
     xlab = "T",
     ylab = "Y",
     main = "regression estimate")

abline(reg_list$param[1],
       reg_list$param[2],
       lty = 2,
       col = "blue",
       lwd = 2)

legend('bottomright',
      "regression estimate",
      lty = 2,
      bty = 'Y',
      cex = 1,
      col = "blue",
      lwd = 2)

rm(example_data, reg_list, sample_index)
```

---

 scalar\_wts

*This function calculates scalar weights for use in other models*


---

### Description

This function calculates the scalar weights

### Usage

```
scalar_wts(treat,
           treat_formula,
           numerator_formula,
           data,
           treat_mod,
           link_function,
           ...)
```

### Arguments

treat	is the name of the treatment variable contained in data.
treat_formula	an object of class "formula" (or one that can be coerced to that class) that regresses treat on a linear combination of X: a symbolic description of the model to be fitted.
numerator_formula	an object of class "formula" (or one that can be coerced to that class) that regresses treat on a linear combination of X: a symbolic description of the model to be fitted. i.e. $treat \sim 1$ .
data	is a dataframe containing treat, and X.
treat_mod	a description of the error distribution to be used in the model for treatment. Options include: "Normal" for normal model, "LogNormal" for lognormal model, "Poisson" for Poisson model, "Sqrt" for square-root transformation to a normal treatment, "NegBinom" for negative binomial model, "Gamma" for gamma model.
link_function	is either "log", "inverse", or "identity" for the "Gamma" treat_mod.
...	additional arguments to be passed to the treatment regression fitting function.

### Value

scalar\_wts returns an object of class "causaldrf\_wts", a list that contains the following components:

param	summary of estimated weights.
t_mod	the result of the treatment model fit.
num_mod	the result of the numerator model fit.
weights	estimated weights for each unit.
call	the matched call.

**References**

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

**See Also**

[iptw\\_est](#), [ismw\\_est](#), [reg\\_est](#), [aipwee\\_est](#), [wtrg\\_est](#), etc. for other estimates.

[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

**Examples**

```
## Example from Schafer (2015).

example_data <- sim_data

scalar_wts_list <- scalar_wts(treat = T,
                             treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
                             numerator_formula = T ~ 1,
                             data = example_data,
                             treat_mod = "Normal")

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
      scalar_wts_list$weights[sample_index],
      xlab = "T",
      ylab = "weights",
      main = "scalar_wts")

rm(example_data, scalar_wts_list, sample_index)
```

---

sim\_data

*Simulated data from Schafer and Galagate (2015)*

---

**Description**

Simulated data used in the paper "Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models".

**Usage**

```
data(sim_data)
```

**Format**

A data frame with 1000 rows and 20 variables:

**Details**

A dataset containing sim\_data.

**Value**

(A.1, A.2, A.3, A.4, A.5, A.6, A.7, A.8) are the true measured covariates.

(B.1, B.2, B.3, B.4, B.5, B.6, B.7, B.8) are the transformed covariates.

T	treatment
Theta.1	unit level intercept
Theta.2	unit level slope
Y	outcome

**Source**

use the draw\_sample function

**References**

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

**Examples**

```
## Example from Schafer (2015).
data(sim_data)
head(sim_data)
```

---

t_mod	<i>A function to estimate conditional expected values and higher order moments</i>
-------	--

---

**Description**

This function fits a glm regression specified by the user to estimate conditional moments.

**Usage**

```
t_mod(treat,
      treat_formula,
      data,
      treat_mod,
      link_function,
      ...)
```

**Arguments**

treat	is the name of the treatment variable contained in data.
treat_formula	an object of class "formula" (or one that can be coerced to that class) that regresses treat on a linear combination of X: a symbolic description of the model to be fitted.
data	is a dataframe containing Y, treat, and X.
treat_mod	a description of the error distribution to be used in the model for treatment. Options include: "Normal" for normal model, "LogNormal" for lognormal model, "Sqrt" for square-root transformation to a normal treatment, "Poisson" for Poisson model, "NegBinom" for negative binomial model, "Gamma" for gamma model.
link_function	is either "log", "inverse", or "identity" for the "Gamma" treat_mod.
...	additional arguments to be passed to the low level treatment regression fitting functions.

**Value**

t\_mod returns a list containing the following elements:

T_data	a dataframe containing estimated treatment, estimated treatment squared, estimated treatment cube, estimated treatment quartic, and estimated gps.
T_result	the result of the treatment model fit.

**References**

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

**See Also**

[ismw\\_est](#), [reg\\_est](#), [wtrg\\_est](#), [aipwee\\_est](#), etc. for other estimates.  
[overlap\\_fun](#) to prepare the data for use in the different estimates.

**Examples**

```
## Example from Schafer (2015).

example_data <- sim_data

t_mod_list <- t_mod(treat = T,
                   treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
                   data = example_data,
                   treat_mod = "Normal")

cond_exp_data <- t_mod_list$T_data

full_data <- cbind(example_data, cond_exp_data)

rm(example_data, t_mod_list, cond_exp_data, full_data)
```

---

wtrg\_est

*The weighted regression estimator*


---

### Description

This method uses weight matrices to estimate parameters for an ADRF with quadratic or linear fits.

### Usage

```
wtrg_est(Y,
         treat,
         covar_formula,
         data,
         e_treat_1,
         e_treat_2,
         e_treat_3,
         e_treat_4,
         degree)
```

### Arguments

Y	is the output
treat	is the treatment variable
covar_formula	is the formula for the covariates model of the form: $\sim X.1 + \dots$
data	will contain all the data: X, treat, and Y
e_treat_1	is estimated treatment
e_treat_2	is estimated treatment squared
e_treat_3	is estimated treatment cubed
e_treat_4	is estimated treatment to the fourth
degree	is 1 for linear fit and 2 for quadratic fit

### Details

This function estimates the ADRF by the method described in Schafer and Galagate (2015) which uses weight matrices to adjust for possible bias.

### Value

wtrg\_est returns an object of class "causaldrf", a list that contains the following components:

param	the estimated parameters.
call	the matched call.

## References

Schafer, J.L., Galagate, D.L. (2015). Causal inference with a continuous treatment and outcome: alternative estimators for parametric dose-response models. *Manuscript in preparation*.

## See Also

[iptw\\_est](#), [ismw\\_est](#), [reg\\_est](#), [aipwee\\_est](#), [wtrg\\_est](#), etc. for other estimates.  
[t\\_mod](#), [overlap\\_fun](#) to prepare the data for use in the different estimates.

## Examples

```
## Example from Schafer (2015).

example_data <- sim_data

t_mod_list <- t_mod(treat = T,
  treat_formula = T ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data = example_data,
  treat_mod = "Normal")

cond_exp_data <- t_mod_list$T_data
full_data <- cbind(example_data, cond_exp_data)

wtrg_list <- wtrg_est(Y = Y,
  treat = T,
  covar_formula = ~ B.1 + B.2 + B.3 + B.4 + B.5 + B.6 + B.7 + B.8,
  data = example_data,
  e_treat_1 = full_data$est_treat,
  e_treat_2 = full_data$est_treat_sq,
  e_treat_3 = full_data$est_treat_cube,
  e_treat_4 = full_data$est_treat_quartic,
  degree = 1)

sample_index <- sample(1:1000, 100)

plot(example_data$T[sample_index],
  example_data$Y[sample_index],
  xlab = "T",
  ylab = "Y",
  main = "weighted regression estimate")

abline(wtrg_list$param[1],
  wtrg_list$param[2],
  lty = 2,
  lwd = 2,
  col = "blue")

legend('bottomright',
  "weighted regression estimate",
  lty = 2,
```

*wtrg\_est*

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```
lwd = 2,  
col = "blue",  
bty='Y',  
cex=1)
```

```
rm(example_data, t_mod_list, cond_exp_data, full_data, wtrg_list, sample_index)
```

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