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Cross-referencing the documentation

When reading this manual, you will find references to other Stata manuals. For example,

[U] 26 Overview of Stata estimation commands
[R] regress
[D] reshape

The first example is a reference to chapter 26, *Overview of Stata estimation commands*, in the *User’s Guide*; the second is a reference to the *regress* entry in the *Base Reference Manual*; and the third is a reference to the *reshape* entry in the *Data Management Reference Manual*.

All the manuals in the Stata Documentation have a shorthand notation:

[GSM] Getting Started with Stata for Mac
[GSU] Getting Started with Stata for Unix
[GSW] Getting Started with Stata for Windows
[U] Stata User’s Guide
[R] Stata Base Reference Manual
[BAYES] Stata Bayesian Analysis Reference Manual
[FN] Stata Functions Reference Manual
[XT] Stata Longitudinal-Data/Panel-Data Reference Manual
[M] Stata Multiple-Imputation Reference Manual
[PSS] Stata Power and Sample-Size Reference Manual
[SVY] Stata Survey Data Reference Manual
[I] Stata Glossary and Index
This manual documents commands for the analysis of treatment effects and is referred to as [TE] in cross-references.

After this entry, [TE] treatment effects provides an overview of the treatment-effects estimation commands. The other parts of this manual are arranged alphabetically. If you are new to Stata’s treatment-effects commands, we recommend that you read the following sections first:

- [TE] tefects intro  Introduction to treatment effects for observational data
- [TE] tefects intro advanced  Advanced introduction to treatment effects for observational data
- [TE] tefects multivalued  Multivalued treatment effects

If you are interested in survival analysis, we also recommend that you read the following section first:

- [TE] stteffects intro  Introduction to treatment effects for observational survival-time data

Stata is continually being updated, and Stata users are always writing new commands. To find out about the latest treatment-effects features, type `search treatment effects`.

Reference


Also see

- [U] 1.3 What’s new
- [R] intro  Introduction to base reference manual
treatment effects — Introduction to treatment-effects commands

Description

This manual documents commands that use observational data to estimate the effect caused by getting one treatment instead of another. In observational data, treatment assignment is not controlled by those who collect the data; thus some common variables affect treatment assignment and treatment-specific outcomes. Observational data is sometimes called retrospective data or nonexperimental data, but to avoid confusion, we will always use the term “observational data”.

When all the variables that affect both treatment assignment and outcomes are observable, the outcomes are said to be conditionally independent of the treatment, and the t/effects and st/effects estimators may be used.

When not all of these variables common to both treatment assignment and outcomes are observable, the outcomes are not conditionally independent of the treatment, and et/effects, et/poisson, or et/regress may be used.

t/effects and st/effects offer much flexibility in estimators and functional forms for the treatment-assignment models. t/effects provides models for continuous, binary, count, fractional, and nonnegative outcome variables. st/effects provides many functional forms for survival-time outcomes. See [TE] t/effects intro, [TE] t/effects intro advanced, and [TE] st/effects intro for more information.

et/effects, et/poisson, and et/regress offer less flexibility than t/effects because more structure must be imposed when conditional independence is not assumed. et/effects is for continuous, binary, count, fractional, and nonnegative outcomes and uses a probit model for binary treatments; see [TE] et/effects. et/poisson is for count outcomes and uses a normal distribution to model treatment assignment; see [TE] et/poisson. et/regress is for linear outcomes and uses a normal distribution to model treatment assignment; see [TE] et/regress.

Treatment effects

<table>
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<th>Command</th>
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<td>t/effects ipw</td>
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<td>t/effects ipwra</td>
<td>Inverse-probability-weighted regression adjustment</td>
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<td>t/effects nnmatch</td>
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<tr>
<td>t/effects ra</td>
<td>Regression adjustment</td>
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Survival treatment effects

[TE] stteffects ipw  Survival-time inverse-probability weighting
[TE] stteffects ipwra  Survival-time inverse-probability-weighted regression adjustment
[TE] stteffects ra  Survival-time regression adjustment
[TE] stteffects wra  Survival-time weighted regression adjustment

Endogenous treatment effects

[TE] eteffects  Endogenous treatment-effects estimation
[TE] etpoisson  Poisson regression with endogenous treatment effects
[TE] etregress  Linear regression with endogenous treatment effects

Treatment effects with sample selection and endogenous covariates

[ERM] eregress  Extended linear regression
[ERM] eintreg  Extended interval regression
[ERM] eprobit  Extended probit regression
[ERM] eoprobit  Extended ordered probit regression

Postestimation tools

[TE] tebalance  Check balance after teffects or stteffects estimation
[TE] tebalance box  Covariate balance box
[TE] tebalance density  Covariate balance density
[TE] tebalance overid  Test for covariate balance
[TE] tebalance summarize  Covariate-balance summary statistics
[TE] teffects overlap  Overlap plots
[TE] eteffects postestimation  Postestimation tools for eteffects
[TE] etpoisson postestimation  Postestimation tools for etpoisson
[TE] etregress postestimation  Postestimation tools for etregress
[TE] stteffects postestimation  Postestimation tools for stteffects

Also see

[U] 1.3 What’s new
[TE] teffects intro  — Introduction to treatment effects for observational data
[TE] teffects intro advanced  — Advanced introduction to treatment effects for observational data
[TE] teffects multivalued  — Multivalued treatment effects
[TE] stteffects intro  — Introduction to treatment effects for observational survival-time data
[TE] Glossary
**Description**

*eteffects* estimates the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs) from observational data when treatment assignment is correlated with the potential outcomes. It allows for continuous, binary, count, fractional, and nonnegative outcomes and requires a binary treatment. To control for the endogeneity of the treatment assignment, the estimator includes residuals from the treatment model in the models for the potential outcomes, known as a control-function approach.

**Quick start**

ATE of binary treatment *treat* using a linear model for outcome *y1* on *x* and the residuals from a probit model for *treat* on *x* and *z*

\[ \text{eteffects (y1 x) (treat x z)} \]

As above, but estimate ATET

\[ \text{eteffects (y1 x) (treat x z), atet} \]

As above, but estimate POMs

\[ \text{eteffects (y1 x) (treat x z), pomeans} \]

As above, and show parameters from auxiliary equations

\[ \text{eteffects (y1 x) (treat x z), pomeans aequations} \]

ATE of *treat* using an exponential-mean model for *y1*

\[ \text{eteffects (y1 x, exponential) (treat x z)} \]

Same as above, but for count outcome *y2*

\[ \text{eteffects (y2 x, exponential) (treat x z)} \]

As above, but use a probit model for binary outcome *y3*

\[ \text{eteffects (y3 x, probit) (treat x z)} \]

As above, but use a fractional probit model for *y4* ranging from 0 to 1

\[ \text{eteffects (y4 x, fractional) (treat x z)} \]

**Menu**

Statistics > Treatment effects > Endogenous treatment > Control function estimator > Continuous outcomes
Statistics > Treatment effects > Endogenous treatment > Control function estimator > Binary outcomes
Statistics > Treatment effects > Endogenous treatment > Control function estimator > Count outcomes
Statistics > Treatment effects > Endogenous treatment > Control function estimator > Fractional outcomes
Statistics > Treatment effects > Endogenous treatment > Control function estimator > Nonnegative outcomes
et effects — Endogenous treatment-effects estimation  5

Syntax

et effects (ovar omvarlist [, omodel noconstant])

(tvar tmvarlist [, noconstant]) [if] [in] [weight] [, stat options]

ovar is the depvar of the outcome model.

omvarlist is the list of exogenous indepvars in the outcome model.

tvar is the binary treatment variable.

tmvarlist is the list of covariates that predict treatment assignment.

omodel  Description

Model
linear  linear outcome model; the default
fractional  fractional probit outcome model
probit  probit outcome model
exponential  exponential-mean outcome model

stat  Description

Model
ate  estimate average treatment effect in population; the default
atet  estimate average treatment effect on the treated
pomeans  estimate potential-outcome means

options  Description

Model
noconstant  suppress constant term

SE/Robust
vce(vcetype)  vcetype may be robust, cluster clustvar, bootstrap, or jackknife

Reporting
level(#)  set confidence level; default is level(95)
agredonations  display auxiliary-equation results
display_options  control columns and column formats, row spacing, line width, display of omitted variables and base and empty cells, and factor-variable labeling

Maximization
maximize_options  control the maximization process; seldom used

Advanced
pstolerance(#)  set tolerance for overlap assumption
osample(newvar)  generate newvar to mark observations that violate the overlap assumption
coefflegend  display legend instead of statistics
**eteffects** — Endogenous treatment-effects estimation

`omvarlist` and `tmvarlist` may contain factor variables; see [U] 11.4.3 Factor variables.

`bootstrap`, `by`, `jackknife`, and `statsby` are allowed; see [U] 11.1.10 Prefix commands.

Weights are not allowed with the `bootstrap` prefix; see [R] `bootstrap`.

`fweights`, `iweights`, and `pweights` are allowed; see [U] 11.1.6 weight.

`coeflegend` does not appear in the dialog box.

See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.

### Options

#### Model

- **noconstant**; see [R] estimation options.

  `stat` is one of three statistics: `ate`, `atet`, or `pomeans`. `ate` is the default.

  - `ate` specifies that the average treatment effect be estimated.
  - `atet` specifies that the average treatment effect on the treated be estimated.
  - `pomeans` specifies that the potential-outcome means for each treatment level be estimated.

#### SE/Robust

- **vce(vcetype)** specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (`robust`), that allow for intragroup correlation (`cluster clustvar`), and that use bootstrap or jackknife methods (`bootstrap`, `jackknife`); see [R] `vce_option`.

#### Reporting

- **level(#)**; see [R] estimation options.

  `aequations` specifies that the results for the outcome-model or the treatment-model parameters be displayed. By default, the results for these auxiliary parameters are not displayed.

  `display_options`: `noci`, `nopvalues`, `noomitted`, `vsquish`, `noemptycells`, `baselevels`, `allevels`, `nofvlabel`, `fvwrap(#)`, `fvwrapon(style)`, `cformat(%)`, `pformat(%)`, `sformat(%)`, and `nolstretch`; see [R] estimation options.

#### Maximization

- **maximize_options**: `iterate(#)`, `[no]log`, and `from(init_specs)`; see [R] `maximize`. These options are seldom used.

  - `init_specs` is one of

    - `matname [ , skip copy ]`
    - `# [ , # ...], copy`

#### Advanced

- **pstolerance(#)** specifies the tolerance used to check the overlap assumption. The default value is `pstolerance(1e-5)`. `eteffects` will exit with an error if an observation has an estimated propensity score smaller than that specified by `pstolerance()`.

  `osample(newvar)` specifies that indicator variable `newvar` be created to identify observations that violate the overlap assumption.

The following option is available with `eteffects` but is not shown in the dialog box:

- **coeflegend**; see [R] estimation options.
Remarks and examples

If you are unfamiliar with treatment-effects estimators for observational data or the teffects commands, we recommend that you look at [TE] teffects intro. For the intuition behind some of the concepts discussed below, we recommend that you read Defining treatment effects in [TE] teffects intro advanced.

The estimators implemented in eteffects extend the regression adjustment (RA) estimators implemented in teffects ra to allow for endogenous treatments, that is, when treatment assignment is not independent of outcomes. This endogeneity is a violation of the conditional mean independence assumption used by teffects ra, as discussed in The potential-outcome model in [TE] teffects intro advanced.

eteffects estimates the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs). It uses a linear, a probit, a fractional probit, or an exponential-mean model for the potential outcomes and a probit model for treatment assignment. After conditioning on the observable covariates, eteffects allows some remaining unobservable components to affect both treatment assignment and the potential outcomes. The treatment assignment process is endogenous because these unobservable components affect both treatment assignment and the potential outcomes.

To control for the endogeneity of the treatment assignment, eteffects uses a control-function approach. This method controls for endogeneity by including the residuals from the treatment-assignment model as a regressor in the models for the potential outcome. The implementation in eteffects follows Wooldridge (2010), who provides an excellent discussion of the control-function approach that addresses endogeneity problems in a treatment-effects context.

The control-function approach estimates the parameters of the conditional means of the potential outcomes. Sample averages of the conditional means are used to estimate the unconditional ATE, ATET, or POMs. This method is known as RA.

Taken collectively, the estimators implemented in eteffects are control-function RA estimators. See Methods and formulas below for details about the estimation procedure.

Example 1: Linear outcome estimates for ATE

Suppose we want to know the effect of a mother smoking while pregnant on the birthweight of her infant. We use an extract from Cattaneo (2010) in which bweight records the baby’s birthweight and mbsmoke is the variable (0 or 1) indicating whether a mother smoked while pregnant.

We may believe that birthweight (the potential outcome) is influenced by whether the mother had a prenatal exam in the first trimester, whether the mother is married, the mother’s age, whether this is the first birth, and the education level of the father. We may also believe that the smoking decision (the treatment) is influenced by the mother’s marital status, the education level of the mother, her age, whether she had a prenatal exam in the first trimester, and whether this baby is her first baby.

Thus we condition on different sets of covariates in the models for treatment assignment and the potential outcomes. In the probit model for smoking status (mbsmoke), we condition on marital status (mmarried), age (mage), mother’s education level (medu), father’s education level (fedu), and whether it was the mother’s first baby (fbaby). We model birthweight (bweight) as a linear function of whether the mother had a first-trimester prenatal exam (prenatal1), mmarried, mage, and fbaby. We can estimate the ATE of smoking status using one of the teffects estimators if we believe that there are no unobservable components that affect both the decision to smoke while pregnant and the potential birthweights.
If we believe there is some unobservable factor that affects both assignment to treatment and the potential outcome, we must select another estimator. For example, we do not observe a mother’s health consciousness, which affects both the smoking decision and each potential birthweight through other behaviors such as intake of prenatal vitamins. Under these assumptions, the estimators in `eteffects` consistently estimate the ATE, but the estimators in `TE` `teffects` yield inconsistent estimates.

```
. use http://www.stata-press.com/data/r15/cattaneo2
. eteffects (bweight i.prenatal1 i.mmarried mage i.fbaby)
   > (mbsmoke i.mmarried mage i.fbaby medu fedu)
Iteration 0:   EE criterion = 4.704e-24
Iteration 1:   EE criterion = 1.223e-25
Endogenous treatment-effects estimation Number of obs = 4,642
Outcome model : linear
Treatment model: probit

        bweight |      Coef.     Std. Err.      z    P>|z|     [95% Conf. Interval]
-------------+----------------------------------
        ATE   |  mbsmoke (smoker vs nonsmoker)  
          | -455.9119    212.4393   -2.15   0.032    -872.2853    -39.53852
        POmean  |  mbsmoke nonsmoker               
          |  3437.964     31.21145  110.15   0.000     3376.791    3499.138
```

When no mother smokes, the average birthweight is 3,438 grams. The average birthweight is 456 grams less when all mothers smoke than when no mother smokes.

We can compare these results with those obtained if we ignore the endogeneity of the smoking decision. Below we estimate the ATE using the inverse-probability-weighted regression-adjustment estimator in `TE` `teffects ipwra`.

```
. teffects ipwra (bweight i.prenatal1 i.mmarried mage i.fbaby)
   > (mbsmoke i.mmarried mage i.fbaby medu fedu)
Iteration 0:   EE criterion = 3.036e-22
Iteration 1:   EE criterion = 3.755e-26
Treatment-effects estimation Number of obs = 4,642
Estimator : IPW regression adjustment
Outcome model : linear
Treatment model: logit

        bweight |      Coef.     Std. Err.      z    P>|z|     [95% Conf. Interval]
-------------+----------------------------------
        ATE   |  mbsmoke (smoker vs nonsmoker)  
          |  -233.6835    25.07695   -9.32   0.000    -282.8335   -184.5336
        POmean  |  mbsmoke nonsmoker               
          |   3403.191     9.529709  357.11   0.000     3384.513     3421.869
```

When no mother smokes, the average birthweight is 3,438 grams. The average birthweight is 456 grams less when all mothers smoke than when no mother smokes.
In magnitude, the estimated ATE is more than half the estimate that allows for endogenous treatment assignment. If there is endogeneity, disregarding it underestimates the effect of smoking on birthweight. We show how to test for endogeneity in example 1 of \texttt{eteffects postestimation}.

\section*{Example 2: Estimating the \textit{ATET}}

Continuing \texttt{example 1}, we can use the \texttt{atet} option to estimate the \textit{ATET}.

\begin{verbatim}
. eteffects (bweight i.prenatal1 i.mmarried mage i.fbaby) > (mbsmoke i.mmarried mage i.fbaby medu fedu), atet

Iteration 0:   EE criterion =  4.688e-24
Iteration 1:   EE criterion =  8.479e-26

Endogenous treatment-effects estimation       Number of obs = 4,642
Outcome model : linear
Treatment model: probit

|          | Coef.  | Std. Err. | z     | P>|z|    | [95% Conf. Interval] |
|----------|--------|-----------|-------|--------|---------------------|
| ATET mbsmoke smoker vs nonsmoker | -409.8527 | 161.4816  | -2.54 | 0.011  | (-726.3507, -93.35466) |
|          | POmean mbsmoke nonsmoker | 3547.512 | 160.0595 | 22.16 | 0.000  | (3233.801, 3861.223) |
\end{verbatim}

In the population of mothers who smoke, the average infant birthweight would be 3,548 grams if none of these mothers smoked. For the mothers who smoke, the average infant birthweight is 410 grams less than if none of these mothers smoked.

\section*{Example 3: Exponential-mean outcomes}

We estimate the ATE of living in an urban area on monthly earnings (\textit{wage}), using a subset of the National Longitudinal Survey in 1980 found in \textit{Wooldridge (2010)}. We assume that once we condition on work experience (\textit{exper}), whether education level attained is college or higher (\textit{college}), and IQ (\textit{iq}), individual wages follow an exponential mean. The variables used to predict residence in an urban area (\textit{urban}) are \textit{college} and whether the respondent’s father attained a bachelor’s degree or higher (\textit{fcollege}).
. use http://www.stata-press.com/data/r15/nlsy80
. eteffects (wage exper iq i.college, exponential nocons)
> (urban i.college fcollege)
Iteration 0:  EE criterion = 2.903e-25
Iteration 1:  EE criterion = 2.903e-25  (backed up)
Endogenous treatment-effects estimation  Number of obs =  935
Outcome model :  exponential
Treatment model:  probit

|          | Coef.  |  Robust Std. Err. |      z    |  P>|z|   | [95% Conf. Interval] |
|-----------|--------|-------------------|-----------|-------|---------------------|
|         ATE |        |                   |           |       |                     |
| urban     |        |                   |           |       |                     |
| (1 vs 0)  | 481.0465  |  31.74882        |  15.15    | 0.000 |  418.82 - 543.2731  |
| POmean    |        |                   |           |       |                     |
| urban     |        |                   |           |       |                     |
| 0         |  233.8083  |  13.51028        |  17.31    | 0.000 |  207.3286 - 260.288 |

When everyone lives outside urban areas, wages are $234 a month on average. Wages are $481 a month greater, on average, when everyone lives in urban areas.

Stored results

eteffects stores the following in e():

Scalars

e(N)            number of observations

Scalars

e(nj)           number of observations for treatment level j

e(k_eq)         number of equations in e(b)

e(k_levels)     number of levels in treatment variable

e(rank)         rank of e(V)

e(converged)    1 if converged, 0 otherwise

Macros

e(cmd)          eteffects

e(cmdline)      command as typed

e(depvar)       name of outcome variable

e(tvar)         name of treatment variable

e(omodel)       fractional, linear, probit, or exponential

e(stat)         statistic estimated, ate, atet, or pomeans

e(wtype)        weight type

e(wexp)         weight expression

e(title)        title in estimation output

e(tlevels)      levels of treatment variable

e(vce)          vctype specified in vce()

e(vcetype)      title used to label Std. Err.

e(properties)   b V

e(estat_cmd)    program used to implement estat

e(predict)      program used to implement predict

e(marginsnotok) predictions disallowed by margins

e(asbalanced)   factor variables fvset as asbalanced

e(asobserved)   factor variables fvset as asobserved

Matrices

e(b)            coefficient vector

e(V)            variance–covariance matrix of the estimators

Functions

e(sample)       marks estimation sample
Methods and formulas

The treatment-effects models considered in eteffects are given by

\[ y_{i0} = E(y_{i0}|x_i) + \epsilon_{i0} \]  
\[ y_{i1} = E(y_{i1}|x_i) + \epsilon_{i1} \]  
\[ t_i = E(t_i|z_i) + \nu_i \]  
\[ y_i = t_i y_{i1} + (1 - t_i) y_{i0} \]  
\[ E(\epsilon_{ij}|x_i, z_i) = E(\epsilon_{ij}|z_i) = E(\epsilon_{ij}|x_i) = 0 \text{ for } j \in \{0,1\} \]  
\[ E(\epsilon_{ij}|t_i) \neq 0 \text{ for } j \in \{0,1\} \]

where the subscript \( i \) denotes individual level observations, \( y_{i1} \) is the potential outcome of receiving the treatment, \( y_{i0} \) is the potential outcome when the treatment is not received, \( t_i \) is the observed binary treatment, and \( y_i \) is the observed outcome. Each one of the potential outcomes is determined by its expected value conditional on a set of regressors \( x_i \) and an unobserved random component \( \epsilon_{ij} \), for \( j \in \{0,1\} \). Similarly, the treatment is given by its expectation conditional on a set of regressors \( z_i \), which does not need to differ from \( x_i \), and an unobserved component \( \nu_i \).

Equations (1)–(5) describe the parametric treatment-effects models in [TE] eteffects. Equation (6) adds endogeneity to the framework. It states that the unobservables in the potential-outcome equations are correlated to treatment status. For our birthweight example, this would happen if mothers who do not smoke are more health conscious than those who smoke and if we do not observe health awareness in our data. If we do not observe health awareness, the decision to smoke or not to smoke is not independent of the infant’s birthweight.

Equations (3), (5), and (6) are the basis of the control-function estimator implemented by eteffects. Equation (5) states that the unobserved components in the potential outcome are independent of \( z_i \). Therefore, the correlation between \( t_i \) and the unobserved components must be equivalent to the correlation between \( \epsilon_{ij} \) and \( \nu_i \). Another way of stating this is

from (3) \[ E(\epsilon_{ij}|t_i) = E(\epsilon_{ij}|E(t_i|z_i) + \nu_i) \]  
from (5) \[ = E(\epsilon_{ij}|\nu_i) \]  
\[ = \nu_i \beta_{2j} \]

We fit (3) using a probit estimator. We then obtain \( \hat{\nu}_i \) as the difference between the treatment and our estimate of \( E(t_i|z_i) \) and use this statistic to compute an estimate of \( E(y_{ij}|x_i, \nu_i, t_i) \) for \( j \in \{0,1\} \). If the outcome is linear, for instance,

\[ E(y_{ij}|x_i, \nu_i, t_i = j) = x_i' \beta_{1j} + \nu_i \beta_{2j} \text{ for } j \in \{0,1\} \]
For the probit and exponential-mean cases, respectively, we have the following:

\[
E(y_{ij}|x_i, \nu_i, t_i = j) = \Phi(x_i'\beta_{1j} + \nu_i\beta_{2j})
\]

(8)

\[
E(y_{ij}|x_i, \nu_i, t_i = j) = \exp(x_i'\beta_{1j} + \nu_i\beta_{2j})
\]

(9)

The parameters of (3) and (7)–(9), and the ATE, ATET, and POMs are estimated using the generalized method of moments (GMM). The moment equations used in GMM are the sample analogs of \( E \{ w'_i \epsilon_i(\theta) \} = 0 \), where \( w_i \) are the instruments, \( \epsilon_i(\theta) \) are residuals, and \( \theta \) are the parameters of the model (see \([R] gmm\)). The moment conditions in the GMM estimation for the linear model are given by

\[
\frac{1}{n} \sum_{i=1}^{n} x_i'(y_i - x_i'\hat{\beta}_{1j} + \hat{\nu}_i\hat{\beta}_{2j})t_i = 0
\]

(10)

\[
\frac{1}{n} \sum_{i=1}^{n} x_i'(y_i - x_i'\hat{\beta}_{1j} + \hat{\nu}_i\hat{\beta}_{2j})(1 - t_i) = 0
\]

(11)

\[
\frac{1}{n} \sum_{i=1}^{n} z_i' \left\{ t_i \frac{\phi(z_i'\hat{\pi})}{\Phi(z_i'\hat{\pi})} - (1 - t_i) \frac{\phi(z_i'\hat{\pi})}{1 - \Phi(z_i'\hat{\pi})} \right\} = 0
\]

(12)

\[
\frac{1}{n} \sum_{i=1}^{n} \left\{ (x_i'\hat{\beta}_{10} + \hat{\nu}_i\hat{\beta}_{20}) - \text{POM0} \right\} = 0
\]

(13)

\[
\frac{1}{n} \sum_{i=1}^{n} \left\{ (x_i'\hat{\beta}_{11} + \hat{\nu}_i\hat{\beta}_{21}) - \text{POM0} - \hat{\text{ATE}} \right\} = 0
\]

(14)

where \( \hat{\nu}_i = t_i - \Phi(z_i'\hat{\pi}) \), \( n \) is the number of observations, and \( \hat{\beta}_{11}, \hat{\beta}_{10}, \hat{\beta}_{21}, \hat{\beta}_{20}, \hat{\pi}, \hat{\text{ATE}}, \text{and} \hat{\text{POM0}} \) are the parameters. If we want to estimate the ATET, we replace (14) with

\[
\frac{1}{n} \sum_{i=1}^{n} \left\{ (x_i'\hat{\beta}_{11} + \hat{\nu}_i\hat{\beta}_{21}) \frac{n}{n_t} - \text{POM0} \frac{n}{n_t} - \hat{\text{ATET}} \right\} = 0
\]

(15)

and if we want to estimate the potential-outcome means, we replace (14) with

\[
\frac{1}{n} \sum_{i=1}^{n} \left\{ (x_i'\hat{\beta}_{11} + \hat{\nu}_i\hat{\beta}_{21}) - \text{POM1} \right\} = 0
\]

(16)

where \( \hat{\text{ATET}} \) and \( \hat{\text{POM1}} \) are the parameters of the model, and \( n_t \) is the number of treated units.
For the exponential-mean outcome model, we replace \( x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j} \) with \( \exp(x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j}) \) to obtain the residual equations in (10)–(16). For the probit outcome model, we replace (10) and (11) with the following:

\[
\frac{1}{n} \sum_{i=1}^{n} t_i x_i' \begin{cases} 
\phi \left( x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j} \right) - (1 - y_i) \frac{\phi \left( x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j} \right)}{1 - \Phi \left( x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j} \right)} \\
\frac{1}{n} \sum_{i=1}^{n} (1 - t_i) x_i' \begin{cases} 
\phi \left( x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j} \right) - (1 - y_i) \frac{\phi \left( x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j} \right)}{1 - \Phi \left( x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j} \right)} 
\end{cases}
\end{cases}
\]

For the remaining equations, \( x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j} \) is replaced with \( \Phi(x_i' \hat{\beta}_{1j} + \hat{\nu}_i \hat{\beta}_{2j}) \). The fractional probit model uses the same moment conditions as the probit model.

### Acknowledgment

We thank Jeffrey Wooldridge of Michigan State University for helpful conversations and advice about the literature and methods in this area.

### References


### Also see

[TE] eteffects — Postestimation tools for eteffects

[TE] etregress — Linear regression with endogenous treatment effects

[TE] teffects — Treatment-effects estimation for observational data

[R] gmm — Generalized method of moments estimation

[R] probit — Probit regression

[R] regress — Linear regression

[U] 20 Estimation and postestimation commands
Postestimation commands

The following postestimation command is of special interest after `eteffects`:

<table>
<thead>
<tr>
<th>Command</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>estat endogenous</code></td>
<td>perform tests of endogeneity</td>
</tr>
</tbody>
</table>

The following standard postestimation commands are available after `eteffects`:

<table>
<thead>
<tr>
<th>Command</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>estat summarize</code></td>
<td>summary statistics for the estimation sample</td>
</tr>
<tr>
<td><code>estat vce</code></td>
<td>variance–covariance matrix of the estimators (VCE)</td>
</tr>
<tr>
<td><code>estimates</code></td>
<td>cataloging estimation results</td>
</tr>
<tr>
<td><code>hausman</code></td>
<td>Hausman’s specification test</td>
</tr>
<tr>
<td><code>lincom</code></td>
<td>point estimates, standard errors, testing, and inference for linear combinations of coefficients</td>
</tr>
<tr>
<td><code>nlcom</code></td>
<td>point estimates, standard errors, testing, and inference for nonlinear combinations of coefficients</td>
</tr>
<tr>
<td><code>predict</code></td>
<td>predictions, residuals, influence statistics, and other diagnostic measures</td>
</tr>
<tr>
<td><code>predictnl</code></td>
<td>point estimates, standard errors, testing, and inference for generalized predictions</td>
</tr>
<tr>
<td><code>test</code></td>
<td>Wald tests of simple and composite linear hypotheses</td>
</tr>
<tr>
<td><code>testnl</code></td>
<td>Wald tests of nonlinear hypotheses</td>
</tr>
</tbody>
</table>

predict

**Description for predict**

`predict` creates a new variable containing predictions such as treatment effects, conditional means, propensity scores, and linear predictions.

**Menu for predict**

Statistics  >  Postestimation
Syntax for predict

predict [type] { stub* | newvar | newvarlist} [if] [in] [, statistic tlevel]

predict [type] { stub* | newvarlist} [if] [in], scores

**statistic**          Description

<table>
<thead>
<tr>
<th>Main</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>te</td>
<td>treatment effect; the default</td>
</tr>
<tr>
<td>cmean</td>
<td>conditional mean at treatment level</td>
</tr>
<tr>
<td>ps</td>
<td>propensity score</td>
</tr>
<tr>
<td>xb</td>
<td>linear prediction</td>
</tr>
<tr>
<td>psxb</td>
<td>linear prediction for propensity score</td>
</tr>
<tr>
<td>xbtotall</td>
<td>linear prediction, using residuals from treatment model</td>
</tr>
</tbody>
</table>

Specify one new variable with te; specify one or two new variables with cmean, ps, and xb.

**Options for predict**

**Main**

te, the default, calculates the treatment effect.

cmean calculates the conditional mean for the control group. To also obtain the conditional mean for the treatment group, specify two variables. If you want the conditional mean for only the treatment group, specify the tlevel option.

ps calculates the probability of being in the control group. To also obtain the probability of being in the treatment group, specify two variables. If you want the probability of being in the treatment group only, specify the tlevel option.

xb calculates the linear prediction for the control group. To also obtain the linear prediction for the treatment group, specify two variables. If you want the linear prediction for only the treatment group, specify the tlevel option.

psxb calculates the linear prediction for the propensity score.

xbtotal calculates the linear prediction for the control group, including the residuals from the treatment model as regressors. To also obtain the linear prediction for the treatment group, specify two variables. If you want the linear prediction, including the residuals from the treatment model as regressors, only for the treatment group, specify the tlevel option.

tlevel specifies that the statistic be calculated for the treatment group; the default is to calculate the statistic for the control group.

scores calculates the score variables. For eteffects, this is the same as the residuals in the moment conditions used by the generalized method of moments (see [R] gmm). For the average treatment effect, the average treatment effect on the treated, and the potential-outcome means, parameter-level scores are computed. For the auxiliary equations, equation-level scores are computed.
estat

Description for estat

estat endogenous performs a Wald test to determine whether the estimated correlations between the treatment-assignment and potential-outcome models are different from zero. The null hypothesis is that the correlations are jointly zero. Rejection of the null hypothesis suggests endogeneity.

Menu for estat

Statistics > Postestimation

Syntax for estat

estat endogenous

Remarks and examples

Example 1: Testing for endogeneity

In example 3 of [TE] eteffects, endogeneity could arise if unobservable factors that determine wages are correlated with the decision to live in an urban area. If there is no endogeneity, we would prefer to use one of the eteffects estimators because they will give us the correct standard errors. The control-function approach used by eteffects allows us to test for endogeneity.

The control-function approach estimates the correlation between the unobservables of the treatment-assignment and potential-outcome models. If there is no correlation between the unobservables, then there is no endogeneity. We test for correlation, and thus for endogeneity, by typing

. use http://www.stata-press.com/data/r15/nlsy80
. eteffects (wage exper iq i.college, exponential nocons)
> (urban i.college fcollege)
(output omitted)
. estat endogenous

Test of endogeneity
Ho: treatment and outcome unobservables are uncorrelated

    chi2(  2) =  275.36
Prob > chi2 =    0.0000

We reject the null hypothesis of no endogeneity. This suggests that unobservable factors that determine wages mediate the decision to live in an urban area.

Technical note

The estimated correlations between the unobservables of the treatment-assignment and potential-outcome models are auxiliary parameters. They appear under the headings TEOM0 and TEOM1, which refer to treatment residuals (TE) for outcome model 0 (OM0) and outcome model 1 (OM1), when the option aequations is specified.
For the model in example 3 of [TE] eteffects with the aequations option, the results are the following:

```
. eteffects (wage exper iq i.college, exponential nocons)
> (urban i.college fcollege), aequations
Iteration 0: EE criterion = 2.903e-25
Iteration 1: EE criterion = 2.903e-25 (backed up)
Endogenous treatment-effects estimation Number of obs = 935
Outcome model : exponential
Treatment model: probit

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coef.</td>
</tr>
<tr>
<td>ATE</td>
<td></td>
</tr>
<tr>
<td>urban (1 vs 0)</td>
<td></td>
</tr>
<tr>
<td>POmean</td>
<td></td>
</tr>
<tr>
<td>urban 0</td>
<td></td>
</tr>
<tr>
<td>TME1</td>
<td></td>
</tr>
<tr>
<td>college 1</td>
<td></td>
</tr>
<tr>
<td>fcollege</td>
<td></td>
</tr>
<tr>
<td>_cons</td>
<td></td>
</tr>
<tr>
<td>OME0</td>
<td></td>
</tr>
<tr>
<td>exper</td>
<td></td>
</tr>
<tr>
<td>iq</td>
<td></td>
</tr>
<tr>
<td>college 1</td>
<td></td>
</tr>
<tr>
<td>OME1</td>
<td></td>
</tr>
<tr>
<td>exper</td>
<td></td>
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<tr>
<td>iq</td>
<td></td>
</tr>
<tr>
<td>college 1</td>
<td></td>
</tr>
<tr>
<td>TEOM0</td>
<td></td>
</tr>
<tr>
<td>_cons</td>
<td></td>
</tr>
<tr>
<td>TEOM1</td>
<td></td>
</tr>
<tr>
<td>_cons</td>
<td></td>
</tr>
</tbody>
</table>
```

Among other things, we can use these correlations to test the joint significance of the coefficients on the residuals from the treatment-assignment models. This is equivalent to the endogeneity test in example 1. We type

```
. test [TEOM0]_cons [TEOM1]_cons
( 1) [TEOM0]_cons = 0
( 2) [TEOM1]_cons = 0
   chi2( 2) = 275.36
   Prob > chi2 = 0.0000
```
Also see

[TE] eteffects — Endogenous treatment-effects estimation

[U] 20 Estimation and postestimation commands
etpoisson estimates the parameters of a Poisson regression model in which one of the regressors is an endogenous binary treatment. Both the average treatment effect and the average treatment effect on the treated can be estimated with etpoisson.

Quick start

Poisson model of \( y \) on \( x \) and endogenous binary treatment \( \text{treat} \) modeled by \( x \) and \( w \)

\[
etpoisson y \ x, \ \text{treat}(\text{treat} = x \ w)
\]

With robust standard errors

\[
etpoisson y \ x, \ \text{treat}(\text{treat} = x \ w) \ \text{vce(robust)}
\]

Average treatment effect after etpoisson with the required \text{vce(robust)} option

\[
margins \ r.treat, \ \text{vce(unconditional)}
\]

As above, but calculate average treatment effect on the treated

\[
margins, \ \text{vce(unconditional)} \ \text{predict(cte)} \ \text{subpop(if} \ \text{treat==1)}
\]

Menu

Statistics > Treatment effects > Endogenous treatment > Maximum likelihood estimator > Count outcomes
Syntax

\texttt{etpoisson depvar \ [indepvars] \ [if] \ [in] \ [weight],}
\[ \text{treat}(\text{depvar}_t = \text{indepvars}_t \ [, \ \text{noconstant} \ \text{offset(\text{varname}_o)})] \ [\text{options}] \]

<table>
<thead>
<tr>
<th>\textit{options}</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>\textbf{Model}</td>
<td></td>
</tr>
<tr>
<td>\texttt{treat()}</td>
<td>equation for treatment effects</td>
</tr>
<tr>
<td>\texttt{noconstant}</td>
<td>suppress constant term</td>
</tr>
<tr>
<td>\texttt{exposure(\text{varname}_e)}</td>
<td>include $\ln(\text{varname}_e)$ in model with coefficient constrained to 1</td>
</tr>
<tr>
<td>\texttt{offset(\text{varname}_o)}</td>
<td>include $\text{varname}_o$ in model with coefficient constrained to 1</td>
</tr>
<tr>
<td>\texttt{constraints(\text{constraints})}</td>
<td>apply specified linear constraints</td>
</tr>
<tr>
<td>\texttt{collinear}</td>
<td>keep collinear variables</td>
</tr>
</tbody>
</table>

| \textbf{SE/Robust}        |                                                  |
| \texttt{vce(vcetype)}     | \text{vcetype} may be \text{oim}, \text{robust}, \text{cluster} \text{ clustvar}, \text{opg}, \text{bootstrap}, or \text{jackknife} |

| \textbf{Reporting}        |                                                  |
| \texttt{level(#)}         | set confidence level; default is \text{level(95)} |
| \texttt{irr}              | report incidence-rate ratios                     |
| \texttt{noconsreport}     | do not display constraints                       |
| \texttt{display_options}  | control columns and column formats, row spacing, line width, display of omitted variables and base and empty cells, and factor-variable labeling |

| \textbf{Integration}      |                                                  |
| \texttt{intpoints(#)}     | use \# Gauss–Hermite quadrature points; default is \text{intpoints(24)} |

| \textbf{Maximization}     |                                                  |
| \texttt{maximize_options} | control the maximization process; seldom used    |
| \texttt{coeflegend}       | display legend instead of statistics             |

*\texttt{treat()}* is required.

The full specification is \texttt{treat(\text{depvar}_t = \text{indepvars}_t \ [, \ \text{noconstant} \ \text{offset(\text{varname}_o)})]}.

\textit{indepvars} and \textit{indepvars}_t may contain factor variables; see \cite{factorvariables}.

\textit{depvar}, \textit{depvar}_t, \textit{indepvars}, and \textit{indepvars}_t may contain time-series operators; see \cite{timeseries}.

\texttt{bootstrap}, \texttt{by}, \texttt{jackknife}, \texttt{rolling}, \texttt{statsby}, and \texttt{svy} are allowed; see \cite{prefix}

Weights are not allowed with the \texttt{bootstrap} prefix; see \cite{bootstrap}.

\texttt{aweights} are not allowed with the \texttt{jackknife} prefix; see \cite{jackknife}.

\texttt{vce()} and weights are not allowed with the \texttt{svy} prefix; see \cite{svy}.

\texttt{fweights}, \texttt{aweights}, \texttt{iweights}, and \texttt{pweights} are allowed; see \cite{weight}.

\texttt{coeflegend} does not appear in the dialog box.

See \cite{estimation} for more capabilities of estimation commands.
**Options**

**Model**

treat\( (\text{depvar}_t = \text{indepvars}_t \ [, \text{noconstant offset(varname}_o]) \)\) specifies the variables and options for the treatment equation. It is an integral part of specifying a treatment-effects model and is required.

The indicator of treatment, \( \text{depvar}_t \), should be coded as 0 or 1.

\text{noconstant}, \text{exposure(varname}_e), \text{offset(varname}_o), \text{constraints(constraints)}, \text{collinear}; see [R] estimation options.

**SE/Robust**

\text{vce(vcetype)} specifies the type of standard error reported, which includes types that are derived from asymptotic theory (\text{oim}, \text{opg}), that are robust to some kinds of misspecification (\text{robust}), that allow for intragroup correlation (\text{cluster clustvar}), and that use bootstrap or jackknife methods (\text{bootstrap}, \text{jackknife}); see [R] vce_option.

**Reporting**

\text{level(#)}; see [R] estimation options.

\text{irr} reports estimated coefficients transformed to incidence-rate ratios, that is, \( e^{\beta_i} \) rather than \( \beta_i \). Standard errors and confidence intervals are similarly transformed. This option affects how results are displayed, not how they are estimated or stored. \text{irr} may be specified at estimation or when replaying previously estimated results.

\text{nocnsreport}; see [R] estimation options.

\text{display_options: noci, nopvalues, noomitted, vsquish, noemptycells, baselevels, allbaselevels, nofvlabel, fvwrap(#), fvwrapon(style), cformat(\%fmt), pformat(\%fmt), sformat(\%fmt), and nolstretch}; see [R] estimation options.

**Integration**

\text{intpoints(#)} specifies the number of integration points to use for integration by quadrature. The default is \text{intpoints(24)}; the maximum is \text{intpoints(128)}. Increasing this value improves the accuracy but also increases computation time. Computation time is roughly proportional to its value.

**Maximization**

\text{maximize_options: difficult, technique(algorithm_spec), iterate(#), [no]log, trace, gradient, showstep, hessian, showtolerance, tolerance(#), ltolerance(#), nrtolerance(#), nonrtolerance, and from(init_specs)}; see [R] maximize. These options are seldom used.

Setting the optimization type to \text{technique(bhhh)} resets the default \text{vcetype} to \text{vce(opg)}.

The following option is available with \text{etpoisson} but is not shown in the dialog box:

\text{coeflegend}; see [R] estimation options.
etpoisson estimates the parameters of a Poisson regression model that includes an endogenous binary-treatment variable. The dependent variable must be a Poisson distributed count. The parameters estimated by etpoisson can be used to estimate the average treatment effect (ATE) and average treatment effect on the treated (ATET).

We call the model fit by etpoisson an endogenous treatment-regression model, although it is also known as an endogenous binary-variable model or as an endogenous dummy-variable model. The endogenous treatment-regression model fit by etpoisson is a specific endogenous treatment-effects model; it uses a nonlinear model for the outcome and a constrained normal distribution to model the deviation from the conditional independence assumption imposed by the estimators implemented by teffects; see [TE] teffects intro. In treatment-effects jargon, the endogenous binary-variable model fit by etpoisson is a nonlinear potential-outcome model that allows for a specific correlation structure between the unobservables that affect the treatment and the unobservables that affect the potential outcomes. See [TE] etregress for an estimator that allows for a linear-outcome model and a similar model for the endogeneity of the treatment.

More formally, we have an equation for outcome $y_j$ and an equation for treatment $t_j$:  

$$E(y_j|x_j, t_j, \epsilon_j) = \exp(x_j \beta + \delta t_j + \epsilon_j)$$

$$t_j = \begin{cases} 
1, & w_j \gamma + u_j > 0 \\
0, & \text{otherwise} 
\end{cases}$$

The $x_j$ are the covariates used to model the outcome, $w_j$ are the covariates used to model treatment assignment, and error terms $\epsilon_j$ and $u_j$ are bivariate normal with mean 0 and covariance matrix

$$\begin{bmatrix}
\sigma^2 & \sigma \rho \\
\sigma \rho & 1
\end{bmatrix}$$

The covariates $x_j$ and $w_j$ are unrelated to the error terms; in other words, they are exogenous. Note that $y_j$ may be a count or continuous and nonnegative in this specification.

Terza (1998) describes the maximum likelihood estimator used in etpoisson. Terza (1998) categorized the model fit by etpoisson as an endogenous-switching model. These models involve a binary switch that is endogenous for the outcome. Calculation of the maximum likelihood estimate involves numeric approximation of integrals via Gauss–Hermite quadrature. This is computationally intensive, but the computational costs are reasonable on modern computers.
Basic example

Example 1

In this example, we observe a simulated random sample of 5,000 households. The outcome of interest is the number of trips taken by members of the household in the 24-hour period immediately prior to the interview time.

We have fictional household level data on the following variables: number of trips taken in the past 24 hours (trips), distance to the central business district from the household (cbd), distance from the household to a public transit node (ptn), an indicator of whether there is a full-time worker in the household (worker), an indicator of whether the examined period is on a weekend (weekend), the ratio of the household income to the median income of the census tract (realinc), and an indicator of car ownership (owncar). We suspect that unobservables that affect the number of trips also affect the household’s propensity to own a car.

We use etpoisson to estimate the parameters of a Poisson regression model for the number of trips with car ownership as an endogenous treatment. In subsequent examples, we will use margins (see [R] margins) to estimate the ATE and the ATET of car ownership on the number of trips taken by the household. In the etpoisson command below, we specify the vce(robust) option because we need to specify vce(unconditional) when we use margins later.
. use http://www.stata-press.com/data/r15/trip1
(Household trips, car ownership)
. etpoisson trips cbd ptn worker weekend,
   > treat(owncar = cbd ptn worker realinc) vce(robust)

Iteration 0: log pseudolikelihood = -14845.147 (not concave)
Iteration 1: log pseudolikelihood = -14562.997 (not concave)
Iteration 2: log pseudolikelihood = -13655.592 (not concave)
Iteration 3: log pseudolikelihood = -12847.219 (not concave)
Iteration 4: log pseudolikelihood = -12566.037
Iteration 5: log pseudolikelihood = -12440.974
Iteration 6: log pseudolikelihood = -12413.485
Iteration 7: log pseudolikelihood = -12412.699
Iteration 8: log pseudolikelihood = -12412.696
Iteration 9: log pseudolikelihood = -12412.696

Poisson regression with endogenous treatment
Number of obs = 5,000
(24 quadrature points)
Log pseudolikelihood = -12412.696

Robust

|                | Robust | Std. Err. | z     | P>|z| | [95% Conf. Interval] |
|----------------|--------|-----------|-------|------|----------------------|
| trips          |        |           |       |      |                      |
| cbd            | -.0100919 | .0020071 | -5.03 | 0.000 | -.0140258 to -.006158 |
| ptn            | -.0204038 | .0020289 | -10.06 | 0.000 | -.0243805 to -.0164272 |
| worker         | .692301  | .0548559 | 12.62 | 0.000 | .5847854 to .7998166  |
| weekend        | .0930517 | .034538  | 2.69  | 0.007 | .0253585 to .160745   |
| 1.owncar       | .5264713 | .1124157 | 4.68  | 0.000 | .3061406 to .746802   |
| _cons          | -.2340772 | .0810812 | -2.89 | 0.004 | -.3929934 to -.0751609 |
| owncar         |        |           |       |      |                      |
| cbd            | .007218  | .00239   | 3.02  | 0.003 | .0025337 to .0119023  |
| ptn            | .0084769 | .0024518 | 3.46  | 0.001 | .0063714 to .0132824  |
| worker         | .543643  | .0504267 | 10.78 | 0.000 | .4448085 to .6424774  |
| realinc        | .176479  | .0108746 | 16.23 | 0.000 | .1551652 to .1977928  |
| _cons          | -.4611246 | .0592161 | -7.79 | 0.000 | -.5771859 to -.3450633 |
| /athrho        | .5741169 | .0957832 | 5.99  | 0.000 | .3863852 to .7618486  |
| /lnsigma       | -.2182037 | .0256046 | -8.51 | 0.000 | -.2884338 to -.1679735 |
| rho            | .5183763 | .0700449 | 7.42  | 0.000 | .3862398 to .6421645  |
| sigma          | .8039617 | .020604  | 3.88  | 0.000 | .764576 to .8453762   |

Wald test of indep. eqns. (rho = 0): chi2(1) = 35.93  Prob > chi2 = 0.0000

The Wald test in the header is highly significant, indicating a good model fit. All the covariates are statistically significant, and the Wald test in the footer indicates that we can reject the null hypothesis of no correlation between the treatment errors and the outcome errors.

We can interpret the coefficient on 1.owncar as the logarithm of the ratio of the treatment potential-outcome mean to the control potential-outcome mean. The treatment variable did not interact with any of the outcome covariates, so the effect of each regressor is the same in the two regimes and will cancel from the ratio of potential-outcome means. This means the ratio is equivalent to the exponentiated coefficient on 1.owncar. After discussing the other parameters, we will use lincom to obtain this ratio. See [R] lincom for more information.

The estimated correlation between the treatment-assignment errors and the outcome errors is 0.518, indicating that unobservables that increase the number of trips tend to occur with unobservables that increase the chance of car ownership.
The results for the two ancillary parameters require explanation. \texttt{etpoisson} estimates the inverse hyperbolic tangent of $\rho$,

$$\text{atanh} \rho = \frac{1}{2} \ln \left( \frac{1 + \rho}{1 - \rho} \right)$$

and $\ln \sigma$ rather than $\rho$ and $\sigma$. For numerical stability during optimization, \texttt{etpoisson} does not directly estimate $\rho$ and $\sigma$.

Now we use \texttt{lincom} and the \texttt{eform} option to estimate the exponentiated coefficient for \texttt{1.owncar}. This corresponds to the ratio of the treatment regime potential-outcome mean to the control regime potential-outcome mean.

\[
\text{. lincom [trips]_b[1.owncar], eform ( 1) [trips]1.owncar = 0}
\]

\begin{table}[h]
\centering
\begin{tabular}{l|cccc}
\hline
 & exp(b) & Std. Err. & z & P>|z| & [95\% Conf. Interval] \\
\hline
(1) & 1.692948 & .1903139 & 4.68 & 0.000 & 1.358173 & 2.110241 \\
\hline
\end{tabular}
\end{table}

The potential-outcome mean for the treatment regime is 1.69 times the potential-outcome mean for the control regime. So the average number of trips in the treatment regime is over one and a half times the average number of trips in the control regime.

By interacting the treatment, \texttt{owncar}, with the other regressors, we could estimate different coefficients for the regressors in the treatment and control regimes. In the current model, there are no treatment interactions, so the coefficients are the same in each regime.

---

**Average treatment effect (ATE)**

The parameter estimates from \texttt{etpoisson} can be used by \texttt{margins} to estimate the ATE, the average difference of the treatment and control potential outcomes.

> **Example 2**

Continuing with example 1, we use \texttt{margins} to estimate the ATE of car ownership on the number of trips taken in a 24-hour period.

We can estimate the ATE of car ownership by using the potential-outcome means obtained through the \texttt{predict, pomean} command and the \texttt{margins} command; see \textit{Methods and formulas} below and \texttt{etpoisson postestimation} for more details about the use of \texttt{predict} after \texttt{etpoisson}.

The \texttt{r.} notation indicates that the potential-outcome means for treatment and control will be contrasted. We specify the \texttt{contrast(nowald)} option to suppress the Wald tests that \texttt{margins} displays by default for contrasts.

\[
\text{. margins r.owncar, vce(unconditional) contrast(nowald)}
\]

### Contrasts of predictive margins

<table>
<thead>
<tr>
<th>Expression</th>
<th>Potential-outcome mean, predict()</th>
</tr>
</thead>
<tbody>
<tr>
<td>owncar</td>
<td>1.058914 .1922909 .6820309 1.435797</td>
</tr>
</tbody>
</table>

---
The estimated ATE of car ownership on the number of trips taken is 1.06. The average household will take 1.06 more trips when it owns a car.

### Average treatment effect on the treated (ATET)

The parameter estimates from `etpoisson` can be used by `margins` to estimate the ATET, the average difference of the treatment and control potential outcomes in the treated population.

#### Example 3

Continuing with the previous example, we use `margins` to estimate the ATET of car ownership on the number of trips taken in a 24-hour period.

We can estimate the ATET of car ownership by using the conditional treatment effect (conditional on exogenous covariates and treatment level) obtained through the `predict, cte` command and the `margins` command; see *Methods and formulas* below and [TE] `etpoisson postestimation` for more details about the use of `predict` after `etpoisson`.

We estimate the ATET with `margins`. We specify `cte` in the `predict()` option. Estimation is restricted to the treated subpopulation by specifying `owncar` in the `subpop()` option.

```
. margins, predict(cte) vce(unconditional) subpop(owncar)
Predictive margins                     Number of obs  =   5,000
                                               Subpop. no. obs  =   3,504
Expression : Conditional treatment effect, predict(cte)

<table>
<thead>
<tr>
<th>Unconditional</th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Margin</td>
<td>Std. Err.</td>
<td>z</td>
</tr>
<tr>
<td>_cons</td>
<td>1.251971</td>
<td>.2059201</td>
<td>6.08</td>
</tr>
</tbody>
</table>
```

The estimated ATET of car ownership on the number of trips taken is 1.25. Thus the average household in the treated population will take 1.25 more trips than it would if it did not own a car. This number is higher than the ATE. In this model, the ATE and ATET will only coincide when there is no correlation between the treatment errors and outcome errors and the exogenous covariates $x$ have the same distribution in the general population and treated subpopulation. See *Methods and formulas* for more details.
etpoisson — Poisson regression with endogenous treatment effects

Stored results

etpoisson stores the following in e():

Scalars

- `e(N)` number of observations
- `e(k)` number of parameters
- `e(k_eq)` number of equations in `e(b)`
- `e(k_eq_model)` number of equations in overall model test
- `e(k_aux)` number of auxiliary parameters
- `e(k_df)` number of dependent variables
- `e(df_m)` model degrees of freedom
- `e(ll)` log likelihood
- `e(N_clust)` number of clusters
- `e(chi2)` $\chi^2$
- `e(chi2_c)` $\chi^2$ for comparison, $\rho=0$ test
- `e(n_quad)` number of quadrature points
- `e(p)` $p$-value for model test
- `e(p_c)` $p$-value for comparison test
- `e(ranking)` rank of `e(V)`
- `e(ic)` number of iterations
- `e(rc)` return code
- `e(converged)` 1 if converged, 0 otherwise

Macros

- `e(cmd)` etpoisson
- `e(cmdline)` command as typed
- `e(depvar)` name of dependent variable
- `e(wtype)` weight type
- `e(wexp)` weight expression
- `e(title)` title in estimation output
- `e(title2)` secondary title in estimation output
- `e(clustvar)` name of cluster variable
- `e(offset1)` offset for regression equation
- `e(offset2)` offset for treatment equation
- `e(chi2type)` Wald; type of model $\chi^2$ test
- `e(chi2_ct)` Wald; type of comparison $\chi^2$ test
- `e(vce)` vcetype specified in `vce()`
- `e(vcetype)` title used to label Std. Err.
- `e(opt)` type of optimization
- `e(which)` max or min; whether optimizer is to perform maximization or minimization
- `e(ml_method)` type of ml method
- `e(user)` name of likelihood-evaluator program
- `e(technique)` maximization technique
- `e(properties)` b V
- `e(predict)` program used to implement predict
- `e(marginsok)` predictions allowed by margins
- `e(asbalanced)` factor variables fvset as asbalanced
- `e(asobserved)` factor variables fvset as asobserved

Matrices

- `e(b)` coefficient vector
- `e(Cns)` constraints matrix
- `e(ilog)` iteration log (up to 20 iterations)
- `e(gradient)` gradient vector
- `e(V)` variance–covariance matrix of the estimators
- `e(V_modelbased)` model-based variance

Functions

- `e(sample)` marks estimation sample
Methods and formulas

Terza (1998) derives the maximum likelihood estimator implemented here. We provide some details of the derivation and then explain how the model is nested in a more general potential-outcomes model. Then the ATE and ATET are derived.

Let \( x_j \) be the covariates used to model the outcome, and let \( w_j \) be the covariates used to model treatment assignment. Define \( z_j = (w_j, x_j) \). The vector \( z_j \) contains all the exogenous covariates in the model. When offsets \( o \beta_j \) are used in the outcome variable equation, the following formulas apply with \( x_j \beta \) changed to \( x_j \beta + o \beta_j \). Similarly, when offsets \( o \gamma_j \) are used in the endogenous treatment equation, the following formulas apply with \( w_j \gamma \) changed to \( w_j \gamma + o \gamma_j \). If offsets are used in either equation, they are included in the vector of exogenous covariates \( z_j \).

For treatment \( t_j \), \( z_j \), and \( \epsilon_j \), outcome \( y_j \) of this model has conditional mean

\[
E(y_j|x_j, t_j, \epsilon_j) = \exp(x_j \beta + \delta t_j + \epsilon_j)
\]

The probability density function of \( y_j \) for this model, conditioned on treatment \( t_j \), \( z_j \), and \( \epsilon_j \), is given by

\[
f(y_j|z_j, t_j, \epsilon_j) = \exp\{-\exp(x_j \beta + \delta t_j + \epsilon_j)\} \frac{\exp(x_j \beta + \delta t_j + \epsilon_j)^{y_j}}{y_j!}
\]

The treatment \( t_j \) is determined by

\[
t_j = \begin{cases} 
1, & \text{if } w_j \gamma + u_j > 0 \\
0, & \text{otherwise}
\end{cases}
\]

The error terms \( \epsilon_j \) and \( u_j \) are bivariate normal with mean zero and covariance matrix

\[
\begin{bmatrix} 
\sigma^2 & \sigma \rho \\
\sigma \rho & 1
\end{bmatrix}
\]

Conditional on \( \epsilon_j \), \( u_j \) is normal with mean \( \epsilon_j \rho/\sigma \) and variance \((1 - \rho^2)\); thus we obtain the following conditional probability density for \( t_j \):

\[
Pr(t_j|z_j, \epsilon_j) = t_j \Phi \left( \frac{w_j \gamma + (\rho/\sigma) \epsilon_j}{\sqrt{1 - \rho^2}} \right) + (1 - t_j) \left[ 1 - \Phi \left( \frac{w_j \gamma + (\rho/\sigma) \epsilon_j}{\sqrt{1 - \rho^2}} \right) \right]
\]

\( \Phi \) denotes the standard normal cumulative distribution function. This leads to the following joint density of \( y_j \), \( t_j \), and \( \epsilon_j \):

\[
f(y_j, t_j, \epsilon_j|z_j) = f(y_j|z_j, t_j, \epsilon_j)P(t_j|z_j, \epsilon_j)f(\epsilon_j)
\]

The density of \( y_j \) and \( t_j \), conditioned on \( z_j \), is obtained by integrating the above with respect to \( \epsilon_j \). Recall that \( \epsilon_j \) is normal with mean 0 and variance \( \sigma^2 \).

\[
f(y_j, t_j|z_j) = \int_{-\infty}^{\infty} f(y_j|z_j, t_j, \epsilon_j)P(t_j|z_j, \epsilon_j)f(\epsilon_j)\frac{1}{\sigma \sqrt{2\pi}} \exp \left\{ -\left( \frac{\epsilon_j}{\sigma \sqrt{2}} \right)^2 \right\} d\epsilon_j
\]
\( f(y_j, t_j|z_j) \) cannot be evaluated in a closed form. We change the variable of integration from \( \epsilon_j \) to \( \eta_j = \epsilon_j/(\sigma \sqrt{2}) \), which yields

\[
f(y_j, t_j|z_j) = \frac{1}{\sqrt{\pi}} \int_{-\infty}^{\infty} f(y_j|z_j, t_j, \sqrt{2}\sigma \eta_j) P(t_j|z_j, \sqrt{2}\sigma \eta_j) \exp(-\eta_j^2) \, d\eta_j
\]

We approximate this integral by Gauss–Hermite quadrature. Observing a sample of \( t_j, y_j, \) and \( z_j \), we calculate the log likelihood as the following:

\[
\ln L = \sum_{j=1}^{n} w_j \ln \{f(y_j, t_j|z_j)\}
\]

The \( w_j \) terms denote optional weights.

In the maximum likelihood estimation, \( \sigma \) and \( \rho \) are not directly estimated. Directly estimated are \( \ln \sigma \) and \( \text{atanh} \rho \):

\[
\text{atanh} \rho = \frac{1}{2} \ln \left( \frac{1 + \rho}{1 - \rho} \right)
\]

Now we present formulas for the ATE and ATET. First, we nest the endogenous-treatment Poisson regression model in a potential-outcome model. A potential-outcome model specifies what each individual would obtain in each treatment level.

A potential-outcome model that nests the endogenous-treatment Poisson regression fit by \texttt{etpoisson} is:

\[
\begin{align*}
E(y_{0j}|x_j, \epsilon_j) &= \exp(x_j \beta_0 + \epsilon_{0j}) \\
E(y_{1j}|x_j, \epsilon_j) &= \exp(x_j \beta_1 + \epsilon_{1j}) \\
t_j &= \begin{cases} 1, & \text{if } w_j \gamma + u_j > 0 \\ 0, & \text{otherwise} \end{cases}
\end{align*}
\]

where \( y_{0j} \) is the outcome that person \( j \) obtains if person \( j \) selects treatment 0, and \( y_{1j} \) is the outcome that person \( j \) obtains if person \( j \) selects treatment 1. This formulation allows differing coefficients for the control (\( \beta_0 \)) and treatment (\( \beta_1 \)) regimes. The constant intercept for the control group is \( \beta_{00} \). The constant intercept for the treatment group is \( \beta_{11} = \beta_{00} + \delta \), where \( \delta \) is the coefficient for treatment \( t_j \) in the outcome (1). The remaining notation was defined above.

We may allow other coefficients to differ across regimes in the outcome (1) by adding interactions between the treatment \( t_j \) and covariates \( x_j \) to the model. To be concise, we use two coefficient vectors \( \beta_0 \) and \( \beta_1 \) here rather than a single coefficient vector with interactions between the treatment \( t_j \) and covariates \( x_j \). The two formulations are equivalent.

We never observe both \( y_{0j} \) and \( y_{1j} \), only one or the other. We observe

\[
y_j = t_j y_{1j} + (1-t_j) y_{0j}
\]

The vector of error terms \( (\epsilon_{0j}, \epsilon_{1j}, u_j)' \) comes from a mean zero trivariate normal distribution with covariance matrix

\[
\begin{bmatrix}
\sigma^2 & \theta & \sigma \rho \\
\theta & \sigma^2 & \sigma \rho \\
\sigma \rho & \sigma \rho & 1
\end{bmatrix}
\]
The parameters $\sigma$ and $\rho$ were discussed earlier. The parameter $\theta$ is the covariance between the two potential outcomes. We cannot estimate $\theta$ because we have no observations in which an individual is observed in both potential outcomes. Fortunately, $\theta$ is not required for the calculations that we present.

The ATE is the difference in means of the potential outcomes. The mean of each potential outcome accounts for each individual’s contribution, regardless of whether that individual selects that treatment level.

The conditional means of the potential outcomes $y_{tj}$, $t \in (0, 1)$ for exogenous covariates $z_j$ are

$$E(y_{tj}|z_j) = \exp \left( x_j \beta_t + \frac{\sigma^2}{2} \right)$$

We can see that when the coefficients are the same across the regimes, the ratio of potential-outcome means will be equal to $\exp(\delta)$; this is true of the conditional and marginal potential-outcome means.

The difference in potential-outcome means or treatment effect at exogenous covariates $z_j$ is

$$E(y_{1j} - y_{0j}|z_j) = \{ \exp (x_j \beta_1) - \exp (x_j \beta_0) \} \exp \left( \frac{\sigma^2}{2} \right)$$

By the law of iterated expectations, the ATE is

$$E(y_{1j} - y_{0j}) = E\{E(y_{1j} - y_{0j}|z_j)\}$$

$$= E \left[ \{ \exp (x_j \beta_1) - \exp (x_j \beta_0) \} \exp \left( \frac{\sigma^2}{2} \right) \right]$$

This expectation can be estimated as a predictive margin.

Now we will derive an expression for the ATET.

The conditional means of the potential outcomes $y_{tj}$, $t \in (0, 1)$ for exogenous covariates $z_j$ and treatment $t_j$ are

$$E(y_{tj}|z_j, t_j) = \exp \left( x_j \beta_t + \frac{\sigma^2}{2} \right) \left\{ \frac{\Phi (\rho \sigma + w_j \gamma)}{\Phi (w_j \gamma)} \right\}^{t_j} \left\{ \frac{1 - \Phi (\rho \sigma + w_j \gamma)}{1 - \Phi (w_j \gamma)} \right\}^{1-t_j}$$

Rather than the conditional potential-outcome means, the conditional mean of the observed outcome may be of interest. The conditional mean of the observed outcome $y_j$ for endogenous treatment indicator $t_j$ and exogenous covariates $z_j$ is given by

$$E(y_j|z_j, t_j) = t_j \exp \left( x_j \beta_1 + \frac{\sigma^2}{2} \right) \frac{\Phi (\rho \sigma + w_j \gamma)}{\Phi (w_j \gamma)}$$

$$+ (1 - t_j) \exp \left( x_j \beta_0 + \frac{\sigma^2}{2} \right) \frac{1 - \Phi (\rho \sigma + w_j \gamma)}{1 - \Phi (w_j \gamma)}$$
The treatment effect at exogenous covariates $z_j$ and treatment $t_j$ is

$$E(y_{1j} - y_{0j}|z_j, t_j) =$$

$$\{ \exp(x_j \beta_1) - \exp(x_j \beta_0) \} \exp \left( \frac{\sigma^2}{2} \right) \left\{ \frac{\Phi(\rho \sigma + w'_j \gamma)}{\Phi(w_j \gamma)} \right\}^{t_j} \left\{ \frac{1 - \Phi(\rho \sigma + w_j \gamma)}{1 - \Phi(w'_j \gamma)} \right\}^{1-t_j}$$

By the law of iterated expectations, the ATET is

$$E(y_{1j} - y_{0j}|t_j = 1) = E\{E(y_{1j} - y_{0j}|z_j, t_j = 1)|t_j = 1\}$$

$$= E \left[ \{ \exp(x_j \beta_1) - \exp(x_j \beta_0) \} \exp \left( \frac{\sigma^2}{2} \right) \frac{\Phi(\rho \sigma + w_j \gamma)}{\Phi(w_j \gamma)} \right|_{t_j = 1}$$

This can be estimated as a predictive margin on the treated subpopulation.

We note that when $\rho = 0$, the correction factor involving $\Phi$ will disappear from the ATET. Then the ATE and ATET will be equivalent if the distribution of $x_j$ under the treated population is identical to the distribution over the entire population.

The probability of $y_j$ conditional on $t_j$ and $z_j$ is

$$\Pr(y_j = n|z_j, t_j) = \frac{f(y_j = n, t_j|z_j)}{\Phi(w_j \gamma)^{t_j} \Phi(-w_j \gamma)^{1-t_j}}$$

As discussed earlier, we approximate $f(y_j, t_j|z_j)$ using Gauss–Hermite quadrature.

References

Also see
[TE] etpoisson postestimation — Postestimation tools for etpoisson
[TE] etregress — Linear regression with endogenous treatment effects
[R] heckpoisson — Poisson regression with sample selection
[R] ivpoisson — Poisson model with continuous endogenous covariates
[R] ivprobit — Probit model with continuous endogenous covariates
[R] ivregress — Single-equation instrumental-variables regression
[R] ivtobit — Tobit model with continuous endogenous covariates
[R] poisson — Poisson regression
[SVY] svy estimation — Estimation commands for survey data
[U] 20 Estimation and postestimation commands
Postestimation commands

The following standard postestimation commands are available after `etpoisson`:

<table>
<thead>
<tr>
<th>Command</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>contrast</td>
<td>contrasts and ANOVA-style joint tests of estimates</td>
</tr>
<tr>
<td>estat ic</td>
<td>Akaike’s and Schwarz’s Bayesian information criteria (AIC and BIC)</td>
</tr>
<tr>
<td>estat summarize</td>
<td>summary statistics for the estimation sample</td>
</tr>
<tr>
<td>estat vce</td>
<td>variance–covariance matrix of the estimators (VCE)</td>
</tr>
<tr>
<td>estat (svy)</td>
<td>postestimation statistics for survey data</td>
</tr>
<tr>
<td>estimates</td>
<td>cataloging estimation results</td>
</tr>
<tr>
<td>*hausman</td>
<td>Hausman’s specification test</td>
</tr>
<tr>
<td>lincom</td>
<td>point estimates, standard errors, testing, and inference for linear combinations of coefficients</td>
</tr>
<tr>
<td>*lrtest</td>
<td>likelihood-ratio test</td>
</tr>
<tr>
<td>margins</td>
<td>marginal means, predictive margins, marginal effects, and average marginal effects</td>
</tr>
<tr>
<td>marginsplot</td>
<td>graph the results from margins (profile plots, interaction plots, etc.)</td>
</tr>
<tr>
<td>nlcom</td>
<td>point estimates, standard errors, testing, and inference for nonlinear combinations of coefficients</td>
</tr>
<tr>
<td>predict</td>
<td>predictions, probabilities, and treatment effects</td>
</tr>
<tr>
<td>predictnl</td>
<td>point estimates, standard errors, testing, and inference for generalized predictions</td>
</tr>
<tr>
<td>pwcompare</td>
<td>pairwise comparisons of estimates</td>
</tr>
<tr>
<td>suest</td>
<td>seemingly unrelated estimation</td>
</tr>
<tr>
<td>test</td>
<td>Wald tests of simple and composite linear hypotheses</td>
</tr>
<tr>
<td>testnl</td>
<td>Wald tests of nonlinear hypotheses</td>
</tr>
</tbody>
</table>

* `hausman` and `lrtest` are not appropriate with `svy` estimation results.
predict

Description for predict

`predict` creates a new variable containing predictions such as counts, conditional treatment effects, probabilities, and linear predictions.

Menu for predict

Statistics > Postestimation

Syntax for predict

```
predict [type] newvar [if] [in] [, statistic nooffset]
```

```
predict [type] { stub* | newvar_reg newvar_treat newvar_athrho newvar_lnsigma } [if] [in], scores
```

<table>
<thead>
<tr>
<th>statistic</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Main</strong></td>
<td></td>
</tr>
<tr>
<td><code>pomean</code></td>
<td>potential-outcome mean (the predicted count); the default</td>
</tr>
<tr>
<td><code>omean</code></td>
<td>observed-outcome mean (the predicted count)</td>
</tr>
<tr>
<td><code>cte</code></td>
<td>conditional treatment effect at treatment level</td>
</tr>
<tr>
<td><code>pr(n)</code></td>
<td>probability ( \text{Pr}(y_j = n) )</td>
</tr>
<tr>
<td><code>pr(a,b)</code></td>
<td>probability ( \text{Pr}(a \leq y_j \leq b) )</td>
</tr>
<tr>
<td><code>xb</code></td>
<td>linear prediction</td>
</tr>
<tr>
<td><code>xbtreat</code></td>
<td>linear prediction for treatment equation</td>
</tr>
</tbody>
</table>

These statistics are available both in and out of sample; type `predict ... if e(sample) ... if wanted` only for the estimation sample.

Options for predict

- `pomean`, the default, calculates the potential-outcome mean.
- `omean` calculates the observed-outcome mean.
- `cte` calculates the treatment effect, the difference of potential-outcome means, conditioned on treatment level.
pr(n) calculates the probability Pr(yj = n), where n is a nonnegative integer that may be specified as a number or a variable.

pr(a,b) calculates the probability Pr(a ≤ yj ≤ b), where a and b are nonnegative integers that may be specified as numbers or variables;

b missing (b ≥ .) means +∞;
pr(20,.) calculates Pr(yj ≥ 20);
pr(20,b) calculates Pr(yj ≥ 20) in observations for which b ≥ . and calculates Pr(20 ≤ yj ≤ b) elsewhere.

pr(.,b) produces a syntax error. A missing value in an observation of the variable a causes a missing value in that observation for pr(a,b).

xb calculates the linear prediction for the dependent count variable, which is xjβ if neither offset() nor exposure() was specified; xjβ + offsetjβ if offset() was specified; or xjβ + ln(exposurej) if exposure() was specified.

xbtreat calculates the linear prediction for the endogenous treatment equation, which is wjγ if offset() was not specified in treat() and wjγ + offsetjα if offset() was specified in treat().

nooffset is relevant only if you specified offset() or exposure() when you fit the model. It modifies the calculations made by predict so that they ignore the offset or exposure variable. nooffset removes the offset from calculations involving both the treat() equation and the dependent count variable.

scores calculates equation-level score variables.

The first new variable will contain $\partial \ln L / \partial (x_j \beta)$.

The second new variable will contain $\partial \ln L / \partial (w_j \gamma)$.

The third new variable will contain $\partial \ln L / \partial \tanh \rho$.

The fourth new variable will contain $\partial \ln L / \partial \ln \sigma$. 
margins

Description for margins

margins estimates margins of response for counts, conditional treatment effects, probabilities, and linear predictions.

Menu for margins

Statistics > Postestimation

Syntax for margins

margins [marginlist] [, options]

margins [marginlist], predict(statistic ...) [predict(statistic ...) ...] [options]

<table>
<thead>
<tr>
<th>statistic</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>pomean</td>
<td>potential-outcome mean (the predicted count); the default</td>
</tr>
<tr>
<td>omean</td>
<td>observed-outcome mean (the predicted count)</td>
</tr>
<tr>
<td>cte</td>
<td>conditional treatment effect at treatment level</td>
</tr>
<tr>
<td>pr((n))</td>
<td>probability ( \Pr(y_j = n) )</td>
</tr>
<tr>
<td>pr(a, b)</td>
<td>probability ( \Pr(a \leq y_j \leq b) )</td>
</tr>
<tr>
<td>xb</td>
<td>linear prediction</td>
</tr>
<tr>
<td>xbtreat</td>
<td>linear prediction for treatment equation</td>
</tr>
</tbody>
</table>

Statistics not allowed with margins are functions of stochastic quantities other than \( \mathbf{e}(\mathbf{b}) \).

For the full syntax, see [R] margins.

Remarks and examples

The average treatment effect (ATE) and the average treatment effect on the treated (ATET) are the parameters most frequently estimated by postestimation techniques after etpoisson.

You can use the margins command (see [R] margins) after etpoisson to estimate the ATE or ATET. See example 2 of [TE] etpoisson for an example of ATE estimation. See example 3 of [TE] etpoisson for an example of ATET estimation.

See example 1 of [TE] etpoisson for an example using lincom after etpoisson.

Methods and formulas

See Methods and formulas of [TE] etpoisson for details.

Also see

[TE] etpoisson — Poisson regression with endogenous treatment effects

[U] 20 Estimation and postestimation commands
etregress estimates an average treatment effect (ATE) and the other parameters of a linear regression model augmented with an endogenous binary-treatment variable. Estimation is by full maximum likelihood, a two-step consistent estimator, or a control-function estimator.

In addition to the ATE, etregress can be used to estimate the average treatment effect on the treated (ATET) when the outcome may not be conditionally independent of the treatment.

etreg is a synonym for etregress.

Quick start

ATE and ATET from a linear regression model of $y$ on $x$ and endogenous binary treatment $treat$ modeled by $x$ and $w$

etregress $y$ $x$, treat($treat = x$ $w$)

As above, but use a control-function estimator

etregress $y$ $x$, treat($treat = x$ $w$) cfunction

With robust standard errors

etregress $y$ $x$, treat($treat = x$ $w$) vce(robust)

Add the interaction between $treat$ and continuous covariate $x$ using factor variables

etregress $y$ $x$ i.treat#c.x, treat($treat = x$ $w$) vce(robust)

ATE after etregress with the required vce(robust) option and endogenous treatment interaction terms

margins r.treat, vce(unconditional)

As above, but calculate ATET

margins, vce(unconditional) predict(cte) subpop(if treat==1)

Menu

Statistics > Treatment effects > Endogenous treatment > Maximum likelihood estimator > Continuous outcomes
**Syntax**

**Basic syntax**

```
etregress depvar [ indepvars ], treat(depvar_t = indepvar_{t}) [ twostep | cfunction ]
```

**Full syntax for maximum likelihood estimates only**

```
etregress depvar [ indepvars ] [ if ] [ in ] [ weight ],
   treat(depvar_t = indepvar_{t} [, noconstant]) [ etregress_ml_options ]
```

**Full syntax for two-step consistent estimates only**

```
etregress depvar [ indepvars ] [ if ] [ in ],
   treat(depvar_t = indepvar_{t} [, noconstant]) twostep [ etregress_ts_options ]
```

**Full syntax for control-function estimates only**

```
etregress depvar [ indepvars ] [ if ] [ in ],
   treat(depvar_t = indepvar_{t} [, noconstant]) cfunction [ etregress_cf_options ]
```
### etregress_ml_options

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>treat()</strong></td>
</tr>
<tr>
<td>noconstant</td>
</tr>
<tr>
<td>poutcomes</td>
</tr>
<tr>
<td>constraints(constants)</td>
</tr>
<tr>
<td>collinear</td>
</tr>
</tbody>
</table>

### SE/Robust

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>vce(vcetype)</td>
</tr>
</tbody>
</table>

### Reporting

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>level(#)</td>
</tr>
<tr>
<td>first</td>
</tr>
<tr>
<td>hazard(newvar)</td>
</tr>
<tr>
<td>lrmodel</td>
</tr>
<tr>
<td>noconsreport</td>
</tr>
<tr>
<td>display_options</td>
</tr>
</tbody>
</table>

### Maximization

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>maximize_options</td>
</tr>
<tr>
<td>coeflegend</td>
</tr>
</tbody>
</table>

* `treat(depvar_t = indepvars_t [, noconstant])` is required.

### etregress_ts_options

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>treat()</strong></td>
</tr>
<tr>
<td><strong>twostep</strong></td>
</tr>
<tr>
<td>noconstant</td>
</tr>
</tbody>
</table>

### SE

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>vce(vcetype)</td>
</tr>
</tbody>
</table>

### Reporting

<table>
<thead>
<tr>
<th>Description</th>
</tr>
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<tbody>
<tr>
<td>level(#)</td>
</tr>
<tr>
<td>first</td>
</tr>
<tr>
<td>hazard(newvar)</td>
</tr>
<tr>
<td>display_options</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>coeflegend</td>
</tr>
</tbody>
</table>

* `treat(depvar_t = indepvars_t [, noconstant])` and `twostep` are required.
**etregress — Linear regression with endogenous treatment effects**

<table>
<thead>
<tr>
<th>etregress_cf_options</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>*treat()</td>
<td>equation for treatment effects</td>
</tr>
<tr>
<td>*cfunction</td>
<td>produce control-function estimate</td>
</tr>
<tr>
<td>noconstant</td>
<td>suppress constant term</td>
</tr>
<tr>
<td>poutcomes</td>
<td>use potential-outcome model with separate treatment and control group variance and correlation parameters</td>
</tr>
<tr>
<td>SE</td>
<td>vctype may be robust, bootstrap, or jackknife</td>
</tr>
<tr>
<td>vce(vcetype)</td>
<td>vcetype may be robust, bootstrap, or jackknife</td>
</tr>
<tr>
<td>Reporting</td>
<td></td>
</tr>
<tr>
<td><em>level(#)</em></td>
<td>set confidence level; default is level(95)</td>
</tr>
<tr>
<td><em>first</em></td>
<td>report first-step probit estimates</td>
</tr>
<tr>
<td>_hazard(newvar)</td>
<td>create newvar containing hazard from treatment equation</td>
</tr>
<tr>
<td><em>display_options</em></td>
<td>control columns and column formats, row spacing, line width, display of omitted variables and base and empty cells, and factor-variable labeling</td>
</tr>
<tr>
<td>Maximization</td>
<td></td>
</tr>
<tr>
<td><em>maximize_options</em></td>
<td>control the maximization process; seldom used</td>
</tr>
<tr>
<td><em>coeflegend</em></td>
<td>display legend instead of statistics</td>
</tr>
</tbody>
</table>

*treat(depvar = indepvars_t [ , noconstant ] ) and cfunction are required.

indepvars and indepvars_t may contain factor variables; see [U] 11.4.3 Factor variables.

depvar, indepvars, depvar_t, and indepvars_t may contain time-series operators; see [U] 11.4.4 Time-series varlists.
twostep, cfunction, vce(), first, hazard(), lrmodel, and weights are not allowed with the svy prefix; see [SVY] svy.

See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.
Options for maximum likelihood estimates

treat(\(depvar_t = indepvars_t\[, \text{noconstant}\]\) specifies the variables and options for the treatment equation. It is an integral part of specifying a treatment-effects model and is required.

\text{noconstant}; see \[R\] estimation options.

poutcomes specifies that a potential-outcome model with separate variance and correlation parameters for each of the treatment and control groups be used.

\text{constraints}(\text{constraints}), \text{collinear}; see \[R\] estimation options.

\text{vce}(\text{vcetype}) specifies the type of standard error reported, which includes types that are derived from asymptotic theory (\text{oim}, \text{opg}), that are robust to some kinds of misspecification (\text{robust}), that allow for intragroup correlation (\text{cluster clustvar}), and that use bootstrap or jackknife methods (\text{bootstrap}, \text{jackknife}); see \[R\] \text{vce option}.

\text{level}(#)\); see \[R\] estimation options.

\text{first} specifies that the first-step probit estimates of the treatment equation be displayed before estimation.

\text{hazard}(\text{newvar}) will create a new variable containing the hazard from the treatment equation. The hazard is computed from the estimated parameters of the treatment equation.

\text{lrmodel}, \text{nocnsreport}; see \[R\] estimation options.

\text{display_options}: \text{noci}, \text{nopvalues}, \text{noomitted}, \text{vsquish}, \text{noemptycells}, \text{baselevels}, \text{allbaselevels}, \text{nofvlabel}, \text{fwrap(#)}, \text{fwrapon(style)}, \text{cformat(\%fmt)}, \text{pformat(\%fmt)}, \text{sformat(\%fmt)}, \text{and nolstretch}; see \[R\] estimation options.

\text{maximize_options}: \text{difficult}, \text{technique(algorithm_spec)}, \text{iterate(#)}, [\text{no}] \text{log}, \text{trace}, \text{gradient}, \text{showstep}, \text{hessian}, \text{showtolerance}, \text{tolerance(#)}, \text{ltolerance(#)}, \text{nrtolerance(#)}, \text{nonrtolerance}, \text{and from(init_specs)}; see \[R\] \text{maximize}. These options are seldom used.

Setting the optimization type to \text{technique(bhhh)} resets the default \text{vcetype} to \text{vce(opg)}.

The following option is available with \text{etregress} but is not shown in the dialog box:

\text{coeflegend}; see \[R\] estimation options.
Options for two-step consistent estimates

Model

\texttt{treat(\textit{depvar}_t = \textit{indepvars}_t[, \text{noconstant}])} specifies the variables and options for the treatment equation. It is an integral part of specifying a treatment-effects model and is required.

two\text{step} specifies that two-step consistent estimates of the parameters, standard errors, and covariance matrix be produced, instead of the default maximum likelihood estimates.

\text{noconstant}; see [R] estimation options.

SE

\texttt{vce(\textit{vcetype})} specifies the type of standard error reported, which includes types that are derived from asymptotic theory (conventional) and that use bootstrap or jackknife methods (bootstrap, jackknife); see [R] vce_option.

\texttt{vce(\text{conventional})}, the default, uses the conventionally derived variance estimator for the two-step estimator of the treatment-effects model.

Reporting

\texttt{level(\#)}; see [R] estimation options.

\texttt{first} specifies that the first-step probit estimates of the treatment equation be displayed before estimation.

\texttt{hazard(\textit{newvar})} will create a new variable containing the hazard from the treatment equation. The hazard is computed from the estimated parameters of the treatment equation.

\textit{display_options}: noci, nopvalues, noomitted, vsquish, noemptycells, baselevels, allbaselevels, noreflabel, fvwrap(\#), fvwrapon(style), cformat(\%fmt), pformat(\%fmt), sformat(\%fmt), and nolstretch; see [R] estimation options.

The following option is available with etregress but is not shown in the dialog box:

\texttt{coeflegend}; see [R] estimation options.

Options for control-function estimates

Model

\texttt{treat(\textit{depvar}_t = \textit{indepvars}_t[, \text{noconstant}])} specifies the variables and options for the treatment equation. It is an integral part of specifying a treatment-effects model and is required.

cfunction specifies that control-function estimates of the parameters, standard errors, and covariance matrix be produced instead of the default maximum likelihood estimates. cfunction is required.

\text{noconstant}; see [R] estimation options.

poutcomes specifies that a potential-outcome model with separate variance and correlation parameters for each of the treatment and control groups be used.

SE

\texttt{vce(\textit{vcetype})} specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (robust) and that use bootstrap or jackknife methods (bootstrap, jackknife); see [R] vce_option.
Reporting

level(#); see [R] estimation options.

first specifies that the first-step probit estimates of the treatment equation be displayed before estimation.

hazard(newvar) will create a new variable containing the hazard from the treatment equation. The hazard is computed from the estimated parameters of the treatment equation.

display_options: noci, nopvalues, noomitted, vsquish, noemptycells, baselevels, allbaselevels, nofvlabel, fvwrap(#), fvwrapon(style), cformat(%,fmt), pformat(%,fmt), sformat(%,fmt), and nolstretch; see [R] estimation options.

Maximization

maximize_options: iterate(#), [no]log, and from(init_specs); see [R] maximize. These options are seldom used.

init_specs is one of

matname [, skip copy]

# [# ...] copy

The following option is available with etregress but is not shown in the dialog box:

coefflegend; see [R] estimation options.

Remarks and examples

Remarks are presented under the following headings:

Overview
Basic examples
Average treatment effect (ATE)
Average treatment effect on the treated (ATET)

Overview

etregress estimates an ATE and the other parameters of a linear regression model that also includes an endogenous binary-treatment variable. In addition to the ATE, the parameters estimated by etregress can be used to estimate the ATET when the outcome is not conditionally independent of the treatment.

We call the model fit by etregress an endogenous treatment-regression model, although it is also known as an endogenous binary-variable model or as an endogenous dummy-variable model. The endogenous treatment-regression model is a specific endogenous treatment-effects model; it uses a linear model for the outcome and a normal distribution to model the deviation from the conditional independence assumption imposed by the estimators implemented in teffects; see [TE] teffects intro. In treatment-effects jargon, the endogenous binary-variable model is a linear potential-outcome model that allows for a specific correlation structure between the unobservables that affect the treatment and the unobservables that affect the potential outcomes. See [TE] etpoisson for an estimator that allows for a nonlinear outcome model and a similar model for the endogeneity of the treatment.
Heckman (1976, 1978) brought this model into the modern literature. Maddala (1983) derives the maximum likelihood and the control-function (CF) estimators of the model. Maddala (1983) also reviews some empirical applications and describes it as an endogenous-switching model. Barnow, Cain, and Goldberger (1981) provide another useful derivation of this model. They concentrate on deriving the conditions for which the self-selection bias of the simple OLS estimator of the treatment effect, $\delta$, is nonzero and of a specific sign. Cameron and Trivedi (2005, sec. 16.7 and 25.3.4) and Wooldridge (2010, sec. 21.4.1) discuss the endogenous binary-variable model as an endogenous treatment-effects model and link it to recent work.

*etregress* performs CF estimation in one step by using the generalized method of moments (GMM) with stacked moments. See Newey (1984) and Wooldridge (2010, sec. 14.2) for a description of this technique. Many econometric and statistical models can be expressed as conditions on the population moments. The parameter estimates produced by GMM estimators make the sample-moment conditions as true as possible given the data. See [R] gmm for further information on GMM estimation and how Stata performs it. Two-step CF estimation is also supported by *etregress*.

Formally, the endogenous treatment-regression model is composed of an equation for the outcome $y_j$ and an equation for the endogenous treatment $t_j$. The variables $x_j$ are used to model the outcome. When there are no interactions between $t_j$ and $x_j$, we have

$$y_j = x_j \beta + \delta t_j + \epsilon_j$$

$$t_j = \begin{cases} 1, & \text{if } w_j \gamma + u_j > 0 \\ 0, & \text{otherwise} \end{cases}$$

where $w_j$ are the covariates used to model treatment assignment, and the error terms $\epsilon_j$ and $u_j$ are bivariate normal with mean zero and covariance matrix

$$\begin{bmatrix} \sigma^2 & \rho \sigma \\ \rho \sigma & 1 \end{bmatrix}$$

The covariates $x_j$ and $w_j$ are unrelated to the error terms; in other words, they are exogenous. We call this the constrained model because the variance and correlation parameters are identical across the treatment and control groups.

This model can be generalized to a potential-outcome model with separate variance and correlation parameters for the treatment and control groups. The generalized model is

$$y_{0j} = x_j \beta_0 + \epsilon_{0j}$$

$$y_{1j} = x_j \beta_1 + \epsilon_{1j}$$

$$t_j = \begin{cases} 1, & \text{if } w_j \gamma + u_j > 0 \\ 0, & \text{otherwise} \end{cases}$$

where $y_{0j}$ is the outcome that person $j$ obtains if person $j$ selects treatment 0, and $y_{1j}$ is the outcome that person $j$ obtains if person $j$ selects treatment 1. We never observe both $y_{0j}$ and $y_{1j}$, only one or the other. We observe

$$y_j = t_j y_{1j} + (1 - t_j) y_{0j}$$
In this unconstrained model, the vector of error terms \( (\epsilon_{0j}, \epsilon_{1j}, u_j)' \) comes from a mean zero trivariate normal distribution with covariance matrix

\[
\begin{pmatrix}
\sigma_0^2 & \sigma_{01} & \sigma_{0}\rho_0 \\
\sigma_{01} & \sigma_1^2 & \sigma_{1}\rho_1 \\
\sigma_{0}\rho_0 & \sigma_{1}\rho_1 & 1
\end{pmatrix}
\]

The covariance \( \sigma_{01} \) cannot be identified because we never observe both \( y_{1j} \) and \( y_{0j} \). However, identification of \( \sigma_{01} \) is not necessary to estimate the other parameters because all covariates and the outcome are observed in observations from each group. We normalize the treatment error variance to be 1 because we observe only whether an outcome occurs under treatment. More details are found in Methods and formulas.

Rather than showing two separate regression equations, etregress reports one outcome equation with interaction terms between the treatment and outcome covariates. etregress can fit the constrained and generalized potential-outcome models using either the default maximum likelihood estimator or the one-step CF estimator obtained with option cfunction. The two-step CF estimator provides consistent estimates for the constrained model.

**Basic examples**

When there are no interactions between the treatment variable and the outcome covariates in the constrained model, etregress directly estimates the ATE and the ATET.

> **Example 1: Basic example**

We estimate the ATE of being a union member on wages of women in 1972 from a nonrepresentative extract of the National Longitudinal Survey on young women who were ages 14–26 in 1968. We will use the variables wage (wage), grade (years of schooling completed), smsa (an indicator for living in an SMSA—standard metropolitan statistical area), black (an indicator for being African-American), tenure (tenure at current job), and south (an indicator for living in the South).
We use `etregress` to estimate the parameters of the endogenous treatment-regression model.

```
. use http://www.stata-press.com/data/r15/union3
(National Longitudinal Survey. Young Women 14-26 years of age in 1968)
. etregress wage age grade smsa black tenure, treat(union = south black tenure)
Iteration 0: log likelihood = -3140.811
Iteration 1: log likelihood = -3053.6629
Iteration 2: log likelihood = -3051.5847
Iteration 3: log likelihood = -3051.575
Iteration 4: log likelihood = -3051.575
```

Linear regression with endogenous treatment

| Coef. | Std. Err. | z    | P>|z| | 95% Conf. Interval |
|-------|-----------|-----|-----|------------------|
| wage  |           |     |     |                  |
| age   | 0.1487409 | 0.0193291 | 7.70 | 0.000 | 0.1108566 - 0.1866252 |
| grade | 0.4205658 | 0.0293577 | 14.33 | 0.000 | 0.3630258 - 0.4781058 |
| smsa  | 0.9117044 | 0.1249041 | 7.30 | 0.000 | 0.6668969 - 1.156512 |
| black | -0.7882471 | 0.1367078 | -5.77 | 0.000 | -1.05619 - -0.5203048 |
| tenure| 0.1524015 | 0.0369596 | 4.12 | 0.000 | 0.0879621 - 0.2168409 |
| 1.union| 2.945815 | 0.7249621 | 10.71 | 0.000 | 2.4069 - 3.484731 |
| _cons | -4.351572 | 0.5283952 | -8.24 | 0.000 | -5.387208 - -3.315936 |

union

| Coef. | Std. Err. | z    | P>|z| | 95% Conf. Interval |
|-------|-----------|-----|-----|------------------|
| south| -0.5807419 | 0.0851111 | -6.82 | 0.000 | -0.7475566 - -0.4139271 |
| black| 0.4557499 | 0.0958042 | 4.76 | 0.000 | 0.2679771 - 0.6435226 |
| tenure| 0.0871536 | 0.0232483 | 3.75 | 0.000 | 0.0415878 - 0.1327195 |
| _cons| -0.8855758 | 0.0724506 | -12.22 | 0.000 | -1.027576 - -0.7435753 |

/athrho | -0.6544347 | 0.0910314 | -7.19 | 0.000 | -0.832853 - -0.4760164 |
/lnsigma| 0.7026769 | 0.0293372 | 23.95 | 0.000 | 0.645177 - 0.7601767 |

rho | -0.574678 | 0.060971 | -9.51 | 0.000 | -0.706545 - -0.4430476 |
sigma | 2.019151 | 0.0592362 | 34.42 | 0.000 | 2.006325 - 2.031978 |
lambda | -1.1603 | 0.1495097 | -7.63 | 0.000 | -1.453334 - -0.8672668 |

LR test of indep. eqns. (rho = 0): chi2(1) = 19.84 Prob > chi2 = 0.0000

The likelihood-ratio test in the footer indicates that we can reject the null hypothesis of no correlation between the treatment-assignment errors and the outcome errors. The estimated correlation between the treatment-assignment errors and the outcome errors, $\rho$, is $-0.575$. The negative relationship indicates that unobservables that raise observed wages tend to occur with unobservables that lower union membership. We discuss some details about this parameter in the technical note below.

The estimated ATE of being a union member is 2.95. The ATET is the same as the ATE in this case because the treatment indicator variable has not been interacted with any of the outcome covariates, and the correlation and variance parameters are identical across the control and treatment groups.
Technical note

The results for the ancillary parameters $\rho$ and $\sigma$ require explanation. For numerical stability during optimization, `etregress` does not directly estimate $\rho$ or $\sigma$. Instead, `etregress` estimates the inverse hyperbolic tangent of $\rho$,

$$\text{atanh} \rho = \frac{1}{2} \ln \left( \frac{1 + \rho}{1 - \rho} \right)$$

and $\ln \sigma$. Also `etregress` reports $\lambda = \rho \sigma$, along with an estimate of the standard error of the estimate and the confidence interval.

In contrast to the constrained model, `etregress` directly estimates the ATE only when there are no interactions between the treatment variable and the outcome covariates in the unconstrained model.

Example 2: Allowing group-specific variance and correlation

We estimate the ATE of having health insurance on the natural logarithm of total out-of-pocket prescription drug expenditures from a simulated random sample of individuals between the ages of 26 and 64. We will use the variables `lndrug` (natural logarithm of spending on prescription drugs), `age` (age of the individual), `chron` (whether the individual has a chronic condition), `lninc` (natural logarithm of income), `married` (marriage status), and `work` (employment status). Our treatment is whether the person has health insurance, `ins`. We allow the outcome error variance and correlation parameters to vary between the treated (insured) and control (uninsured) groups in this example, rather than constraining them to be equal as in example 1.

We use `etregress` to estimate the parameters of the endogenous treatment-effects model. To estimate separate variance and correlation parameters for each of the control and treatment groups, we specify the `poutcomes` option. We specify the `cfunction` option to use the CF estimator.
. use http://www.stata-press.com/data/r15/drugexp
(Prescription drug expenditures)
. etregress lndrug chron age lninc, treat(ins=age married lninc work) poutcomes > cfunction

Iteration 0:  GMM criterion Q(b) =  2.279e-15
Iteration 1:  GMM criterion Q(b) =  6.358e-30

Linear regression with endogenous treatment  Number of obs =  6,000
Estimator:  control-function

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<th></th>
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<th>[95% Conf. Interval]</th>
</tr>
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</table>

Wald test of indep. (rho0 = rho1 = 0):  chi2(2) = 8.88  Prob > chi2 = 0.0118

The Wald test reported in the footer indicates that we can reject the null hypothesis of no correlation between the treatment-assignment errors and the outcome errors for the control and treatment groups. The estimate of the correlation of the treatment-assignment errors for the control group ($\rho_0$) is positive, indicating that unobservables that increase spending on prescription drugs tend to occur with unobservables that increase health insurance coverage. Because $\rho_1$ is also positive, we make the same interpretation for individuals with insurance. The estimate $\rho_1$ is larger than the estimate $\rho_0$, indicating a stronger relationship between the unobservables and treatment outcomes in the treated group.

The estimated ATE of having health insurance is $-0.86$. Note that while the ATE and ATET were the same in example 1, that is not the case here. We show how to calculate the ATET for a potential-outcome model in example 6.

The estimate of the outcome error standard-deviation parameter for the control group ($\sigma_0$) is slightly larger than that of the treatment group parameter ($\sigma_1$), indicating a greater variability in the unobservables among the untreated group.
Average treatment effect (ATE)

When there is a treatment variable and outcome covariate interaction, the parameter estimates from `etregress` can be used by `margins` to estimate the ATE, the average difference of the treatment potential outcomes and the control potential outcomes.

Example 3: Allowing interactions between treatment and outcome covariates, ATE

In example 1, the coefficients on the outcome covariates do not vary by treatment level. The differences in wages between union members and nonmembers are modeled as a level shift captured by the coefficient on the indicator for union membership. In this example, we use factor-variable notation to allow some of the coefficients to vary over treatment level and then use `margins` (see [R] `margins`) to estimate the ATE. (See [U] 11.4.3 Factor variables for an introduction to factor-variable notation.)

We begin by estimating the parameters of the model in which the coefficients on `black` and `tenure` differ for union members and nonmembers. We specify the `vce(robust)` option because we need to specify `vce(unconditional)` when we use `margins` below.
The results indicate that the coefficients on `black` differ by union membership and that the coefficient on `tenure` for nonmembers is positive, while the coefficient on `tenure` for members is 0. The model fits well overall, so we proceed with interpretation. Because we interacted the treatment variable with two of the covariates, the estimated coefficient on the treatment level is not an estimate of the ATE. Below we use `margins` to estimate the ATE from these results. We specify the `vce(unconditional)` option to obtain the standard errors for the population ATE instead of the sample ATE. We specify the `contrast(nowald)` option to suppress the Wald tests, which `margins` displays by default for contrasts.
We can use the same methods above to obtain the ATE in an unconstrained model.

Example 4: Treatment interactions and group-specific variance and correlation, ATE

In example 2, the coefficients on the outcome covariates do not vary by treatment level. Suppose we believe that the effect of having a chronic condition on out-of-pocket spending differs between the insured and uninsured. Again, we use an interaction term. Because we are using a CF estimator, the variance–covariance of the estimator (VCE) is already robust so we do not specify vce(robust).
. use http://www.stata-press.com/data/r15/drugexp
(Prescription drug expenditures)
. etregress lndrug i.ins#i.chron age lninc, treat(ins=age married lninc work) > poutcomes cfunction
Iteration 0:  GMM criterion Q(b) = 2.279e-15
Iteration 1:  GMM criterion Q(b) = 1.561e-28
Linear regression with endogenous treatment  Number of obs = 6,000
Estimator: control-function

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</tbody>
</table>

Wald test of indep. (rho0 = rho1 = 0): chi2(2) = 8.90  Prob > chi2 = 0.0117

The results indicate that the coefficient on chron differs by whether an individual has insurance. The model fits well overall, so we proceed with interpretation.

Because we interacted the treatment variable with one of the covariates, the estimated coefficient on the treatment level is not an estimate of the ATE. Below we use margins to estimate the ATE from these results. We specify the vce(unconditional) option to obtain the standard errors for the population ATE instead of the sample ATE. We specify the contrast(nowald) option to suppress the Wald tests.
. margins r.ins, vce(unconditional) contrast(nowald)

Contrasts of predictive margins
Expression : Linear prediction, predict()

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<tr>
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</thead>
<tbody>
<tr>
<td>Contrast</td>
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<tr>
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<tr>
<td>ins</td>
</tr>
<tr>
<td>(1 vs 0)</td>
</tr>
</tbody>
</table>

The ATE estimate is similar to the one produced by the constrained model in example 2.

**Average treatment effect on the treated (ATET)**

When there is a treatment variable and outcome covariate interaction, the parameter estimates from `etregress` can be used by `margins` to estimate the ATET, the average difference of the treatment potential outcomes and the control potential outcomes on the treated population.

**Example 5: Allowing interactions between treatment and outcome covariates, ATET**

The ATET may differ from the ATE in example 3 because the interaction between the treatment variable and some outcome covariates makes the ATE and the ATET vary over outcome covariate values. Below we use `margins` to estimate the ATET by specifying the `subpop(union)` option, which restricts the sample used by `margins` to union members.

. use http://www.stata-press.com/data/r15/union3
(National Longitudinal Survey. Young Women 14-26 years of age in 1968)
. etregress wage age grade smsa i.union#c.(black tenure), treat(union = south black tenure) vce(robust)
(output omitted)
. margins r.union, vce(unconditional) contrast(nowald) subpop(union)

Contrasts of predictive margins
Expression : Linear prediction, predict()

<table>
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<td>(1 vs 0)</td>
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The estimated ATET and ATE are close, indicating that the average predicted outcome for the treatment group is similar to the average predicted outcome for the whole population.
Example 6: Treatment interactions and group-specific variance and correlation, ATET

The ATET may differ from the ATE in example 4 because the interaction between the treatment variable and some outcome covariates makes the ATE and the ATET vary over values of the covariate in the outcome equation. Even if there is no interaction between treatment assignment and a covariate in the outcome equation, the estimated ATE and ATET will differ if the variances of the outcome errors and their correlations with the treatment-assignment errors differ across the control and treatment groups.

We can estimate the ATET of having health insurance by using the conditional treatment effect (conditional on exogenous covariates and treatment level) obtained using the predict, cte and the margins commands; see Methods and formulas below and [TE] etregress postestimation for more details about the use of predict after etregress.

We restrict estimation to the treated subpopulation by specifying the subpop(ins) option with margins.

```
. use http://www.stata-press.com/data/r15/drugexp
   (Prescription drug expenditures)
. etregress lndrug i.ins#i.chron age lninc,
   > treat(ins = age married lninc work) poutcomes cfunction
   (output omitted)
. margins, predict(cte) subpop(ins) vce(unconditional)
Predictive margins Number of obs = 6,000
   Subpop. no. obs = 4,556
Expression : Conditional treatment effect, predict(cte)

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```

In absolute value, the treatment effect on the treated of $-0.76$ is smaller than the population average effect of $-0.86$ that we found in example 4.
etregress (maximum likelihood) stores the following in e():

Scalars
- e(N) number of observations
- e(k) number of parameters
- e(k_eq) number of equations in e(b)
- e(k_eq_model) number of equations in overall model test
- e(k_aux) number of auxiliary parameters
- e(k_dv) number of dependent variables
- e(df_m) model degrees of freedom
- e(ll) log likelihood
- e(ll0) log likelihood, constant-only model (lrmodel only)
- e(N_clust) number of clusters
- e(lambda) estimate of $\lambda$ in constrained model
- e(lambda0) estimate of $\lambda_0$ in potential-outcome model
- e(lambda0) standard error of $\lambda$ in constrained model
- e(sigma0) estimate of $\sigma_0$ in potential-outcome model
- e(lambda1) estimate of $\lambda_1$ in potential-outcome model
- e(sigma1) estimate of $\sigma_1$ in potential-outcome model
- e(chi2) $\chi^2$
- e(chi2_c) $\chi^2$ for comparison test
- e(p) $p$-value for model test
- e(p_c) $p$-value for comparison test
- e(rho) estimate of $\rho$ in constrained model
- e(rho0) estimate of $\rho_0$ in potential-outcome model
- e(rho1) estimate of $\rho_1$ in potential-outcome model
- e(rank) rank of e(V)
- e(rank0) rank of e(V) for constant-only model
- e(ic) number of iterations
- e(rc) return code
- e(converged) 1 if converged, 0 otherwise

Macros
- e(cmd) etregress
- e(cmdline) command as typed
- e(depvar) name of dependent variable
- e(hazard) variable containing hazard
- e(ctype) weight type
- e(wexp) weight expression
- e(title) title in estimation output
- e(title2) secondary title in estimation output
- e(clustvar) name of cluster variable
- e(chi2type) Wald or LR; type of model $\chi^2$ test
- e(chi2_ct) Wald or LR; type of model $\chi^2$ test corresponding to e(chi2_c)
- e(vcetype) vcetype specified in vce() or
- e(vcetype) title used to label Std. Err.
- e(opt) type of optimization
- e(which) max or min; whether optimizer is to perform maximization or minimization
- e(method) ml
- e(ml_method) type of ml method
- e(user) name of likelihood-evaluator program
- e(technique) maximization technique
- e(properties) b V
- e(predict) program used to implement predict
- e(footnote) program used to implement the footnote display
- e(marginsok) predictions allowed by margins
- e(asbalanced) factor variables fvset as asbalanced
- e(asobserved) factor variables fvset as asobserved
etregress — Linear regression with endogenous treatment effects

Matrices
- e(b) — coefficient vector
- e(Cns) — constraints matrix
- e(i) — iteration log (up to 20 iterations)
- e(gradient) — gradient vector
- e(V) — variance–covariance matrix of the estimators
- e(V_modelbased) — model-based variance

Functions
- e(sample) — marks estimation sample

etregress (two-step) stores the following in e():

Scalars
- e(N) — number of observations
- e(df_m) — model degrees of freedom
- e(lambda) — $\lambda$
- e(selambda) — standard error of $\lambda$
- e(sigma) — estimate of sigma
- e(chi2) — $\chi^2$
- e(p) — $p$-value for model test
- e(rho) — $\rho$
- e(rank) — rank of e(V)

Macros
- e(cmd) — etregress
- e(cmdline) — command as typed
- e(depvar) — name of dependent variable
- e(hazard) — variable containing hazard
- e(title) — title in estimation output
- e(chi2type) — Wald or LR; type of model $\chi^2$ test
- e(vce) — vcetype specified in vce()
- e(method) — twostep
- e(properties) — b V
- e(predict) — program used to implement predict
- e(footnote) — program used to implement the footnote display
- e(marginsok) — predictions allowed by margins
- e(marginsnotok) — predictions disallowed by margins
- e(asbalanced) — factor variables fvset as asbalanced
- e(asobserved) — factor variables fvset as asobserved

Matrices
- e(b) — coefficient vector
- e(V) — variance–covariance matrix of the estimators

Functions
- e(sample) — marks estimation sample
etregress (control-function) stores the following in e():

Scalars
- \( e(N) \): number of observations
- \( e(k) \): number of parameters
- \( e(k_{-eq}) \): number of equations in \( e(b) \)
- \( e(k_{-aux}) \): number of auxiliary parameters
- \( e(k_{-dv}) \): number of dependent variables
- \( e(lambda) \): estimate of \( \lambda \) in constrained model
- \( e(sigma) \): estimate of \( \sigma \) in constrained model
- \( e(lambda0) \): estimate of \( \lambda_0 \) in potential-outcome model
- \( e(sigma0) \): estimate of \( \sigma_0 \) in potential-outcome model
- \( e(lambda1) \): estimate of \( \lambda_1 \) in potential-outcome model
- \( e(sigma1) \): estimate of \( \sigma_1 \) in potential-outcome model
- \( e(chi2_{-c}) \): \( \chi^2 \) for comparison test
- \( e(p_{-c}) \): \( p \)-value for comparison test
- \( e(rho) \): estimate of \( \rho \) in constrained model
- \( e(rho0) \): estimate of \( \rho_0 \) in potential-outcome model
- \( e(rho1) \): estimate of \( \rho_1 \) in potential-outcome model
- \( e(rank) \): rank of \( e(V) \)
- \( e(converged) \): 1 if converged, 0 otherwise

Macros
- \( e(cmd) \): etregress
- \( e(cmdline) \): command as typed
- \( e(depvar) \): name of dependent variable
- \( e(hazard) \): variable containing hazard
- \( e(wtype) \): weight type
- \( e(wexp) \): weight expression
- \( e(title) \): title in estimation output
- \( e(title2) \): secondary title in estimation output
- \( e(chi2_{-ct}) \): Wald; type of model \( \chi^2 \) test corresponding to \( e(chi2_{-c}) \)
- \( e(vce) \): vcetype specified in vce()
- \( e(vcetype) \): title used to label Std. Err.
- \( e(method) \): cfunction
- \( e(properties) \): b V
- \( e(predict) \): program used to implement predict
- \( e(footnote) \): program used to implement the footnote display
- \( e(marginsok) \): predictions allowed by margins
- \( e(asbalanced) \): factor variables fvset as asbalanced
- \( e(asobserved) \): factor variables fvset as asobserved

Matrices
- \( e(b) \): coefficient vector
- \( e(V) \): variance–covariance matrix of the estimators

Functions
- \( e(sample) \): marks estimation sample

Methods and formulas
Maddala (1983, 117–122 and 223–228) derives both the maximum likelihood and the CF estimators implemented here. Greene (2012, 890–894) also provides an introduction to the treatment-effects model. Cameron and Trivedi (2005, sections 16.7 and 25.3.4) and Wooldridge (2010, section 21.4.1) discuss the endogenous binary-variable model as an endogenous treatment-effects model and link it to recent work.
Methods and formulas are presented under the following headings:

- Constrained model
- General potential-outcome model
- Average treatment effect
- Average treatment effect on the treated

### Constrained model

The primary regression equation of interest is

\[ y_j = x_j \beta + \delta t_j + \epsilon_j \]  \hspace{1cm} (1)

where \( t_j \) is a binary-treatment variable that is assumed to stem from an unobservable latent variable:

\[ t_j^* = w_j \gamma + u_j \]

The decision to obtain the treatment is made according to the rule

\[ t_j = \begin{cases} 
1, & \text{if } t_j^* > 0 \\
0, & \text{otherwise}
\end{cases} \]

where \( \epsilon \) and \( u \) are bivariate normal with mean zero and covariance matrix

\[
\begin{bmatrix}
\sigma^2 & \rho \sigma \\
\rho \sigma & 1
\end{bmatrix}
\]

Interactions between \( x_j \) and the treatment \( t_j \) are also allowed in (1). The likelihood function for this model is given in Maddala (1983, 122). Greene (2000, 180) discusses the standard method of reducing a bivariate normal to a function of a univariate normal and the correlation \( \rho \). The following is the log likelihood for observation \( j \),

\[
\ln L_j = \begin{cases} 
\ln \Phi \left( \frac{w_j \gamma + (y_j - x_j \beta - \delta) \rho \sigma}{\sqrt{1 - \rho^2}} \right) - \frac{1}{2} \left( \frac{y_j - x_j \beta - \delta}{\sigma} \right)^2 - \ln(\sqrt{2\pi} \sigma) & t_j = 1 \\
\ln \Phi \left( \frac{-w_j \gamma - (y_j - x_j \beta) \rho \sigma}{\sqrt{1 - \rho^2}} \right) - \frac{1}{2} \left( \frac{y_j - x_j \beta}{\sigma} \right)^2 - \ln(\sqrt{2\pi} \sigma) & t_j = 0
\end{cases}
\]

where \( \Phi(\cdot) \) is the cumulative distribution function of the standard normal distribution.

In the maximum likelihood estimation, \( \sigma \) and \( \rho \) are not directly estimated. Rather \( \ln \sigma \) and \( \text{atanh} \rho \) are directly estimated, where

\[
\text{atanh} \rho = \frac{1}{2} \ln \left( \frac{1 + \rho}{1 - \rho} \right)
\]

The standard error of \( \lambda = \rho \sigma \) is approximated through the delta method, which is given by

\[
\text{Var}(\lambda) \approx D \text{Var} \left\{ \left( \text{atanh} \rho \hspace{0.5cm} \ln \sigma \right) \right\} D'
\]

where \( D \) is the Jacobian of \( \lambda \) with respect to \( \text{atanh} \rho \) and \( \ln \sigma \).
Maddala (1983, 120–122) also derives the CF estimator as a two-step estimator. This estimator is implemented here. We will discuss it and then discuss the one-step CF estimator that is also implemented.

For the two-step estimator, probit estimates of the treatment equation

\[ \Pr(t_j = 1 \mid w_j) = \Phi(w_j \gamma) \]

are obtained in the first stage. From these estimates, the hazard, \( h_j \), for each observation \( j \) is computed as

\[
\begin{align*}
    h_j &= \begin{cases} 
        \frac{\phi(w_j \hat{\gamma})}{\Phi(w_j \hat{\gamma})} & t_j = 1 \\
        -\frac{\phi(w_j \hat{\gamma})}{\{1 - \Phi(w_j \hat{\gamma})\}} & t_j = 0 
    \end{cases}
\end{align*}
\]

where \( \phi \) is the standard normal density function. If \( d_j = h_j (h_j + w_j \hat{\gamma}) \) then

\[
\begin{align*}
    E(y_j \mid t_j, x_j, w_j) &= x_j \beta + \delta t_j + \rho \sigma h_j \\
    \text{Var}(y_j \mid t_j, x_j, w_j) &= \sigma^2 (1 - \rho^2 d_j)
\end{align*}
\]

The two-step parameter estimates of \( \beta \) and \( \delta \) are obtained by augmenting the regression equation with the hazard \( h \). Thus the regressors become \( [x \ t \ h] \), and the additional parameter estimate \( \beta_h \) is obtained on the variable containing the hazard. A consistent estimate of the regression disturbance variance is obtained using the residuals from the augmented regression and the parameter estimate on the hazard

\[
\hat{\sigma}^2 = \frac{e'e + \beta_h^2 \sum_{j=1}^{N} d_j}{N}
\]

The two-step estimate of \( \rho \) is then

\[
\hat{\rho} = \frac{\beta_h}{\hat{\sigma}}
\]

To understand how the consistent estimates of the coefficient covariance matrix based on the augmented regression are derived, let \( A = [x \ t \ h] \) and \( D \) be a square diagonal matrix of size \( N \) with \( (1 - \hat{\rho}^2 d_j) \) on the diagonal elements. The conventional VCE is

\[
V_{\text{twostep}} = \hat{\sigma}^2 (A'A)^{-1}(A'DA + Q)(A'A)^{-1}
\]

where

\[
Q = \hat{\rho}^2 (A'DA) V_p (A'DA)
\]

and \( V_p \) is the variance–covariance estimate from the probit estimation of the treatment equation.

The one-step CF estimator is a GMM estimator with stacked moments. See Newey (1984) and Wooldridge (2010, sec. 14.2) for a description of this technique. Many econometric and statistical models can be expressed as conditions on the population moments. The parameter estimates produced by GMM estimators make the sample-moment conditions as true as possible given the data.

Under CF estimation, as in maximum likelihood estimation, we directly estimate \( \text{atanh} \ \rho \) and \( \ln \ \sigma \) rather than \( \rho \) and \( \sigma \), so the parameter vector is

\[
\theta = (\beta', \delta, \gamma', \text{atanh} \ \rho, \ln \sigma)'
\]
In this case, we have separate error functions for the treatment assignment

\[ u_t(t_j, w_j, \theta) = \begin{cases} \phi(w_j \gamma)/\Phi(w_j \gamma) & t_j = 1 \\ -\phi(w_j \gamma)/(1 - \Phi(w_j \gamma)) & t_j = 0 \end{cases} \]

for the outcome mean

\[ u_m(y_j, t_j, x_j, w_j, \theta) = y_i - x_j \beta - \delta t_j - \rho \sigma u_{t,j} \]

and for the outcome variance

\[ u_v(y_j, t_j, x_j, w_j, \theta) = u_{m,j}^2 - \sigma^2 \left[ 1 - \rho^2 \{ u_{t,j}(u_{t,j} + w_j \gamma) \} \right] \]

We calculate the hazard, \( h_j \), prior to estimation from a probit regression of the treatment \( t_j \) on the treatment covariates \( w_j \). Let \( \tilde{z}_j = (x_j, t_j, h_j) \). Now we define

\[ Z_j = \begin{bmatrix} \tilde{z}_j & 0 & 0 \\ 0 & w_j & 0 \\ 0 & 0 & 1 \end{bmatrix} \]

and

\[ s_j(y_j, t_j, x_j, w_j, \theta) = Z_j' \begin{bmatrix} u_{m,j} \\ u_{t,j} \\ u_{v,j} \end{bmatrix} \]

The CF estimator \( \hat{\theta} \) is the value of \( \theta \) that satisfies the sample-moment conditions

\[ 0 = \frac{1}{N} \sum_i s_j(y_j, t_j, x_j, w_j, \theta) \]

The Huber/White/robust sandwich estimator is consistent for the VCE. See Wooldridge (2010, chap. 14), Cameron and Trivedi (2005, chap. 6), and Newey and McFadden (1994).

The formula is

\[ \hat{V} = (1/N) S \tilde{S} \tilde{G}' \]

where

\[ \tilde{G} = \left\{ (1/N) \sum_i \frac{\partial s_j(y_j, t_j, x_j, w_j, \theta)}{\partial \theta} \right\}^{-1} \]

and

\[ \tilde{S} = (1/N) \sum_i s_j(y_j, t_j, x_j, w_j, \theta) s_j(y_j, t_j, x_j, w_j, \theta)' \]

The matrix \( \tilde{G} \) is not symmetric because our estimator comes from stacking the moment conditions instead of optimizing one objective function. The implication is that the robust formula should always be used because, even under correct specification, the nonsymmetric \( \tilde{G} \) and the symmetric \( \tilde{S} \) converge to different matrices.
General potential-outcome model

Equation (1) can be generalized to a potential-outcome model with separate variance and correlation parameters for the control and treatment groups.

The generalized model is

\[ y_{0j} = x_j \beta_0 + \epsilon_{0j} \]
\[ y_{1j} = x_j \beta_1 + \epsilon_{1j} \]
\[ t_j = \begin{cases} 1, & \text{if } w_j \gamma + u_j > 0 \\ 0, & \text{otherwise} \end{cases} \]

where \( y_{0j} \) is the outcome that person \( j \) obtains if person \( j \) selects treatment 0, and \( y_{1j} \) is the outcome that person \( j \) obtains if person \( j \) selects treatment 1. We never observe both \( y_{0j} \) and \( y_{1j} \), only one or the other. We observe

\[ y_j = t_j y_{1j} + (1 - t_j) y_{0j} \]

In this unconstrained model, the vector of error terms \((\epsilon_{0j}, \epsilon_{1j}, u_j)'\) comes from a mean zero trivariate normal distribution with covariance matrix

\[
\begin{bmatrix}
\sigma_0^2 & \sigma_{01} & \sigma_0 \rho_0 \\
\sigma_{01} & \sigma_1^2 & \sigma_1 \rho_1 \\
\sigma_0 \rho_0 & \sigma_1 \rho_1 & 1
\end{bmatrix}
\]

The likelihood function for this model is given in Maddala (1983, 224).

\[
\ln f_j = \begin{cases} 
\ln \Phi \left( \frac{w_j \gamma + (y_j - x_j \beta_1) \rho_1 / \sigma_1}{\sqrt{1 - \rho_1^2}} \right) - \frac{1}{2} \left( \frac{y_j - x_j \beta_1}{\sigma_1} \right)^2 - \ln(\sqrt{2\pi \sigma_1}), & t_j = 1 \\
\ln \Phi \left( \frac{-w_j \gamma - (y_j - x_j \beta_0) \rho_0 / \sigma_0}{\sqrt{1 - \rho_0^2}} \right) - \frac{1}{2} \left( \frac{y_j - x_j \beta_0}{\sigma_0} \right)^2 - \ln(\sqrt{2\pi \sigma_0}), & t_j = 0
\end{cases}
\]

\[
\ln L = \sum_{j=1}^n w_j \ln f_j
\]

where \( \Phi(\cdot) \) is the cumulative distribution function of the standard normal distribution, and \( w_j \) is an optional weight. The covariance between \( \epsilon_{0j} \) and \( \epsilon_{1j}, \sigma_{01} \), cannot be estimated because the potential outcomes \( y_{0j} \) and \( y_{1j} \) are never observed simultaneously.

As in the constrained model, \( \sigma_0 \) and \( \sigma_1 \) are not directly estimated in the maximum likelihood estimation; rather, \( \ln \sigma_0 \) and \( \ln \sigma_1 \) are estimated.

The parameters \( \rho_0 \) and \( \rho_1 \) are also not directly estimated; rather, \( \text{atanh} \rho_0 \) and \( \text{atanh} \rho_1 \) are directly estimated.

The new parameter vector is

\[ \theta = (\beta'_0, \beta'_1, \gamma', \text{atanh} \rho_0, \ln \sigma_0, \text{atanh} \rho_1, \ln \sigma_1)' \]

The CF estimator for this potential-outcome model uses new error functions for the outcome mean

\[
u_m(y_j, t_j, x_j, w_j, \theta) = y_i - t_j (x_j \beta_1 + \rho_1 \sigma_1 u_{i,j}) - (1 - t_j) (x_j \beta_0 + \rho_0 \sigma_0 u_{i,j})
\]
and for the outcome variances
\[ u_{v,0}(y_j, t_j, x_j, w_j, \theta) = (1 - t_j) \left( u_{m,j}^2 - \sigma_0^2 \left[ 1 - \rho_0^2 \{ u_{t,j}(u_{t,j} + w_j \gamma) \} \right] \right) \]
\[ u_{v,1}(y_j, t_j, x_j, w_j, \theta) = t_j \left( u_{m,j}^2 - \sigma_1^2 \left[ 1 - \rho_1^2 \{ u_{t,j}(u_{t,j} + w_j \gamma) \} \right] \right) \]

These error functions are derived based on the identities
\[
E(y_j | t_j, x_j, w_j) = t_j(x_j \beta_1 + \rho_1 \sigma_1 u_{t,j}) + (1 - t_j)(x_j \beta_0 + \rho_0 \sigma_0 u_{t,j})
\]
\[
\text{Var}(y_j | t_j = 0, x_j, w_j) = \sigma_0^2 \left[ 1 - \rho_0^2 \{ u_{t,j}(u_{t,j} + w_j \gamma) \} \right]
\]
\[
\text{Var}(y_j | t_j = 1, x_j, w_j) = \sigma_1^2 \left[ 1 - \rho_1^2 \{ u_{t,j}(u_{t,j} + w_j \gamma) \} \right]
\]

We calculate the hazard, \( h_j \), prior to estimation from a probit regression of the treatment, \( t_j \), on the treatment covariates, \( w_j \). Let \( \tilde{z}_j = \{ x_j, t_j, h_j, (1 - t_j)h_j \} \). Now we define
\[
Z_j = \begin{bmatrix}
\tilde{z}_j & 0 & 0 & 0 \\
0 & w_j & 0 & 0 \\
0 & 0 & 1 & 0 \\
0 & 0 & 0 & 1
\end{bmatrix}
\]

and
\[
s_j(y_j, t_j, x_j, w_j, \theta) = Z_j' \begin{bmatrix}
u_{m,j} \\
u_{t,j} \\
u_{v,0,j} \\
u_{v,1,j}
\end{bmatrix}
\]

The CF estimator \( \hat{\theta} \) is the value of \( \theta \) that satisfies the sample-moment conditions
\[
0 = \frac{1}{N} \sum_i s_j(y_j, t_j, x_j, w_j, \theta)
\]

The Huber/White/robust sandwich estimator is consistent for the VCE. See Wooldridge (2010, chap. 14), Cameron and Trivedi (2005, chap. 6), and Newey and McFadden (1994).

The formula is
\[
\hat{V} = (1/N) \overline{G} \overline{S} \overline{G}'
\]
where
\[
\overline{G} = \left\{ (1/N) \sum_i \frac{\partial s_j(y_j, t_j, x_j, w_j, \theta)}{\partial \theta} \right\}^{-1}
\]
and
\[
\overline{S} = (1/N) \sum_i s_j(y_j, t_j, x_j, w_j, \theta) s_j(y_j, t_j, x_j, w_j, \theta)'
\]

The matrix \( \overline{G} \) is not symmetric because our estimator comes from stacking the moment conditions instead of optimizing one objective function. The implication is that the robust formula should always be used because, even under correct specification, the nonsymmetric \( \overline{G} \) and the symmetric \( \overline{S} \) converge to different matrices.
Average treatment effect

The ATE is the average difference of the treated potential outcomes and the control potential outcomes.

By the law of iterated expectations, the ATE is

\[ E(y_{1j} - y_{0j}) = E\{E(y_{1j} - y_{0j}|x_j, \epsilon_{0j}, \epsilon_{1j})\} \]
\[ = E(x_j\beta_1 + \epsilon_1 - x_j\beta_0 - \epsilon_0) \]
\[ = E\{x_j(\beta_1 - \beta_0)\} \]

This expectation can be estimated as a predictive margin when \( x_j(\beta_1 - \beta_0) \) varies in \( x_j \). Otherwise, the ATE is estimated as the coefficient of \( t_j \) in the model.

Average treatment effect on the treated

The ATET is the average difference of the treated potential outcomes and the control potential outcomes on the treated population.

The conditional means of the potential outcomes \( y_{tj}, t \in (0, 1) \) for exogenous covariates \( x_j \) and treatment covariates \( w_j \) at treatment \( t_j = 1 \) are

\[ E(y_{tj}|x_j, w_j, t_j = 1) = x_j\beta_t + \rho_t\sigma_t\phi(w_j\gamma)/\Phi(w_j\gamma) \]

By the law of iterated expectations, the ATET is

\[ E(y_{1j} - y_{0j}|t_j = 1) = E\{E(y_{1j} - y_{0j}|x_j, w_j, t_j = 1)\} \]
\[ = E\{x_j(\beta_1 - \beta_0) + (\rho_1\sigma_1 - \rho_0\sigma_0)\phi(w_j\gamma)/\Phi(w_j\gamma)|t_j = 1\} \]

This expectation can be estimated as a predictive margin on the treated population when \( x_j(\beta_1 - \beta_0) \) varies in \( x_j \) or when the variance and correlation parameters differ by treatment group. Otherwise, the ATET is estimated as the coefficient of \( t_j \) in the model.

References


etregress — Linear regression with endogenous treatment effects


Heckman, J. 1976. The common structure of statistical models of truncation, sample selection and limited dependent variables and a simple estimator for such models. *Annals of Economic and Social Measurement* 5: 475–492.


Also see

[TE] etregress postestimation — Postestimation tools for etregress

[TE] etpoisson — Poisson regression with endogenous treatment effects

[ERM] eregress — Extended linear regression

[R] heckman — Heckman selection model

[R] probit — Probit regression

[R] regress — Linear regression

[SVY] svy estimation — Estimation commands for survey data

[U] 20 Estimation and postestimation commands
# Postestimation commands

The following postestimation commands are available after `etregress`:

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<td>Akaike’s and Schwarz’s Bayesian information criteria (AIC and BIC)</td>
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<td><code>estat summarize</code></td>
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<td><code>marginsplot</code></td>
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<td><code>predict</code></td>
<td>predictions, residuals, influence statistics, and other diagnostic measures</td>
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<td><code>predictnl</code></td>
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<td><code>testnl</code></td>
<td>Wald tests of nonlinear hypotheses</td>
</tr>
</tbody>
</table>

* `estat ic`, `lrtest`, and `suest` are not appropriate after `etregress`, `twostep` or `etregress, cfunction`.  
  `hausman` and `lrtest` are not appropriate with `svy` estimation results.
predict

Description for predict

predict creates a new variable containing predictions such as linear predictions, conditional treatment effects, standard errors, expected values, and probabilities.

Menu for predict

Statistics > Postestimation

Syntax for predict

After ML, twostep, or cfunction

predict [type] newvar [if] [in] [ , statistic ]

After ML or cfunction for constrained model

predict [type] { stub* | newvar_reg newvar_treat newvar_athrho newvar_lnsigma } [if] [in] , scores

After ML or cfunction for general potential-outcome model

predict [type] { stub* | newvar_reg newvar_treat newvar_athrho0 newvar_lnsigma0 
newvar_athrho1 newvar_lnsigma1 } [if] [in] , scores

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<td>conditional treatment effect at treatment level</td>
</tr>
<tr>
<td>stdp</td>
<td>standard error of the prediction</td>
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<tr>
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<td>standard error of the forecast</td>
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<tr>
<td>yctr</td>
<td>$E(y_j</td>
</tr>
<tr>
<td>ycntrt</td>
<td>$E(y_j</td>
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<tr>
<td>ptrt</td>
<td>$Pr(\text{treatment} = 1)$</td>
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<tr>
<td>xbrt</td>
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</tr>
<tr>
<td>stdptrt</td>
<td>standard error of the linear prediction for treatment equation</td>
</tr>
</tbody>
</table>

These statistics are available both in and out of sample; type predict ... if e(sample) ... if wanted only for the estimation sample.

stdf is not allowed with svy estimation results.
Options for predict

**xb**, the default, calculates the linear prediction, \(x_j b\).

**cte** calculates the treatment effect, the difference of potential-outcome means, conditioned on treatment level.

**stdp** calculates the standard error of the prediction, which can be thought of as the standard error of the predicted expected value or mean for the observation’s covariate pattern. The standard error of the prediction is also referred to as the standard error of the fitted value.

**stdf** calculates the standard error of the forecast, which is the standard error of the point prediction for one observation. It is commonly referred to as the standard error of the future or forecast value. By construction, the standard errors produced by **stdf** are always larger than those produced by **stdp**; see *Methods and formulas* in [R] *regress postestimation*.

**yctrt** calculates the expected value of the dependent variable conditional on the presence of the treatment: \(E(y_j | \text{treatment} = 1)\).

**ycntrt** calculates the expected value of the dependent variable conditional on the absence of the treatment: \(E(y_j | \text{treatment} = 0)\).

**ptrt** calculates the probability of the presence of the treatment:
\[ \Pr(\text{treatment} = 1) = \Pr(w_j \gamma + u_j > 0). \]

**xbtrt** calculates the linear prediction for the treatment equation.

**stdptrt** calculates the standard error of the linear prediction for the treatment equation.

**scores**, not available with **twostep**, calculates equation-level score variables.

The first new variable will contain \(\partial \ln L/\partial (x_j \beta)\).

The second new variable will contain \(\partial \ln L/\partial (w_j \gamma)\).

Under the constrained model, the third new variable will contain \(\partial \ln L/\partial \tanh \rho\).

Under the constrained model, the fourth new variable will contain \(\partial \ln L/\partial \ln \sigma\).

Under the general potential-outcome model, the third new variable will contain \(\partial \ln L/\partial \tanh \rho_0\).

Under the general potential-outcome model, the fourth new variable will contain \(\partial \ln L/\partial \ln \sigma_0\).

Under the general potential-outcome model, the fifth new variable will contain \(\partial \ln L/\partial \tanh \rho_1\).

Under the general potential-outcome model, the sixth new variable will contain \(\partial \ln L/\partial \ln \sigma_1\).
margins

Description for margins

margins estimates margins of response for linear predictions, conditional treatment effects, expected values, and probabilities.

Menu for margins

Statistics > Postestimation

Syntax for margins

margins [marginlist] [ , options ]
margins [marginlist], predict(statistic ...) [ predict(statistic ...) ... ] [options ]

Maximum likelihood and control-function estimation results

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<tr>
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</tr>
<tr>
<td>ycnt</td>
<td>$E(y_j \mid \text{treatment} = 0)$</td>
</tr>
<tr>
<td>ptrt</td>
<td>$Pr(\text{treatment} = 1)$</td>
</tr>
<tr>
<td>xbt</td>
<td>linear prediction for treatment equation</td>
</tr>
<tr>
<td>stdp</td>
<td>not allowed with margins</td>
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<td>stdptrt</td>
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</table>

Two-step estimation results

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<tr>
<th>statistic</th>
<th>Description</th>
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<tbody>
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<td>linear prediction; the default</td>
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<td>not allowed with margins</td>
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<td>stdf</td>
<td>not allowed with margins</td>
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<tr>
<td>stdptrt</td>
<td>not allowed with margins</td>
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</tbody>
</table>

Statistics not allowed with margins are functions of stochastic quantities other than $e(b)$. For the full syntax, see [R] margins.
Remarks and examples

The average treatment effect (ATE) and the average treatment effect on the treated (ATET) are the parameters most frequently estimated by postestimation techniques after etregress.

When there are no interactions between the treatment variable and the outcome covariates in the constrained model, etregress directly estimates the ATE and the ATET; see example 1 of [TE] etregress.

When there are no interactions between the treatment variable and the outcome covariates in the general potential-outcome model, etregress directly estimates the ATE; see example 2 of [TE] etregress.

When there are interactions between the treatment variable and the outcome covariates, you can use margins after etregress to estimate the ATE. See example 3 and example 4 of [TE] etregress for examples of ATE estimation.

When there are interactions between the treatment variable and the outcome covariates in the constrained model, you can use margins after etregress to estimate the ATET. See example 5 of [TE] etregress for an example of ATET estimation in the constrained model.

In the general potential-outcome model, you can use margins after etregress to estimate the ATET. See example 6 of [TE] etregress for an example of ATET estimation in the general potential-outcome model.

Also see

[TE] etregress — Linear regression with endogenous treatment effects

[U] 20 Estimation and postestimation commands
stteffects — Treatment-effects estimation for observational survival-time data

Description

stteffects estimates average treatment effects, average treatment effects on the treated, and potential-outcome means using observational survival-time data. The available estimators are regression adjustment, inverse-probability weighting, and more efficient methods that combine regression adjustment and inverse-probability weighting.

For a brief description and example of each estimator, see Remarks and examples in [TE] stteffects intro.

Syntax

stteffects subcommand ... [, options]

subcommand Description
ra regression adjustment
ipw inverse-probability weighting
ipwra inverse-probability-weighted regression adjustment
wra weighted regression adjustment

Also see

[TE] stteffects intro — Introduction to treatment effects for observational survival-time data
This entry provides an overview of the treatment-effects estimators that use observational survival-time data and are implemented in *stteffects*. It also provides an overview of the potential-outcomes framework and its application to survival-time data and to the interpretation of the treatment-effects parameters estimated.

The *stteffects* command estimates average treatment effects (ATEs), average treatment effects on the treated (ATETs), and potential-outcome means (POMs). Each of these effect parameters is discussed in this entry. *stteffects* implements a variety of estimators for the ATE, ATET, and POM. The treatment effects can be estimated using regression adjustment (RA), inverse-probability weights (IPW), inverse-probability-weighted regression adjustment (IPWRA), and weighted regression adjustment (WRA). This entry also provides some intuition for the estimators and discusses the trade-offs between them.

Remarks and examples

Remarks are presented under the following headings:

- Introduction
- A quick tour of the estimators
  - Regression adjustment
  - Inverse-probability weighting
  - Combinations of RA and IPW
  - Weighted regression adjustment
- Average treatment effect on the treated
- Comparison of treatment-effects estimators
- Assumptions and trade-offs
  - The conditional independence assumption
  - The sufficient overlap assumption
  - The correct adjustment for censoring assumption
  - Assumptions for the ATET
- Specification diagnostics and tests
- Multivalued treatments

Introduction

The *stteffects* command estimates treatment effects using observational survival-time data.

For some intuition about the methods implemented in the *stteffects* command, consider the following question: Does smoking decrease the time to a second heart attack in the population of women aged 45–55 who have had one heart attack? Three aspects of this question stand out.

1. For ethical reasons, these data will be observational.
2. This question is about the time to an event, and such data are commonly known as survival-time data or time-to-event data. These data are nonnegative and, frequently, right-censored.
3. Many researchers and practitioners want an effect estimate in easy-to-understand units of time.
Aspects of treatment effects are introduced in treatment effects for observational survival-time data.

We are most concerned with aspect 3 because it helps us define and understand the effect of interest. In particular, we would like to know the average change in time to a second heart attack that would occur in the population if all women smoked instead of if no women smoked. This effect is an ATE.

We must solve a missing-data problem to estimate the ATE. The ATE is the population average of the contrast in outcomes when everyone gets the treatment and when no one gets the treatment. Formally, we write this as

\[ ATE = E(t_1 - t_0) \]

where \( t_1 \) is the survival time when a subject gets the treatment and \( t_0 \) is the survival time when a subject does not get the treatment. For each treatment level, there is a potential outcome that would be observed if a subject received that treatment level: \( t_1 \) is the potential outcome that would occur if someone gets the treatment and \( t_0 \) is the potential outcome that would occur if someone does not get the treatment. The missing-data problem arises because each subject receives only one treatment level, and so we observe only one of the two potential outcomes.

Much of the survival-time literature uses a hazard ratio as the effect of interest. The ATE has three advantages over the hazard ratio as an effect measure.

1. The ATE measures the effect in the same time units as the outcome instead of in relative conditional probabilities.
2. The ATE is much easier to explain to nontechnical audiences.
3. The models used to estimate the ATE can be much more flexible. Hazard ratios are useful for population effects when they are constant, which occurs when the treatment enters linearly and the distribution of the outcome has a proportional-hazards form. Neither linearity in treatment nor proportional-hazards form is required for the ATE, and neither is imposed on the models fit by the estimators implemented in stteffects.

The estimators implemented in stteffects use the common missing-data techniques of regression modeling, weighting, and combinations thereof to account for data lost to censoring and to unobserved potential outcomes.

Here we note only a few contributions and entry points to the vast literature on estimating ATEs. The use of potential outcomes to define treatment effects has proved extraordinarily useful; see Holland (1986), Rubin (1974), and Heckman (1997). Cameron and Trivedi (2005, chap. 25), Wooldridge (2010, chap. 21), and Vittinghoff et al. (2012, chap. 9) provide excellent general introductions to estimating ATEs.

Technical note

Left-truncation would be another type of missing data. The estimators implemented in stteffects do not adjust for left-truncation, so stteffects cannot be used with delayed-entry data.

stteffects cannot be used with time-varying covariates or multiple-record data because these add a repeated-measure structure that significantly complicates the estimation problem.
A quick tour of the estimators

The `stteffects` command implements five estimators of treatment effects. We introduce each one by showing the basic syntax used to apply it to a common example dataset. See each command’s entry for detailed information.

We have some fictional data on the time to a second heart attack among women aged 45–55 years. The treatment, smoking, is stored in the 0/1 indicator `smoke`. These data also contain each woman’s age at the time of her first heart attack (`age`), and indices of her exercise level (`exercise`), diet quality (`diet`), and education attainment (`education`) prior to her first heart attack.

Like `streg` and other survival-time commands, `stteffects` uses the outcome variable and the failure indicator computed by `stset`. In this dataset, `atime` is the observed time in years to the second heart attack, and `fail` is the 0/1 indicator that a second heart attack was observed and recorded in `atime`. (When `fail` is 1, `atime` records the time to the second attack; when `fail` is 0, `atime` records a censored observation of the time to the second attack.)

We begin our examples by first reading in the data and then specifying the raw outcome and failure variables to `stset`.

```
. use http://www.stata-press.com/data/r15/sheart
   (Time to second heart attack (fictional))
. stset atime, failure(fail)
    failure event:  fail != 0 & fail <.
    obs. time interval:  (0, atime]
    exit on or before:  failure

         2,000  total observations
                0  exclusions
2,000  observations remaining, representing
1,208  failures in single-record/single-failure data
3,795.226  total analysis time at risk and under observation
             at risk from t =          0
              earliest observed entry t =          0
              last observed exit t =  34.17743
```

The output indicates that 1,208 of the 2,000 observations record actual time to a second heart attack. The remaining observations were censored. Now that we have `stset` the data, we can use `stteffects`.

Regression adjustment

Regression modeling of the outcome variable is a venerable approach to solving the missing-data problem in treatment-effects estimation. Known as the regression-adjustment (RA) estimator, this method uses averages of predicted outcomes to estimate the ATE. If the outcome model is well specified, this approach is surprisingly robust.
Example 1: RA estimation

We now use `stteffects ra` to estimate the ATE by RA. We model the outcome as a function of age, exercise, diet, and education, and we specify that smoke is the treatment variable.

```
. stteffects ra (age exercise diet education) (smoke)
    failure _d: fail
    analysis time _t: atime
```

Iteration 0:   EE criterion = 1.525e-19
Iteration 1:   EE criterion = 1.931e-30

Survival treatment-effects estimation
Number of obs = 2,000

Estimator : regression adjustment
Outcome model : Weibull
Treatment model: none
Censoring model: none

|       | Coef.  | Std. Err. | z     | P>|z|    | [95% Conf. Interval] |
|-------|--------|-----------|-------|--------|---------------------|
| ATE   |        |           |       |        |                     |
| smoke | -1.956657 | 0.3331787 | -5.87 | 0.000  | -2.609676 to -1.303639 |
| (Smoker vs Non smoker) |       |           |       |        |                     |
| POmean|        |           |       |        |                     |
| smoke | 4.243974 | 0.2620538 | 16.20 | 0.000  | 3.730358 to 4.75759  |
| Non smoker |       |           |       |        |                     |

When all women in the population smoke, the average time to a second heart attack is estimated to be 1.96 years less than when no women smoke. The estimated average time to a second heart attack when no women smoke is 4.24 years.

The output reports that a Weibull model was used for the outcome. The other outcome models available are exponential, gamma, and log normal. See example 2 in [TE] stteffects ra for an application of the gamma parameterization to this model.

The ratio of the ATE to control-level POM measures the importance of the effect. In this example, when all women smoke, the time to a second heart attack falls by an estimated 46% relative to the case in which none of them smoke. See example 3 in [TE] stteffects ra for an example that uses `nlcom` to compute a point estimate and a confidence interval for this ratio.

Unlike the IPW estimator discussed in the next section, RA does not model treatment assignment or the censoring process. Treatment assignment is handled by fitting separate models for each treatment level and averaging the predicted outcomes. As is standard in the survival-time literature, the censoring term in the log-likelihood function accounts for censoring; see Kalbfleisch and Prentice (2002, chap. 3), Cameron and Trivedi (2005, chap. 17), Cleves, Gould, and Marchenko (2016, chap. 13), and Wooldridge (2010, chap. 22).

See [TE] stteffects ra for further discussion of this command and the RA estimator.
Inverse-probability weighting

Sometimes researchers are more comfortable modeling treatment assignment than the outcome. Inverse-probability-weighted (IPW) estimators use weighted averages of the observed outcome to estimate the POMs and the ATE. The weights correct for the missing data. When there is no censoring, the missing potential outcome is the only missing data, and the weights are constructed from a model of treatment assignment. When the data may be censored, the weights must control for censoring and the missing potential outcome. In this case, IPW estimators construct the weights from two models, one for the censoring time and one for treatment assignment.

Example 2: IPW estimation

Here we use `stteffects ipw` to estimate the effect of smoking on the time to a second heart attack. The model of assignment to the treatment `smoke` depends on age, exercise, diet, and education. The time-to-censoring model also depends on age, exercise, diet, and education.

```
. stteffects ipw (smoke age exercise diet education)
> (age exercise diet education)
   failure _d: fail
   analysis time _t: atime
```

| _t | Coef. | Std. Err. | z | P>|z| | [95% Conf. Interval] |
|----|-------|-----------|---|------|---------------------|
| ATE | smoke (Smoker vs Nonsmoker) | -2.187297 | .6319837 | -3.46 | 0.001 | -3.425962 to -.9486314 |
| POmean | smoke Nonsmoker | 4.225331 | .517501 | 8.16 | 0.000 | 3.211047 to 5.239614 |

When all women in the population smoke, the average time to a second heart attack is estimated to be 2.19 years less than when no women smoke. The estimated average time to a second heart attack when no women smoke is 4.23 years. When all women smoke, the average time to a second heart attack falls by an estimated 52% relative to the case when no women smoke.

The estimates have changed; however, the interpretation is the same as for the RA estimator because the IPW and RA estimators are estimating the same population effects. Under correct model specification, the estimates will differ in finite samples, but the size of these differences will decrease as the sample size gets larger. For the case at hand, the estimated ATE and control-level POM are roughly similar to those produced by the RA estimator using the Weibull model for the outcome.

Recall that IPW estimators are weighted averages of observed outcomes and that the weights control for the missing outcomes. Weights in survival-time data have two components: one for the missing potential outcome and one for data lost to censoring. We used a logit model for treatment assignment, so the component of the weights that controls for the missing potential outcome comes from the
estimated logit parameters. We used a Weibull model for the time to censoring, so the component of
the weights that controls for data lost to censoring comes from the estimated Weibull parameters.

Using weighting from an estimated treatment-assignment model to control for the missing potential
outcome is standard in the treatment-effects literature; for example, see [TE] teffects intro advanced,
Wooldridge (2010, chap. 21), Vittinghoff et al. (2012, chap. 9), Hirano, Imbens, and Ridder (2003),
Cattaneo (2010), and Cattaneo, Drukker, and Holland (2013). Modeling the time to censoring is
specific to the survival-time treatment-effects literature; see Bai, Tsiatis, and O’Brien (2013) and

See [TE] stteffects ipw for further discussion of this command and the IPW estimator.

Combinations of RA and IPW

More efficient estimators are obtained by combining IPW and RA, due to Wooldridge (2007)
and Wooldridge (2010, chap. 21) and denoted by IPWRA. Unlike the estimators discussed in
Wooldridge (2010, chap. 21), both the treatment and the outcome models must be correctly specified
to estimate the ATE.

The IPWRA estimator uses estimated weights that control for missing data to obtain missingness-
adjusted regression coefficients that are used to compute averages of predicted outcomes to estimate
the POMs. The estimated ATE is a contrast of the estimated POMs. These weights always involve a
model for treatment assignment. You choose whether to account for censoring by including a term in
the log-likelihood function or whether to use weights that also account for the data lost to censoring.

Example 3: Likelihood-adjusted-censoring IPWRA estimation

We model the outcome (time to a second heart attack) as a function of age, exercise, diet, and
education. We model assignment to the treatment smoke as a function of the same covariates.

```
. stteffects ipwra (age exercise diet education)
   > (smoke age exercise diet education)
     failure _d: fail
     analysis time _t: atime
Iteration 0: EE criterion = 2.153e-16
Iteration 1: EE criterion = 2.940e-30
Survival treatment-effects estimation
   Number of obs = 2,000
   Robust
   Estimator : IPW regression adjustment
   Outcome model : Weibull
   Treatment model: logit
   Censoring model: none

                  Coef.  Std. Err.     z   P>|z|   [95% Conf. Interval]
-------------  -------------  --------  -------  ---------------------
       _t
  ATE      smoke
  (Smoker   vs       Nonsmoker)
         -1.592494   .4872777  -3.27    0.001   -2.54754   -.637447
     POmean
      smoke
    Nonsmoker       4.214523   .2600165   16.21    0.000       3.7049     4.724146
-------------  -------------  --------  -------  ---------------------
```

The estimated ATE of −1.59 and control-level POM of 4.21 are similar to the reported values of
−1.96 and 4.24 in example 1.
We did not specify a model for the time to censoring, so censoring is handled by including a term in the log-likelihood function in the Weibull outcome model. We denote this likelihood-adjusted-censoring (LAC) version of the IPWRA estimator by LAC-IPWRA.

Example 4: Weighted-adjusted-censoring IPWRA estimation

Instead of including a term in the log-likelihood function, the weighted-adjusted-censoring IPWRA (WAC-IPWRA) estimator uses estimated weights to adjust for censoring. We model the time to a second heart attack as a function of age, exercise, diet, and education; we model assignment to the treatment smoke as a function of the same covariates; and we model the time to censoring as a function of age, exercise, and diet.

```
. stteffects ipwra (age exercise diet education) > (smoke age exercise diet education) (age exercise diet)
   failure _d: fail
   analysis time _t: atime
Iteration 0:   EE criterion = 1.632e-16
Iteration 1:   EE criterion = 2.367e-30
Survival treatment-effects estimation
   Number of obs = 2,000
   Estimator : IPW regression adjustment
   Outcome model : Weibull
   Treatment model: logit
   Censoring model: Weibull

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
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<tbody>
<tr>
<td></td>
<td>Coef.</td>
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<td>_t</td>
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<tr>
<td>ATE</td>
<td></td>
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<tr>
<td></td>
<td>smoke</td>
</tr>
<tr>
<td></td>
<td>(Smoker vs Nonsmoker)</td>
</tr>
<tr>
<td></td>
<td>-2.037944  .6032549</td>
</tr>
<tr>
<td>POM mean</td>
<td>smoke</td>
</tr>
<tr>
<td></td>
<td>Nonsmoker</td>
</tr>
<tr>
<td></td>
<td>4.14284  .4811052</td>
</tr>
</tbody>
</table>
```

The estimated ATE of $-2.04$ and control-level POM of $4.14$ are similar to the reported values of $-1.96$ and $4.24$ in example 1.

The weights for censoring are constructed from the estimated parameters because we specified a time-to-censoring model.

Under correct specification, both versions of the IPWRA estimator estimate the same ATE and control-level POM as estimated by RA and IPW.

The addition of the time-to-censoring model makes the WAC-IPWRA somewhat less robust than the LAC-IPWRA estimator. Weighting methods to control for censoring also place more restrictive assumptions on the censoring process. For example, the censoring time must be random, otherwise it would be impossible to construct the weights. In Assumptions and trade-offs below, we discuss the trade-offs among the estimators and the assumptions that each requires. For the moment, we note that we believe the LAC-IPWRA estimator is more robust than the WAC-IPWRA estimator.

See [TE] stteffects ipwra for further discussion of this command and the IPWRA estimator.
Weighted regression adjustment

When estimating the parameters of an outcome model, the weighted regression-adjustment (WRA) estimator uses weights instead of a term in the log-likelihood function to adjust for censoring. These weights are constructed from a model for the censoring process. The estimated parameters are subsequently used to compute averages of predicted outcomes that estimate the POMs. A contrast of the estimated POMs estimates the ATE.

Example 5: WRA estimation

We model the time to a second heart attack as a function of age, exercise, diet, and education; we specify that smoke is the treatment; and we model the time to censoring as a function of age, exercise, and diet.

```
. stteffects wra (age exercise diet education) (smoke) (age exercise diet) 
   failure _d: fail 
   analysis time _t: atime 
Iteration 0: EE criterion = 7.037e-19 
Iteration 1: EE criterion = 1.110e-30 
Survival treatment-effects estimation Number of obs = 2,000 
Estimator : weighted regression adjustment 
Outcome model : Weibull 
Treatment model: none 
Censoring model: Weibull
```

|              | Robust Coef. | Std. Err. | z   | P>|z| | [95% Conf. Interval] |
|--------------|--------------|-----------|-----|------|----------------------|
| ATE          |              |           |     |      |                      |
| smoke (Smoker vs Nonsmoker) | -2.152014 | 0.4986005 | -4.32 | 0.000 | -3.129253, -1.174775 |
| POmean       |              |           |     |      |                      |
| smoke Nonsmoker | 4.079273 | 0.4379517 | 9.31 | 0.000 | 3.220903, 4.937642   |

The estimated ATE of −2.15 and control-level POM of 4.08 are similar to the reported values of −1.96 and 4.24 in example 1. Like the other estimators discussed, the WRA estimators estimate the same effect parameters as the RA estimator, so the interpretation is the same.

In many survival-time applications, using weights to adjust for censoring is probably less robust than just including a term in the log-likelihood function for the outcome model. The model used to construct the weights is just as complicated as the outcome model, and including the term in the log-likelihood function places fewer restrictions on the censoring process, as discussed in The correct adjustment for censoring assumption below.

See [TE] stteffects wra for further discussion of this command and the WRA estimator.
Average treatment effect on the treated

Intuitively, the average treatment effect on the treated (ATET) is the effect in a well-defined, at-risk subpopulation. Sometimes the subpopulation that gets the treatment defines such an at-risk subpopulation. For example, we may want to know the average change in time to a second heart attack among female smokers aged 45–55 who have had a heart attack if they all became nonsmokers. This effect is the ATET.

Below, we use `stteffects ra` to estimate the ATET by RA.

```
. stteffects ra (age exercise diet education) (smoke), atet
    failure _d: fail
    analysis time _t: atime
Iteration 0:  EE criterion = 1.525e-19
Iteration 1:  EE criterion = 2.002e-31
Survival treatment-effects estimation  Number of obs = 2,000
Estimator : regression adjustment
Outcome model : Weibull
Treatment model: none
Censoring model: none

|            | Coef.    | Std. Err. | z    | P>|z| | [95% Conf. Interval] |
|------------|----------|-----------|------|-----|---------------------|
| ATET       |          |           |      |     |                     |
| smoke      |          |           |      |     |                     |
| (Smoker vs Non smoker) | -1.527476 | .2489203  | -6.14| 0.000 | -2.015351 -1.039602 |
| POmean     |          |           |      |     |                     |
| smoke      |          |           |      |     |                     |
| Non smoker | 3.436937 | .2217808  | 15.50| 0.000 | 3.002255 3.87162   |
```

Now, all effects are calculated only for the subpopulation of women aged 45–55 years who smoke after their first heart attack. If no women in the subpopulation were to smoke, the average time to a second heart attack would be 3.44 years. When all women in the subpopulation smoke (the observed behavior), the average time to a second heart attack is estimated to be 1.53 years less than if no women in the subpopulation had smoked. In other words, if we could somehow turn all smokers in the subpopulation into nonsmokers, the average time to a second heart attack would be 3.44 years instead of 1.91 years (3.44 − 1.53 = 1.91).

These point estimates are a little different than those for the ATE and the control-level POM in the full population of women aged 45–55 years who have had one heart attack. The difference indicates that this particular health cost of smoking may be smaller among women who choose to smoke than in the full population.

Comparison of treatment-effects estimators

We can classify the estimators implemented in `stteffects` into five categories: 1) estimators based on a model for the outcome variable; 2) estimators based on models for the treatment assignment and the censoring time; 3) estimators based on models for the outcome variable and the treatment assignment; 4) estimators based on models for the outcome variable, the treatment assignment, and the censoring time; and 5) estimators based on models for the outcome variable and the censoring time.
Because there are several categories of estimators, the user must decide whether to model the outcome, the probability of treatment, the time to censoring, or some combination thereof.

Each category of estimator contains a variety of choices about the functional forms for the models. We now provide some intuition behind each category of estimator and discuss the relationships.

1. When modeling only the outcome, separate outcome models for each treatment level account for treatment assignment, and censoring is adjusted for in the log-likelihood function. This approach is used in the RA estimators.

2. Some researchers would rather avoid modeling the outcome. Some estimators use weighted averages of the observed outcome to estimate the effect. When estimating treatment effects from observational survival-time data, the weights used must account for treatment assignment and censoring. Models for treatment assignment and time to censoring are used to construct the weights. This approach is used in the IPW estimators.

3. When seeking a more efficient estimator, it is natural to model both the outcome and the treatment and to adjust for censoring in the outcome model. This approach is used in the LAC-IPWRA estimators.

4. When seeking a more efficient estimator, another natural approach is to model both the outcome and the treatment and to adjust for censoring by weights that come from a time-to-censoring model. This approach is used in the WAC-IPWRA estimators.

5. We could modify approach 1 to model the outcome and the time to censoring so that censoring is handled by weighting and its own model instead of by likelihood adjustment. This approach is used in the WRA estimators.

While researcher preferences over what to model largely dictate the approach selected, we quickly note two points that could affect which approach works best. First, we can adjust for censoring by weighting only when censoring time is random. Second, weighting estimators become unstable if the weights get too large.

In the next section, we elaborate on the assumptions needed and the trade-offs among the approaches to estimation.

**Assumptions and trade-offs**

The estimators implemented in `stteffects` require three assumptions: conditional independence, sufficient overlap, and correct adjustment for censoring.

**The conditional independence assumption**

All estimators implemented in `stteffects` require the potential outcomes to be independent of the treatment assignment after conditioning on the covariates. Randomized experiments and the Heckman selection model are two motivating frameworks for the conditional independence assumption.

When the treatment is assigned randomly, the randomization ensures that the potential outcomes are independent of the treatment assignment. In observational data, the treatment is not randomly assigned. However, many important questions can only be answered using observational data because it would be unethical to randomly allocate hazardous treatments, for example, smoking. The conditional independence assumption in observational data says that treatment assignment is as good as random after conditioning on the covariates.
We can also understand conditional independence from a modeling framework. The Heckman selection model specifies that each of the potential outcomes and the treatment assignment process are functions of observable covariates and unobservable errors. The potential outcomes are conditionally independent of the treatment assignment when the unobservable errors in the treatment-assignment process are independent of the unobservable errors in each of the potential-outcome processes. See The CI assumption in [TE] teffects intro advanced for a detailed example.

Both frameworks lead to the same conclusion: we need to observe and to condition on a sufficient number of covariates.

Essentially, all the estimators in stteffects are equally susceptible to violations of the conditional independence assumption. No one estimator is any more robust to the conditional independence assumption than any other one.

Estimating the ATE among the subpopulation of those who get the treatment requires a significantly weaker version of the CI assumption; see Assumptions for the ATET below.

For more details about the conditional independence assumption, see The CI assumption in [TE] teffects intro advanced, and see Rosenbaum and Rubin (1983), Heckman (1997), Imbens and Wooldridge (2009), Cameron and Trivedi (2005, sec. 25.2), Wooldridge (2010, chap. 21), and Vittinghoff et al. (2012, chap. 9).

The sufficient overlap assumption

The sufficient overlap assumption requires that each individual have a sufficiently positive probability of being assigned to each treatment level. We believe that the RA estimator is more robust than the other estimators to near violations of the sufficient overlap condition, under correct model specification.

The overlap condition has no specification test, but using teffects overlap and then summarizing the predicted treatment probabilities presents good diagnostics of overlap problems.

The correct adjustment for censoring assumption

The correct adjustment for censoring assumption has two parts. First, either the censoring time must be fixed or the process must be conditionally-on-covariates independent of the potential outcomes and the treatment-assignment process. This assumption is standard in survival analysis; see, for example, Kalbfleisch and Prentice (2002, chap. 3).

Second, the method used to adjust to censoring must be correct. For the RA and LAC-IPWRA estimators, which use likelihood-adjusted censoring, the second assumption is no more restrictive than assuming correct specification of the outcome model. For the IPW, WAC-IPWRA, and WRA estimators, which adjust by weighting, the second assumption requires that the censoring be random and that the censoring process be correctly modeled.

Under correct specification, all the estimators in stteffects perform well. However, we believe that estimators that use likelihood adjustment instead of weighting are more robust for three reasons.

1. The estimators that use weighting to adjust for censoring cannot handle fixed censoring processes. If the censoring process is not random, the weights are not well defined.
2. The estimators that use weighting to adjust for censoring do not allow the random censoring process to vary by treatment level.
3. The estimators that use weighting to adjust for censoring require an additional sufficient overlap condition: the probability of not being censored must be sufficiently greater than 0 or else the weights that adjust for censoring get too large.
While the estimators that use \textit{WAC} instead of \textit{LAC} require a few more assumptions, some researchers are more comfortable modeling the treatment and censoring than the outcome. In this case, the \textit{IPW} or \textit{WAC-IPWRA} estimator would be the estimator of choice.

See \textit{Specification diagnostics and tests} below for information about testing these assumptions.

\textbf{Assumptions for the ATET}

We noted in \textit{Average treatment effect on the treated} that the \textit{ATET} is sometimes more interesting than the \textit{ATE}. We can also estimate the \textit{ATET} under less restrictive versions of the conditional independence assumption and the sufficient overlap assumption than those required for the \textit{ATE}.

While \textit{ATE} estimation requires that the potential outcomes for both the treated and the not treated be conditionally independent of treatment assignment, \textit{ATET} estimation requires that only the not treated potential outcome be conditionally independent of treatment assignment.

This weaker version of conditional independence allows the gains from the treatment to be related to treatment assignment, after conditioning on the covariates. We can estimate the \textit{ATET}, but not the \textit{ATE}, if some unobserved factor increases (or decreases) the likelihood of assignment to the treatment, increases (or decreases) the time to event in the treatment group, and has no effect on the time to event when not in the treatment group.

For example, suppose that smoking is an acquired taste and that individuals who acquire the taste for smoking more easily are less adversely affected by smoking and otherwise similar to everyone else when not smoking. Taste for smoking is unobservable, and our data have no measure of this variable. In this case, we could estimate the \textit{ATET} but not the \textit{ATE}.

The weaker version of the sufficient overlap assumption only requires that each individual in the treated subpopulation have a positive probability of not getting treated. In contrast, \textit{ATE} estimation requires that each individual in the population have a positive probability of getting each treatment level. In particular, we can estimate the \textit{ATET}, but not the \textit{ATE}, when some individuals in the population have zero chance of getting the treatment. For example, we could estimate the \textit{ATET}, but not the \textit{ATE}, if some women will never smoke for religious reasons.

Even when the conditions for \textit{ATE} estimation hold, the \textit{ATE} and \textit{ATET} may differ. Finding that the \textit{ATET} is significantly different from the \textit{ATE} does not mean that the \textit{ATE} is incorrectly estimated.

See \textit{Heckman (1997)} and \textit{Wooldridge (2010, 911–912)} for more information about the assumptions necessary to estimate the \textit{ATET}.

\textbf{Specification diagnostics and tests}

After \texttt{stteffects ipw} and \texttt{stteffects ipwra}, some specification checks for the treatment-assignment model and the overlap condition are available.

The checks for the treatment-assignment model are known as balance checks. When the covariate distributions are invariant to the treatment level, the covariates are said to be balanced. The concept of balanced covariates comes from the experimental literature, in which random treatment assignment ensures that the covariates are balanced.

In observational data, the covariates are almost never balanced in the raw data. Weighting methods can be viewed as using a treatment-assignment model to balance the covariates. If the treatment-assignment model is well specified, the weights constructed from this model will balance the covariates. One of the nice features of balance checks is that they do not depend on the outcome or its distribution. This fact is especially useful for survival-time outcomes because censoring of the outcome has no effect on the balance checks, so the balance checks implemented in \texttt{tebalance} work without modification.
Conditional on the treatment-assignment model being well specified, we can use the estimated probabilities of treatment, known as the propensity scores, to look for signs that the overlap condition is violated. These checks depend only on the estimated treatment probabilities and are not affected by any censoring of the outcome, so the methods implemented in `teffects overlap` work without modification.

We begin examining our model by using `tebalance summarize` after refitting the models used by the LAC-IPWRA estimator.

```
. quietly stteffects ipwra (age exercise diet education)
    > (smoke age exercise diet education)
. tebalance summarize
```

Covariate balance summary

<table>
<thead>
<tr>
<th></th>
<th>Raw Weighted</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of obs</td>
<td>2,000</td>
</tr>
<tr>
<td>Treated obs</td>
<td>738</td>
</tr>
<tr>
<td>Control obs</td>
<td>1,262</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Covariate</th>
<th>Standardized differences</th>
<th>Variance ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Raw</td>
<td>Weighted</td>
</tr>
<tr>
<td>age</td>
<td>-.3122094</td>
<td>-.0184574</td>
</tr>
<tr>
<td>exercise</td>
<td>-.4975269</td>
<td>-.0458412</td>
</tr>
<tr>
<td>diet</td>
<td>-.2479756</td>
<td>.0021802</td>
</tr>
<tr>
<td>education</td>
<td>-.4801442</td>
<td>-.0216366</td>
</tr>
</tbody>
</table>

The weighted standardized differences are much closer to 0 than the raw standardized differences, and the weighted variance ratios are much closer to 1 than the raw variance ratios. These results indicate that the model-based treatment weights balanced the covariates; see [TE] `tebalance` and [TE] `tebalance summarize` for details.

The diagnostics presented by `tebalance summarize` are not a formal test. However, we can use `tebalance overid` to conduct a formal test of the hypothesis that the weights constructed from the treatment-assignment model balanced the covariates.

```
. tebalance overid
```

Iteration 0: criterion = .22681884
Iteration 1: criterion = .22692316 (backed up)
Iteration 2: criterion = .23090158
Iteration 3: criterion = .2311461
Iteration 4: criterion = .23256285
Iteration 5: criterion = .23286304
Iteration 6: criterion = .23335858
Iteration 7: criterion = .2335567
Iteration 8: criterion = .2335671
Iteration 9: criterion = .23356711

Overidentification test for covariate balance

H0: Covariates are balanced:

<table>
<thead>
<tr>
<th>chi2(5)</th>
<th>Prob &gt; chi2</th>
</tr>
</thead>
<tbody>
<tr>
<td>3.28142</td>
<td>0.6567</td>
</tr>
</tbody>
</table>

There is no significant evidence against the null hypothesis. The interpretation is that we do not reject the null hypothesis that the treatment-assignment model is well specified; see [TE] `tebalance` and [TE] `tebalance overid` for details.

Given that we do not reject the treatment-assignment model, we can use this model to look for evidence that the overlap condition is violated. We begin by using `teffects overlap`.
. tteffects overlap, plevel(Smoker)

The densities of the propensity scores for the smokers and nonsmokers appear to have the same support, indicating that there is no violation of the overlap condition. The only indicator of a possible problem is that the support of the density for nonsmokers gets very close to 0. This problem would affect ATE estimation but not ATET estimation, as discussed in Assumptions and trade-offs. To further investigate, we compute and summarize the predicted propensity score by treatment level.

. predict ps1, p tlevel(Smoker)
. summarize ps1 if smoke == 0

<table>
<thead>
<tr>
<th>Variable</th>
<th>Obs</th>
<th>Mean</th>
<th>Std. Dev.</th>
<th>Min</th>
<th>Max</th>
</tr>
</thead>
<tbody>
<tr>
<td>ps1</td>
<td>1,262</td>
<td>.3410001</td>
<td>.1381673</td>
<td>.014819</td>
<td>.6161401</td>
</tr>
</tbody>
</table>

. summarize ps1 if smoke == 1

<table>
<thead>
<tr>
<th>Variable</th>
<th>Obs</th>
<th>Mean</th>
<th>Std. Dev.</th>
<th>Min</th>
<th>Max</th>
</tr>
</thead>
<tbody>
<tr>
<td>ps1</td>
<td>738</td>
<td>.4168805</td>
<td>.1107557</td>
<td>.0454891</td>
<td>.6216282</td>
</tr>
</tbody>
</table>

To interpret these results, recall that ATE estimation requires that the minimum propensity score for each treatment level be sufficiently greater than 0 and that the maximum propensity score for each treatment level be sufficiently less than 1. Also recall that ATET estimation only requires that the maximum propensity score for each treatment level be sufficiently less than 1.

For ATE estimation, only the minimum predicted propensity score for nonsmokers presents a challenge, and 0.015 is probably not too small. For ATET estimation, neither maximum causes concern.

For information about choosing among the stteffects estimators and their functional forms for the different models, see Model choice under Remarks and examples in [TE] tteffects intro advanced.

**Multivalued treatments**

stteffects can estimate treatment effects for multivalued treatments; here we provide some examples. See [TE] tteffects multivalued for an introduction to interpreting effects from multivalued treatments.
Example 6: Multivalued ATE estimation

We have another fictional dataset that records the time to a second heart attack among women aged 45–55 years. In this dataset, \( \text{atime} \) is the observed time in years to the second heart attack, and \( \text{fail} \) is the 0/1 indicator that a second heart attack was observed and recorded in \( \text{atime} \). (When \( \text{fail} = 1 \), \( \text{atime} \) records the time to the second attack; when \( \text{fail} = 0 \), \( \text{atime} \) records a censored observation of the time to the second attack.)

These data also contain the age at the time of the first heart attack (\( \text{age} \)), and indices of each woman’s exercise level (\( \text{exercise} \)), diet quality (\( \text{diet} \)), and education attainment (\( \text{education} \)) prior to her first heart attack.

The treatment, smoking, is stored in the categorical variable \( \text{smoke} \), which has the following value labels. The women who never smoked are labeled as \( \text{N} \); the women who previously smoked but quit before their first heart attack are labeled as \( \text{B} \); the women who previously smoked but quit after their first heart attack are labeled as \( \text{A} \); and the women who continued to smoke after their first heart attack are labeled as \( \text{S} \).

We begin by first reading in the data and then reviewing information previously stored using \text{stset}.

```
. use http://www.stata-press.com/data/r15/sheartm, clear
   (Time to second heart attack (fictional))
. stset atime, failure(fail)
    -> stset atime, failure(fail)
      failure event:  fail != 0 & fail < .
      obs. time interval: (0, atime]
      exit on or before:  failure

  10,000  total observations
       0  exclusions

   10,000  observations remaining, representing
   9,741  failures in single-record/single-failure data
         27,999.155  total analysis time at risk and under observation
                      at risk from t = 0
                      earliest observed entry t = 0
                      last observed exit t = 17.40826
```

We continue by tabulating the treatment variable \( \text{smoke} \).

```
. tabulate smoke

<table>
<thead>
<tr>
<th>Smoking level</th>
<th>Freq.</th>
<th>Percent</th>
<th>Cum.</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>3,167</td>
<td>31.67</td>
<td>31.67</td>
</tr>
<tr>
<td>B</td>
<td>2,263</td>
<td>22.63</td>
<td>54.30</td>
</tr>
<tr>
<td>A</td>
<td>1,924</td>
<td>19.24</td>
<td>73.54</td>
</tr>
<tr>
<td>S</td>
<td>2,646</td>
<td>26.46</td>
<td>100.00</td>
</tr>
</tbody>
</table>

Total 10,000 100.00
```

We see that 31.67% of the women never smoked, 22.63% of the women previously smoked but quit before their first heart attack, 19.24% of the women previously smoked but quit after their first heart attack, and 26.46% of the women continued to smoke after their first heart attack.
We now use `stteffects ra` to estimate the ATE by RA. We model the outcome as a function of age, exercise, diet, and education, and we specify that smoke is the treatment variable.

```
. stteffects ra (age exercise diet education) (smoke)
    failure _d: fail
    analysis time _t: atime
Iteration 0:   EE criterion = 6.709e-21
Iteration 1:   EE criterion = 8.284e-30
```

Survival treatment-effects estimation  Number of obs = 10,000
Estimator : regression adjustment
Outcome model : Weibull
Treatment model: none
Censoring model: none

|   _t  | Coef. | Std. Err. | z    | P>|z|  | [95% Conf. Interval] |
|-------|-------|-----------|------|------|----------------------|
| ATE   | smoke |           |      |      |                      |
| (B vs N) | -.4129793 | .0317    | -13.03 | 0.000 | -.47511 -.3508485    |
| (A vs N) | -1.281031  | .032866  | -38.98 | 0.000 | -1.345447 -1.216614   |
| (S vs N) | -2.167359  | .0338994 | -63.93 | 0.000 | -2.233801 -2.100917   |

POmean

| smoke | N     | Coef. | Std. Err. | z    | P>|z|  | [95% Conf. Interval] |
|-------|-------|-------|-----------|------|------|----------------------|
| N     | 3.745919 | .0289014 | 129.61 | 0.000 | 3.689273 3.802565 |

The average time to a second heart attack is 0.41 years sooner when all the women smoked at some point but quit before their first heart attack than when all the women never smoked. The average time to a second heart attack is 1.28 years sooner when all the women smoked at some point but quit after their first heart attack than when all the women never smoked. The average time to a second heart attack is 2.17 years sooner when all the women continued to smoke after their first heart attack than when all the women never smoked.
Example 7: Multivalued ATET estimation

In the at-risk subpopulation of women who continued to smoke, we want to estimate the effect of continuing to smoke (S) versus quitting after the first heart attack (A). Below we estimate the ATETs by RA, specifying A to be the control level and S to be the treatment level.

```
stteffects ra (age exercise diet education) (smoke), atet control(A) tlevel(S)
   failure _d: fail
   analysis time _t: atime
```

Iteration 0: EE criterion = 6.709e-21
Iteration 1: EE criterion = 6.836e-30

Survival treatment-effects estimation
Number of obs = 10,000
Estimator : regression adjustment
Outcome model : Weibull
Treatment model: none
Censoring model: none

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>_t</td>
<td>Coef.</td>
<td>Std. Err.</td>
<td>z</td>
</tr>
<tr>
<td>ATET</td>
<td>smoke</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>(N vs A)</td>
<td>1.290123</td>
<td>.0377552</td>
<td>34.17</td>
</tr>
<tr>
<td></td>
<td>(B vs A)</td>
<td>.8748349</td>
<td>.0239595</td>
<td>36.51</td>
</tr>
<tr>
<td></td>
<td>(S vs A)</td>
<td>-.8869257</td>
<td>.0272301</td>
<td>-32.57</td>
</tr>
<tr>
<td>POmean</td>
<td>smoke</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>A</td>
<td>2.500108</td>
<td>.0217833</td>
<td>114.77</td>
</tr>
</tbody>
</table>

The parameter (S vs A) is the one of interest. The estimate implies that the average time to a second heart attack among women who continue to smoke is 0.89 years sooner when they all continue to smoke than when they all quit smoking after their first heart attack.

Acknowledgments

We thank Xiaofei Bai of Boehringer Ingelheim, Matias Cattaneo of the University of Michigan, Barbara Sianesi of the Institute for Fiscal Studies, and Jeff Wooldridge of Michigan State University for helpful conversations about methods and the literature.

References


Also see

[TE] stteffects postestimation — Postestimation tools for stteffects
[TE] tteffects intro advanced — Advanced introduction to treatment effects for observational data
[ST] streg — Parametric survival models
[ST] stset — Declare data to be survival-time data
[U] 20 Estimation and postestimation commands
Description

stteffects ipw estimates the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs) from observational survival-time data with random time to censoring. Estimation is by inverse-probability weighting (IPW). IPW estimators use weighted averages of the observed outcome. The estimated weights correct for missing data on the potential outcomes and for censored survival times. stteffects ipw offers several choices for the functional forms of the treatment model and the time-to-censoring model. Binary and multivalued treatments are accommodated.

See [TE] stteffects intro for an overview of estimating treatment effects from observational survival-time data.

Quick start

Specify time as observed failure time and fail as failure indicator

```
stset time, failure(fail)
```

ATE of binary treat2 on time by IPW using a logistic model of treat2 on x and w and using x and w in a Weibull model for the censoring time

```
stteffects ipw (treat2 x w) (x w)
```

As above, but estimate the ATET

```
seffects ipw (treat2 x w) (x w), atet
```

ATE of treat2 on time by IPW using a probit model of treat2 on x and w and using x and w in a gamma model for the censoring time

```
seffects ipw (treat2 x w, probit) (x w, gamma)
```

ATE for treatment levels 2 and 3 of three-valued treatment treat3

```
seffects ipw (treat3 x w) (x w)
```

As above, and specify that treat3 = 3 is the control level using the value label “MyControl” for 3

```
seffects ipw (treat3 x w) (x w), control("MyControl")
```

Menu

Statistics > Treatment effects > Survival outcomes > Inverse-probability weighting (IPW)
Syntax

```
stteffects ipw (tvar tmvarlist [ , tmoptions ]) (cmvarlist [ , cmoptions ])  
    [ if ] [ in ] [ , stat options ]
```

tvar must contain integer values representing the treatment levels.

tmvarlist specifies the variables that predict treatment assignment in the treatment model.

cmvarlist specifies the variables that predict censoring in the censoring model.

### tmoptions

<table>
<thead>
<tr>
<th>Model</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>logit</td>
<td>logistic treatment model; the default</td>
</tr>
<tr>
<td>probit</td>
<td>probit treatment model</td>
</tr>
<tr>
<td>hetprobit(varlist)</td>
<td>heteroskedastic probit treatment model</td>
</tr>
<tr>
<td>noconstant</td>
<td>suppress constant from treatment model</td>
</tr>
</tbody>
</table>

### cmoptions

<table>
<thead>
<tr>
<th>Model</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>weibull</td>
<td>Weibull; the default</td>
</tr>
<tr>
<td>exponential</td>
<td>exponential</td>
</tr>
<tr>
<td>gamma</td>
<td>two-parameter gamma</td>
</tr>
<tr>
<td>lnormal</td>
<td>lognormal</td>
</tr>
<tr>
<td>ancillary(avarlist [ , noconstant ])</td>
<td>specify variables used to model ancillary parameter</td>
</tr>
<tr>
<td>noconstant</td>
<td>suppress constant from censoring model</td>
</tr>
</tbody>
</table>

### stat

<table>
<thead>
<tr>
<th>Stat</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>ate</td>
<td>estimate average treatment effect in population; the default</td>
</tr>
<tr>
<td>atet</td>
<td>estimate average treatment effect on the treated</td>
</tr>
<tr>
<td>pomeans</td>
<td>estimate potential-outcome means</td>
</tr>
</tbody>
</table>
### Options

<table>
<thead>
<tr>
<th>Model</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>ancillary</strong></td>
<td>specifies the variables used to model the ancillary parameter. By default, the ancillary parameter does not depend on covariates. Specifying <code>ancillary(avarlist, noconstant)</code> causes the constant to be suppressed in the model for the ancillary parameter.</td>
</tr>
<tr>
<td><strong>noconstant</strong></td>
<td>see [R] estimation options.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Stat</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>stat</strong></td>
<td>is one of three statistics: <code>ate</code>, <code>atet</code>, or <code>pomeans</code>. <code>ate</code> is the default.</td>
</tr>
<tr>
<td><code>ate</code></td>
<td>specifies that the average treatment effect be estimated.</td>
</tr>
<tr>
<td><code>atet</code></td>
<td>specifies that the average treatment effect on the treated be estimated.</td>
</tr>
<tr>
<td><code>pomeans</code></td>
<td>specifies that the potential-outcome means for each treatment level be estimated.</td>
</tr>
</tbody>
</table>
vce(vcetype) specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (robust), that allow for intragroup correlation (cluster clustvar), and that use bootstrap or jackknife methods (bootstrap, jackknife); see [R] vce_option.

Reporting

level(#) ; see [R] estimation options.
aequations specifies that the results for the outcome-model or the treatment-model parameters be displayed. By default, the results for these auxiliary parameters are not displayed.
noshow prevents stteffects ipw from showing the key st variables. This option is rarely used because most people type stset, show or stset, noshow to permanently set whether they want to see these variables mentioned at the top of the output of every st command; see [ST] stset.

display_options: noci, nopvalues, nomitted, vsquish, noemptycells, baselevels, allbaselevels, nofvlabel, fvwrap(#), fvwrapon(style), cformat(%fmt), pformat(%fmt), sformat(%fmt), and nolstretch; see [R] estimation options.

Maximization

maximize_options: iterate(#), [no] log, and from(init_specs); see [R] maximize. These options are seldom used.

init_specs is one of

matname [, skip copy]

# [, # ...], copy

iterinit(#) specifies the maximum number of iterations used to calculate the starting values. This option is seldom used.

Advanced

pstolerance(#) specifies the tolerance used to check the overlap assumption. The default value is pstolerance(1e-5). stteffects will exit with an error if an observation has an estimated propensity score smaller than that specified by pstolerance().

osample(newvar) specifies that indicator variable newvar be created to identify observations that violate the overlap assumption.

control(# | label) specifies the level of tvar that is the control. The default is the first treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. control() may not be specified with the statistic mass means. control() and tlevel() may not specify the same treatment level.

tlevel(# | label) specifies the level of tvar that is the treatment for the statistic atet. The default is the second treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. tlevel() may only be specified with statistic atet. tlevel() and control() may not specify the same treatment level.

The following option is available with stteffects but is not shown in the dialog box:

coefflegend; see [R] estimation options.
Remarks and examples

If you are not familiar with the framework for treatment-effects estimation from observational survival-time data, please see [TE] stteffects intro.

IPW estimators use contrasts of weighted averages of observed outcomes to estimate treatment effects. The estimated weights correct for data that are missing because each subject is only observed after receiving one of the possible treatment levels and because some survival-time outcomes are censored.

The IPW estimators implemented in stteffects ipw use a three-step approach to estimating the ATE:

1. Estimate the parameters of a treatment-assignment model, and compute the component of the estimated weights that accounts for data missing because each subject is only observed after receiving one of the possible treatment levels.
2. Estimate the parameters of a time-to-censoring model, and compute the component of the estimated weights that accounts for data lost to censoring.
3. Use the estimated weights to compute weighted averages of the outcomes for each treatment level.

To estimate the ATET, we use different weights in step 2.

The time to censoring must be random to use stteffects ipw because the model in step 2 is not well defined if the time to censoring is fixed. See [TE] stteffects intro for more details. For information about estimators that accommodate a fixed time to censoring, see [TE] stteffects ra and [TE] stteffects ipwra.

Here we note only a few entry points to the vast literature on IPW estimators. Hirano, Imbens, and Ridder (2003), Imbens (2000, 2004), Imbens and Wooldridge (2009), Rosenbaum and Rubin (1983), Robins and Rotnitzky (2006), Wooldridge (2002, 2007), Cameron and Trivedi (2005, chap. 25), Wooldridge (2010, chap. 21), and Vittinghoff et al. (2012, chap. 9) provide excellent general introductions to estimating ATEs and to the IPW estimators in particular.

Like streg and other survival-time commands, stteffects ipw uses the outcome variable and the failure indicator computed by, and optionally weights specified with, stset. stteffects ipw is not appropriate for data with time-varying covariates, also known as multiple-record survival-time data, or for delayed-entry data.

Example 1: Estimating the ATE

Suppose we wish to study the effect of smoking on the time to a second heart attack among women aged 45–55 years. In our fictional sheart dataset, atime is the observed time in years to a second heart attack or censoring, and fail is the 0/1 indicator that a second heart attack was observed. (When fail is 1, atime records the time to the second heart attack; when fail is 0, atime records a censored observation of the time to a second heart attack.) We previously stset these data; see A quick tour of the estimators in [TE] stteffects intro.

The treatment, smoking, is stored in the 0/1 indicator smoke. These data also contain age at the time of the first heart attack (age), and indices of the level of exercise (exercise), diet quality (diet), and education (education) prior to the first heart attack.

We can use stteffects ipw to estimate the ATE. We model treatment assignment using the default logit model with covariates on age, exercise, and education. We model the time to censoring using the default Weibull model with covariates on age, exercise, diet, and education.
. use http://www.stata-press.com/data/r15/sheart
  (Time to second heart attack (fictional))
. stteffects ipw (smoke age exercise education) (age exercise diet education)
          failure _d:  fail
          analysis time _t:  atime
Iteration 0:  EE criterion =  2.042e-18
Iteration 1:  EE criterion =  5.191e-31
Survival treatment-effects estimation  Number of obs   =    2,000
Estimator         : inverse-probability weights
Outcome model     : weighted mean
Treatment model: logit
Censoring model: Weibull

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>t</td>
<td>Coef.</td>
<td>Std. Err.</td>
<td>z</td>
<td>P&gt;</td>
</tr>
<tr>
<td>ATE</td>
<td>smoke</td>
<td>-2.22226</td>
<td>.6307573</td>
<td>-3.52</td>
<td>0.000</td>
</tr>
<tr>
<td></td>
<td>(Smoker vs Nonsmoker)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>POmean</td>
<td>smoke</td>
<td>4.235569</td>
<td>.5210937</td>
<td>8.13</td>
<td>0.000</td>
</tr>
<tr>
<td></td>
<td>Nonsmoker</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

When every woman smoked in the population of women aged 45–55 years who have had a heart attack, the average time to a second heart attack is estimated to be 2.22 years less than when no women in the population of interest smoked. The estimated average time to a second heart attack when no women in the population of interest smoked is 4.24 years.

The ratio of the ATE to the control-level POM measures the importance of the effect. In this example, when every woman smoked, the average time to a second heart attack falls by an estimated 52% relative to the case when none of them smoked. See example 3 in [TE] stteffects ra for an example that uses nlcom to compute a point estimate and a confidence interval for this ratio.

Example 2: Different treatment and censoring models

Instead of a logit model for the treatment assignment, we could have used a probit or a heteroskedastic probit model. Instead of a Weibull model for the censoring time, we could have used an exponential, a gamma, or a lognormal model. For a quick comparison, we now estimate the ATE using a probit model for the treatment assignment and using a gamma model for the censoring time.
. stteffects ipw (smoke age exercise education, probit) 
> (age exercise diet education, gamma)
    failure _d: fail
    analysis time _t: atime
Iteration 0:   EE criterion = 3.534e-15
Iteration 1:   EE criterion = 5.263e-27
Survival treatment-effects estimation  Number of obs =   2,000
Estimator : inverse-probability weights
Outcome model : weighted mean
Treatment model: probit
Censoring model: gamma

| _t | Robust Coef. | Std. Err. | z    | P>|z|  | [95% Conf. Interval]   |
|----|--------------|-----------|------|------|------------------------|
| ATE| smoke        |           |      |      |                        |
|    | (Smoker vs   |           |      |      |                        |
|    | Nonsmoker)   |           |      |      |                        |
|    | -2.646808    | .8368254  | -3.16| 0.002| -4.286956 -1.006661    |
| POM| smoke        |           |      |      |                        |
|    | Nonsmoker    |           |      |      |                        |
|    | 4.702301     | .7404567  | 6.35 | 0.000| 3.251033 6.15357      |

The estimated ATE of $-2.65$ and control-level POM of $4.70$ are similar to the values of $-2.22$ and $4.24$ reported in example 1.

---

Example 3: Estimating the ATET

Intuitively, the ATET measures the effect of the treatment on an at-risk subpopulation. Sometimes the subpopulation that gets the treatment defines such an at-risk subpopulation. The ATET has the added benefit that it can be estimated under weaker conditions than the ATE; see Assumptions and trade-offs under Remarks and examples in [TE] stteffects intro.
. stteffects ipw (smoke age exercise education) (age exercise diet education),
  > atet
     failure _d: fail
     analysis time _t: atime
Iteration 0: EE criterion = 2.042e-18
Iteration 1: EE criterion = 1.248e-32
Survival treatment-effects estimation Number of obs = 2,000
Estimator : inverse-probability weights
Outcome model : weighted mean
Treatment model: logit
Censoring model: Weibull

| _t | Robust Coef. | Std. Err. | z  | P>|z| | [95% Conf. Interval] |
|----|-------------|------------|----|-------|---------------------|
| ATET | smoke (Smoker vs Nonsmoker) | -1.846136 | 0.5076872 | -3.64 | 0.000 | -2.841185 -0.8510877 |
| POmean | smoke Nonsmoker | 3.543788 | 0.474395 | 7.47 | 0.000 | 2.613991 4.473585 |

When every woman in the subpopulation smoked, the average time to a second heart attack is estimated to be 1.85 years less than when no women in the subpopulation smoked. The estimated average time to a second heart attack when no women in the subpopulation smoked is 3.54 years.

Stored results

stteffects ipw stores the following in e():

Scalars
- e(N) number of observations
- e(n_j) number of observations for treatment level j
- e(N_clust) number of clusters
- e(k_eq) number of equations in e(b)
- e(k_levels) number of levels in treatment variable
- e(treated) level of treatment variable defined as treated
- e(control) level of treatment variable defined as control
- e(converged) 1 if converged, 0 otherwise

Macros
- e(cmd) stteffects
- e(cmdline) command as typed
- e(dead) _d
- e(depvar) _t
- e(tvar) name of treatment variable
- e(subcmd) ipw
- e(tmodel) treatment model: logit, probit, or hetprobit
- e(cmodel) censoring model: weibull, exponential, gamma, or lognormal
- e(stat) statistic estimated: ate, atet, or pomeans
- e(wtype) weight type
- e(wexp) weight expression
- e(title) title in estimation output
- e(clustvar) name of cluster variable
- e(tlevels) levels of treatment variable
Methods and formulas

The methods and formulas for the IPW estimators implemented in `stteffects ipw` are given in Methods and formulas of [TE] stteffects ipwra.

References


Also see

[TE] \texttt{stteffects postestimation} — Postestimation tools for \texttt{stteffects}

[TE] \texttt{stteffects intro} — Introduction to treatment effects for observational survival-time data

[ST] \texttt{streg} — Parametric survival models

[ST] \texttt{stset} — Declare data to be survival-time data

[U] 20 Estimation and postestimation commands
**Description**

`stteffects ipwra` estimates the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs) from observational survival-time data by inverse-probability-weighted regression adjustment (IPWRA). IPWRA estimators use missingness-adjusted regression coefficients to compute averages of treatment-level predicted outcomes. Contrasts of these averages estimate the treatment effects. `stteffects ipwra` offers several choices for the functional forms of the outcome model, of the treatment model, and of the optional time-to-censoring model. Binary and multivalued treatments are accommodated.

See [TE] `stteffects intro` for an overview of estimating treatment effects from observational survival-time data.

**Quick start**

Specify `time` as observed failure time and `fail` as failure indicator

```
stset time, failure(fail)
```

ATE of binary treatment `treat2` estimated by IPWRA using a Weibull model for `time` on `x1` and `x2` and a logistic model for `treat2` on `x1` and `w`

```
stteffects ipwra (x1 x2) (treat2 x1 w)
```

As above, but estimate the ATET

```
stteffects ipwra (x1 x2) (treat2 x1 w), atet
```

Gamma model for time and probit model for `treat2`

```
stteffects ipwra (x1 x2, gamma) (treat2 x1 w, probit)
```

ATE for each level of three-valued treatment `treat3`

```
stteffects ipwra (x1 x2) (treat3 x1 w)
```

As above, and specify that `treat3 = 3` is the control level using the value label “MyControl” for 3

```
stteffects ipwra (x1 x2) (treat3 x1 w), control("MyControl")
```

ATE of `treat2` estimated by IPWRA using a Weibull model for `time` on `x1` and `x2`, a logistic model for `treat2` on `x1` and `w`, and a Weibull model for the time to censoring with covariates `x1` and `x2`

```
stteffects ipwra (x1 x2) (treat2 x1 w) (x1 x2)
```

Gamma model for time, probit model for `treat2`, and gamma model for censoring

```
stteffects ipwra (x1 x2, gamma) (treat2 x1 w, probit) (x1 x2, gamma)
```
Menu

Statistics > Treatment effects > Survival outcomes > Regression adjustment with IPW

Syntax

```
stteffects ipwra (omvarlist [, omoptions]) (tvar tmvarlist [, tmoptions])
    [(cmvarlist [, cmoptions]) [if] [in] [, stat options]]
```

`omvarlist` specifies the variables that predict the survival-time variable in the outcome model.
`tvar` must contain integer values representing the treatment levels.
`tmvarlist` specifies the variables that predict treatment assignment in the treatment model.
`cmvarlist` specifies the variables that predict censoring in the censoring model.

### omoptions

**Model**

- `weibull`          Weibull; the default
- `exponential`      exponential
- `gamma`            two-parameter gamma
- `lnormal`          lognormal
- `ancillary(avarlist [, noconstant])` specify variables used to model ancillary parameter
- `noconstant`       suppress constant from outcome model

### tmoptions

**Model**

- `logit`             logistic treatment model; the default
- `probit`            probit treatment model
- `hetprobit(varlist)` heteroskedastic probit treatment model
- `noconstant`        suppress constant from treatment model

### cmoptions

**Model**

- `weibull`          Weibull; the default
- `exponential`      exponential
- `gamma`            two-parameter gamma
- `lnormal`          lognormal
- `ancillary(avarlist [, noconstant])` specify variables used to model ancillary parameter
- `noconstant`       suppress constant from censoring model
### Options

<table>
<thead>
<tr>
<th>Stat</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>ate</td>
<td>estimate average treatment effect in population; the default</td>
</tr>
<tr>
<td>atet</td>
<td>estimate average treatment effect on the treated</td>
</tr>
<tr>
<td>pomeans</td>
<td>estimate potential-outcome means</td>
</tr>
</tbody>
</table>

#### Options Description

- **SE/Robust**
  - `vce(vcetype)`
    - `vcetype` may be `robust`, `cluster clustvar`, `bootstrap`, or `jackknife`

- **Reporting**
  - `level(#)`
    - set confidence level; default is `level(95)`
  - `aequations`
    - display auxiliary-equation results
  - `noshow`
    - do not show `st` setting information
  - `display_options`
    - control columns and column formats, row spacing, line width, display of omitted variables and base and empty cells, and factor-variable labeling

- **Maximization**
  - `maximize_options`
    - control the maximization process; seldom used
  - `iterinit(#)`
    - specify starting-value iterations; seldom used

- **Advanced**
  - `pstolerance(#)`
    - set the tolerance for the overlap assumption
  - `osample(newvar)`
    - identify observations that violate the overlap assumption
  - `control(# | label)`
    - specify the level of `tvar` that is the control
  - `tlevel(# | label)`
    - specify the level of `tvar` that is the treatment
  - `coeflegend`
    - display legend instead of statistics

You must `stset` your data before using `stteffects`; see [ST] `stset`.
`omvarlist`, `tmvarlist`, `cmvarlist`, and `avarlist` may contain factor variables; see [U] 11.4.3 Factor variables.
`bootstrap`, `by`, `jackknife`, and `statsby` are allowed; see [U] 11.1.10 Prefix commands.
Weights are not allowed with the `bootstrap` prefix; see [R] `bootstrap`.
`fweights`, `iwights`, and `pweights` may be specified using `stset`; see Weights under Remarks and examples in [ST] `stset`. However, weights may not be specified if you are using the `bootstrap` prefix.
`coeflegend` does not appear in the dialog box.
See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.

**Options**

- **Model**
  - `ancillary(avarlist [, noconstant ])`
    - specifies the variables list used to model the ancillary parameter.
    - By default, the ancillary parameter does not depend on covariates. Specifying `ancillary(avarlist, noconstant)` causes the constant to be suppressed in the model for the ancillary parameter.
ancillary() may be specified for the model for survival-time outcome, for the model for the censoring variable, or for both. If ancillary() is specified for both, the varlist used for each model may be different.

noconstant; see [R] estimation options.

Stat

stat is one of three statistics: ate, atet, or pomeans. ate is the default.
ate specifies that the average treatment effect be estimated.
atet specifies that the average treatment effect on the treated be estimated.
pomeans specifies that the potential-outcome means for each treatment level be estimated.

SE/Robust

vce(vcetype) specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (robust), that allow for intragroup correlation (cluster clustvar), and that use bootstrap or jackknife methods (bootstrap, jackknife); see [R] vce_option.

level(#) ; see [R] estimation options.
aequations specifies that the results for the outcome-model or treatment-model parameters be displayed. By default, the results for these auxiliary parameters are not displayed.
noshow prevents stteffects ipwra from showing the key st variables. This option is rarely used because most people type stset, show or stset, noshow to permanently set whether they want to see these variables mentioned at the top of the output of every st command; see [ST] stset.
display_options: noci, nopvalues, nomitted, vsquish, nomemptycells, baselevels, allbaselevels, nofvlabel, fvwrap(#), fvwrapon(style), cformat(%fmt), pformat(%fmt), sformat(%fmt), and nolstretch; see [R] estimation options.

Maximization

maximize_options: iterate(#), [no]log, and from(init_specs); see [R] maximize. These options are seldom used.
init_specs is one of
matname [ , skip copy ]
# [, # ... ] , copy
iterinit(#) specifies the maximum number of iterations used to calculate the starting values. This option is seldom used.

Advanced

pstolerance(#) specifies the tolerance used to check the overlap assumption. The default value is pstolerance(1e-5). stteffects will exit with an error if an observation has an estimated propensity score smaller than that specified by pstolerance().

osample(newvar) specifies that indicator variable newvar be created to identify observations that violate the overlap assumption.
control(\# | label) specifies the level of \texttt{tvar} that is the control. The default is the first treatment level. You may specify the numeric level \# (a nonnegative integer) or the label associated with the numeric level. \texttt{control()} may not be specified with the statistic \texttt{pmeans}. \texttt{control()} and \texttt{tlevel()} may not specify the same treatment level.

tlevel(\# | label) specifies the level of \texttt{tvar} that is the treatment for the statistic \texttt{atet}. The default is the second treatment level. You may specify the numeric level \# (a nonnegative integer) or the label associated with the numeric level. \texttt{tlevel()} may only be specified with statistic \texttt{atet}. \texttt{tlevel()} and \texttt{control()} may not specify the same treatment level.

The following option is available with \texttt{stteffects} but is not shown in the dialog box: \texttt{coeflegend}; see \cite{R} \texttt{estimation options}.

Remarks and examples

If you are not familiar with the framework for treatment-effects estimation from observational survival-time data, please see \cite[TE]{stteffects intro}.

IPWRA estimators use estimated weights to obtain missingness-adjusted outcome-regression parameters. The missingness-adjusted outcome-regression parameters are used to compute averages of treatment-level predicted outcomes. Contrasts of these averages estimate the treatment effects.

The estimated weights account for the missing potential outcome and, optionally, for data lost to censoring. The weights are estimated using a treatment-assignment model and, optionally, a model for the censoring time. A term in the estimator for the outcome-regression parameters accounts for data lost to censoring when estimated weights are not used.

There are two versions of the IPWRA estimator because there are two methods of accounting for the data lost to censoring.

1. IPWRA estimators that adjust for censoring by including a term in the likelihood function for the outcome-model parameters are known as likelihood-adjusted-censoring IPWRA (LAC-IPWRA) estimators.

2. IPWRA estimators that adjust for censoring by weighting the likelihood function for the outcome-model parameters by estimated inverse-probability-of-censoring weights are known as weighted-adjusted-censoring IPWRA (WAC-IPWRA) estimators.

The LAC-IPWRA estimators require fewer assumptions than the WAC-IPWRA estimators. Outlining the steps performed by LAC-IPWRA and WAC-IPWRA estimators allows us to be more specific about the trade-offs between the estimators.

LAC-IPWRA estimators use a three-step approach to estimating treatment effects:

1. Estimate the parameters of a treatment-assignment model and compute inverse-probability-of-treatment weights.

2. Obtain the treatment-specific predicted mean outcomes for each subject by using the weighted maximum likelihood estimators. Estimated inverse-probability-of-treatment weights are used to weight the maximum likelihood estimator. A term in the likelihood function adjusts for right-censored survival times.

3. Compute the means of the treatment-specific predicted mean outcomes. Contrasts of these averages provide the estimates of the ATEs. By restricting the computations of the means to the subset of treated subjects, we can obtain the ATETs.
WAC-IPWRA estimators use a four-step approach to estimating treatment effects:

1. Estimate the parameters of a treatment-assignment model and compute inverse-probability-of-treatment weights.
2. Estimate the parameters of a time-to-censoring model and compute inverse-probability-of-censoring weights.
3. Obtain the treatment-specific predicted mean outcomes for each subject by using the weighted maximum likelihood estimators. Estimated inverse-probability-of-treatment weights and inverse-probability-of-censoring weights are used to weight the maximum likelihood estimator. The inverse-probability-of-censoring weights account for right-censored survival times.
4. Compute the means of the treatment-specific predicted mean outcomes. Contrasts of these averages provide the estimates of the ATEs. By restricting the computations of the means to the subset of treated subjects, we can obtain the ATETs.

The WAC-IPWRA estimators require that the censoring time be random and that the time-to-censoring model be well specified. The implemented WAC-IPWRA estimators also require that the time-to-censoring process not vary by treatment level. The LAC-IPWRA estimators do not require these extra assumptions because they use a likelihood term instead of weights to adjust for the data lost to censoring.


Like streg and other survival-time commands, stteffects ipwra uses the outcome variable and the failure indicator computed by, and optionally weights specified with, stset. stteffects ipwra is not appropriate for data with time-varying covariates, also known as multiple-record survival-time data, or for delayed-entry data.

Example 1: Estimating the ATE by LAC-IPWRA

Suppose we wish to study the effect of smoking on the time to a second heart attack among women aged 45–55 years. In our fictional sheart dataset, atime is the observed time in years to a second heart attack or censoring, and fail is the 0/1 indicator that a second heart attack was observed. (When fail is 1, atime records the time to the second heart attack; when fail is 0, atime records a censored observation of the time to a second heart attack.) We previously stset these data; see A quick tour of the estimators in [TE] stteffects intro.

The treatment, smoking, is stored in the 0/1 indicator smoke. These data also contain age at the time of the first heart attack (age), and indices of the level of exercise (exercise), diet quality (diet), and education (education) prior to the first heart attack.

We can use stteffects ipwra to estimate the ATE. We model the mean survival time using the default Weibull model, controlling for age, exercise, diet, and education. We model treatment assignment using the default logit model with covariates age, exercise, and education. We do not specify a time-to-censoring model so that we obtain the LAC estimator.
**use http://www.stata-press.com/data/r15/sheart**

(Time to second heart attack (fictional))

```
. stteffects ipwra (age exercise diet education) (smoke age exercise education)
    failure _d: fail
    analysis time _t: atime
Iteration 0:   EE criterion =  2.432e-16
Iteration 1:   EE criterion =  1.021e-29
Survival treatment-effects estimation  Number of obs   =    2,000
Estimator : IPW regression adjustment
Outcome model : Weibull
Treatment model: logit
Censoring model: none
```

| _t  | Robust Coef. | Std. Err. | z   | P>|z|  | [95% Conf. Interval] |
|-----|--------------|-----------|-----|------|----------------------|
| ATE | smoke        | -1.591874 | .4837332 | -3.29 | 0.001 | -2.539973 to -.643774 |
|     | (Smoker vs Nonsmoker) |          |         |       |       |                      |
| POmean | smoke | 4.214263 | .2598689 | 16.22 | 0.000 | 3.704929 to 4.723597  |
|     | Nonsmoker   |           |         |       |       |                      |

When every woman smoked in the population of women aged 45–55 years who have had a heart attack, the average time to a second heart attack is estimated to be 1.59 years less than when no women in the population of interest smoked. The estimated average time to a second heart attack when no women in the population of interest smoked is 4.21 years.

The ratio of the ATE to the control-level potential-outcome mean (POM) measures the importance of the effect. In this example, when all women smoked, the time to the second heart attack falls by an estimated 38% relative to the case in which no women smoked. See example 3 in [TE] stteffects ra for an example that uses nlcom to compute a point estimate and a confidence interval for this ratio.
Example 2: Different outcome and treatment models

Instead of a Weibull model for the outcome model, we could have used an exponential, a gamma, or a lognormal model. Instead of a logit model for the treatment assignment, we could have used a probit or a heteroskedastic probit model. This example uses a gamma model for the outcome and a probit model for the treatment assignment.

```
   . stteffects ipwra (age exercise diet education, gamma)
   > (smoke age exercise education, probit)
       failure _d: fail
       analysis time _t: atime
Iteration 0: EE criterion = 2.644e-13
Iteration 1: EE criterion = 2.153e-23
Survival treatment-effects estimation  Number of obs = 2,000
   Estimator : IPW regression adjustment
   Outcome model : gamma
   Treatment model: probit
   Censoring model: none

  t        Coef.   Std. Err.     z  P>|z|     [95% Conf. Interval]
----------- ------- -------- -------- -------- ------------------
   ATE       smoke        (Smoker vs Nonsmoker)
                 -1.387303 .4786032  -2.90  0.004  -2.325348 -.4492583
   POmean     smoke        Nonsmoker
                 3.97986  .2258474   17.62  0.000   3.537207 4.422512
```

The estimated ATE of \(-1.39\) and control-level POM of 3.98 are similar to the values of \(-1.59\) and 4.21 that we obtained in example 1.
Example 3: Estimating the ATE by WAC-IPWRA

Rather than using LAC, we may want to specify a time-to-censoring model. We now use `stteffects ipwra` to estimate the ATE by WAC-IPWRA. We use the same specification of the outcome and treatment models that we used in example 1. However, now we specify a time-to-censoring model, using the default Weibull model with covariates `age`, `exercise`, `diet`, and `education`.

```
. stteffects ipwra (age exercise diet education) (smoke age exercise education)>
    > (age exercise diet education)
    failure _d: fail
    analysis time _t: atime
Iteration 0: EE criterion = 1.217e-17
Iteration 1: EE criterion = 9.176e-31
```

Survival treatment-effects estimation Number of obs = 2,000
Estimator : IPW regression adjustment
Outcome model : Weibull
Treatment model: logit
Censoring model: Weibull

|             | Coef.     | Std. Err. | z    | P>|z|   | [95% Conf. Interval] |
|-------------|-----------|-----------|------|-------|---------------------|
| _t          |           | Robust    |      |       |                     |
| ATE         |           |           |      |       |                     |
| smoke       | -2.285057 | .7318456  | -3.12| 0.002 | -3.719448 -.8506656 |
| (Smoker vs  |           |           |      |       |                     |
| Nonsmoker)  |           |           |      |       |                     |
| POmean      |           |           |      |       |                     |
| smoke       | 4.385841  | .6427521  | 6.82 | 0.000 | 3.12607  5.645612   |
| Nonsmoker   |           |           |      |       |                     |

The estimated ATE of \(-2.29\) differs from the ATE of \(-1.59\) estimated by LAC-IPWRA, but the estimates of the control-level POM are similar between the two models: 4.39 for the WAC compared with 4.21 for the LAC.
Example 4: Estimating the ATET by LAC-IPWRA

Intuitively, the ATET measures the effect of the treatment on an at-risk subpopulation. Sometimes the subpopulation that gets the treatment defines such an at-risk subpopulation. The ATET has the added benefit that it can be estimated under weaker conditions than the ATE; see Assumptions and trade-offs under Remarks and examples in [TE] stteffects intro.

```
stteffects ipwra (age exercise diet education)
> (smoke age exercise education), atet

failure _d: fail
analysis time _t: atime

Iteration 0: EE criterion = 2.671e-18
Iteration 1: EE criterion = 1.638e-30

Survival treatment-effects estimation
Number of obs = 2,000
Estimator : IPW regression adjustment
Outcome model : Weibull
Treatment model: logit
Censoring model: none

| _t | Robust Coef. | Std. Err. | z   | P>|z|       | [95% Conf. Interval] |
|----|--------------|-----------|-----|----------|---------------------|
| ATET | smoke | (Smoker vs Non-smoker) | -1.775107     | .3437506  | -5.16   | 0.000    | -2.448846 | -1.101368 |
| POmean | smoke | Non-smoker | 4.062424       | .2779877  | 14.61   | 0.000    | 3.517578  | 4.60727 |
```

When all women in the subpopulation smoked, the average time to a second heart attack is estimated to be 1.78 years less than when no women in the subpopulation of interest smoked. If no women in the subpopulation of interest smoked, the average time to a second heart attack is 4.06 years.

Stored results

`stteffects ipwra` stores the following in `e()`:

Scalars

- `e(N)`: number of observations
- `e(nj)`: number of observations for treatment level `j`
- `e(N_clust)`: number of clusters
- `e(k_eq)`: number of equations in `e(b)`
- `e(k_levels)`: number of levels in treatment variable
- `e(treated)`: level of treatment variable defined as treated
- `e(control)`: level of treatment variable defined as control
- `e(converged)`: 1 if converged, 0 otherwise

Macros

- `e(cmd)`: `stteffects`
- `e(cmdline)`: command as typed
- `e(dead)`: `_d`
- `e(depvar)`: `_t`
- `e(tvar)`: name of treatment variable
Methods and formulas

Methods and formulas are presented under the following headings:

Introduction
Regression-adjusted estimators
Weighted-adjusted-censoring assumptions
Inverse-probability-weighted estimators
Uncensored data
Inverse-probability-weighted regression-adjustment estimators
Weighted-adjusted-censoring IPWRA
Likelihood-adjusted-censoring IPWRA
Functional-form details

Introduction

This section presents the methods and formulas used by the estimators implemented in stteffects ra, stteffects wra, stteffects ipw, and stteffects ipwra. This section assumes that you are familiar with the concepts and intuition from the estimators discussed in [TE] teffects intro advanced.

Each of the estimators implemented in stteffects has a multistep logic but is implemented as one step by simultaneously solving the estimating equations that define each step. This one-step estimating-equation approach provides consistent point estimates and a consistent variance–covariance of the estimator (VCE); see Newey (1984), Wooldridge (2010), and Drukker (2014).

Survival-time treatment-effects estimators handle two types of missing data. First, only one of the potential outcomes is observed, as is standard in causal inference. Second, the potential outcome for the received treatment may be censored. The data missing because of censoring may be handled by an outcome model, a censoring model, or both, just like the data missing due to observing only one potential outcome.
Technical note

Delayed entry would be a third type of missing data. The left-truncation process caused by delayed entry would also need to be modeled to estimate ATE parameters. The estimators implement in stteffects do not allow for delayed entry because they do not have a method for modeling how the left-truncation process selects the sample, conditional on the covariates.

All the implemented estimators are combinations of regression-adjustment (RA) and inverse-probability-weighted (IPW) techniques. RA estimators use an outcome model to account for the missing potential outcome and for censoring. IPW estimators use models for treatment assignment and censoring to construct weights that account for the missing potential outcome and for censoring.

The remainder of this section provides technical details about how the estimators in stteffects were implemented. We provide details only for the two-treatment-level case to simplify the formulas. We provide outlines for how the extensions to the multiple-treatment-level case were implemented.

Regression-adjusted estimators

We begin with the RA estimators implemented in stteffects ra. The RA estimators have the following logic:

RA1. For each treatment level \( \tau \in \{0, 1\} \), estimate by maximum likelihood (ML) the parameters \( \beta_\tau \) of a parametric model for the survival-time outcome \( t \) in which \( F(t|x, \tau, \beta_\tau) \) is the distribution of \( t \) conditional on covariates \( x \) and treatment level \( \tau \). Denote the estimates \( \hat{\beta}_\tau \) by \( \hat{\beta}_{ra, \tau} \).

RA2. Use the estimated \( \hat{\beta}_{ra, \tau} \) and the functional form implied by \( F(t|x, \tau, \beta_\tau) \) to estimate the mean survival time, conditional on \( x \) and treatment level \( \tau \), for each sample observation, denoted by \( \hat{E}(t_i|x_i, \tau, \hat{\beta}_{ra, \tau}) \). Conditional independence of the treatment and the survival-time potential outcomes ensures that \( E(t|x, \tau, \beta_\tau) = E(t_\tau|x, \beta_\tau) \), where \( t_\tau \) is the potential survival-time outcome corresponding to treatment level \( \tau \). Under correct model specification, sample averages of \( \hat{E}(t_i|x_i, \tau, \hat{\beta}_{ra, \tau}) \) consistently estimate the POM for treatment level \( \tau \), denoted by \( POM_\tau \).

RA3. A contrast of the estimated POMs estimates the ATE.

If estimating an ATET, step RA2 is modified to use only the treated observations when estimating the POMs. A contrast of these POMs then estimates the ATET.

The contribution of the \( i \)th observation to the log likelihood that is maximized in step RA1 is

\[
L_{ra}(t_i, x_i, \tau, \hat{\beta}_{ra, \tau}) = \bar{\omega}_i(\tau_i == \tau) \left[ (1 - c_i) \ln \{ f(t_i|x_i, \tau, \hat{\beta}_{ra, \tau}) \} + c_i \ln \{ 1 - F(t_i|x_i, \tau, \hat{\beta}_{ra, \tau}) \} \right]
\]

where \( \bar{\omega}_i \) is the observation-level weight, \( c_i \) is the 0/1 indicator for whether the survival-time observation on person \( i \) was censored, and \( f(t_i|x_i, \tau, \hat{\beta}_{ra, \tau}) \) is the density corresponding to distribution \( F(t_i|x_i, \tau, \hat{\beta}_{ra, \tau}) \). The first term inside the curly braces in (1) accounts for the noncensored observations, and the second term inside the curly braces accounts for the censored observations.
The RA estimators for the POMs simultaneously solve estimating equations (2a) through (2d) for \( \hat{\beta}_{ra,0}, \hat{\beta}_{ra,1}, \hat{\text{POM}}_{ra,0}, \) and \( \hat{\text{POM}}_{ra,1} \).

\[
\frac{1}{N} \sum_{i=1}^{N} s_{ra}(t_i, x_i, 0, \hat{\beta}_{ra,0}, F) = 0 \tag{2a}
\]

\[
\frac{1}{N} \sum_{i=1}^{N} s_{ra}(t_i, x_i, 1, \hat{\beta}_{ra,1}, F) = 0 \tag{2b}
\]

\[
\frac{1}{N} \sum_{i=1}^{N} \varpi_i \left\{ \hat{E}(t_i|\mathbf{x}_i, \tau = 0, \hat{\beta}_{ra,0}) - \hat{\text{POM}}_{ra,0} \right\} = 0 \tag{2c}
\]

\[
\frac{1}{N} \sum_{i=1}^{N} \varpi_i \left\{ \hat{E}(t_i|\mathbf{x}_i, \tau = 1, \hat{\beta}_{ra,1}) - \hat{\text{POM}}_{ra,1} \right\} = 0 \tag{2d}
\]

where

\[
s_{ra}(t_i, x_i, 0, \hat{\beta}_{ra,0}, F) = \frac{\partial L_{ra}(t_i, x_i, 0, \hat{\beta}_{ra,0})}{\partial \beta_{ra,0}}
\]

is the vector of score equations from the ML estimator for \( \hat{\beta}_{ra,0} \) based on survival-time model \( F \),

\[
s_{ra}(t_i, x_i, 1, \hat{\beta}_{ra,1}, F) = \frac{\partial L_{ra}(t_i, x_i, 1, \hat{\beta}_{ra,1})}{\partial \beta_{ra,1}}
\]

is the vector of score equations from the ML estimator for \( \hat{\beta}_{ra,1} \) based on survival-time model \( F \),

\( \hat{E}(t_i|\mathbf{x}_i, \tau = 0, \hat{\beta}_{ra,0}) \) is the predicted mean survival time assuming treatment level 0 for observation \( i \) conditional on \( \mathbf{x}_i \), and

\( \hat{E}(t_i|\mathbf{x}_i, \tau = 1, \hat{\beta}_{ra,1}) \) is the predicted mean survival time assuming treatment level 1 for observation \( i \) conditional on \( \mathbf{x}_i \).

The ATE is estimated by replacing (2d) with

\[
\frac{1}{N_1} \sum_{i=1}^{N} \varpi_i \left\{ \hat{E}(t_i|\mathbf{x}_i, \tau = 1, \hat{\beta}_{ra,1}) - \hat{\text{POM}}_{ra,0} - \hat{\text{ATE}}_{ra} \right\} = 0 \tag{3}
\]

and the ATET is estimated by replacing (2c) and (3) with

\[
\frac{1}{N_1} \sum_{i=1}^{N} \varpi_i(\tau_i == 1) \left\{ \hat{E}(t_i|\mathbf{x}_i, \tau = 0, \hat{\beta}_{ra,0}) - \hat{\text{POM}}_{ra,cot,0} \right\} = 0
\]

\[
\frac{1}{N_1} \sum_{i=1}^{N} \varpi_i(\tau_i == 1) \left\{ \hat{E}(t_i|\mathbf{x}_i, \tau = 1, \hat{\beta}_{ra,1}) - \hat{\text{POM}}_{ra,cot,0} - \hat{\text{ATE}}_{ra} \right\} = 0
\]

where \( N_1 = \sum_{i=1}^{N} (\tau_i == 1) \) and \( \hat{\text{POM}}_{ra,cot,0} \) is the estimated conditional-on-treatment POM for treatment level 0.
Asymptotic standard errors for estimating equation estimators, also known as exactly identified generalized method of moments estimators, are standard in the literature; see Newey (1984), Newey and McFadden (1994), Tsiatis (2006), and Wooldridge (2010). These standard errors always have a robust structure and have been generalized to cluster–robust standard errors (see Wooldridge [2010]).

The score equations and the functional form for the predicted mean survival time depend on the model for survival-time outcome $F$. We provide these details below, under *Functional-form details*. 

**Weighted-adjusted-censoring assumptions**

All estimators that permit you to model the time to censoring are subject to three assumptions:

1. The censoring time must be random.
2. The censoring time must be from a known distribution.
3. The distribution of the censoring time cannot vary by treatment level.

We call these three requirements the WAC assumptions. If the WAC assumptions are violated, you can use either an RA estimator or the LAC version of the IPWRA estimator.

---

### Technical note

We now describe how the observed survival-time outcome $t$ is generated from the random censoring time $t_c$, the received treatment $\tau$, and the potential-outcome survival times $t_0$ and $t_1$ under the WAC assumptions. First, each potential outcome is either censored or not censored.

$$
\tilde{t}_0 = t_c(t_0 \geq t_c) + t_0\{1 - (t_0 \geq t_c)\}
$$

$$
\tilde{t}_1 = t_c(t_1 \geq t_c) + t_1\{1 - (t_1 \geq t_c)\}
$$

Under the WAC assumptions, $t_c$ is a random variable from a known distribution, and $t_c$ does not vary by treatment level.

Next, the received treatment $\tau \in \{0, 1\}$ determines which, possibly censored, potential outcome is observed.

$$
t = (1 - \tau)\tilde{t}_0 + \tau\tilde{t}_1
$$

The 0/1 indicator for whether the observed $t$ was censored, denoted by $c$, is given by

$$
c = (1 - \tau)(t_0 \geq t_c) + \tau(t_1 \geq t_c)
$$

---

### Weighted regression-adjusted estimators

As is standard in the survival literature, the RA estimators account for censored survival times by adding a term to the log-likelihood function for censored observations [see (1)]. In contrast, weighted regression-adjustment (WRA) estimators use weights to account for censored observations and are subject to the WAC assumptions.

Wooldridge (2007) and Lin (2000) derived estimators for the regression parameters that maximize a weighted objective function of the uncensored observations. Each observation-level weight is the inverse of the probability of not being censored. Like the RA estimators, the WRA estimators use averages of the predicted mean survival times to estimate treatment-effect parameters.
The WRA estimators have the following logic.

WRA1. Estimate by ML the parameters $\gamma$ of a parametric survival-time model for the time to censoring $t_c$, in which $F_c(t_c | w, \gamma)$ is the distribution of $t_c$ conditional on covariates $w$. Note that the censoring process does not vary by treatment level and that we only observe $t_c$ when the observed potential outcome was censored. Denote the estimates of $\gamma$ by $\hat{\gamma}$.

WRA2. For each treatment level $\tau \in \{0, 1\}$, estimate by weighted maximum likelihood (WML) the $\beta_\tau$ parameters of a parametric survival-time model, denoted by $F(t|x, \tau, \beta_\tau)$, where $t$ is the survival-time outcome and $x$ are the covariates. The weights are the inverse of the estimated probabilities of not being censored, $1/[1 - F_c(t_c | w, \hat{\gamma})]$, and only the uncensored observations are used. Denote the estimates of $\beta_\tau$ by $\hat{\beta}_{wra, \tau}$.

WRA3. Use the estimated $\hat{\beta}_{wra, \tau}$ and the functional form implied by $F(t | x, \tau, \beta_\tau)$ to estimate the mean survival time, conditional on $x$ and treatment level $\tau$, for each sample observation, denoted by $\hat{E}(t_i | x_i, \tau, \hat{\beta}_{wra, \tau})$. Conditional independence of the treatment and the survival-time potential outcomes ensures that $E(t_i | x_i, \tau, \beta_\tau) = E(t_\tau | x, \beta_\tau)$, where $t_\tau$ is the potential survival-time outcome corresponding to treatment level $\tau$. Under correct model specification, sample averages of $\hat{E}(t_i | x_i, \tau, \hat{\beta}_{wra, \tau})$ consistently estimate the POM for treatment level $\tau$, denoted by $\text{POM}_\tau$.

WRA4. A contrast of the estimated POMs estimates the ATE.

If estimating an ATET, step WRA3 is modified to use only the treated observations when estimating the POMs. A contrast of these POMs then estimates the ATET.

The contribution of the $i$th observation to the log likelihood that is maximized in step WRA1 is

$$L_{c,wra}(t_i, w_i, \hat{\gamma}) = \omega_i [c_i \ln \{f_c(t_i | w_i, \hat{\gamma})\} + (1 - c_i) \ln \{1 - F_c(t_i | w_i, \hat{\gamma})\}]$$

(4)

where $\omega_i$ is the observation-level weight, $c_i$ is the 0/1 indicator for whether the survival-time observation on person $i$ was censored, $t_i$ is the observed failure time, and $f_c(t_i | w_i, \hat{\gamma})$ is the density corresponding to conditional time-to-censoring distribution $F_c(t_i | w_i, \hat{\gamma})$. When $c_i = 1$, $t_i$ is the time to censoring. When $c_i = 0$, the censoring time is not observed; we only know that it is greater than the observed $t_i$. The first term accounts for the observations in which $t_i$ is observed to be the censoring time, and the second term accounts for the observations in which the censoring time is greater than the observed $t_i$.

The contribution of the $i$th observation to the log likelihood that is maximized in step WRA2 is

$$L_{wra}(t_i, x_i, \tau, \hat{\beta}_{wra, \tau}) = \omega_i (\tau_i = \tau) \left[ \frac{(1 - c_i)}{\{1 - F_c(t_i | w_i, \hat{\gamma})\}} \right] \ln \{f(t_i | x_i, \tau, \hat{\beta}_{wra, \tau})\}$$

(5)

where $f(t_i | x_i, \tau, \hat{\beta}_{wra, \tau})$ is the density corresponding to distribution $F(t_i | x_i, \tau, \hat{\beta}_{wra, \tau})$. Equation (5) does not contain a term that adjusts for censoring; see (1) for a comparison. Rather, it uses inverse-probability weights to account for both the censored and the uncensored observations.

The WRA estimators for the POMs simultaneously solve estimating equations (6a) through (6e) for $\hat{\gamma}$, $\hat{\beta}_{wra, 0}$, $\hat{\beta}_{wra, 1}$, $\text{POM}_{wra, 0}$, and $\text{POM}_{wra, 1}$.

$$1/N \sum_{i=1}^{N} s_{wra}(t_i, w_i, \hat{\gamma}, F_c) = 0$$

(6a)

$$1/N \sum_{i=1}^{N} s_{wra}(t_i, x_i, 0, \hat{\beta}_{wra, 0}, F) = 0$$

(6b)
\[
1/N \sum_{i=1}^{N} s_{wra}(t_i, x_i, 1, \hat{\beta}_{wra,1}, F) = 0 \tag{6c}
\]

\[
1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i | x_i, \tau = 0, \hat{\beta}_{wra,0}) - \widehat{\text{POM}}_{wra,0} \right\} = 0 \tag{6d}
\]

\[
1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i | x_i, \tau = 1, \hat{\beta}_{wra,1}) - \widehat{\text{POM}}_{wra,1} \right\} = 0 \tag{6e}
\]

where

\[ s_{wra}(t_i, w_i, \gamma, F_c) = \frac{\partial L_{c,wra}(t_i, w_i, \gamma)}{\partial \gamma} \]

is the vector of score equations from the ML estimator for \( \gamma \) based on survival-time model \( F_c \),

\[ s_{wra}(t_i, x_i, 0, \hat{\beta}_{wra,0}, F) = \frac{\partial L(t_i, x_i, 0, \hat{\beta}_{wra,0})}{\partial \hat{\beta}_{wra,0}} \]

is the vector of score equations from the WML estimator for \( \hat{\beta}_{wra,0} \) based on survival-time model \( F \),

\[ s_{wra}(t_i, x_i, 1, \hat{\beta}_{wra,1}, F) = \frac{\partial L(t_i, x_i, 1, \hat{\beta}_{wra,1})}{\partial \hat{\beta}_{wra,1}} \]

is the vector of score equations from the WML estimator for \( \hat{\beta}_{wra,1} \) based on survival-time model \( F \),

\[ \hat{E}(t_i | x_i, \tau = 0, \hat{\beta}_{wra,0}) \]

is the predicted mean survival time assuming treatment level 0 for observation \( i \) conditional on \( x_i \), and

\[ \hat{E}(t_i | x_i, \tau = 1, \hat{\beta}_{wra,1}) \]

is the predicted mean survival time assuming treatment level 1 for observation \( i \) conditional on \( x_i \).

The observation-level scores \( s_{wra}(t_i, x_i, 0, \hat{\beta}_{wra,0}, F) \) and \( s_{wra}(t_i, x_i, 1, \hat{\beta}_{wra,1}, F) \) also depend on \( c_i, w_i, \gamma \), and \( F_c \), but we ignored this dependence to simplify the notation.

The ATE is estimated by replacing (6e) with

\[
1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i | x_i, \tau = 1, \hat{\beta}_{wra,1}) - \widehat{\text{POM}}_{wra,0} - \hat{\text{ATE}}_{wra} \right\} = 0 \tag{7}
\]

and the ATET is estimated by replacing (6d) and (7) with

\[
1/N_1 \sum_{i=1}^{N} \omega_i(\tau_i = 1) \left\{ \hat{E}(t_i | x_i, \tau = 0, \hat{\beta}_{wra,0}) - \widehat{\text{POM}}_{wra,cot,0} \right\} = 0
\]

\[
1/N_1 \sum_{i=1}^{N} \omega_i(\tau_i = 1) \left\{ \hat{E}(t_i | x_i, \tau = 1, \hat{\beta}_{wra,1}) - \widehat{\text{POM}}_{wra,cot,0} - \hat{\text{ATE}}_{wra} \right\} = 0
\]

where \( \widehat{\text{POM}}_{wra,cot,0} \) is the estimated conditional-on-treatment POM.
Inverse-probability-weighted estimators

IPW estimators are weighted averages of the observed outcome. The weights correct for missing data due to unobserved potential outcomes and censoring. Each weight is the inverse of the probability that a given value is observed. Observed values that were not likely to be observed have higher weights.

When the outcome variable is never censored, the missing data is the unobserved potential outcome and an observation’s weight is the inverse of a treatment probability. When the outcome may be censored, the censoring is an additional source of missing data. In this case, an observation’s weight is the inverse of the joint probability that an observation is uncensored and has a particular treatment level.

To define this joint probability, the censoring time must be random. In practice, we make the WAC assumptions.

As is standard in the survival-time literature, we assume that the censoring-time process is independent of treatment assignment after conditioning on the covariates. This conditional independence assumption implies that the probability that observation $i$ receives treatment level 1 and is not censored is the product of the probability that $i$ gets treatment level 1 and the probability that $i$ is not censored at time $t_i$, which we denote by

$$p(z_i, \alpha)\{1 - F_c(t_i|w_i, \gamma)\}$$

where

$p(z_i, \alpha)$ is the modeled probability that $i$ gets treatment level 1, conditional on covariates $z_i$ with parameters $\alpha$, and

$F_c(t_i|w_i, \gamma)$ is the survival-time model for the censoring time, conditional on covariates $w_i$ with parameters $\gamma$, and evaluated at time $t_i$.

Bai, Tsiatis, and O’Brien (2013) formally derive these weights to control jointly for the missing potential outcome and censoring.

The IPW estimators have the following logic.

IPW1. Estimate by ML the parameters $\gamma$ of a parametric survival-time model for the time to censoring, in which $F_c(t_c|w, \gamma)$ is the distribution of censoring time, conditional on covariates $w$. Denote the estimates of $\gamma$ by $\hat{\gamma}$.

IPW2. Estimate by ML the parameters $\alpha$ of a parametric model for the probability of treatment model $p(z_i, \alpha)$. Denote the estimates of $\alpha$ by $\hat{\alpha}$.

IPW3. Use the $\hat{\gamma}$ estimated in IPW1 and the $\hat{\alpha}$ estimated in IPW2 to construct inverse-probability weights by (8a) for treatment level 1 and by (8b) for treatment level 0.

$$\omega_{i,1} = \frac{(\tau_i == 1)(c_i == 0)}{[p(z_i, \hat{\alpha})\{1 - F_c(t_i|w_i, \hat{\gamma})\}]} \quad (8a)$$

$$\omega_{i,0} = \frac{(\tau_i == 0)(c_i == 0)}{[\{1 - p(z_i, \hat{\alpha})\}\{1 - F_c(t_i|w_i, \hat{\gamma})\}]} \quad (8b)$$

IPW4. Use the estimated weights to estimate each $POM_\tau$ by a weighted average of the uncensored observations on the observed potential outcome.
The contribution of the \( i \)th observation to the log likelihood that is maximized in step IPW1 is

\[
L_{c, \text{ipw}}(t_i, w_i, \hat{\gamma}) = c_i \ln \{ f_c(t_i | w_i, \hat{\gamma}) \} + (1 - c_i) \ln \{ 1 - F_c(t_i | w_i, \hat{\gamma}) \}
\]

where the definitions and intuition are as described after (4).

The contribution of the \( i \)th observation to the log likelihood that is maximized in step IPW2 is

\[
L_{p, \text{ipw}}(\tau_i, z_i, \hat{\alpha}) = c_i \ln \{ p(z_i, \hat{\alpha}) \} + (1 - c_i) \ln \{ 1 - p(z_i, \hat{\alpha}) \}
\]

where \( p(z_i, \hat{\alpha}) \) is the model for the probability that \( i \) gets treatment level 1.

The IPW estimators for the POMs simultaneously solve estimating equations (9a) through (9d) for \( \hat{\gamma}, \hat{\alpha}, \text{POM}_{\text{ipw}, 0}, \) and \( \text{POM}_{\text{ipw}, 1} \).

\[
\frac{1}{N} \sum_{i=1}^{N} s_{\text{ipw}}(t_i, w_i, \hat{\gamma}, F_c) = 0 \quad (9a)
\]

\[
\frac{1}{N} \sum_{i=1}^{N} s_{\text{ipw}}(\tau_i, z_i, \hat{\alpha}, p) = 0 \quad (9b)
\]

\[
\frac{1}{N} \sum_{i=1}^{N} \omega_i \omega_{i, 0} (t_i - \text{POM}_{\text{ipw}, 0}) = 0 \quad (9c)
\]

\[
\frac{1}{N} \sum_{i=1}^{N} \omega_i \omega_{i, 1} (t_i - \text{POM}_{\text{ipw}, 1}) = 0 \quad (9d)
\]

where

\[
s_{\text{ipw}}(t_i, w_i, \hat{\gamma}, F_c) = \frac{\partial L_{c, \text{ipw}}(t_i, w_i, \hat{\gamma})}{\partial \hat{\gamma}}
\]

is the vector of score equations from the ML estimator for \( \hat{\gamma} \) based on survival-time model \( F_c \), and

\[
s_{\text{ipw}}(\tau_i, z_i, \hat{\alpha}, p) = \frac{\partial L_{p, \text{ipw}}(\tau_i, z_i, \hat{\alpha})}{\partial \hat{\alpha}}
\]

is the vector of score equations from the ML estimator for \( \hat{\alpha} \) based on probability model \( p \).

The literature on IPW estimators discusses using normalized versus unnormalized weights, with normalized weights doing better in simulation studies; see Busso, DiNardo, and McCrary (2014) for example. The way that weights enter moment equations (9c) and (9d) implies that they are normalized, because the scale of the weights does not affect the estimates.

The estimated ATE is computed as

\[
\text{POM}_{\text{ipw}, 1} - \text{POM}_{\text{ipw}, 0} = \hat{\text{ATE}}_{\text{ipw}}
\]

The estimated ATET uses weights

\[
\omega_{i, \text{cot}, 1} = \frac{(\tau_i == 1)(c_i == 0)}{\{1 - F_c(t_i | w_i, \hat{\gamma})\}}
\]

for treatment level 1 and

\[
\omega_{i, \text{cot}, 0} = \frac{p(z_i, \hat{\alpha})(\tau_i == 0)(c_i == 0)}{\{1 - p(z_i, \hat{\alpha})\}\{1 - F_c(t_i | w_i, \hat{\gamma})\}}
\]
stteffects ipwra — Survival-time inverse-probability-weighted regression adjustment

for treatment level 0, and replaces (9c) and (9d) with

\[
\frac{1}{N_1} \sum_{i=1}^{N} \omega_{i,\text{cot},0} (t_i - \hat{\text{POM}}_{\text{ipw},\text{cot},0}) = 0 \quad (11a)
\]

\[
\frac{1}{N_1} \sum_{i=1}^{N} \omega_{i,\text{cot},1} (t_i - \hat{\text{POM}}_{\text{ipw},\text{cot},1}) = 0 \quad (11b)
\]

and then computes

\[
\hat{\text{POM}}_{\text{ipw},\text{cot},1} - \hat{\text{POM}}_{\text{ipw},\text{cot},0} = \hat{\text{ATET}}_{\text{ipw}}
\]

These IPW estimators can be viewed as weighted IPW estimators and are thus related to those in Hirano, Imbens, and Ridder (2003).

Uncensored data

As mentioned, when the outcome variable is never censored, the missing data is the unobserved potential outcome and an observation’s weight is the inverse of a treatment probability. In the never-censored case, the IPW estimators are identical to those implemented in teffects ipw; see IPW estimators under Methods and formulas in [TE] teffects aipw.

stteffects ipw computes the estimator for never-censored data when a censoring model is not specified and there are no censored observations in the sample. In the never-censored case, the following changes are made to the IPW estimator for the POMs and the ATE.

1. Step IPW1 is not performed.
2. The weights in (8a) and (8b) for the POMs and the ATE are replaced with (12a) for treatment level 1 and (12b) for treatment level 0.

\[
\omega_{i,1} = \frac{(\tau_i = 1)}{p(z_i, \hat{\alpha})} \quad (12a)
\]

\[
\omega_{i,0} = \frac{(\tau_i = 0)}{\{1 - p(z_i, \hat{\alpha})\}} \quad (12b)
\]

3. Only moment conditions (9b), (9c), and (9d) are used.

The following changes also are made to the IPW estimator for the ATET.

1. Step IPW1 is not performed.
2. The weights in (10a) and (10b) are replaced with (13a) for treatment level 1 and (13b) for treatment level 0.

\[
\omega_{i,\text{cot},1} = (\tau_i = 1) \quad (13a)
\]

\[
\omega_{i,\text{cot},0} = \frac{p(z_i, \hat{\alpha})(\tau_i = 0)}{\{1 - p(z_i, \hat{\alpha})\}} \quad (13b)
\]

3. Only moment conditions (9b), (11a), and (11b) are used.
Inverse-probability-weighted regression-adjustment estimators

IPWRA estimators are averages of treatment-specific predicted conditional means that were made using missingness-adjusted regression parameters. These estimators are Wooldridge’s IPWRA for survival-time outcomes; see Wooldridge (2010, chap. 21) and Wooldridge (2007).

The censored observations can be handled either by weighting under the WAC assumptions to obtain the WAC-IPWRA estimator or by adding a term to the log-likelihood function (which we call likelihood-adjusted censoring) to obtain the LAC-IPWRA estimator. Correspondingly, there are two versions of formulas for the IPWRA estimator.

1. When a censoring model is specified, stteffects ipwra uses the formulas for the WAC-IPWRA estimator given in Weighted-adjusted-censoring IPWRA.

2. When a censoring model is not specified, stteffects ipwra uses the formulas for the LAC-IPWRA given in Likelihood-adjusted-censoring IPWRA, below.

The WAC-IPWRA estimator requires that some observations be censored and that the WAC assumptions hold; see Weighted-adjusted-censoring assumptions, above. The LAC-IPWRA estimator handles the case in which no observations are censored and requires the weaker independent censoring assumptions, which allows for fixed censoring times.

Weighted-adjusted-censoring IPWRA

When a censoring model is specified, stteffects ipwra uses the formulas for the WAC-IPWRA estimator to obtain the model-based weights that account for censoring. For notational conciseness and to reinforce its dependence on random censoring, we denote the WAC-IPWRA estimator by IPWRAR. The WAC-IPWRA estimators have the following logic.

IPWRAR1. Estimate by ML the parameters $\gamma$ of a parametric survival-time model for the time to censoring, in which $F_c(t_c|w, \gamma)$ is the censoring-time distribution, conditional on covariates $w$. We denote the estimates of $\gamma$ by $\hat{\gamma}$.

IPWRAR2. Estimate by ML the parameters $\alpha$ of a parametric model for the probability of treatment model $p(z_i, \alpha)$. We denote the estimates of $\alpha$ by $\hat{\alpha}$.

IPWRAR3. For each treatment level $t \in \{0, 1\}$, estimate by WML the parameters $\beta_t$ of a parametric model for the survival-time outcome $t$, in which $F(t|x, \tau, \beta_t)$ is the distribution of $t$ conditional on covariates $x$ and treatment level $\tau$. For the ATE, the weights are those in equations (8a) and (8b). For the ATET, the weights are those in equations (10a) and (10b). We denote the estimates of $\beta_{ipwra, \tau}$ by $\hat{\beta}_t$.

IPWRAR4. Use the estimated $\hat{\beta}_{ipwra, \tau}$ and the functional form implied by $F(t|x, \tau, \beta_t)$ to estimate the mean survival time, conditional on $x$ and treatment level $\tau$, for each sample observation, denoted by $\hat{E}(t_i|x_i, \tau, \hat{\beta}_{ipwra, \tau})$. Conditional independence of the treatment and the survival-time potential outcomes ensures that $E(t|x, \tau, \beta_t) = E(t|\tau|x, \beta_t)$, where $t_\tau$ is the potential survival-time outcome corresponding to treatment level $\tau$. Under correct model specification, sample averages of $\hat{E}(t_i|x_i, \tau, \hat{\beta}_{ipwra, \tau})$ consistently estimate the POM for treatment level $\tau$, denoted by $POM_\tau$.

The contribution of the $i$th observation to the log likelihood that is maximized in step IPWRAR1 is

$$L_{c,ipwra}(t_i, w_i, \hat{\gamma}) = \omega_i [c_i \ln \{f_c(t_i|w_i, \hat{\gamma})\} + (1 - c_i) \ln \{1 - F_c(t_i|w_i, \hat{\gamma})\}]$$

where the definitions and intuition are as described after (4).
The contribution of the $i$th observation to the log likelihood that is maximized in step IPWRAR2 is

$$L_{p, ipwrar}(\tau_i, z_i, \hat{\alpha}) = \omega_i \left\{ (\tau_i == 1) \ln \{ p(z_i, \hat{\alpha}) \} + \{ 1 - (\tau_i == 1) \} \ln \{ 1 - p(z_i, \hat{\alpha}) \} \right\}$$

where $p(z_i, \hat{\alpha})$ is the model for the probability that $i$ gets treatment level 1.

The weights and the parameters in step IPWRAR3 used to estimate the ATE differ from those used to estimate the ATET. For the ATE, the contribution of the $i$th observation to the log likelihood that is maximized in step IPWRAR3 is

$$L_{i, ipwrar}(t_i, x_i, \tau, \hat{\beta}_{ipwrar, ate, \tau}) = \omega_i \omega_i, \tau \ln \{ f(t_i | x_i, \tau, \hat{\beta}_{ipwrar, ate, \tau}) \}$$

where $\omega_{i,1}$ is given in (8a), $\omega_{i,0}$ is given in (8b), and $f(t_i | x_i, \tau, \hat{\beta}_{ipwrar, ate, \tau})$ is the density corresponding to distribution $F(t_i | x_i, \tau, \hat{\beta}_{ipwrar, ate, \tau})$. Like WRA, only the uncensored observations are used because the weights account for censoring.

The IPWRAR estimators for the POMs simultaneously solve estimating equations (14a) through (14f) for $\hat{\gamma}$, $\hat{\alpha}$, $\hat{\beta}_{ipwrar, ate, 0}$, $\hat{\beta}_{ipwrar, ate, 1}$, $\hat{\beta}_{ipwrar, ate, 0}$, $\hat{\beta}_{ipwrar, ate, 1}$, $\hat{\beta}_{ipwrar, ate, 0}$, and $\hat{\beta}_{ipwrar, ate, 1}$.

$$
\begin{align*}
1/N \sum_{i=1}^{N} s_{ipwrar}(t_i, w_i, \hat{\gamma}, F_c) &= 0 \quad (14a) \\
1/N \sum_{i=1}^{N} s_{ipwrar}(\tau_i, z_i, \hat{\alpha}, p) &= 0 \quad (14b) \\
1/N \sum_{i=1}^{N} s_{ipwrar}(t_i, x_i, 0, \hat{\beta}_{ipwrar, ate, 0}, F) &= 0 \quad (14c) \\
1/N \sum_{i=1}^{N} s_{ipwrar}(t_i, x_i, 1, \hat{\beta}_{ipwrar, ate, 1}, F) &= 0 \quad (14d) \\
1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i | x_i, \tau = 0, \hat{\beta}_{ipwrar, ate, 0}) - \hat{\beta}_{ipwrar, ate, 0} \right\} &= 0 \quad (14e) \\
1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i | x_i, \tau = 1, \hat{\beta}_{ipwrar, ate, 1}) - \hat{\beta}_{ipwrar, ate, 1} \right\} &= 0 \quad (14f)
\end{align*}
$$

where

$$s_{ipwrar}(t_i, w_i, \hat{\gamma}, F_c) = \frac{\partial L_{c, ipwrar}(t_i, w_i, \hat{\gamma})}{\partial \hat{\gamma}}$$

is the vector of score equations from the ML estimator for $\hat{\gamma}$ based on survival-time model $F_c$,

$$s_{ipwrar}(\tau_i, z_i, \hat{\alpha}, p) = \frac{\partial L_{p, ipwrar}(\tau_i, z_i, \hat{\alpha})}{\partial \hat{\alpha}}$$

is the vector of score equations from the ML estimator for $\hat{\alpha}$ based on probability model $p$,

$$s_{ipwrar}(t_i, x_i, 0, \hat{\beta}_{ipwrar, ate, 0}, F) = \frac{\partial L_{ipwrar}(t_i, x_i, 0, \hat{\beta}_{ipwrar, ate, 0})}{\partial \hat{\beta}_{ipwrar, ate, 0}}$$

is the vector of score equations from the ML estimator for $\hat{\beta}_{ipwrar, ate, 0}$ based on survival-time model $F$. 
\[ s_{\text{ipwrar}}(t_i, x, 1, \hat{\beta}_{\text{ipwrar,ate}}, F) = \frac{\partial L_{\text{ipwrar}}(t_i, x, 1, \hat{\beta}_{\text{ipwrar,ate}}, 1)}{\partial \hat{\beta}_{\text{ipwrar,ate}}, 1} \]

is the vector of score equations from the ML estimator for \( \hat{\beta}_{\text{ipwrar,ate}}, 1 \) based on survival-time model \( F \),

\[ \hat{E}(t_i | x, \tau = 0, \hat{\beta}_{\text{ipwrar,ate}, 0}) \]

is the predicted mean survival time assuming treatment level 0 for observation \( i \) conditional on \( x \), and

\[ \hat{E}(t_i | x, \tau = 1, \hat{\beta}_{\text{ipwrar,ate}, 1}) \]

is the predicted mean survival time assuming treatment level 1 for observation \( i \) conditional on \( x \).

The ATE is estimated by replacing (14f) with

\[ 1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i | x, \tau = 1, \hat{\beta}_{\text{ipwrar,ate}, 1}) - \hat{POM}_{\text{ipwrar}, 0} - \hat{ATE}_{\text{ipwrar}} \right\} = 0 \]

For the ATET, the contribution of the \( i \)th observation to the weighted log likelihood that is maximized in step IPWRAR3 is

\[ L_{\text{ipwrar}}(t_i, x, \tau, \hat{\beta}_{\text{ipwrar,ate}, \tau}) = \omega_i \omega_{i, \text{cot}, \tau} (\tau_i = \tau) \ln \{ f(t_i | x, \tau, \hat{\beta}_{\text{ipwrar,atet}, \tau}) \} \]

where \( \omega_{i, \text{cot}, 1} \) is given in (10a), \( \omega_{i, \text{cot}, 0} \) is given in (10b), and \( f(t_i | x, \tau, \hat{\beta}_{\text{ipwrar,atet}, \tau}) \) is the density corresponding to distribution \( F(t_i | x, \tau, \hat{\beta}_{\text{ipwrar,atet}, \tau}) \).

The WAC-IPWRA estimators for the conditional-on-treatment POMs simultaneously solve estimating equations (15a) through (15f) for \( \hat{\beta}_{\text{ipwrar,atet}, 0}, \hat{\beta}_{\text{ipwrar,atet}, 0}, \hat{\gamma}, \hat{\alpha}, \hat{\beta}_{\text{ipwrar,atet}, \tau}, \beta_{\text{ipwrar,atet}, \tau}, \) and \( \hat{POM}_{\text{ipwrar,atet}, 0}, \hat{POM}_{\text{ipwrar,atet}, 0}, \) and \( \hat{POM}_{\text{ipwrar,atet}, 0} \).

\[
\begin{align*}
1/N \sum_{i=1}^{N} s_{\text{ipwrar}}(t_i, w, \hat{\gamma}, F_c) &= 0 \quad (15a) \\
1/N \sum_{i=1}^{N} s_{\text{ipwrar}}(\tau_i, z, \hat{\alpha}, p) &= 0 \quad (15b) \\
1/N \sum_{i=1}^{N} s_{\text{ipwrar}}(t_i, x_i, 0, \hat{\beta}_{\text{ipwrar,atet}, 0}, F) &= 0 \quad (15c) \\
1/N \sum_{i=1}^{N} s_{\text{ipwrar}}(t_i, x_i, 1, \hat{\beta}_{\text{ipwrar,atet}, 1}, F) &= 0 \quad (15d) \\
1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i | x, \tau = 0, \hat{\beta}_{\text{ipwrar,atet}, 0}) - \hat{POM}_{\text{ipwrar,atet}, 0} \right\} &= 0 \quad (15e) \\
1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i | x, \tau = 1, \hat{\beta}_{\text{ipwrar,atet}, 1}) - \hat{POM}_{\text{ipwrar,atet}, 1} \right\} &= 0 \quad (15f)
\end{align*}
\]

where

\[ s_{\text{ipwrar}}(t_i, w, \hat{\gamma}, F_c) = \frac{\partial L_{\text{ipwrar}}(t_i, w, \hat{\gamma})}{\partial \hat{\gamma}} \]

is the vector of score equations from the ML estimator for \( \hat{\gamma} \) based on survival-time model \( F_c \).
\( s_{\text{ipwrar}}(\tau, z_i, \hat{\alpha}, p) = \frac{\partial L_{\text{ipwrar}}(\tau, z_i, \hat{\alpha})}{\partial \alpha} \) is the vector of score equations from the ML estimator for \( \hat{\alpha} \) based on probability model \( p \),

\[ s_{\text{ipwrar}}(t_i, x_i, 0, \hat{\beta}_{\text{ipwrar}, \text{atet}, 0}, F) = \frac{\partial L_{\text{ipwrar}}(t_i, x_i, 0, \hat{\beta}_{\text{ipwrar}, \text{atet}, 0})}{\partial \beta_{\text{ipwrar}, \text{atet}, 0}} \] is the vector of score equations from the WML estimator for \( \hat{\beta}_{\text{ipwrar}, \text{atet}, 0} \) based on survival-time model \( F \),

\[ s_{\text{ipwrar}}(t_i, x_i, 1, \hat{\beta}_{\text{ipwrar}, \text{atet}, 1}, F) = \frac{\partial L_{\text{ipwrar}}(t_i, x_i, 1, \hat{\beta}_{\text{ipwrar}, \text{atet}, 1})}{\partial \beta_{\text{ipwrar}, \text{atet}, 1}} \] is the vector of score equations from the WML estimator for \( \hat{\beta}_{\text{ipwrar}, \text{atet}, 1} \) based on survival-time model \( F \),

\( \hat{E}(t_i | x_i, \tau = 0, \hat{\beta}_{\text{ipwrar}, \text{atet}, 0}) \) is the predicted mean survival time assuming treatment level 0 for observation \( i \) conditional on \( x_i \), and

\( \hat{E}(t_i | x_i, \tau = 1, \hat{\beta}_{\text{ipwrar}, \text{atet}, 1}) \) is the predicted mean survival time assuming treatment level 1 for observation \( i \) conditional on \( x_i \).

The ATET is estimated by replacing (15f) with

\[ 1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i | x_i, \tau = 1, \hat{\beta}_{\text{ipwrar}, \text{atet}, 1}) - \hat{\text{POM}}_{\text{ipwrar}, \text{cot}, 0} - \text{ATET}_{\text{ipwrar}} \right\} = 0 \]

**Likelihood-adjusted-censoring IPRA**

When a censoring model is not specified, \texttt{stteffects ipwr} uses the formulas for the LAC-IPRA estimator that add a term to the log-likelihood function. For notational conciseness and to reinforce its use of an additional term in the log likelihood, we denote the LAC-IPRA estimator by IPRA in lists and formulas.

The methods and formulas for the LAC-IPRA estimator differ in three ways from those for the WAC-IPRA estimator.

1. No censoring model is specified, so LAC-IPRA does not perform a version of \texttt{step IPWRAR1} and it does not use the moment equations (14a).
2. The weights only depend on the treatment level and treatment assignment probabilities, not on the censoring.
3. The WML estimator for \( \beta_\tau \) includes a term for censored observations and censored observations are used. Recall that for the WAC-IPRA estimator, the weights used in the WML estimator for \( \beta_\tau \) account for the censoring, and the censored observations are not used in the WML estimator.

The LAC-IPRA estimators have the following logic.

**IPWAL1.** Estimate by ML the parameters \( \alpha \) of a parametric model for the probability of treatment model \( p(z_i, \alpha) \).

**IPWAL2.** For each treatment level \( \tau \in \{0, 1\} \), estimate by WML the parameters \( \beta_\tau \) of a parametric model for the survival-time outcome \( t \) in which \( F(t | x, \tau, \beta_\tau) \) is the distribution of \( t \) conditional on covariates \( x \) and treatment level \( \tau \). The weights depend only on the treatment level and the treatment-assignment probabilities. For the ATE, the weights are those in (12a) and (12b). For the ATET, the weights are those in (13a) and (13b). We denote the estimates of \( \beta_\tau \) by \( \hat{\beta}_{\text{ipwral}, \tau} \).
IPWRAL3. Use the estimated \( \hat{\beta}_{ipwral, \tau} \) and the functional form implied by \( F(t|x, \tau, \beta_{\tau}) \) to estimate the mean survival time, conditional on \( x \) and treatment level \( \tau \), for each sample observation, denoted by \( \hat{E}(t_i|x_i, \tau, \hat{\beta}_{ipwral, \tau}) \). Conditional independence of the treatment and the survival-time potential outcomes ensures that \( E(t|x, \tau, \beta_{\tau}) = E(t_{\tau}|x, \beta_{\tau}) \), where \( t_{\tau} \) is the potential survival-time outcome corresponding to treatment level \( \tau \). Under correct model specification, sample averages of \( \hat{E}(t_i|x_i, \tau, \hat{\beta}_{ipwral, \tau}) \) consistently estimate the POM for treatment level \( \tau \), denoted by \( POM_{\tau} \).

The contribution of the \( i \)th observation to the log likelihood that is maximized in step \( \text{IPWRAL1} \) is

\[
L_{p, \text{ipwral}}(\tau_i, z_i, \hat{\alpha}) = \omega_i \left[ (\tau_i == 1) \ln \left\{ p(z_i, \hat{\alpha}) \right\} \right.
\]

where \( p(z_i, \hat{\alpha}) \) is the model for the probability that \( i \) gets treatment level 1.

The weights and the parameters in step \( \text{IPWRAL2} \) used to estimate the ATE differ from those used to estimate the ATET. For the ATE, the contribution of the \( i \)th observation to the log likelihood that is maximized in step \( \text{IPWRAL2} \) is

\[
L_{\text{ipwral}}(t_i, x_i, \tau, \hat{\beta}_{\text{ipwral,ate,} \tau}) = (\tau_i == \tau) \omega_i \omega_{i, \tau} \left\{ (1 - c_i) \ln \left\{ f(t_i|x_i, \tau, \hat{\beta}_{\text{ipwral,ate,} \tau}) \right\} \right.
\]

\[
\left. + c_i \ln \left\{ 1 - \hat{E}(t_i|x_i, \tau, \hat{\beta}_{\text{ipwral,ate,} \tau}) \right\} \right\}
\]

where \( \omega_{i,1} \) is given in (12a), \( \omega_{i,0} \) is given in (12b), and \( f(t_i|x_i, \tau, \hat{\beta}_{\text{ipwral,ate,} \tau}) \) is the density corresponding to distribution \( F(t_i|x_i, \tau, \hat{\beta}_{\text{ipwral,ate,} \tau}) \). Unlike the WRA estimator, the censored observations are used, and there is a term in the likelihood function to account for censoring.

The LAC-IPWRA estimators for the POMs simultaneously solve estimating equations (16a) through (16e) for \( \hat{\alpha}, \hat{\beta}_{\text{ipwral,ate,} 0}, \hat{\beta}_{\text{ipwral,ate,} 1}, \text{POM}_{\text{ipwral,0}}, \) and \( \text{POM}_{\text{ipwral,1}} \).

\[
1/N \sum_{i=1}^{N} s_{\text{ipwral}}(\tau_i, z_i, \hat{\alpha}, p) = 0 \quad (16a)
\]

\[
1/N \sum_{i=1}^{N} s_{\text{ipwral}}(t_i, x_i, 0, \hat{\beta}_{\text{ipwral,ate,} 0}, F) = 0 \quad (16b)
\]

\[
1/N \sum_{i=1}^{N} s_{\text{ipwral}}(t_i, x_i, 1, \hat{\beta}_{\text{ipwral,ate,} 1}, F) = 0 \quad (16c)
\]

\[
1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i|x_i, \tau = 0, \hat{\beta}_{\text{ipwral,ate,} 0}) - \text{POM}_{\text{ipwral,0}} \right\} = 0 \quad (16d)
\]

\[
1/N \sum_{i=1}^{N} \omega_i \left\{ \hat{E}(t_i|x_i, \tau = 1, \hat{\beta}_{\text{ipwral,ate,} 1}) - \text{POM}_{\text{ipwral,1}} \right\} = 0 \quad (16e)
\]

where

\[
s_{\text{ipwral}}(\tau_i, z_i, \hat{\alpha}, p) = \frac{\partial L_{p, \text{ipwral}}(\tau_i, z_i, \hat{\alpha})}{\partial \hat{\alpha}}
\]

is the vector of score equations from the ML estimator for \( \hat{\alpha} \) based on probability model \( p \).
The ATE is estimated by replacing \( \widehat{\beta}_{\text{ipwral}, \text{ate}, 0} \) with \( \widehat{\beta}_{\text{ipwral}, \text{ate}, 1} \) based on survival-time model \( F \), and \( \widehat{\beta}_{\text{ipwral}, \text{ate}, 1} \) is the predicted mean survival time assuming treatment level 1 for observation \( i \) conditional on \( x_i \).

The LAC-IPWRA estimators for the conditional-on-treatment POMs simultaneously solve estimating equations (17a) through (17e) for \( \hat{\alpha} \), \( \hat{\beta}_{\text{ipwral}, \text{ate}, 0} \), \( \hat{\beta}_{\text{ipwral}, \text{ate}, 1} \), \( \widehat{\text{POM}}_{\text{ipwral}, \text{cot}, 0} \), and \( \widehat{\text{POM}}_{\text{ipwral}, \text{cot}, 1} \).
In this section, we specify the functional forms for the conditional distribution function used in the survival-time outcome model $F$, the conditional distribution function used in the survival-time censoring model $F_c$, and the conditional distribution used to model the treatment probabilities $p$.

You may choose among the same set of conditional distribution functions for either $F$ or $F_c$: exponential, Weibull, log normal, or gamma.

<table>
<thead>
<tr>
<th>Name</th>
<th>Cumulative Function</th>
<th>Density Function</th>
<th>Mean</th>
</tr>
</thead>
<tbody>
<tr>
<td>exponential</td>
<td>$1 - \exp(-\lambda_i t_i)$</td>
<td>$\lambda_i \exp(-\lambda_i t_i)$</td>
<td>$1/\lambda_i$</td>
</tr>
<tr>
<td>Weibull</td>
<td>$1 - \exp{-(-\lambda_i t_i)^{s_i}}$</td>
<td>$s_i t_i^{s_i - 1} \lambda_i^{s_i} \exp{-(-\lambda_i t_i)^{s_i}}$</td>
<td>$(1/\lambda_i) \Gamma{(s_i + 1)/s_i}$</td>
</tr>
<tr>
<td>log normal</td>
<td>$\Phi{\ln(t_i) - \lambda_i/s_i}$</td>
<td>$(1/(s_i t_i)) \phi{\ln(t_i) - \lambda_i/s_i}$</td>
<td>exp($\lambda_i + s_i^2/2$)</td>
</tr>
<tr>
<td>gamma</td>
<td>$\gammamap{s_i, (s_i t_i/\lambda_i)}$</td>
<td>$(s_i^{s_i} t_i^{s_i - 1})/{\lambda_i^{s_i} \Gamma(s_i)} \exp(-s_i t_i/\lambda_i)$</td>
<td>$\lambda_i$</td>
</tr>
</tbody>
</table>

where the following table specifies how $\lambda_i$ and $s_i$ are parameterized in terms of the covariates $x_i$ and the ancillary covariates $\tilde{x}_i$, respectively.

<table>
<thead>
<tr>
<th>Name</th>
<th>$\lambda_i$</th>
<th>$s_i$</th>
</tr>
</thead>
<tbody>
<tr>
<td>exponential</td>
<td>$\exp(-x_i \beta)$</td>
<td>$s_i$</td>
</tr>
<tr>
<td>Weibull</td>
<td>$\exp(-x_i \beta)$</td>
<td>$\exp(\tilde{x}_i \tilde{\beta})$</td>
</tr>
<tr>
<td>log normal</td>
<td>$x_i \beta$</td>
<td>$\exp(\tilde{x}_i \tilde{\beta})$</td>
</tr>
<tr>
<td>gamma</td>
<td>$\exp(x_i \beta)$</td>
<td>$\exp(-2\tilde{x}_i \tilde{\beta})$</td>
</tr>
</tbody>
</table>
For the treatment-assignment models, the probit model uses the standard normal distribution, the logit uses the standard logistic distribution, the hetprobit model uses

$$\Phi(z_1\alpha_1/\exp(z_2\alpha_2))$$

and the multinomial logit uses

$$p(z, t) = \exp(z\alpha_t) / \{1 + \sum_{k=1}^{q} \exp(z\alpha_k)\}$$

where the notation is defined below.

In the hetprobit model, $z_1$ are the covariates specified in the treatment-assignment specification, $z_2$ are the covariates specified in the hetprobit() option, and $\alpha_1$ and $\alpha_2$ are the corresponding coefficients.

In the multinomial logit model, $z$ are the covariates specified in the treatment-assignment specification and $\alpha_k$ are the coefficients; see [R] mlogit for further details.

References


Also see

[TE] `stteffects postestimation` — Postestimation tools for `stteffects`

[TE] `stteffects intro` — Introduction to treatment effects for observational survival-time data

[ST] `streg` — Parametric survival models

[ST] `stset` — Declare data to be survival-time data

[U] 20 Estimation and postestimation commands
## Postestimation commands

The following postestimation commands are of special interest after `stteffects`:

<table>
<thead>
<tr>
<th>Command</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>teffects overlap</code></td>
<td>overlap plots</td>
</tr>
<tr>
<td><code>tebalance</code></td>
<td>check balance of covariates</td>
</tr>
</tbody>
</table>

The following standard postestimation commands are also available:

<table>
<thead>
<tr>
<th>Command</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>estat summarize</code></td>
<td>summary statistics for the estimation sample</td>
</tr>
<tr>
<td><code>estat vce</code></td>
<td>variance–covariance matrix of the estimators (VCE)</td>
</tr>
<tr>
<td><code>estimates</code></td>
<td>cataloging estimation results</td>
</tr>
<tr>
<td><code>hausman</code></td>
<td>Hausman’s specification test</td>
</tr>
<tr>
<td><code>lincom</code></td>
<td>point estimates, standard errors, testing, and inference for linear combinations of coefficients</td>
</tr>
<tr>
<td><code>nlcom</code></td>
<td>point estimates, standard errors, testing, and inference for nonlinear combinations of coefficients</td>
</tr>
<tr>
<td><code>predict</code></td>
<td>predictions, residuals, influence statistics, and other diagnostic measures</td>
</tr>
<tr>
<td><code>predictnl</code></td>
<td>point estimates, standard errors, testing, and inference for generalized predictions</td>
</tr>
<tr>
<td><code>test</code></td>
<td>Wald tests of simple and composite linear hypotheses</td>
</tr>
<tr>
<td><code>testnl</code></td>
<td>Wald tests of nonlinear hypotheses</td>
</tr>
</tbody>
</table>
predict

Description for predict

predict creates a new variable containing predictions such as treatment effects, conditional means, propensity scores, linear predictions, and log square roots of latent variances.

Menu for predict

Statistics ➤ Postestimation

Syntaxes for predict

Syntaxes are presented under the following headings:

- Syntax for predict after stteffects ipw
- Syntax for predict after stteffects ipwra
- Syntax for predict after stteffects ra
- Syntax for predict after stteffects wra

Syntax for predict after stteffects ipw

```
predict [type] { stub* | newvar | newvarlist } [if] [in]
[ , statistic tlevel(treat_level) ]
```

```
predict [type] { stub* | newvarlist } [if] [in], scores
```

<table>
<thead>
<tr>
<th>statistic</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>ps</td>
<td>propensity score; the default</td>
</tr>
<tr>
<td>censurv</td>
<td>censored survival probability</td>
</tr>
<tr>
<td>xb</td>
<td>linear prediction for propensity score</td>
</tr>
<tr>
<td>cb</td>
<td>linear prediction for censoring model</td>
</tr>
<tr>
<td>lnsigma</td>
<td>log square root of latent variance (for treatment model hetprobit())</td>
</tr>
<tr>
<td>clnshape</td>
<td>log of conditional latent shape (for censoring distribution Weibull, log normal, or gamma)</td>
</tr>
</tbody>
</table>

If you do not specify tlevel() and only specify one new variable, ps assumes tlevel() specifies the first treatment level.

If you do not specify tlevel() and only specify one new variable, xb and lnsigma assume tlevel() specifies the first noncontrol treatment level.

You specify one or \( t \) new variables with ps, where \( t \) is the number of treatment levels.

You specify one or \( t-1 \) new variables with xb and lnsigma.

You specify one new variable with censurv, cb, and clnshape.
## Syntax for predict after stteffects ipwra

```
predict [\{type\}] \{stub* | newvar | newvarlist\} [if] [in]
[\, statistic tlevel(\text{treat\_level})]
```

```
predict [\{type\}] \{stub* | newvarlist\} [if] [in], scores
```

<table>
<thead>
<tr>
<th>statistic</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>\text{te}</td>
<td>treatment effect; the default</td>
</tr>
<tr>
<td>\text{cmean}</td>
<td>conditional mean at treatment level</td>
</tr>
<tr>
<td>\text{ps}</td>
<td>propensity score</td>
</tr>
<tr>
<td>\text{censurv}</td>
<td>censored survival probability</td>
</tr>
<tr>
<td>\text{xb}</td>
<td>linear prediction for outcome model</td>
</tr>
<tr>
<td>\text{cxb}</td>
<td>linear prediction for censoring model</td>
</tr>
<tr>
<td>\text{psxb}</td>
<td>linear prediction for propensity score</td>
</tr>
<tr>
<td>\text{lnshape}</td>
<td>log of conditional latent shape (for outcome distribution Weibull, log normal, or gamma) at treatment level</td>
</tr>
<tr>
<td>\text{clnshape}</td>
<td>log of conditional latent shape (for censoring distribution Weibull, log normal, or gamma)</td>
</tr>
<tr>
<td>\text{pslnsigma}</td>
<td>log square root of latent variance (for treatment model \text{hetprobit}) for propensity score</td>
</tr>
</tbody>
</table>

If you do not specify \text{tlevel}() and only specify one new variable, \text{te} and \text{psxb} assume \text{tlevel}() specifies the first noncontrol treatment level.

If you do not specify \text{tlevel}() and only specify one new variable, \text{cmean}, \text{ps}, \text{xb}, and \text{pslnsigma} assume \text{tlevel}() specifies the first treatment level.

You specify one or \text{t} new variables with \text{cmean}, \text{ps}, \text{xb}, and \text{lnshape}, where \text{t} is the number of treatment levels.

You specify one or \text{t}–\text{1} new variables with \text{te}, \text{psxb}, and \text{pslnsigma}.

You specify one new variable with \text{censurv}, \text{cxb}, and \text{clnshape}. 
### Syntax for predict after stteffects ra

```
predict [type] { stub* | newvar | newvarlist } [if] [in]
    [, statistic tlevel(treat_level) ]
```

```
predict [type] { stub* | newvarlist } [if] [in], scores
```

<table>
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<tr>
<th>statistic</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>te</code></td>
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</tr>
<tr>
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<td>conditional mean at treatment level</td>
</tr>
<tr>
<td><code>xb</code></td>
<td>linear prediction for outcome model</td>
</tr>
<tr>
<td><code>lnshape</code></td>
<td>log of conditional latent shape (for outcome distribution Weibull, log normal, or gamma) at treatment level</td>
</tr>
</tbody>
</table>

If you do not specify `tlevel()` and only specify one new variable, `te` assumes `tlevel()` specifies the first noncontrol treatment level.

If you do not specify `tlevel()` and only specify one new variable, `cmean`, `xb`, and `lnshape` assume `tlevel()` specifies the first treatment level.

You specify one or `t` new variables with `cmean`, `xb`, and `lnshape`, where `t` is the number of treatment levels.

You specify one or `t-1` new variables with `te`.

### Syntax for predict after stteffects wra

```
predict [type] { stub* | newvar | newvarlist } [if] [in]
    [, statistic tlevel(treat_level) ]
```

```
predict [type] { stub* | newvarlist } [if] [in], scores
```

<table>
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If you do not specify `tlevel()` and only specify one new variable, `te` assumes `tlevel()` specifies the first noncontrol treatment level.

If you do not specify `tlevel()` and only specify one new variable, `cmean`, `xb`, and `lnshape` assume `tlevel()` specifies the first treatment level.

You specify one or `t` new variables with `cmean`, `xb`, and `lnshape`, where `t` is the number of treatment levels.

You specify one or `t-1` new variables with `te`.

You specify one new variable with `censurv`, `cxb`, and `clnshape`. 
Options for predict

Options are presented under the following headings:

- Options for predict after stteffects ipw
- Options for predict after stteffects ipwra
- Options for predict after stteffects ra
- Options for predict after stteffects wra

Options for predict after stteffects ipw

- **ps**, the default, calculates the propensity score of each treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

- **censurv** calculates the survivor probability from the time-to-censoring model. (In other words, it calculates the probability that an outcome is not censored.) This option is allowed only if a censoring model is specified at estimation time. You need to specify only one new variable.

- **xb** calculates the propensity score linear prediction at each noncontrol level of the treatment or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level (except the control level).

- **cxb** calculates the linear prediction of the censoring model. This option is allowed only if a censoring model is specified at estimation time. You need to specify only one new variable.

- **lnsigma** calculates the log square root of the latent variance. This option is valid only when treatment model `hetprobit()` is used. You need to specify only one new variable.

- **clnshape** calculates the log of the conditional latent shape parameter of the censoring distribution. This option is valid when censoring distribution Weibull, log normal, or gamma is used. You need to specify only one new variable.

- **tlevel(treat_level)** specifies the treatment level for prediction.

- **scores** calculates the score variables. Parameter-level scores are computed for the treatment mean and average treatment-effect equations. Equation-level scores are computed for the censoring and propensity-score equations.

   The jth new variable will contain the scores for the jth parameter in the coefficient table if \( j \leq t \), where \( t \) is the number of treatment levels. Otherwise, it will contain the scores for fitted equation \( j - t \) following the first \( t \) parameters in the coefficient table.

Options for predict after stteffects ipwra

- **te**, the default, calculates the treatment effect for each noncontrol treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level (except the control level).

- **cmean** calculates the conditional mean for each treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.
ps calculates the propensity score of each treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

censurv calculates the survivor probability from the time-to-censoring model. (In other words, it calculates the probability that an outcome is not censored.) This option is allowed only if a censoring model is specified at estimation time. You need to specify only one new variable.

xb calculates the outcome model linear prediction at each treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

cxb calculates the linear prediction of the censoring model. This option is allowed only if a censoring model is specified at estimation time. You need to specify only one new variable.

psxb calculates the propensity score linear prediction at each noncontrol level of the treatment or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level (except the control level).

lnshape calculates the log of the conditional latent shape parameter for each treatment level or the treatment level specified in `tlevel()`. This option is valid when outcome distribution Weibull, log normal, or gamma is used. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

clnshape calculates the log of the conditional latent shape parameter for the censoring distribution. This option is valid when censoring distribution Weibull, log normal, or gamma is used. You need to specify only one new variable.

pslnsigma calculates the log square root of the latent variance for the propensity score. This option is valid only when treatment model `hetprobit()` is used. You need to specify only one new variable.

`tlevel(treat_level)` specifies the treatment level for prediction.

scores calculates the score variables. Parameter-level scores are computed for the treatment mean and average treatment-effect equations. Equation-level scores are computed for the outcome, censoring, and propensity-score equations.

The `j`th new variable will contain the scores for the `j`th parameter in the coefficient table if `j ≤ t`, where `t` is the number of treatment levels. Otherwise, it will contain the scores for fitted equation `j − t` following the first `t` parameters in the coefficient table.

Options for `predict` after `stteffects ra`:

- **Main**
  - **te**, the default, calculates the treatment effect for each noncontrol treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.
  - **cmean** calculates the conditional mean for each treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.
  - **xb** calculates the outcome model linear prediction at each treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.
\texttt{lnshape} calculates the log of the conditional latent shape parameter for each treatment level or the treatment level specified in \texttt{tlevel()}. This option is valid when the outcome distribution Weibull, log normal, or gamma is used. If you specify the \texttt{tlevel()} option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

\texttt{tlevel(treat\_level)} specifies the treatment level for prediction.

\texttt{scores} calculates the score variables. Parameter-level scores are computed for the treatment mean and average treatment-effect equations. Equation-level scores are computed for the outcome equations. The \textit{j}th new variable will contain the scores for the \textit{j}th parameter in the coefficient table if \textit{j} \leq \textit{t}, where \textit{t} is the number of treatment levels. Otherwise, it will contain the scores for fitted equation \textit{j} – \textit{t} following the first \textit{t} parameters in the coefficient table.

\section*{Options for predict after \texttt{stteffects wra}}

- \texttt{te}, the default, calculates the treatment effect for each noncontrol treatment level or the treatment level specified in \texttt{tlevel()}. If you specify the \texttt{tlevel()} option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level (except the control level).

- \texttt{cmean} calculates the conditional mean for each treatment level or the treatment level specified in \texttt{tlevel()}. If you specify the \texttt{tlevel()} option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

- \texttt{censurv} calculates the survivor probability from the time-to-censoring model. (In other words, it calculates the probability that an outcome is not censored.) This option is allowed only if a censoring model is specified at estimation time. You need to specify only one new variable.

- \texttt{xb} calculates the outcome model linear prediction at each treatment level or the treatment level specified in \texttt{tlevel()}. If you specify the \texttt{tlevel()} option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

- \texttt{lnshape} calculates the log of the conditional latent shape parameter for each treatment level or the treatment level specified in \texttt{tlevel()}. This option is valid when the outcome distribution Weibull, log normal, or gamma is used. If you specify the \texttt{tlevel()} option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

- \texttt{clnshape} calculates the log of the conditional latent shape parameter of the censoring distribution. This option is valid when the censoring distribution Weibull, log normal, or gamma is used. You need to specify only one new variable.

- \texttt{tlevel(treat\_level)} specifies the treatment level for prediction.

- \texttt{scores} calculates the score variables. Parameter-level scores are computed for the treatment mean and average treatment-effect equations. Equation-level scores are computed for the outcome and censoring equations. The \textit{j}th new variable will contain the scores for the \textit{j}th parameter in the coefficient table if \textit{j} \leq \textit{t}, where \textit{t} is the number of treatment levels. Otherwise, it will contain the scores for fitted equation \textit{j} – \textit{t} following the first \textit{t} parameters in the coefficient table.
Remarks and examples

Checking model specification is the most frequent reason for postestimation computation after \texttt{stteffects}. \texttt{teffects overlap} provides a graphical method for checking the overlap assumption; see \texttt{[TE] teffects overlap}. Summarizing the estimated probabilities provides another check. Recall that the reciprocals of these estimated probabilities are used as weights by some of the estimators. If the estimated probabilities are too small, the weights get too large and the estimators become unstable.

We estimate the average treatment effect of smoking on the time to a second heart attack by inverse-probability weighting; see example 1 of \texttt{[TE] stteffects ipw} for background.

```
. use http://www.stata-press.com/data/r15/sheart
   (Time to second heart attack (fictional))
. stteffects ipw (smoke age exercise education) (age exercise diet education)
   failure _d: fail
   analysis time _t: atime
Iteration 0:   EE criterion = 2.042e-18
Iteration 1:   EE criterion = 5.191e-31
Survival treatment-effects estimation Number of obs    =     2,000
   Estimator : inverse-probability weights
   Outcome model : weighted mean
   Treatment model: logit
   Censoring model: Weibull

                     _t | Coef.  Std. Err.     z  P>|z|     [95% Conf. Interval]
-----------|------------------|------------------|------|---------|------------------|------------------|
        ATE |               |
   smoke   | (Smoker vs Nonsmoker) | -2.22226   .6307573   -3.52     0.000    -3.458522    -.9859983 |
        POmean |               |
   smoke   | Nonsmoker     |  4.235569   .5210937    8.13     0.000     3.214244     5.256894 |
```

Below, we compute the estimated probabilities of being a Nonsmoker and store them in \texttt{ps0}. Likewise, the estimated probabilities of being a Smoker are stored in \texttt{ps1}.

```
. predict ps0 ps1, ps
```

The overlap condition requires that each of these probabilities be sufficiently greater than 0 and less than 1 for every individual; see \textit{Assumptions and trade-offs} under Remarks and examples in \texttt{[TE] stteffects intro}.

In practice, we know that weighting estimators perform poorly when the weights become too large. This approach requires that the probability of being a Nonsmoker not be too small among Nonsmokers and that the probability of being a Smoker not be too small among Smokers. Below, we summarize these probabilities.
The minimum probability of being a Non-smoker among Non-smokers is 0.39. The minimum probability of being a Smoker among Smokers is 0.09. Neither minimum seems too small.

Estimating survival-time treatment effects also uses weights to adjust for censored outcomes; see [TE] stteffects intro. Thus we require that the probability of an uncensored failure also be sufficiently greater than 0. Below, we compute the estimated probabilities of failure and summarize them among those that fail.

```
. predict fprob2, censurv
. summarize fprob if fail==1
```

<table>
<thead>
<tr>
<th>Variable</th>
<th>Obs</th>
<th>Mean</th>
<th>Std. Dev.</th>
<th>Min</th>
<th>Max</th>
</tr>
</thead>
<tbody>
<tr>
<td>fprob2</td>
<td>1,208</td>
<td>.7246067</td>
<td>.2143543</td>
<td>.0364246</td>
<td>.9999086</td>
</tr>
</tbody>
</table>

The minimum probability of 0.04 does not appear too small.

## Technical note

The previous discussion builds on the intuition that the weights used in a weighting estimator should not be too large.

This technical note goes a little further by explicitly computing the weights and using them to replicate the inverse-probability-weighted point estimate for the Non-smoker potential-outcome mean.

We now compute the weights using the predicted probabilities computed in the examples above and then use `mean` to compute the weighted average that estimates the potential-outcome mean for Non-smokers.

```
. generate double ipw0 = 1/(ps0*fprob)
. mean _t [pw=ipw0] if smoke==0 & fail==1
```

<table>
<thead>
<tr>
<th>Mean estimation</th>
<th>Number of obs = 716</th>
</tr>
</thead>
<tbody>
<tr>
<td>_t</td>
<td>Mean</td>
</tr>
<tr>
<td>_t</td>
<td>4.235569</td>
</tr>
</tbody>
</table>

The weights account for data lost to the Smoker potential outcome or to censoring by increasing the importance of observations that were observed to be Non-smoker failure times even though they were not likely to be observed.

The point estimate matches that reported by `stteffects ipw`; the standard errors differ because `mean` takes the estimated weights as given. See `Inverse-probability-weighted estimators` under Methods and formulas in [TE] stteffects ipwra.
References


Also see

[TE] tetable — Check balance after teffects or stteffects estimation
[TE] teffects overlap — Overlap plots
[TE] stteffects ipw — Survival-time inverse-probability weighting
[TE] stteffects ipwra — Survival-time inverse-probability-weighted regression adjustment
[TE] stteffects ra — Survival-time regression adjustment
[TE] stteffects wra — Survival-time weighted regression adjustment
[U] 20 Estimation and postestimation commands
**Description**

`stteffects ra` estimates the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs) from observational survival-time data by regression adjustment (RA). RA uses averages of treatment-specific predicted mean survival times to estimate mean survival times for each potential outcome. Contrasts of these predicted mean survival times estimate the treatment effects. `stteffects ra` offers several choices for the model used to predict mean survival time. Binary and multivalued treatments are accommodated.

See [TE] stteffects intro for an overview of estimating treatment effects from observational survival-time data.

**Quick start**

Specify `time` as observed failure time and `fail` as failure indicator

```
stset time, failure(fail)
```

ATE from a Weibull model for `time` on `x1` and `x2` with binary treatment `treat2`

```
stteffects ra (x1 x2) (treat2)
```

As above, but estimate the ATET

```
stteffects ra (x1 x2) (treat2), atet
```

As above, but estimate the potential-outcome means

```
stteffects ra (x1 x2) (treat2), pomeans
```

ATE of `treat2` using a gamma model for `time`

```
stteffects ra (x1 x2, gamma) (treat2)
```

ATE for each level of three-valued treatment `treat3`

```
stteffects ra (x1 x2) (treat3)
```

As above, and specify that `treat3 = 3` is the control level using the value label “MyControl” for 3

```
stteffects ra (x1 x2) (treat3), control("MyControl")
```

**Menu**

Statistics > Treatment effects > Survival outcomes > Regression adjustment
**Syntax**

```
stteffects ra (omvarlist [, omoptions]) (tvar) [if] [in] [, stat options]
```

*omvarlist* specifies the variables that predict the survival-time variable in the outcome model.
*tvar* must contain integer values representing the treatment levels.

**omoptions**

<table>
<thead>
<tr>
<th>Description</th>
<th>Model</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td><em>weibull</em></td>
</tr>
<tr>
<td></td>
<td><em>exponential</em></td>
</tr>
<tr>
<td></td>
<td><em>gamma</em></td>
</tr>
<tr>
<td></td>
<td><em>lnormal</em></td>
</tr>
<tr>
<td>ancillary(avarlist [, noconstant])</td>
<td>specify variables used to model ancillary parameter</td>
</tr>
<tr>
<td></td>
<td><em>noconstant</em></td>
</tr>
</tbody>
</table>

**stat**

<table>
<thead>
<tr>
<th>Description</th>
<th>Stat</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td><em>ate</em></td>
</tr>
<tr>
<td></td>
<td><em>atet</em></td>
</tr>
<tr>
<td></td>
<td><em>pomeans</em></td>
</tr>
</tbody>
</table>

**options**

<table>
<thead>
<tr>
<th>Description</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>vctype</td>
</tr>
<tr>
<td></td>
<td>may be <em>robust, cluster clustvar, bootstrap, or jackknife</em></td>
</tr>
</tbody>
</table>

**SE/Robust**

| vce(vcetype)         | vcetype may be *robust, cluster clustvar, bootstrap, or jackknife* |

**Reporting**

| level(#)             | set confidence level; default is *level(95)* |
| aequations           | display auxiliary-equation results |
| noshow               | do not show st setting information |

**display_options**

| control(#) | label | specify the level of tvar that is the control |
| tlevel(#)   | label | specify the level of tvar that is the treatment |
| coeflegend  | display legend instead of statistics |
You must stset your data before using stteffects; see [ST] stset.

omvarlist and avarlist may contain factor variables; see [U] 11.4.3 Factor variables.

bootstrap, by, jackknife, and statsby are allowed; see [U] 11.1.10 Prefix commands.

Weights are not allowed with the bootstrap prefix; see [R] bootstrap.

fweights, iweights, and pweights may be specified using stset; see Weights under Remarks and examples in [ST] stset. However, weights may not be specified if you are using the bootstrap prefix.

coefflegend does not appear in the dialog box.

See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.

Options

ancillary(avarlist [, noconstant]) specifies the variables used to model the ancillary parameter.

By default, the ancillary parameter does not depend on covariates. Specifying ancillary(avarlist, noconstant) causes the constant to be suppressed in the model for the ancillary parameter. noconstant; see [R] estimation options.

stat is one of three statistics: ate, atet, or pomeans. ate is the default.

ate specifies that the average treatment effect be estimated.

atet specifies that the average treatment effect on the treated be estimated.

pomeans specifies that the potential-outcome means for each treatment level be estimated.

vce(vcetype) specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (robust), that allow for intragroup correlation (cluster clustvar), and that use bootstrap or jackknife methods (bootstrap, jackknife); see [R] vce_option.

level(#); see [R] estimation options.

eaequations specifies that the results for the outcome-model or treatment-model parameters be displayed. By default, the results for these auxiliary parameters are not displayed.

no shows prevents stteffects ra from showing the key st variables. This option is rarely used because most people type stset, show or stset, noshow to permanently set whether they want to see these variables mentioned at the top of the output of every st command; see [ST] stset.

display_options: noci, nopvalues, noomitted, vsquish, noemptycells, baselevels, allbaselevels, nofvlabel, fvwrap(#), fvwrapon(style), cformat(%fmt), pformat(%fmt), sformat(%fmt), and nolstretch; see [R] estimation options.

maximize_options: iterate(#), [no]log, and from(init_specs); see [R] maximize. These options are seldom used.

init_specs is one of

matname [, skip copy]

# [, # ...], copy
iterinit(#) specifies the maximum number of iterations used to calculate the starting values. This option is seldom used.

ccontrol(#|label) specifies the level of tvar that is the control. The default is the first treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. control() may not be specified with the statistic pomeans. control() and tlevel() may not specify the same treatment level.

tlevel(#|label) specifies the level of tvar that is the treatment for the statistic atet. The default is the second treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. tlevel() may only be specified with statistic atet. tlevel() and control() may not specify the same treatment level.

The following option is available with stteffects but is not shown in the dialog box:

coefflegend; see [R] estimation options.

Remarks and examples

If you are not familiar with the framework for treatment-effects estimation from observational survival-time data, please see [TE] stteffects intro.

RA estimators use contrasts of the averages of treatment-specific predicted mean outcomes to estimate treatment effects. RA estimators use a two-step approach to estimating treatment effects:

1. For each treatment level, fit a model of the survival-time outcome on the same set of covariates.
2. Compute the averages of the predicted outcomes for each subject within each treatment level.

These averages estimate the potential-outcome means (POMs). Contrasts of these averages estimate the ATEs. By restricting the computations of the averages to the subset of treated subjects, we obtain estimates of the ATETs.

Here we note only a few entry points to the vast literature on RA estimators. Imbens (2004), Imbens and Wooldridge (2009), Cameron and Trivedi (2005, chap. 25), Wooldridge (2010, chap. 21), and Vittinghoff et al. (2012, chap. 9) provide excellent general introductions to estimating ATEs and to RA estimators in particular.

Like streg and other survival-time commands, stteffects ra uses the outcome variable and the failure indicator computed by, and optionally weights specified with, stset. stteffects ra is not appropriate for data with time-varying covariates, also known as multiple-record survival-time data, or for delayed-entry data.

Example 1: Estimating the ATE

Suppose we wish to study the effect of smoking on the time to a second heart attack among women aged 45–55 years. In our fictional sheart dataset, atime is the observed time in years to a second heart attack or censoring, and fail is the 0/1 indicator that a second heart attack was observed. (When fail is 1, atime records the time to the second heart attack; when fail is 0, atime records a censored observation of the time to a second heart attack.) We previously stset these data; see A quick tour of the estimators in [TE] stteffects intro.
The treatment, smoking, is stored in the 0/1 indicator `smoke`. These data also contain age at the time of the first heart attack (`age`), and indices of the level of exercise (`exercise`), diet quality (`diet`), and education (`education`) prior to the first heart attack.

We can use `stteffects ra` to estimate the ATE by RA. We model the mean survival time using the default Weibull model, controlling for `age`, `exercise`, `diet`, and `education`, and we specify that `smoke` is the treatment variable.

```
. use http://www.stata-press.com/data/r15/sheart
   (Time to second heart attack (fictional))
. stteffects ra (age exercise diet education) (smoke)
   failure _d:  fail
   analysis time _t:  atime
Iteration 0:  EE criterion = 1.525e-19
Iteration 1:  EE criterion = 1.931e-30
Survival treatment-effects estimation                  Number of obs =   2,000
Estimator : regression adjustment
Outcome model : Weibull
Treatment model: none
Censoring model: none

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>_t</td>
<td>Coef.</td>
<td>Std. Err.</td>
<td>z</td>
<td>P&gt;</td>
</tr>
<tr>
<td>ATE</td>
<td>smoke</td>
<td>-1.956657</td>
<td>.3331787</td>
<td>-5.87</td>
<td>0.000</td>
</tr>
<tr>
<td></td>
<td>(Smoker vs</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Nonsmoker)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>POmean</td>
<td>smoke</td>
<td>4.243974</td>
<td>.2620538</td>
<td>16.20</td>
<td>0.000</td>
</tr>
<tr>
<td></td>
<td>Nonsmoker</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
```

When every woman smoked in the population of women aged 45–55 years who have had a heart attack, the average time to a second heart attack is estimated to be 1.96 years less than when no women in the population of interest smoked. The estimated average time to a second heart attack when no women in the population of interest smoked is 4.24 years. In other words, if every woman in the population of interest smoked, then the average time to a second heart attack would fall by an estimated 46% relative to the case when no women smoked.

Example 2: Changing the outcome model

Instead of a Weibull model for the outcome model, we could have used an exponential, a gamma, or a lognormal model. By way of comparison, we use a gamma model and the same covariates to estimate the ATE.
The estimated ATE of $-1.80$ and control-level POM of $3.99$ are similar to those of $-1.96$ and $4.24$ obtained from the Weibull model in example 1. The ratio of the estimated ATE to the control-level POM indicates a 45% reduction instead of the 46% reduction obtained from the Weibull model.

Example 3: Estimating the ratio of the ATE to the control-level POM

The ratio of the ATE to the control-level POM measures the importance of the effect. In example 1, we computed the point estimate of this ratio from the output, but we were left without a confidence interval. In this example, we use \texttt{nlcom} to compute a point estimate and a confidence interval.

Below, we refit the model from example 1, specifying the \texttt{coeflegend} option to learn the parameter names. We use the parameter names in \texttt{nlcom} to estimate the ratio of the ATE to the control-level POM.
Survival treatment-effects estimation

Number of obs = 2,000

Estimator : regression adjustment
Outcome model : Weibull
Treatment model: none
Censoring model: none

_t  Coef. Legend

ATE
smoke
(Smoker vs Nonsmoker) -1.956657 _b[ATE:r1vs0.smoke]

POmean
smoke
Nonsmoker 4.243974 _b[POmean:0.smoke]

._nlcom _b[ATE:r1vs0.smoke] / _b[POmean:0.smoke]
_n1_1:  _b[ATE:r1vs0.smoke] / _b[POmean:0.smoke]

| _t | Coef. | Std. Err. | z     | P>|z| | [95% Conf. Interval] |
|----|-------|-----------|-------|-------|-----------------------|
| _n1_1 | -.4610437 | .0598709 | -7.70 | 0.000 | -.5783885 -.3436988 |

The output shows that when every woman smoked, the average time to a second heart attack falls by an estimated 46% relative to the case when no women smoked, as we computed earlier. We also obtain a 95% confidence interval of 34% to 58% for this estimate.

Example 4: Estimating the ATET

Intuitively, the ATET measures the effect of the treatment on an at-risk subpopulation. Sometimes the subpopulation that gets the treatment defines such an at-risk subpopulation. The ATET has the added benefit that it can be estimated under weaker conditions than the ATE; see Assumptions and trade-offs in [TE] stteffects intro.
. stteffects ra (age exercise diet education) (smoke), atet
    failure _d: fail
    analysis time _t: atime

Iteration 0:   EE criterion =  1.525e-19
Iteration 1:   EE criterion =  2.002e-31

Survival treatment-effects estimation
Number of obs  =  2,000

Estimator     : regression adjustment
Outcome model : Weibull
Treatment model: none
Censoring model: none

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
<th></th>
<th></th>
<th>[95% Conf. Interval]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>_t</td>
<td>Coef.</td>
<td>Std. Err.</td>
<td>z</td>
</tr>
<tr>
<td>ATET</td>
<td>smoke</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(Smoker vs</td>
<td></td>
<td>-1.527476</td>
<td>.2489203</td>
<td>-6.14</td>
</tr>
<tr>
<td>Non-smoker)</td>
<td></td>
<td>-1.039602</td>
<td></td>
<td></td>
</tr>
<tr>
<td>POmean</td>
<td>smoke</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Non-smoker</td>
<td>3.436937</td>
<td>.2217808</td>
<td>15.50</td>
<td>0.000</td>
</tr>
<tr>
<td></td>
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</tr>
</tbody>
</table>

When every woman in the subpopulation smoked, the average time to a second heart attack is estimated to be 1.53 years less than when no women in the subpopulation smoked. The estimated average time to a second heart attack when no women in the subpopulation smoked is 3.44 years.

Example 5: Fixed or random censoring time

The time to censoring in survival-time data can be random or deterministic, although it must be independent of treatment assignment and the potential outcomes; see Kalbfleisch and Prentice (2002, chap. 3) for the standard case and see The correct adjustment for censoring assumption under Assumptions and trade-offs in [TE] stteffects intro for the treatment-effects case.

The RA estimator and the likelihood-adjusted-censoring version of the inverse-probability-weighted RA estimator can accommodate a fixed time to censoring; see The correct adjustment for censoring assumption in [TE] stteffects intro. (The estimators that handle censoring by weighting cannot accommodate a fixed time to censoring because the weights are not well defined with a fixed time to censoring.)

We have fictional data on the time to rearrest among men aged 25–35 who were previously in prison for a felony conviction (rtime). The time to censoring is fixed in these data because individuals were followed for a maximum of five years.

Some of the young men chose to enter a vocational training program before release from prison; train is 1 for participants and 0 for nonparticipants. The dataset also contains fail (which is 1 if the observed time is a failure time and 0 if it is time to censoring), age at the time of the first arrest (age), an index of the parents’ socioeconomic level (parental), and the number of years behind in school at the time of the first arrest (edeficit).

We estimate the ATET because we wish to allow the gains from the training program to be related to an unobservable characteristic that affects who self-selects into the program; see Average treatment effect on the treated in [TE] stteffects intro.
We model the outcome as a function of *age*, *parental*, and *edeficit*.

```
. use http://www.stata-press.com/data/r15/recid2, clear
    (Time to rearrest (fictional))
. stteffects ra (age parental edeficit) (train), atet
    failure _d: fail
    analysis time _t: rtime
Iteration 0: EE criterion = 2.769e-19
Iteration 1: EE criterion = 9.441e-34
Survival treatment-effects estimation Number of obs = 2,000
Estimator : regression adjustment
Outcome model : Weibull
Treatment model: none
Censoring model: none

Coef. Std. Err. z P>|z| [95% Conf. Interval]
ATET
    train (Student vs Nonstudent)
          2.440919   .4689057  5.21 0.000   1.52188   3.359957
POmean
    train Nonstudent
          2.062029   .1231492 16.74 0.000   1.820661   2.303397
```

When everyone who selected the training got the training, the average time to rearrest is 2.44 years later than the average rearrest time if none of those who selected the training got the training. The average rearrest time if none of those who selected the training got the training is 2.06 years. In other words, the average time to rearrest increases from about 2.06 years to about 4.50 years for the subpopulation of young men who self-selected into the prerelease vocational training program.

Stored results

`stteffects ra` stores the following in `e()`:

Scalars
- `e(N)` number of observations
- `e(nj)` number of observations for treatment level `j`
- `e(N_clust)` number of clusters
- `e(k_eq)` number of equations in `e(b)`
- `e(k_levels)` number of levels in treatment variable
- `e(treated)` level of treatment variable defined as treated
- `e(control)` level of treatment variable defined as control
- `e(converged)` 1 if converged, 0 otherwise

Macros
- `e(cmd)` `stteffects`
- `e(cmdline)` command as typed
- `e(dead)` `_d`
- `e(depvar)` `_t`
- `e(tvar)` name of treatment variable
- `e(subcmd)` `ra`
- `e(omodel)` outcome model: weibull, exponential, gamma, or lognormal
- `e(stat)` statistic estimated: ate, atet, or pomeans
**Methods and formulas**

The methods and formulas for the RA estimators implemented in `stteffects ra` are given in *Methods and formulas* of [TE] `stteffects ipwra`.

**References**


Also see

[TE] stteffects postestimation — Postestimation tools for stteffects
[TE] stteffects intro — Introduction to treatment effects for observational survival-time data
[ST] streg — Parametric survival models
[ST] stset — Declare data to be survival-time data
[U] 20 Estimation and postestimation commands
Description

`stteffects wra` estimates the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs) from observational survival-time data with random time to censoring. Estimation is by weighted regression adjustment (WRA). WRA estimators use inverse-probability-of-censoring adjusted regression coefficients to compute averages of treatment-level predicted outcomes. Contrasts of these averages estimate the treatment effects. WRA uses estimated weights from a time-to-censoring model to account for censored survival times instead of including a term in the likelihood function. `stteffects wra` offers several choices for the functional forms of the outcome model and the time-to-censoring model. Binary and multivalued treatments are accommodated.

See [TE] `stteffects intro` for an overview of estimating treatment effects from observational survival-time data.

Quick start

Specify `time` as observed failure time and `fail` as failure indicator

```
  stset time, failure(fail)
```

ATE from a Weibull model for `time` on `x1` and `x2` with binary treatment `treat2` and a Weibull model on `x1` and `x2` for censoring

```
  stteffects wra (x1 x2) (treat2) (x1 x2)
```

As above, but estimate the ATET

```
  stteffects wra (x1 x2) (treat2) (x1 x2), atet
```

ATE of `treat2` using a gamma model for `time` and a gamma censoring model

```
  stteffects wra (x1 x2, gamma) (treat2) (x1 x2, gamma)
```

ATE for each level of three-valued treatment `treat3`

```
  stteffects wra (x1 x2) (treat3) (x1 x2)
```

As above, and specify that `treat3 = 3` is the control level using the value label “MyControl” for 3

```
  stteffects wra (x1 x2) (treat3) (x1 x2), control("MyControl")
```

Menu

Statistics > Treatment effects > Survival outcomes > Weighted regression adjustment
### Syntax

```
stteffects wra (omvarlist [, omoptions]) (tvar) (cmvarlist [, cmoptions])
```

*omvarlist* specifies the variables that predict the survival-time variable in the outcome model.
*tvar* must contain integer values representing the treatment levels.
*cmvarlist* specifies the variables that predict censoring in the censoring model.

#### omoptions

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Model</td>
</tr>
<tr>
<td><strong>weibull</strong></td>
</tr>
<tr>
<td><strong>exponential</strong></td>
</tr>
<tr>
<td><strong>gamma</strong></td>
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<tr>
<td><strong>lnormal</strong></td>
</tr>
<tr>
<td><strong>ancillary(avarlist [, noconstant])</strong></td>
</tr>
<tr>
<td><strong>noconstant</strong></td>
</tr>
</tbody>
</table>

#### cmoptions

<table>
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<tr>
<th>Description</th>
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<tbody>
<tr>
<td>Model</td>
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<tr>
<td><strong>weibull</strong></td>
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<tr>
<td><strong>ancillary(avarlist [, noconstant])</strong></td>
</tr>
<tr>
<td><strong>noconstant</strong></td>
</tr>
</tbody>
</table>

#### stat

<table>
<thead>
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<tbody>
<tr>
<td>Stat</td>
</tr>
<tr>
<td><strong>ate</strong></td>
</tr>
<tr>
<td><strong>atet</strong></td>
</tr>
<tr>
<td><strong>pomeans</strong></td>
</tr>
</tbody>
</table>
options Description

SE/Robust

vce(vcetype) \(\text{vcetype \ may \ be \ robust, \ cluster \ clustvar, \ bootstrap, \ or \ jackknife}\)

Reporting

level(#) \(\text{set \ confidence \ level; \ default \ is \ level(95)}\)
equations \(\text{display \ auxiliary-equation \ results}\)
noshow \(\text{do \ not \ show \ st \ setting \ information}\)
display_options \(\text{display \ of \ omitted \ variables \ and \ base \ and \ empty \ cells, \ and}\
\text{factor-variable \ labeling}\)

Maximization

maximize_options \(\text{control \ the \ maximization \ process; \ seldom \ used}\)
iterinit(#) \(\text{specify \ starting-value \ iterations; \ seldom \ used}\)

Advanced

pstolerance(#) \(\text{set \ the \ tolerance \ for \ the \ overlap \ assumption}\)
osample(newvar) \(\text{identify \ observations \ that \ violate \ the \ overlap \ assumption}\)
control(# | label) \(\text{specify \ the \ level \ of \ tvar \ that \ is \ the \ control}\)
tlevel(# | label) \(\text{specify \ the \ level \ of \ tvar \ that \ is \ the \ treatment}\)
coeflegend \(\text{display \ legend \ instead \ of \ statistics}\)

You must stset your data before using stteffects; see [ST] stset.

omvarlist, cmvarlist, and avarlist may contain factor variables; see [U] 11.4.3 Factor variables.

bootstrap, by, jackknife, and statsby are allowed; see [U] 11.1.10 Prefix commands.

Weights are not allowed with the bootstrap prefix; see [R] bootstrap.

fweights, iweights, and pweights may be specified using stset; see Weights under Remarks and examples in [ST] stset. However, weights may not be specified if you are using the bootstrap prefix.

coeflegend does not appear in the dialog box.

See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.

Options

ancillary(avarlist [, noconstant]) \(\text{specifies \ the \ variables \ used \ to \ model \ the \ ancillary \ parameter.}\
\text{By \ default, \ the \ ancillary \ parameter \ does \ not \ depend \ on \ covariates. \ Specifying \ ancillary(avarlist,}\
\text{noconstant) \ causes \ the \ constant \ to \ be \ suppressed \ in \ the \ model \ for \ the \ ancillary \ parameter.}\
\text{ancillary()} \ may \ be \ specified \ for \ the \ model \ for \ survival-time \ outcome, \ for \ the \ model \ for \ the}\
censoring \ variable, \ or \ for \ both. \ If \ ancillary() \ is \ specified \ for \ both, \ the \ varlist \ used \ for \ each}\
\text{model \ may \ be \ different.}\
noconstant; see [R] estimation options.
stat is one of three statistics: \texttt{ate}, \texttt{atet}, or \texttt{pomeans}. \texttt{ate} is the default.

\texttt{ate} specifies that the average treatment effect be estimated.

\texttt{atet} specifies that the average treatment effect on the treated be estimated.

\texttt{pomeans} specifies that the potential-outcome means for each treatment level be estimated.

\texttt{vce(vcetype)} specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (\texttt{robust}), that allow for intragroup correlation (\texttt{cluster clustvar}), and that use bootstrap or jackknife methods (\texttt{bootstrap}, \texttt{jackknife}); see \cite{vce_option}.

\texttt{aequations} specifies that the results for the outcome-model or treatment-model parameters be displayed. By default, the results for these auxiliary parameters are not displayed.

\texttt{noshow} prevents \texttt{stteffects wra} from showing the key \texttt{st} variables. This option is rarely used because most people type \texttt{stset}, \texttt{show} or \texttt{stset, noshow} to permanently set whether they want to see these variables mentioned at the top of the output of every \texttt{st} command; see \cite{stset}.

\texttt{display_options}: \texttt{noci}, \texttt{nopvalues}, \texttt{noomitted}, \texttt{vsquish}, \texttt{noemptycells}, \texttt{baselevels}, \texttt{allbaselevels}, \texttt{nofvlabel}, \texttt{fvwrap(#)}, \texttt{fvwrapon(style)}, \texttt{cformat(\%fmt)}, \texttt{pformat(\%fmt)}, \texttt{sformat(\%fmt)}, and \texttt{nolstretch}; see \cite{estimation_options}.

\texttt{maximize_options}: \texttt{iterate(#)}, \texttt{[no]log}, and \texttt{from(init_specs)}; see \cite{maximize}. These options are seldom used.

\texttt{init_specs} is one of

\begin{verbatim}
matname [ , skip copy ]
# [ , # ... ] , copy
\end{verbatim}

\texttt{iterinit(#)} specifies the maximum number of iterations used to calculate the starting values. This option is seldom used.

\texttt{pstolerance(#)} specifies the tolerance used to check the overlap assumption. The default value is \texttt{pstolerance(1e-5)}. \texttt{stteffects} will exit with an error if an observation has an estimated propensity score smaller than that specified by \texttt{pstolerance(\)}.\texttt{osample(newvar)} specifies that indicator variable \texttt{newvar} be created to identify observations that violate the overlap assumption.

\texttt{control(# | label)} specifies the level of \texttt{tvar} that is the control. The default is the first treatment level. You may specify the numeric level \# (a nonnegative integer) or the label associated with the numeric level. \texttt{control()} may not be specified with the statistic \texttt{pomeans}. \texttt{control()} and \texttt{tlevel()} may not specify the same treatment level.
tlevel(# | label) specifies the level of tvar that is the treatment for the statistic atet. The default is the second treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. tlevel() may only be specified with statistic atet. tlevel() and control() may not specify the same treatment level.

The following option is available with stteffects but is not shown in the dialog box: coeflegend; see [R] estimation options.

Remarks and examples

If you are not familiar with the framework for treatment-effects estimation from observational survival-time data, please see [TE] stteffects intro.

Weighted regression-adjustment (WRA) estimators use estimated weights to account for censoring when estimating outcome-regression parameters. The estimated outcome-regression parameters are used to compute averages of treatment-level predicted outcomes. Contrasts of these averages estimate the treatment effects.

WRA estimators use a three-step approach to estimating treatment effects:

1. They estimate the parameters of a time-to-censoring model and compute inverse-probability-of-censoring weights.
2. Using the estimated inverse-probability-of-censoring weights, they use weighted maximum likelihood estimators for the outcome for each treatment level and obtain the treatment-specific predicted mean outcomes for each subject. The inverse-probability-of-censoring weights account for right-censored survival times.
3. They compute the means of the treatment-specific predicted mean outcomes. Contrasts of these averages provide the estimates of the ATEs. By restricting the computations of the means to the subset of treated subjects, we can obtain the ATETs.

WRA estimators differ from RA estimators in that WRA estimators use weights to account for observations lost to censoring while RA estimators use an additional term in the likelihood function. A model for the time to censoring is used to estimate the weights.

WRA estimators require more assumptions than RA estimators. Specifically, they require that the censoring time be random and that the time-to-censoring model be well specified. The implemented WRA estimators also require that the time-to-censoring process not vary by treatment level. The RA estimator and the likelihood-adjusted-censoring version of the inverse-probability-weighted RA estimator do not require these extra assumptions, because they use a likelihood term instead of weights to adjust for the data lost to censoring; see [TE] stteffects ra and [TE] stteffects ipwra.

Here we note only a few entry points to the vast literature on weighted estimators. Imbens (2004), Imbens and Wooldridge (2009), Robins and Rotnitzky (2006), Wooldridge (2002, 2007), Cameron and Trivedi (2005, chap. 25), Wooldridge (2010, chap. 21), and Vittinghoff et al. (2012, chap. 9) provide excellent general introductions to estimating ATEs and to WRA estimators in particular.

Like streg and other survival-time commands, stteffects wra uses the outcome variable and the failure indicator computed by, and optionally weights specified with, stset. stteffects wra is not appropriate for data with time-varying covariates, also known as multiple-record survival-time data, or for delayed-entry data.
Example 1: Estimating the ATE

Suppose we wish to study the effect of smoking on the time to a second heart attack among women aged 45–55 years. In our fictional `sheart` dataset, `atime` is the observed time in years to a second heart attack or censoring, and `fail` is the 0/1 indicator that a second heart attack was observed. (When `fail` is 1, `atime` records the time to the second heart attack; when `fail` is 0, `atime` records a censored observation of the time to a second heart attack.) We previously `stset` these data; see *A quick tour of the estimators* in [TE] `stteffects intro`.

The treatment, smoking, is stored in the 0/1 indicator `smoke`. These data also contain age at the time of the first heart attack (`age`), and indices of the level of exercise (exercise), diet quality (`diet`), and education (`education`) prior to the first heart attack.

We can use `stteffects wra` to estimate the ATE by WRA. We model the mean survival time using the default Weibull outcome model with `age`, `exercise`, `diet`, and `education` as covariates, and we specify that `smoke` is the treatment variable. We also specify the default Weibull time-to-censoring model and include `age`, square of `age`, `exercise`, and `education`.

```
. use http://www.stata-press.com/data/r15/sheart
   (Time to second heart attack (fictional))
. stteffects wra (age exercise diet education) ///
   > (smoke) ///
   > (age c.age#c.age exercise diet education)
   > failure _d: fail
   > analysis time _t: atime

Iteration 0:   EE criterion = 4.096e-18
Iteration 1:   EE criterion = 1.302e-29

Survival treatment-effects estimation     Number of obs   =    2,000
Estimator       : weighted regression adjustment
Outcome model   : Weibull
Treatment model : none
Censoring model : Weibull

                      Robust                    Robust
       Coef.     Std. Err.     z    P>|z|  [95% Conf. Interval]
---------------------------------------------------------------
     ATE          smoke (Smoker vs Nonsmoker)  -2.374174    .6017498   -3.95  0.000  -3.553582   -1.194766
     POmean       smoke Nonsmoker              4.302131    .5528943    7.78  0.000     3.218478    5.385784
```

When every woman smoked in the population of women aged 45–55 years who have had a heart attack, the average time to a second heart attack is estimated to be 2.37 years less than when no women in the subpopulation of interest smoked. The estimated average time to a second heart attack when no women in the subpopulation of interest smoked is 4.30 years.
Stored results

\texttt{stteffects wra} stores the following in \texttt{e()}: 

\begin{itemize}
  \item \texttt{e(N)} number of observations
  \item \texttt{e(nj)} number of observations for treatment level \( j \)
  \item \texttt{e(N_clust)} number of clusters
  \item \texttt{e(k_eq)} number of equations in \texttt{e(b)}
  \item \texttt{e(k_levels)} number of levels in treatment variable
  \item \texttt{e(treated)} level of treatment variable defined as treated
  \item \texttt{e(control)} level of treatment variable defined as control
  \item \texttt{e(converged)} 1 if converged, 0 otherwise
\end{itemize}

\begin{itemize}
  \item \texttt{e(cmd)} \texttt{stteffects}
  \item \texttt{e(cmdline)} command as typed
  \item \texttt{e(dead)} 	exttt{_d}
  \item \texttt{e(depvar) \_t}
  \item \texttt{e(tvar)} name of treatment variable
  \item \texttt{e(subcmd) wra}
  \item \texttt{e(omodel)} outcome model: \texttt{weibull}, exponential, gamma, or lognormal
  \item \texttt{e(cmodel)} censoring model: \texttt{weibull}, exponential, gamma, or lognormal
  \item \texttt{e(stat)} statistic estimated: \texttt{ate}, \texttt{atet}, or \texttt{pomeans}
  \item \texttt{e(type)} weight type
  \item \texttt{e(wexp)} weight expression
  \item \texttt{e(title)} title in estimation output
  \item \texttt{e(clustvar)} name of cluster variable
  \item \texttt{e(tlevels)} levels of treatment variable
  \item \texttt{e(vce)} \texttt{vctype} specified in \texttt{vce()}
  \item \texttt{e(vctype)} title used to label Std. Err.
  \item \texttt{e(properties) b V}
  \item \texttt{e(estat_cmd)} program used to implement \texttt{estat}
  \item \texttt{e(predict)} program used to implement \texttt{predict}
  \item \texttt{e(marginsnotok)} predictions disallowed by \texttt{margins}
  \item \texttt{e(asbalanced)} factor variables \texttt{fvset} as \texttt{asbalanced}
  \item \texttt{e(asobserved)} factor variables \texttt{fvset} as \texttt{asobserved}
\end{itemize}

Matrices

\begin{itemize}
  \item \texttt{e(b)} coefficient vector
  \item \texttt{e(V)} variance–covariance matrix of the estimators
\end{itemize}

Functions

\begin{itemize}
  \item \texttt{e(sample)} marks estimation sample
\end{itemize}

Methods and formulas

The methods and formulas for the WRA estimators implemented in \texttt{stteffects wra} are given in \textit{Methods and formulas} of [TE] \texttt{stteffects ipwra}.

References


**Also see**

[TE] *stteffects postestimation* — Postestimation tools for stteffects
[TE] *stteffects intro* — Introduction to treatment effects for observational survival-time data
[ST] *streg* — Parametric survival models
[ST] *stset* — Declare data to be survival-time data
[U] *20 Estimation and postestimation commands*
tebalance — Check balance after teffects or stteffects estimation

Description

The tebalance postestimation commands produce diagnostic statistics, test statistics, and diagnostic plots to assess whether a teffects or an stteffects command balanced the covariates over treatment levels.

Syntax

tebalance subcommand ... [, options ]

<table>
<thead>
<tr>
<th>subcommand</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>summarize</td>
<td>compare means and variances in raw and balanced data</td>
</tr>
<tr>
<td>overid</td>
<td>overidentification test</td>
</tr>
<tr>
<td>density</td>
<td>kernel density plots for raw and balanced data</td>
</tr>
<tr>
<td>box</td>
<td>box plots for each treatment level for balanced data</td>
</tr>
</tbody>
</table>

Remarks and examples

This entry provides an overview of the commands in tebalance. We recommend that you read this entry before proceeding to [TE] tebalance summarize, [TE] tebalance overid, [TE] tebalance density, or [TE] tebalance box for command-specific syntax and details.

A covariate is said to be balanced when its distribution does not vary over treatment levels.

Covariates are balanced in experimental data because treatment assignment is independent of the covariates because of the study design. In contrast, covariates must be balanced by weighting or matching in observational data because treatment assignment is related to the covariates that also affect the outcome of interest.

The estimators implemented in teffects and stteffects use a model or matching method to make the outcome conditionally independent of the treatment by conditioning on covariates. If this model or matching method is well specified, it should balance the covariates. Balance diagnostic techniques and tests check the specification of the conditioning method used by a teffects or an stteffects estimator; see [TE] teffects intro advanced for an introduction to teffects, and [TE] stteffects intro for an introduction to stteffects.

tebalance implements four methods to check for balance after teffects and stteffects. Which tebalance methods are available depends on the teffects estimation method, as summarized in the table below.
tebalance — Check balance after tteffects or stteffects estimation

<table>
<thead>
<tr>
<th>Method</th>
<th>Description</th>
<th>ipw</th>
<th>aipw</th>
<th>ipwra</th>
<th>nnmatch</th>
<th>psmatch</th>
<th>ipw</th>
<th>ipwra</th>
</tr>
</thead>
<tbody>
<tr>
<td>summarize</td>
<td>standardized differences and variance ratios</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
</tr>
<tr>
<td>overid</td>
<td>chi-squared test for balance</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td></td>
<td>x</td>
<td></td>
</tr>
<tr>
<td>density</td>
<td>diagnostic kernel-density plots</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
<td>x</td>
</tr>
<tr>
<td>box</td>
<td>diagnostic box plots</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>x</td>
<td>x</td>
</tr>
</tbody>
</table>

Example 1: Balance after estimators that use weighting

Inverse-probability-weighted (IPW) estimators use a model for the treatment to make the outcome conditionally independent of the treatment. If this model is well specified, it will also balance the covariates.

Using an extract from Cattaneo (2010), we use tteffects ipw to estimate the effect of a mother’s smoking behavior (mbsmoke) on the birthweight of her child (bweight), controlling for marital status (mmarried), the mother’s age (mage), whether the mother had a prenatal doctor’s visit in the baby’s first trimester (prenatal1), and whether this baby is the mother’s first child (fbaby).

```
. use http://www.stata-press.com/data/r15/cattaneo2
. tteffects ipw (bweight) (mbsmoke mmarried mage prenatal1 fbaby)
Iteration 0: EE criterion = 1.873e-22
Iteration 1: EE criterion = 3.315e-26
Treatment-effects estimation Number of obs = 4,642
Estimator : inverse-probability weights
Outcome model : weighted mean
Treatment model: logit

|                  | Coef.     | Std. Err. | z     | P>|z|    | [95% Conf. Interval] |
|------------------|-----------|-----------|-------|--------|---------------------|
|                  | Robust    |           |       |        |                     |
| bweight          |           |           |       |        |                     |
| ATE mbsmoke      | -236.1038 | 23.86187  | -9.89 | 0.000  | -282.8722           |
| (smoker vs       |           |           |       |        | -189.3354           |
| nonsmoker)       |           |           |       |        |                     |
| POmean mbsmoke   | 3402.552  | 9.539555  | 356.68| 0.000  | 3383.855            |
| nonsmoker        |           |           |       |        | 3421.249            |
```
Rubin (2008) recommends finding a model that balances the covariates before looking at results for the estimated treatment effect. Thus we do not interpret the above results, and we note that we could pay closer heed to Rubin’s recommendation by preceding the `teffects` command with `quietly` to suppress the output.

Imai and Ratkovic (2014) derived a test for balance by viewing the restrictions imposed by balance as overidentifying conditions. This test is implemented in `tebalance overid`, and we use it to test whether the above treatment model balanced the covariates.

```
. tebalance overid
Iteration 0:  criterion = 0.02146858
Iteration 1:  criterion = 0.02159149  (backed up)
Iteration 2:  criterion = 0.02177783
Iteration 3:  criterion = 0.02260102
Iteration 4:  criterion = 0.02267956
Iteration 5:  criterion = 0.02292367
Iteration 6:  criterion = 0.02431697
Iteration 7:  criterion = 0.02457043
Iteration 8:  criterion = 0.02488579
Iteration 9:  criterion = 0.02529453
Iteration 10: criterion = 0.02545885
Iteration 11: criterion = 0.02554462
Iteration 12: criterion = 0.02554514
Iteration 13: criterion = 0.02554512
Iteration 14: criterion = 0.02554512
Iteration 15: criterion = 0.02554514
```

Overidentification test for covariate balance

<table>
<thead>
<tr>
<th>H0: Covariates are balanced:</th>
</tr>
</thead>
<tbody>
<tr>
<td>chi2(5) = 38.1464</td>
</tr>
<tr>
<td>Prob &gt; chi2 = 0.0000</td>
</tr>
</tbody>
</table>

We reject the null hypothesis that the treatment model balanced the covariates.

Let’s use `tebalance summarize` to see where the problem lies. To get an idea of the extent to which the covariates are unbalanced, we begin by summarizing the covariates by group in the raw data by specifying the `baseline` option.

```
. tebalance summarize, baseline
```

<table>
<thead>
<tr>
<th>Covariate balance summary</th>
<th>Raw</th>
<th>Weighted</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of obs = 4,642</td>
<td>4,642</td>
<td>4,642.0</td>
</tr>
<tr>
<td>Treated obs = 864</td>
<td>2,315.3</td>
<td></td>
</tr>
<tr>
<td>Control obs = 3,778</td>
<td>2,326.7</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Means</th>
<th>Control</th>
<th>Treated</th>
<th>Variances</th>
<th>Control</th>
<th>Treated</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mmmarried</td>
<td>.7514558</td>
<td>.4733796</td>
<td>.1868194</td>
<td>.2495802</td>
<td></td>
</tr>
<tr>
<td>mage</td>
<td>26.81048</td>
<td>25.16667</td>
<td>31.87141</td>
<td>28.10429</td>
<td></td>
</tr>
<tr>
<td>prenatal1</td>
<td>.8268925</td>
<td>.6898148</td>
<td>.1431792</td>
<td>.2142183</td>
<td></td>
</tr>
<tr>
<td>fbaby</td>
<td>.4531498</td>
<td>.3715278</td>
<td>.2478707</td>
<td>.2337654</td>
<td></td>
</tr>
</tbody>
</table>

The output indicates that the covariates may not be balanced in the raw data. For example, the distribution of the mother’s age may differ over the treatment groups. We can investigate the differences further with standardized differences and variance ratios. A perfectly balanced covariate has a standardized difference of zero and variance ratio of one. There are no standard errors on these
statistics, so inference is informal. Austin (2009) provides a recent introduction to these diagnostics, although they have been used at least since Rosenbaum and Rubin (1985).

By omitting the `baseline` option, we obtain these diagnostic statistics for the raw data and the weighted data.

```
. tebalance summarize
Covariate balance summary

<table>
<thead>
<tr>
<th></th>
<th>Raw</th>
<th>Weighted</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of obs</td>
<td>4,642</td>
<td>4,642.0</td>
</tr>
<tr>
<td>Treated obs</td>
<td>864</td>
<td>2,315.3</td>
</tr>
<tr>
<td>Control obs</td>
<td>3,778</td>
<td>2,326.7</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Covariate</th>
<th>Standardized differences</th>
<th>Variance ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Raw</td>
<td>Weighted</td>
</tr>
<tr>
<td>mmarrried</td>
<td>-.5953009</td>
<td>-.0105562</td>
</tr>
<tr>
<td>mage</td>
<td>-.300179</td>
<td>-.0672115</td>
</tr>
<tr>
<td>prenatal1</td>
<td>-.3242695</td>
<td>-.0156339</td>
</tr>
<tr>
<td>fbaby</td>
<td>-.1663271</td>
<td>.0257705</td>
</tr>
</tbody>
</table>
```

Reviewing the output, we see that for `mmarrried`, `prenatal1`, and `fbaby`, our model improved the level of balance. The weighted standardized differences are all close to zero and the variance ratios are all close to one. However, output indicates that `mage` may not be balanced by our model. The weighted standardized difference is close to zero, but the weighted variance ratio still appears to be considerably less than one.

Now, let’s use `tebalance density` to look at how the densities of `mage` for treated and control groups differ.

```
. tebalance density mage
```

The plots also indicate a lack of balance in `mage` between the treatment groups.
To try to achieve better balance, we specify a richer model with interactions between `mage` and the other covariates and look at the resulting standardized differences.

```
. quietly teffects ipw (bweight) (mbsmoke mmarried mage prenatal1 fbaby > c.mage#(c.mage i.mmarried prenatal1))
. tebalance summarize
```

Covariate balance summary

<table>
<thead>
<tr>
<th></th>
<th>Raw</th>
<th>Weighted</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of obs</td>
<td>4,642</td>
<td>4,642.0</td>
</tr>
<tr>
<td>Treated obs</td>
<td>864</td>
<td>2,329.1</td>
</tr>
<tr>
<td>Control obs</td>
<td>3,778</td>
<td>2,312.9</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Covariate</th>
<th>Raw Weighted</th>
<th>Variance ratio Raw Weighted</th>
</tr>
</thead>
<tbody>
<tr>
<td>mmarried</td>
<td>-.5953009</td>
<td>.0053497</td>
</tr>
<tr>
<td>mage</td>
<td>-.300179</td>
<td>.0410889</td>
</tr>
<tr>
<td>prenatal1</td>
<td>-.3242695</td>
<td>.0009807</td>
</tr>
<tr>
<td>fbaby</td>
<td>-.1663271</td>
<td>-.0130638</td>
</tr>
<tr>
<td>mage#mage</td>
<td>-.3028275</td>
<td>.0477465</td>
</tr>
<tr>
<td>mmarried#mage</td>
<td>-.6329701</td>
<td>.0197209</td>
</tr>
<tr>
<td>married</td>
<td></td>
<td></td>
</tr>
<tr>
<td>prenatal1#mage</td>
<td>-.4053969</td>
<td>.0182109</td>
</tr>
</tbody>
</table>
The standardized difference and variance ratio results for `mage` look closer to the expected values of zero and one, so we proceed to the formal test.

```plaintext
. tebalance overid
Iteration 0: criterion = .0602349
Iteration 1: criterion = .06172749 (backed up)
Iteration 2: criterion = .06428588 (backed up)
Iteration 4: criterion = .06527284 (backed up)
Iteration 5: criterion = .06643426
Iteration 6: criterion = .07120383
Iteration 7: criterion = .07647097
Iteration 8: criterion = .07674915
Iteration 9: criterion = .07684127
Iteration 10: criterion = .07703321
Iteration 11: criterion = .0776508
Iteration 12: criterion = .07773156
Iteration 14: criterion = .07773561
Iteration 15: criterion = .07774891
Iteration 16: criterion = .07775314
Iteration 17: criterion = .07775324
Iteration 18: criterion = .07775325
Iteration 19: criterion = .07775325
Iteration 20: criterion = .07775325
Iteration 21: criterion = .07775325
Iteration 22: criterion = .07775325

Overidentification test for covariate balance
H0: Covariates are balanced:
  chi2(8) = 11.8612
  Prob > chi2 = 0.1575
```

We do not reject the null hypothesis that the specified treatment model balances the covariates.

Example 2: Balance after estimators that use matching

Instead of weighting, we might want to use a matching estimator. We can select `teffects nnmatch` or `teffects psmatch` for balance and estimation; in this example, we use `teffects nnmatch`.

```plaintext
. teffects nnmatch (bweight mmarried mage prenatal1 fbaby)
> (mbsmoke), bias(mage) ematch(mmarried prenatal1 fbaby)
Treatment-effects estimation
  Number of obs = 4,642
  Estimator : nearest-neighbor matching
  Matches: requested = 1
  Outcome model : matching
  min = 1
  Distance metric: Mahalanobis
  max = 139

                AI Robust
              Coef. Std. Err.  z  P>|z|  [95% Conf. Interval]
ATE
  mbsmoke
    (smoker vs nonsmoker)  -240.4589   28.43008  -8.46   0.000  -296.1808  -184.7369
```

Again we ignore the estimated effect and first check for balance. We begin by reviewing whether the summary statistics indicate good balance.
We do not have standard errors on these statistics, so we cannot make any formal conclusions, but the summary statistics appear to indicate much better balance than the IPW results. `tebalance` summarize has to refit the model to recover the matched sample because the `generate()` option was not specified on the `teffects nnmatch` command. The reestimation does not affect the results, although the computation takes longer; see example 3 for details.

Because it is a matching estimator, and not an IPW estimator, we cannot use `tebalance overid` after `teffects nnmatch`. The matching estimators, however, provide diagnostic box plots using `tebalance box` that are not available after the IPW estimators.

```
. tebalance box mage
   note: refitting the model using the `generate()` option
```

The box plots of the matched data also indicate covariate balance.
Let's look at the kernel density plots using the matched data.

```
. tebalance density mage
note: refitting the model using the generate() option
```

The plots using the matched data appear to be balanced.

### Technical note

`teffects` implements matching estimators, IPW estimators, regression-adjustment (RA) estimators, and estimators that combine IPW and RA. Matching estimators define a matched sample, and IPW estimators define a weighted sample, each of which can be used to compute covariate balance statistics. RA estimators do not define an adjusted sample that can be used to compute covariate balance statistics, and `tebalance` does not work after `teffects ra`. Only the IPW component of the estimators that combine RA and IPW defines a weighted sample that can be used to compute balance statistics. So, `tebalance` produces the same results after `teffects aipw` or `teffects ipwra` as it does after `teffects ipw`.

### Example 3: Faster results after a matching estimator

The `tebalance` commands run in example 2 executed more slowly than necessary. `tebalance` issued the note

```
note: refitting the model using the generate() option
```

after the commands

```
. tebalance summarize
. tebalance box mage
```

and

```
. tebalance density mage
```
After \texttt{teffects nnmatch} or \texttt{teffects psmatch}, \texttt{tebalance} computes the balance statistics on the matched sample defined by the matching estimator. \texttt{teffects nnmatch} and \texttt{teffects psmatch} leave behind only variables that identify the matched sample when the \texttt{generate()} option is specified. Unless the \texttt{generate()} option is specified with \texttt{teffects nnmatch} or \texttt{teffects psmatch}, each \texttt{tebalance} command must rerun the estimation command to recover the matched sample.

Typing

\begin{verbatim}
. teffects nnmatch (bweight mmarried mage fbaby prenatal1) > (mbsmoke), bias(mage) ematch(mmarried fbaby prenatal1) > generate(matchv)
\end{verbatim}

would generate variables that identify the matched sample that the \texttt{tebalance} commands could use. See \textit{Remarks and examples} in \cite{TE tebalance box}, \cite{TE tebalance density}, and \cite{TE tebalance summarize} for examples using the option \texttt{generate()} on \texttt{teffects psmatch} to speed up the postestimation computations.

### Methods and formulas

Methods and formulas are presented under the following headings:

- \textit{Introduction}
- \textit{Matched samples}
- \textit{IPW samples}
- \textit{Testing the propensity-score model specification}

#### Introduction

For covariate $z$, we observe values $\{z_1, z_2, \ldots, z_N\}$. Define a treatment indicator variable for $J$ treatment levels as $t_i \in \{1, 2, \ldots, J\}$, for $i = 1, \ldots, N$, and frequency weights as $\{w_1, w_2, \ldots, w_N\}$. The sample mean and variance of $z$ for level $t$ are

\[
\hat{\mu}_z(t) = \frac{\sum_i I(t_i = t)w_iz_i}{N_t} \quad \text{and} \quad \hat{\sigma}^2_z(t) = \frac{\sum_i I(t_i = t)w_i\{z_i - \hat{\mu}_z(t)\}^2}{N_t - 1}
\]

where $N_t = \sum_i w_iI(t_i = t)$, and

\[
I(t_i = t) = \begin{cases} 1 & \text{if } t_i = t \\ 0 & \text{otherwise} \end{cases}
\]

As shown in Austin (2011), the standardized differences for covariate $z$ between level $t$ and the control $t_0$ are computed as

\[
\delta_z(t) = \frac{\hat{\mu}_z(t) - \hat{\mu}_z(t_0)}{\sqrt{\hat{\sigma}^2_z(t) + \hat{\sigma}^2_z(t_0)}}
\] (1)

The variance ratio is $\rho_z(t) = \{\hat{\sigma}^2_z(t)\}/\{\hat{\sigma}^2_z(t_0)\}$. 


Matched samples

We now turn our attention to the matched samples for the potential-outcome mean (POM), average treatment effect (ATE), and average treatment effect on the treated (ATET) estimators. We estimate the covariate for the counter-factual treatment by taking the mean of the matched observations

\[ \hat{z}_i = \frac{\sum_{j \in \Omega(i)} w_j z_j}{\sum_{j \in \Omega(i)} w_j} \]

where \( \Omega(i) = (k_1, k_2, \ldots, k_m) \) is the set of observation indices that are matched to observation \( i \) of the opposite treatment condition. The observed covariate and matched covariate pairs, \((z_i, \hat{z}_i)\), \(i = 1, \ldots, N\), are used in the box plot (see [G-2] graph box) and the kernel density plot (see [R] kdensity). The ATET sample is limited to those observations from the conditional treatment, \( \tilde{t} \), and their matched covariate means.

In Methods and formulas of [TE] teffects nnmatch, we define \( K_m(i) \) as the number of times observation \( i \) is used in a match with observation \( j \) of the opposite treatment condition, \( i \in \Omega(j) \), weighted by the total number of matches for observation \( j \). Specifically,

\[ K_m(i) = \sum_{j=1}^{N} I\{i \in \Omega(j)\} \frac{w_j}{\sum_{k \in \Omega(j)} w_k} \]

These weights are used in the estimation of the mean and variance for the matched dataset. For the POM and ATE models, the estimated mean and variance are computed as

\[ \hat{\mu}_z(t) = \frac{\sum_i^{N} I(t_i = t) w_i z_i \{1 + K_m(i)\}}{M_t} \quad \text{and} \quad \hat{\sigma}_z^2(t) = \frac{\sum_i^{N} I(t_i = t) w_i \{z_i - \hat{\mu}_z(t)\}^2}{M_t - 1} \]

where \( M_t = \sum_i^{N} I(t_i = t) w_i \{1 + K_m(i)\} \).

The standardized differences between the control level and all other levels for the matched covariate distribution are computed as in (1), but \( \hat{\mu}_z(t) \) is substituted for \( \hat{\mu}_z(t) \) and \( \hat{\sigma}_z^2(t) \) for \( \hat{\sigma}_z^2(t) \).

For the ATET model, there is no matched sample for the treatment levels other than the conditional treatment \( \tilde{t} \). The covariate mean and variance for the conditional treatment are the same as that of the raw data, \( \mu_z(\tilde{t}) \) and \( \sigma_z(\tilde{t}) \). However, the covariate mean and variance for the sample matched to the conditional treatment, \( t \neq \tilde{t} \), are computed as

\[ \hat{\mu}_z(t) = \frac{\sum_i^{N} I(t_i = t) w_i K_m(i)}{M_t} \quad \text{and} \quad \hat{\sigma}_z^2(t) = \frac{\sum_i^{N} I(t_i = t) w_i K_m(i) \{z_i - \hat{\mu}_z(t)\}^2}{M_t - 1} \]

where \( M_t = \sum_i^{N} I(t_i = t) w_i K_m(i) \).
IPW samples

Computation of the inverse-probability weights is discussed in Methods and formulas of [TE] teffects aipw and in Methods and formulas of [TE] stteffects ipwra. For the POM and ATE estimators, we defined the normalized IPW weights as \( \tilde{d}_i(t) = N_i d_i(t) / \sum_i N_i d_i(t) \), where \( d_i(t) = I(t_i = t) / p(z_i, t, \hat{\gamma}) \) for treatment level \( t \) and individual \( i \).

For the ATET estimator, we use the normalized weights \( \tilde{f}_i = N_i f_i / \sum_i N_i f_i \), where \( f_i = p(z_i, \tilde{t}, \hat{\gamma}) / p(z_i, t_i, \hat{\gamma}) \) are the treatment-adjusted inverse-probability weights, and \( \tilde{t} \) is the conditional treatment.

We will simplify notation by defining a single weight

\[
\bar{w}_i(t) = \begin{cases} 
\tilde{d}_i(t) & \text{if model is ATE or POM} \\
\tilde{f}_i(t) & \text{if model is ATET}
\end{cases}
\]

The covariate mean and variance for treatment level \( t \) are

\[
\tilde{\mu}_z(t) = \frac{\sum_i N_i I(t_i = t) w_i \bar{w}_i x_i}{M_t} \quad \text{and} \quad \tilde{\sigma}^2_z(t) = \frac{I(t_i = t) w_i \bar{w}_i \{ z_i - \tilde{\mu}_z(t) \}^2}{M_t - 1}
\]

where \( M_t = \sum_i N_i I(t_i = t) w_i \bar{w}_i \).

The kernel density is computed by \texttt{kdensity} for each covariate conditioned on each treatment level using the raw covariate with \texttt{iweight}s equal to \( w_i \bar{w}_i \).

Testing the propensity-score model specification

We estimate the probability of treatment conditioned on a set of covariates with a propensity-score model. Imai and Ratkovic (2014) derive a test for whether the estimated propensity score balances the covariates. The score equations for parameters of the propensity-score model define an exactly identified generalized method of moments (GMM) estimator. Imai and Ratkovic (2014) use the conditions imposed by mean balance as overidentifying conditions. A standard GMM test for the validity of the overidentifying conditions is then a test for covariate balance. See [R] \texttt{gmm} for a discussion of this overidentifying test, which is known as Hansen’s \( J \) test in the econometrics literature.

Here are the details about the score equations and the overidentifying balance conditions. Recall from Methods and formulas of [TE] teffects aipw and Methods and formulas of [TE] stteffects ipwra, we have the first-order condition of the treatment model

\[
\frac{1}{N} \sum_{i=1}^{N} s_{tm,i}(z_i, \tilde{\gamma}) = 0
\]

For a two-level treatment-effects model with conditional treatment \( \tilde{t} \) and control \( t_0 \), the score is

\[
s_{tm,i}(z_i, \gamma) = \frac{I(t_i = \tilde{t})}{p(z_i, \tilde{t}, \gamma)} \frac{\partial p(z_i, t, \gamma)}{\partial \gamma'} - \left\{ \frac{I(t_i = t_0)}{1 - p(z_i, \tilde{t}, \gamma)} \right\} \frac{\partial p(z_i, \tilde{t}, \gamma)}{\partial \gamma'} \bigg|_{\gamma = \tilde{\gamma}}
\]
The score reduces to

\[ s_{tm,i}(z_i, \tilde{\gamma}) = \left[ \frac{I(t_i = \tilde{t}) - p(z_i, \tilde{t}, \gamma)}{p(z_i, \tilde{t}, \gamma) \{1 - p(z_i, \tilde{t}, \gamma)\}} \right] \frac{\partial p(z_i, \tilde{t}, \gamma)}{\partial \gamma} \bigg|_{\gamma=\tilde{\gamma}} \]

The corresponding covariate balancing moment conditions are

\[ w_{tm,i}(z_i, \gamma) = \left[ \frac{I(t_i = \tilde{t}) - p(z_i, \tilde{t}, \gamma)}{p(z_i, \tilde{t}, \gamma) \{1 - p(z_i, \tilde{t}, \gamma)\}} \right] z_i \]

for the POM and ATE models. For the ATET model with conditional treatment \( \tilde{t} \), we multiply by \( p(z_i, \tilde{t}, \gamma) \) and scale by \( N/N_{\tilde{t}} \):

\[ w_{tm,i}(z_i, \gamma) = \frac{N}{N_{\tilde{t}}} \left\{ \frac{I(t_i = \tilde{t}) - p(z_i, \tilde{t}, \gamma)}{1 - p(z_i, \tilde{t}, \gamma)} \right\} z_i \]

We stack the moment conditions

\[ g_{tm}(Z, \gamma) = \frac{1}{N} \sum_{i=1}^{N} \left\{ s_{tm,i}(z_i, \gamma) \right\} \]

\[ = \frac{1}{N} \sum_{i=1}^{N} g_{tm,i}(z_i, \gamma) \]

The overidentified GMM estimator is then

\[ \tilde{\gamma} = \text{argmin}_{\gamma} N g_{tm}(Z, \gamma)^{\prime} W_{tm}(Z, \gamma)^{-1} g_{tm}(Z, \gamma) \quad (2) \]

where

\[ W_{tm}(Z, \gamma) = \frac{1}{N} \sum_{i=1}^{N} ET \left\{ g_{tm,i}(z, \gamma) g_{tm,i}(z, \gamma)^{\prime} \right\} \]

and the expectation is taken with respect to treatment distribution. The weight matrix \( W_{tm}(Z, \gamma) \) is computed explicitly (Imai and Ratkovic 2014), and (2), written as a maximization problem, is solved using \texttt{ml}.

Finally, Hansen’s \( J \) statistic is evaluated at its minimum,

\[ J = N g_{tm}(Z, \tilde{\gamma})^{\prime} W_{tm}(Z, \tilde{\gamma})^{-1} g_{tm}(Z, \tilde{\gamma}) \]

and is asymptotically distributed chi-squared with degrees of freedom \( d \),

\[ d = \text{rank} \left\{ W_{tm}(Z, \tilde{\gamma}) \right\} - \text{rank} \left[ \frac{1}{N} \sum_{i=1}^{N} ET \left\{ s_{tm,i}(z_i, \tilde{\gamma}) s_{tm,i}(z_i, \tilde{\gamma})^{\prime} \right\} \right] \]
References


Also see

[TE] **tobalance summarize** — Covariate-balance summary statistics

[TE] **tobalance overid** — Test for covariate balance

[TE] **tobalance density** — Covariate balance density

[TE] **tobalance box** — Covariate balance box

[TE] **tffects** — Treatment-effects estimation for observational data

[TE] **sttffects intro** — Introduction to treatment effects for observational survival-time data
Description

tebalance box produces box plots that are used to check for balance in matched samples after teffects nnmatch and teffects psmatch.

Quick start

Box plot of the propensity score from the last teffects psmatch command

tebalance box

Box plot of values of $x_1$ in the treatment and control groups from raw data and the matched sample from the last teffects nnmatch or teffects psmatch command

tebalance box x1

Menu

Statistics  >  Treatment effects  >  Balance  >  Graphs
Syntax

Box plots for the propensity score

```
tebalance box [, options]
```

Box plots for a covariate

```
tebalance box varname [, options]
```

options Description

<table>
<thead>
<tr>
<th>Main</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td><code>boxlook_options</code></td>
<td>graph box options controlling how the box looks</td>
</tr>
<tr>
<td><code>legending_options</code></td>
<td>graph box options controlling how the variables are labeled</td>
</tr>
<tr>
<td><code>axis_options</code></td>
<td>graph box options controlling how numerical y axis is labeled</td>
</tr>
<tr>
<td><code>title_and_other_options</code></td>
<td>graph box options controlling titles, added text, aspect ratio, etc.</td>
</tr>
<tr>
<td><code>by_options</code></td>
<td>suboptions inside <code>by()</code> controlling plots by raw and matched samples</td>
</tr>
</tbody>
</table>

Options

`boxlook_options` are any of the options documented in `boxlook_options` in [G-2] graph box. `legending_options` are any of the options documented in `legending_options` in [G-2] graph box. `axis_options` are any of the options documented in `axis_options` in [G-2] graph box. `title_and_other_options` are any of the options, except `by()`, documented in `title_and_other_options` in [G-2] graph box. `tebalance box` uses `by()` to differentiate between raw and matched samples, and some `twoway_options` will be repeated for `by graph` and might be better specified as `byopts()`. `by_options` are any of the `byopts` documented in [G-3] `by_option`. `byopts()` generally affects the entire graph, and some `by_options` may be better specified as `twoway_options`; see [G-3] `twoway_options`.

Remarks and examples

When the distribution of a covariate does not vary over the treatment levels, the covariate is said to be balanced. `tebalance box` produces box plots of a covariate over treatment levels for the raw data and for the matched sample produced by `teffects`. If the matched-sample box plots are the same over the treatment levels, the covariate is balanced in the matched sample.

After `teffects nnmatch` and `teffects psmatch`,

```
  . tebalance box varname [, options]
```
produces box plots to check whether `varname` is balanced.
After *teffects psmatch*,

```
  . tebalance box [, options]
```

produces box plots to check whether the propensity score estimated by *teffects* is balanced.

We recommend that you read [TE] *tebalance* before proceeding; it provides an introduction to covariate balance and an overview of the implemented methods.

> **Example 1: Checking covariate balance after psmatch**

Using an extract from the data used by Cattaneo (2010), we use *teffects psmatch* to estimate the effect of a mother’s smoking behavior (*mbsmoke*) on the birthweight of her child (*bweight*), controlling for marital status (*mmarried*), the mother’s age (*mage*), whether the mother had a prenatal doctor’s visit in the baby’s first trimester (*prenatal1*), and whether this baby is the mother’s first child (*fbaby*).

```
  . use http://www.stata-press.com/data/r15/cattaneo2
  . teffects psmatch (bweight) (mbsmoke mmarried mage prenatal1 fbaby),
    > generate(matchv)
```

```
    Treatment-effects estimation
    Estimator : propensity-score matching
    Matches: requested =  1
    Outcome model : matching
    Treatment model: logit
    Number of obs =  4,642
    min =  1
    max =  139

<table>
<thead>
<tr>
<th></th>
<th>AI Robust</th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coef.</td>
<td>Std. Err.</td>
<td>z</td>
<td>P&gt;</td>
<td>z</td>
</tr>
<tr>
<td>bweight</td>
<td></td>
<td></td>
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<td></td>
</tr>
<tr>
<td>ATE</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mbsmoke (smoker vs nonsmoker)</td>
<td>-235.1714</td>
<td>27.74409</td>
<td>-8.48</td>
<td>0.000</td>
<td>-289.5488 -180.794</td>
</tr>
</tbody>
</table>
```

We specified the option `generate(matchv)` to speed up the postestimation command that produces density plots, as discussed in example 3 under *Remarks and examples* of [TE] *tebalance*. We do not interpret the estimated effect produced by this preliminary model but rather check the specification. Now we look at the box plots.
The box plots for the matched sample are very similar. The medians, the 25th percentiles, and the 75th percentiles appear to be the same, although there may be some differences in the tails, the upper adjacent values, the lower adjacent values, and the outliers. Matching on the estimated propensity score appears to have balanced `mage`, except for the tails.

To get an idea of whether `teffects psmatch` balanced all the covariates, we look at the box plots for the estimated propensity score.

The box plots indicate that `teffects psmatch` balanced the estimated propensity scores.
Reference


Also see

[TE] tebalance — Check balance after tefeats or stteffects estimation
[TE] tefeats nnnmatch — Nearest-neighbor matching
[TE] tefeats psmatch — Propensity-score matching
[TE] tefeats overlap — Overlay plots
Description

tebalance density produces kernel density plots that are used to check for covariate balance after estimation by a teffects inverse-probability-weighted estimator, a teffects matching estimator, or an stteffects inverse-probability-weighted estimator.

Quick start

Kernel density plot of the propensity score after teffects psmatch
tebalance density

Kernel density plot of x1 after a teffects command or an stteffects command
tebalance density x1

As above, but rescale the kernel bandwidth by a factor of 2
tebalance density x1, bwidth(*2)

Menu

Statistics > Treatment effects > Balance > Graphs
Syntax

Density plots for the propensity score

tebalance density [, options]

Density plots for a covariate

tebalance density varname [, options]

Options

<table>
<thead>
<tr>
<th>Options</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Main</td>
<td></td>
</tr>
<tr>
<td>kernel(kernel)</td>
<td>specify the kernel function; default is kernel(epanechnikov)</td>
</tr>
<tr>
<td>bwidth(*)</td>
<td>rescale default bandwidth</td>
</tr>
<tr>
<td>line#opts(line_options)</td>
<td>twoway line options for density line number #</td>
</tr>
<tr>
<td>twoway_options</td>
<td>any options other than by() documented in [G-3 twoway_options]</td>
</tr>
<tr>
<td>byopts(byopts)</td>
<td>how subgraphs are combined, labeled, etc.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>kernel</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>triangle</td>
<td>triangle kernel function; the default</td>
</tr>
<tr>
<td>epanechnikov</td>
<td>Epanechnikov kernel function</td>
</tr>
<tr>
<td>epan2</td>
<td>alternative Epanechnikov kernel function</td>
</tr>
<tr>
<td>biweight</td>
<td>biweight kernel function</td>
</tr>
<tr>
<td>cosine</td>
<td>cosine trace kernel function</td>
</tr>
<tr>
<td>gaussian</td>
<td>Gaussian kernel function</td>
</tr>
<tr>
<td>parzen</td>
<td>Parzen kernel function</td>
</tr>
<tr>
<td>rectangle</td>
<td>rectangle kernel function</td>
</tr>
</tbody>
</table>

Options

- **Main**
  - `kernel(kernel)` specifies the kernel function for use in calculating the kernel density estimates. The default kernel is the `kernel(epanechnikov)`.
  - `bwidth(*)` specifies the factor by which the default bandwidths are to be rescaled. A bandwidth is the half-width of the kernel, the width of the density window around each point. Each kernel density plot has its own bandwidth, and by default, each kernel density plot uses its own optimal bandwidth; see [R] `kdensity`. `bwidth()` rescales each plot’s optimal bandwidth by the specified amount.
  - `line#opts(line_options)` specifies the line pattern, width, color, and overall style of density line number #. The line numbers are in the same order as the treatment levels specified in `e(tlevels)`.
  - `twoway_options` are any of the options documented in [G-3 `twoway_options`], excluding by(). These include options for titling the graph (see [G-3 `title_options`]) and for saving the graph to disk (see [G-3 `saving_option`]). `tebalance density` uses by() to differentiate between raw and weighted or matched samples, and some `twoway_options` will be repeated for by graph and might be better specified as `byopts()`.
byopts(by_option) is as documented in [G-3] by_option. byopts() affects how the subgraphs are combined, labeled, etc. byopts() generally affects the entire graph, and some by_option may be better specified as twoway_options; see [G-3] twoway_options.

Remarks and examples

When the distribution of a covariate does not vary over the treatment levels, the covariate is said to be balanced. tebalance density produces kernel density plots of a covariate over treatment levels for the raw data and the weighted or matched sample produced by tefects or stteffects. If the weighted-sample or matched-sample kernel density plots of the covariate are the same over the treatment levels, the covariate is balanced in the weighted or matched sample.

After all tefects commands except tefects ra, stteffects ipw, and stteffects ipwra,

```
. tebalance density varname [, options]
```

produces kernel density plots to check whether varname is balanced.

After all tefects commands except tefects ra, tefects nmmatch, stteffects ipw, and stteffects ipwra,

```
. tebalance density [, options]
```

produces kernel density plots to check whether the propensity score estimated by tefects or tefects is balanced. Our discussion of the use of tebalance density and interpretation of its results for a covariate below also apply to a propensity score.

We recommend that you read [TE] tebalance before proceeding; it provides an introduction to covariate balance and an overview of the implemented methods. See [TE] stteffects intro for a discussion of survival-time features.

Example 1: Checking covariate balance after psmatch

Using an extract from the data used by Cattaneo (2010), we use tefects psmatch to estimate the effect of a mother’s smoking behavior (mbsmoke) on the birthweight of her child (bweight), controlling for marital status (mmarried), the mother’s age (mage), whether the mother had a prenatal doctor’s visit in the baby’s first trimester (prenatal1), and whether this baby is the mother’s first child (fbaby).

```
. use http://www.stata-press.com/data/r15/cattaneo2
. tefects psmatch (bweight) (mbsmoke mmarried mage prenatal1 fbaby),
> generate(matchv)
```

```
Treatment-effects estimation
Number of obs = 4,642
Estimator : propensity-score matching
Matches: requested = 1
Outcome model : matching
min = 1
Treatment model: logit
max = 139

<table>
<thead>
<tr>
<th>bweight</th>
<th>AI Robust</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coef.</td>
</tr>
<tr>
<td>ATE</td>
<td></td>
</tr>
<tr>
<td>mbsmoke (smoker vs nonsmoker)</td>
<td>-235.1714</td>
</tr>
</tbody>
</table>
We specified the option `generate(matchv)` to speed up the postestimation command that produces density plots, as discussed in example 3 under Remarks and examples in [TE] tebalance entry. We do not interpret the estimated effect produced by this preliminary model but rather check the specification. We begin by looking at the default density plots.

```
. tebalance density mage
```

The density plots for the matched sample are nearly indistinguishable, implying that matching on the estimated propensity score balanced the covariates. The density plots are too jagged for presentation, so we oversmooth them by scaling up the bandwidth used for each plot.

```
. tebalance density mage, bwidth(*1.5)
```

Option `bwidth()` rescales the default optimal bandwidths by the specified scale factor. Each of the four density plots has its own sample size and optimal bandwidth. Rescaling each of the four bandwidths by 1.5 produces smoother plots.
Stored results

After `teffects` or `stteffects` fits a binary treatment, `tebalance density` stores the following in `r()`:

Scalars

- `r(bwc_adj)` bandwidth for control in weighted or matched-adjusted sample
- `r(Nc_adj)` observations on control in weighted or matched-adjusted sample
- `r(bwt_adj)` bandwidth for treated in weighted or matched-adjusted sample
- `r(Nt_adj)` observations on treated in weighted or matched-adjusted sample
- `r(bwc_raw)` bandwidth for control in raw sample
- `r(Nc_raw)` observations on control in raw sample
- `r(bwt_raw)` bandwidth for treated in raw sample
- `r(Nt_raw)` observations on treated in raw sample

Macros

- `r(kernel)` name of kernel

After `teffects` or `stteffects` fits a multivalued treatment, `tebalance density` stores the following in `r()`:

Scalars

- `r(bw#_adj)` bandwidth for treatment level # in weighted or matched-adjusted sample
- `r(N#_adj)` observations on treatment level # in weighted or matched-adjusted sample
- `r(bw#_raw)` bandwidth for treatment level # in raw sample
- `r(N#_raw)` observations on treatment level # in raw sample

Macros

- `r(kernel)` name of kernel

Reference


Also see

- `[TE] tebalance` — Check balance after `teffects` or `stteffects` estimation
- `[TE] teeffects aipw` — Augmented inverse-probability weighting
- `[TE] teeffects ipw` — Inverse-probability weighting
- `[TE] teeffects ipwra` — Inverse-probability-weighted regression adjustment
- `[TE] teeffects nnmatch` — Nearest-neighbor matching
- `[TE] teeffects psmatch` — Propensity-score matching
- `[TE] teeffects overlap` — Overlap plots
- `[TE] stteffects intro` — Introduction to treatment effects for observational survival-time data
tebalance overid — Test for covariate balance

Description

tebalance overid performs a test for covariate balance after estimation by a teffects inverse-probability-weighted (IPW) estimator or an stteffects IPW estimator.

Quick start

Test for covariate balance after a teffects or an stteffects IPW estimator

tebalance overid

As above, but test for balance only in base covariates and exclude interaction terms

tebalance overid, bonly

Menu

Statistics > Treatment effects > Balance > Overidentification test
Syntax

tebalance overid [, bconly nolog iterate(#) ]

Options

bconly specifies that only the base covariates be included in the test for balance. By default, the powers and interactions specified by factor-variable notation in the teffects or stteffects model are also included in the test for balance.

nolog suppresses the display of the optimization search log.

iterate(#) sets the maximum number of iterations to # in the generalized method of moments estimator used to compute the test statistic.

Remarks and examples

When the distribution of a covariate is the same for all treatment levels, the covariate is said to be balanced. tebalance overid performs a test to see whether the covariates are balanced after teffects or stteffects. tebalance overid can be executed after teffects ipw, teffects aipw, teffects ipwra, stteffects ipw, or stteffects ipwra, which use the inverse-probability weights predicted by a treatment model to account for how treatment assignment depends on observed covariates. If the treatment model is well specified, IPW functions of the covariates from the model are balanced.

We recommend that you read [TE] tebalance before proceeding; it provides an introduction to covariate balance and an overview of the implemented methods. See [TE] stteffects intro for survival-time discussion and examples.

Example 1: Base covariates and interactions

This example illustrates the interpretation of the bconly option, which excludes powers and interactions when factor variables are included in the propensity-score model.

We frequently use factor variables to include powers of, and interactions between, base covariates in our specification of the propensity-score model. In example 1 under Remarks and examples in [TE] tebalance, we rejected the null hypothesis of balance in a model using only base covariates but not in the richer model that included power and interaction terms. By default, tebalance overid tests whether the model balances the base covariates and the power-and-interaction covariates. When option bconly is specified, tebalance overid tests whether the model balances the base covariates only.

Using an extract from the data used by Cattaneo (2010), we use teffects ipw to estimate the effect of a mother’s smoking behavior (mbsmoke) on the birthweight of her child (bweight), controlling for marital status (mmarried), the mother’s age (mage), whether the mother had a prenatal doctor’s visit in the baby’s first trimester (prenatal1), and whether this baby is the mother’s first child (fbaby). In addition to the base covariates, we include the square of mage, an interaction between mage and mmarried, and an interaction between mage and prenatal1 in the model for the propensity score.
We specified option aequations to see the parameter estimates for the coefficients in the propensity-score model. There are eight coefficients, five on the base covariates (mmarried, mage, fbaby, prenatal1, and _cons) and three on the power-and-interaction covariates (c.mage#c.mage, c.mage#1.mmarried, and c.mage#1.prenatal1). Below we test whether the model balances all eight covariates.
. tebalance overid

Iteration 0:  criterion =  .0602349
Iteration 1:  criterion =  .06172749 (backed up)
Iteration 2:  criterion =  .06428588 (backed up)
Iteration 3:  criterion =  .06489623 (backed up)
Iteration 4:  criterion =  .06527284 (backed up)
Iteration 5:  criterion =  .06643426
Iteration 6:  criterion =  .07120383
Iteration 7:  criterion =  .07647097
Iteration 8:  criterion =  .07647097
Iteration 9:  criterion =  .07703321
Iteration 10: criterion =  .0776508
Iteration 11: criterion =  .07771863
Iteration 12: criterion =  .07773156
Iteration 13: criterion =  .07773561
Iteration 14: criterion =  .07774891
Iteration 15: criterion =  .07775314
Iteration 16: criterion =  .07775324
Iteration 17: criterion =  .07775325
Iteration 18: criterion =  .07775325
Iteration 19: criterion =  .07775325
Iteration 20: criterion =  .07775325
Iteration 21: criterion =  .07775325
Iteration 22: criterion =  .07775325

Overidentification test for covariate balance
H0: Covariates are balanced:
    chi2(8)  =  11.8612
    Prob > chi2 =  0.1575

We cannot reject the null hypothesis that the IPW model balanced all eight covariates.

Below we specify option bconly to test whether the IPW model balanced the five base covariates only.

. tebalance overid, bconly

Iteration 0:  criterion =  .1079977
Iteration 1:  criterion =  .10800825 (backed up)
Iteration 2:  criterion =  .10844177 (backed up)
Iteration 3:  criterion =  .10851228 (backed up)
Iteration 4:  criterion =  .10860856 (backed up)
Iteration 5:  criterion =  .10907494
Iteration 6:  criterion =  .11009793
Iteration 7:  criterion =  .11164037
Iteration 8:  criterion =  .11260665
Iteration 9:  criterion =  .11286445
Iteration 10: criterion =  .11331466
Iteration 11: criterion =  .11333969
Iteration 12: criterion =  .11335601
Iteration 13: criterion =  .11335696
Iteration 14: criterion =  .11335696
Iteration 15: criterion =  .11335696

Overidentification test for covariate balance
H0: Covariates are balanced:
    chi2(5)  =  7.82169
    Prob > chi2 =  0.1663

We cannot reject the null hypothesis that the IPW model balanced the five base covariates.

Each test has a justification.
In a model-based approach, the Imai and Ratkovic (2014) test checks whether the propensity score is correctly specified. We include all eight covariates because they must all be balanced, if the propensity-score model is correctly specified.

A conditional-independence approach can be used to justify only including the base covariates in the test. In this approach, the propensity-score model need only balance the base covariates. Powers and interactions of the base covariates are included to get a propensity-score model that balances the base covariates, but balance of these higher-order terms is more than what needs to be checked.

In large samples, both tests should have nominal coverage under the null hypothesis that the propensity-score model is correctly specified. Under the alternative that the propensity-score model is misspecified, including all the covariates should yield a test with higher power.

The test that includes all the covariates is the default.

### Stored results

tebalance overid stores the following in r():

- Scalars:
  - r(p): \( p \)-value
  - r(df): overidentifying constraints, test degrees of freedom
  - r(chi2): chi-squared statistic

### References


### Also see

- [TE] tebalance — Check balance after teffects or stteffects estimation
- [TE] teffects aipw — Augmented inverse-probability weighting
- [TE] teffects ipw — Inverse-probability weighting
- [TE] teffects ipwra — Inverse-probability-weighted regression adjustment
- [TE] teffects overlap — Overlap plots
- [TE] stteffects intro — Introduction to treatment effects for observational survival-time data
tebalance summarize reports diagnostic statistics that are used to check for covariate balance over treatment groups after estimation by a teffects inverse-probability-weighted (IPW) estimator, a teffects matching estimator, or an stteffects IPW estimator.

**Quick start**

Raw and weighted standardized differences and variance ratios of all covariates from the most recently estimated teffects model or stteffects model

```stata
tebalance summarize
```

As above, but report statistics only for covariates x1 and x2

```stata
tebalance summarize x1 x2
```

Baseline means and variances for treated and control groups

```stata
tebalance summarize, baseline
```

### Menu

- Statistics > Treatment effects > Balance > Summaries
Syntax

```
tebalance summarize [varlist] [, baseline]
```

* varlist may contain factor variables; see [U] 11.4.3 Factor variables.

Option

- **baseline** specifies that tebalance summarize report means and variances by treatment level.

Remarks and examples

When the distribution of a covariate is the same for all treatment levels, the covariate is said to be balanced. tebalance summarize reports diagnostic statistics to check for covariate balance after teffects or stteffects. tebalance summarize can be executed after all teffects estimators with the exception of teffects ra and executed after stteffects ipw and stteffects ipwra.

We recommend that you read [TE] tebalance before proceeding; it provides an introduction to covariate balance and an overview of the implemented methods. See [TE] stteffects intro for survival-time discussion and examples.

➢ Example 1: Checking covariate balance after psmatch

Using an extract from the data used by Cattaneo (2010), we use teffects psmatch to estimate the effect of a mother’s smoking behavior (mbsmoke) on the birthweight of her child (bweight), controlling for marital status (mmarried), the mother’s age (mage), whether the mother had a prenatal doctor’s visit in the baby’s first trimester (prenatal1), and whether this baby is the mother’s first child (fbaby).

```
. use http://www.stata-press.com/data/r15/cattaneo2
. teffects psmatch (bweight) (mbsmoke mmarried mage prenatal1 fbaby),
   > generate(matchv)
```

```
<table>
<thead>
<tr>
<th>Treatment-effects estimation</th>
<th>Number of obs = 4,642</th>
</tr>
</thead>
<tbody>
<tr>
<td>Estimator : propensity-score matching</td>
<td>Matches: requested = 1</td>
</tr>
<tr>
<td>Outcome model : matching</td>
<td>min = 1</td>
</tr>
<tr>
<td>Treatment model: logit</td>
<td>max = 139</td>
</tr>
</tbody>
</table>
```

| bweight  | Coef.  | Std. Err. | z    | P>|z|   | [95% Conf. Interval] |
|----------|--------|-----------|------|-------|---------------------|
| ATE      |        |           |      |       |                     |
| mbsmoke  |        |           |      |       |                     |
| smoker   |        |           |      |       |                     |
| vs       |        |           |      |       |                     |
| nonsmoker|        |           |      |       |                     |

-235.1714  27.74409  -8.48  0.000  -289.5488  -180.794
We specified the option `generate(matchv)` to speed up the postestimation commands that compute balance statistics, as discussed in `example 3` under Remarks and examples in [TE] `tebalance`. We do not interpret the estimated effect produced by this preliminary model but rather check the specification.

We begin by looking at the standardized differences and variance ratios for the raw data and the matched sample.

```
. tebalance summarize
Covariate balance summary

<table>
<thead>
<tr>
<th></th>
<th>Raw</th>
<th>Matched</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of obs</td>
<td>4,642</td>
<td>9,284</td>
</tr>
<tr>
<td>Treated obs</td>
<td>864</td>
<td>4,642</td>
</tr>
<tr>
<td>Control obs</td>
<td>3,778</td>
<td>4,642</td>
</tr>
</tbody>
</table>
```

<table>
<thead>
<tr>
<th></th>
<th>Raw</th>
<th>Matched</th>
</tr>
</thead>
<tbody>
<tr>
<td>mmarried</td>
<td>-.5953009</td>
<td>.0014107</td>
</tr>
<tr>
<td>mage</td>
<td>-.300179</td>
<td>-.0120277</td>
</tr>
<tr>
<td>prenatal1</td>
<td>-.3242695</td>
<td>.0333609</td>
</tr>
<tr>
<td>fbaby</td>
<td>-.1663271</td>
<td>-.0117326</td>
</tr>
</tbody>
</table>

The matched sample results indicate that matching on the estimated propensity score balanced the covariates. The standardized differences are all close to zero, and the variance ratios are all close to one. This inference is informal because we do not have standard errors for these statistics.

We may also wish to see the baseline summary statistics.

```
. tebalance summarize, baseline
Covariate balance summary

<table>
<thead>
<tr>
<th></th>
<th>Raw</th>
<th>Matched</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of obs</td>
<td>4,642</td>
<td>9,284</td>
</tr>
<tr>
<td>Treated obs</td>
<td>864</td>
<td>4,642</td>
</tr>
<tr>
<td>Control obs</td>
<td>3,778</td>
<td>4,642</td>
</tr>
</tbody>
</table>
```

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>Treated</th>
<th>Control</th>
<th>Treated</th>
</tr>
</thead>
<tbody>
<tr>
<td>mmarried</td>
<td>.7514558</td>
<td>.4733796</td>
<td>.1868194</td>
<td>.2495802</td>
</tr>
<tr>
<td>mage</td>
<td>26.81048</td>
<td>25.16667</td>
<td>31.87141</td>
<td>28.10429</td>
</tr>
<tr>
<td>prenatal1</td>
<td>.8268925</td>
<td>.6898148</td>
<td>.1431792</td>
<td>.2142183</td>
</tr>
<tr>
<td>fbaby</td>
<td>.4531498</td>
<td>.3715278</td>
<td>.2478707</td>
<td>.2337654</td>
</tr>
</tbody>
</table>

While we rely on the standardized differences for conclusions about balance in the unmatched sample from this output, the baseline means and variances give us some idea of the scale of the differences.

---

Example 2: Multivalued treatments

In the multivalued-treatment case, `tebalance summarize` produces output grouped by treatment level. In the Cattaneo (2010) extract, the variable `msmoke` is an ordered categorical variable specifying the number of cigarettes smoked. We begin by tabulating `msmoke`. 
All the treatment groups have significantly smaller numbers of observations than the control group of not smoking. Still, each group has at least 200 observations. We continue by quietly fitting a candidate IPW model and reporting the baseline summaries.

The results for the control level of 0 daily are repeated for the treatment group. These results give a sense of the scale of imbalance in the raw data. Now we compute the balance statistics.
. tebalance summarize

Covariate balance summary

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Observations</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Raw</td>
</tr>
<tr>
<td>0 daily</td>
<td>3,778</td>
</tr>
<tr>
<td>1-5 daily</td>
<td>200</td>
</tr>
<tr>
<td>6-10 daily</td>
<td>337</td>
</tr>
<tr>
<td>11+ daily</td>
<td>327</td>
</tr>
<tr>
<td>Total</td>
<td>4,642</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>Raw</th>
<th>Weighted</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-5 daily</td>
<td>-0.634909</td>
<td>-0.0016208</td>
</tr>
<tr>
<td>mmarried</td>
<td>1.3340214</td>
<td>1.0014060</td>
</tr>
<tr>
<td>mage</td>
<td>-0.3857482</td>
<td>-0.0219656</td>
</tr>
<tr>
<td>prenatal1</td>
<td>0.9867038</td>
<td>0.9905584</td>
</tr>
<tr>
<td>fbaby</td>
<td>-0.312519</td>
<td>-0.0012611</td>
</tr>
<tr>
<td>6-10 daily</td>
<td>-0.6506304</td>
<td>-0.0108454</td>
</tr>
<tr>
<td>mmarried</td>
<td>1.3276964</td>
<td>1.0093310</td>
</tr>
<tr>
<td>mage</td>
<td>-0.3220222</td>
<td>-0.0836571</td>
</tr>
<tr>
<td>prenatal1</td>
<td>0.8493666</td>
<td>0.7984901</td>
</tr>
<tr>
<td>fbaby</td>
<td>-0.3465797</td>
<td>-0.0100232</td>
</tr>
<tr>
<td>11+ daily</td>
<td>-0.5147672</td>
<td>-0.0212969</td>
</tr>
<tr>
<td>mmarried</td>
<td>1.3416801</td>
<td>1.0181359</td>
</tr>
<tr>
<td>mage</td>
<td>-0.2239116</td>
<td>-0.0636951</td>
</tr>
<tr>
<td>prenatal1</td>
<td>0.8451058</td>
<td>0.8468934</td>
</tr>
<tr>
<td>fbaby</td>
<td>-0.3077549</td>
<td>-0.0380744</td>
</tr>
</tbody>
</table>

These results indicate that the IPW estimator probably did not fully balance the covariates (the variance ratios for `mage` at the daily levels of 6–10 cigarettes and 11-plus cigarettes are not close to 1). At this point, we would use a richer model and see whether it balanced the covariates.

Note that we cannot use `tebalance overid`, because it has not been implemented for multivalued treatments.

**Stored results**

tebalance summarize stores the following in `r()`:

Matrices

- `r(size)` number of observations in the raw and matched or weighted samples
- `r(table)` table of covariate statistics

**Reference**

Also see

[TE] tebalance — Check balance after teffects or stteffects estimation
[TE] teffects aipw — Augmented inverse-probability weighting
[TE] teffects ipw — Inverse-probability weighting
[TE] teffects ipwra — Inverse-probability-weighted regression adjustment
[TE] teffects nnmatch — Nearest-neighbor matching
[TE] teffects psmatch — Propensity-score matching
[TE] teffects overlap — Overlap plots
[TE] stteffects intro — Introduction to treatment effects for observational survival-time data
[TE] stteffects ipw — Survival-time inverse-probability weighting
[TE] stteffects ipwra — Survival-time inverse-probability-weighted regression adjustment
teffects — Treatment-effects estimation for observational data

Description

teffects estimates potential-outcome means (POMs), average treatment effects (ATEs), and average treatment effects on the treated (ATETs) using observational data. Regression-adjustment, inverse-probability-weighting, and matching estimators are provided, as are doubly robust methods that combine regression adjustment and inverse-probability weighting. teffects overlap plots the estimated densities of the probability of getting each treatment level.

The outcomes can be continuous, binary, count, fractional, or nonnegative. The treatment model can be binary, or it can be multinomial, allowing for multivalued treatments.

For a brief description and example of each estimator, see Remarks and examples in [TE] teffects intro.

Syntax

teffects subcommand ... [, options]

<table>
<thead>
<tr>
<th>subcommand</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>aipw</td>
<td>augmented inverse-probability weighting</td>
</tr>
<tr>
<td>ipw</td>
<td>inverse-probability weighting</td>
</tr>
<tr>
<td>ipwra</td>
<td>inverse-probability-weighted regression adjustment</td>
</tr>
<tr>
<td>nnmatch</td>
<td>nearest-neighbor matching</td>
</tr>
<tr>
<td>overlap</td>
<td>overlap plots</td>
</tr>
<tr>
<td>psmatch</td>
<td>propensity-score matching</td>
</tr>
<tr>
<td>ra</td>
<td>regression adjustment</td>
</tr>
</tbody>
</table>

Also see

[TE] teffects intro — Introduction to treatment effects for observational data
[TE] teffects intro advanced — Advanced introduction to treatment effects for observational data
[TE] teffects multivalued — Multivalued treatment effects
teffects intro — Introduction to treatment effects for observational data

Description

This entry provides a nontechnical introduction to treatment-effects estimators and the teffects command in Stata. Advanced users may want to instead read [TE] teffects intro advanced or skip to the individual commands’ entries.

The teffects command estimates average treatment effects (ATEs), average treatment effects among treated subjects (ATETs), and potential-outcome means (POMs) using observational data.

Treatment effects can be estimated using regression adjustment (RA), inverse-probability weights (IPW), and “doubly robust” methods, including inverse-probability-weighted regression adjustment (IPWRA) and augmented inverse-probability weights (AIPW), and via matching on the propensity score or nearest neighbors.

The outcome can be continuous, binary, count, fractional, or nonnegative. Treatments can be binary or multivalued.

Remarks and examples

This entry presents a nontechnical overview of treatment-effects estimators for those who are new to the subject of treatment-effects estimation or are at least new to Stata’s facilities for estimating treatment effects. More advanced users may want to instead read [TE] teffects intro advanced or skip to the individual commands’ entries.

Remarks are presented under the following headings:

- Introduction
- Defining treatment effects
- Estimating treatment effects
  - Regression adjustment
  - Inverse-probability weighting
  - Doubly robust combinations of RA and IPW
- Matching
- Caveats and assumptions
- A quick tour of the estimators
  - RA
  - IPW
  - IPWRA
  - AIPW
  - Nearest-neighbor matching
  - Propensity-score matching
- Video examples
Introduction

Suppose we have observed a sample of subjects, some of whom received a treatment and the rest of whom did not. As the name suggests, in most applications, the “subjects” are indeed people. A “treatment” could indeed be a medical treatment such as a new drug regimen or surgical procedure. In social science applications, a treatment could be participation in a job-training program or inclusion in a classroom or school in which a new pedagogical method is being used. However, not all applications use individuals as the subjects. For example, a policy analyst might be interested in examining the impact of an experimental program in which a national agency held a lottery to award only some local governments the resources needed to implement the program. Here the subjects are the local governments, and treatment refers to whether a local government received the resources needed to implement the program.

We would like to know if a treatment has an effect on an outcome \( Y \). The outcome could be the cholesterol level of a patient taking either an existing statin or a new experimental drug, or the outcome could be the wage offered to a person who either did or did not participate in a job-training program. In an ideal world, we would observe \( Y \) when a subject is treated (which we denote as \( Y_1 \)), and we would observe \( Y \) when the same subject is not treated (which we denote as \( Y_0 \)). We would be careful to make both observations under identical conditions so that the only difference is the presence or absence of the treatment. We could then average the difference between \( Y_1 \) and \( Y_0 \) across all the subjects in our dataset to obtain a measure of the average impact of the treatment.

Unfortunately, this ideal experiment is almost never available in observational data because it is not possible to observe a specific subject having received the treatment and having not received the treatment. When the outcome is the birthweight of a specific baby and the treatment is the mother smoking while pregnant, it is impossible to observe the baby’s birthweight under both treatments of the mother smoking and the mother not smoking.

A classic solution to this problem is to randomize the treatment. High costs or ethical issues rule out this solution in many observational datasets. For example, we could not ask a random selection of pregnant women to smoke.

The defining characteristic of observational data is that treatment status is not randomized. Moreover, that implies that the outcome and treatment are not necessarily independent. The goal of the estimators implemented by \texttt{teffects} is to utilize covariates to make treatment and outcome independent once we condition on those covariates.

The treatment-effect estimators implemented by \texttt{teffects} allow us to estimate the efficacy of treatments using observational data. The rest of this entry discusses these treatment-effect estimators at an introductory level. For a more technical introduction, see [TE] \texttt{teffects intro advanced}.

Defining treatment effects

We introduce treatment effects more formally by using the potential-outcomes framework, which is also known as the counterfactual framework. What is a potential outcome? Consider a subject that did not receive treatment so that we observe \( Y_0 \). What would \( Y_1 \) be for that same subject if it were exposed to treatment? We call \( Y_1 \) the potential outcome or counterfactual for that subject. For a subject that did receive treatment, we observe \( Y_1 \), so \( Y_0 \) would be the counterfactual outcome for that subject. We can view this as a missing-data problem, and treatment-effect methods can account for that problem.

Treatment-effect estimators allow us to estimate three parameters. The potential-outcome means (POMs) are the means of \( Y_1 \) and \( Y_0 \) in the population. The average treatment effect (ATE) is the mean of the difference \( (Y_1 - Y_0) \). Finally, the average treatment effect on the treated (ATET) is the mean of the difference \( (Y_1 - Y_0) \) among the subjects that actually receive the treatment.
To develop our intuition, suppose we have observed a sample of patients, some of whom received a medication to reduce their blood pressure. Figure 1 plots each of our patient’s systolic blood pressures as a function of weight. We use the color green to indicate patients who did not receive the drug and blue to indicate patients who did receive the drug.

A remarkable feature of our data is that the average blood pressure of patients not taking the drug is 160, and the average blood pressure of patients taking the drug is also 160. Can we therefore conclude that taking the drug has no impact on blood pressure? The answer is no.

Because this is observational data, we could not randomly assign who would receive the drug and who would not. As a result, treatment status could be related to covariates that also affect blood pressure. Heavier patients were more likely to be prescribed the medication, and blood pressure is correlated with weight. The difference in sample means does not estimate the true average treatment effect, because blood pressure depends on weight and weight is correlated with the treatment.

Suppose that we did in fact observe both potential outcomes for all patients. In figure 2, we continue to use solid dots for our observed data points, and we introduce hollow dots to represent the counterfactual outcomes. That is, the green hollow dots represent the blood pressures we would measure if only our treated patients had not taken the drug, and the blue hollow dots represent the blood pressures we would measure if only our untreated patients had taken the drug. The green and blue dashed lines represent the untreated and treated POMs, respectively. That is, the green line represents the mean of all the green dots, and the blue line represents the mean of all the blue dots.
If we did have the data represented by the hollow dots, then we could say that the average treatment effect is the difference between the mean of all the green dots and the mean of all the blue dots. In this ideal scenario, there are no missing data on the other potential outcome, and we have all the data we need to use the difference in means to estimate the ATE.

Looking at figure 2, we can see why a difference in means using only the solid dots does not estimate the ATE. Using only the solid green dots underestimates the average blood pressure for untreated individuals, and using only the solid blue dots overestimates the average blood pressure for treated individuals.

Estimating an ATE is essentially a missing-data problem. When covariates that affect the potential outcomes are related to treatment, we cannot use a difference in sample means, because the missing data are informative.

The treatment-effect estimators implemented in `teffects` allow for covariates like weight to be related to the potential outcomes and the treatment. Essentially, the estimators implemented by `teffects` utilize covariates to fill in the hollow circles or otherwise account for how the missing data depend on covariates that affect the potential outcomes.

**Estimating treatment effects**

We cannot estimate the ATE by simply taking the difference between the sample means for the treated and untreated subjects, because there are covariates that are related to the potential outcomes and the treatment. The estimators implemented by `teffects` require us to specify enough of these covariates so that after we condition on these covariates, any remaining influences on the treatment are not related to the potential outcomes. `teffects` implements several different estimators to accomplish this, including regression adjustment (RA), inverse-probability weighting (IPW), “doubly robust” methods that combine elements of RA and IPW, and matching methods. Here we introduce the methods by using intuition and simple examples.

See [TE] `teffects intro advanced` for a more technical introduction, and see the individual commands’ entries for estimator-specific details.
Regression adjustment

The RA method extends the idea of using sample means to estimate treatment effects by using a regression model to predict potential outcomes adjusted for covariates. In the examples here, we use linear regression, but the `teffects ra` command provides you with the flexibility to use logistic, probit, and heteroskedastic probit regression models for binary outcomes as well as Poisson regression for nonnegative outcomes; see [TE] teffects ra for more information.

`bweightex.dta` is a hypothetical dataset based on Cattaneo (2010) that we have created to illustrate treatment-effects estimators using graphs. The subjects in this dataset are women who were pregnant, some of whom smoked during the pregnancy. The outcome variable is the birthweight of the baby, and we want to know whether smoking during pregnancy affects the birthweight. The dataset also contains other demographic variables that we will use later.

Figure 3 illustrates the relationship between birthweight and smoking status as a function of the mother’s age:

![Figure 3](image)

We see that smokers tend to be older than nonsmokers and that birthweight depends on smoking. Therefore, the difference between the sample means of birthweights of babies born to smokers and nonsmokers will not estimate the true average treatment effect.

We also still have the same problem as in the previous section: we do not observe the counterfactual birthweights of babies. Suppose, however, that we did. In figure 4, we use solid points to represent observed birthweights and the colors green to represent nonsmokers and blue to represent smokers. The hollow points represent the counterfactual birthweights. The hollow blue points represent the birthweights of babies that we would observe if only our young nonsmoking mothers had instead smoked during their pregnancies. Similarly, the hollow green points represent the birthweights of babies that we would observe if only our older smoking mothers had instead not smoked during their pregnancies.
Figure 4 suggests a way to estimate the potential outcomes for each mother:

![Figure 4: Observed (solid) and Unobserved (hollow) Outcomes](image)

We could fit a linear regression of birthweight on mother’s age by using the observed birthweights for nonsmokers, and we could do likewise for smokers. The following graph includes these two regression lines:

![Figure 5: Regression Lines for the Observations](image)

Figure 5 illustrates the principle behind the RA method. We use the green regression line to predict each baby’s birthweight assuming the mother did not smoke, and we use the blue regression line to predict each baby’s birthweight assuming the mother did smoke. The treatment effect of smoking for a mother of a particular age is the vertical difference between the green and blue regression lines.

The three parameters we mentioned in the introduction are now easy to estimate. For each mother, we obtain two values, say, $bw_0$ and $bw_1$, representing our predictions of her baby’s birthweight assuming the mother did not or did smoke, respectively. The means of these variables represent the untreated and treated POMs. The ATE is the sample mean of the difference $(bw_1 - bw_0)$, and the ATET is the sample mean of that difference computed using only the mothers who in fact did smoke during pregnancy.
Adding the circles highlights the fact that the average age is higher for smokers than for nonsmokers. Even though the blue and green lines have different slopes, if the average age was the same for smokers and nonsmokers, a difference in the sample means of birthweights could still estimate the true ATE.

Figure 5 lets us address one more issue. Users who are versed in regression analysis may be inclined to estimate the effect of smoking using a regression model for birthweight as a function of smoking and the mother’s age. We clearly see in figure 5 that regression lines for smokers and nonsmokers have different slopes—the effect of age on birthweight is not the same for smokers and nonsmokers. In regression analysis, we would therefore include an interaction term between smoking and age. The RA method fits separate regression lines for smokers and nonsmokers, which also handles these differential effects of age on smoking.

**Inverse-probability weighting**

As we remarked in our discussion of the RA method, we cannot simply use the sample mean birthweights of babies born to smokers and nonsmokers to estimate the effect of smoking. If we did that, we would conflate the negative effect of smoking with the positive effect of age and the positive relationship between age and smoking. IPW is a treatment-effects estimator that uses weighted means rather than simple unweighted means to disentangle the effects of treatment and other confounders like age.

The concept underlying IPW can be gleaned from figure 2, where, as you will recall, the hollow points represent counterfactual outcomes. As we demonstrated in Defining treatment effects, we could estimate the average treatment effect if we knew the means of all the nonsmoking outcomes and the means of all the smoking outcomes. In the context of figure 4, we need the mean of all the green points, both solid and hollow, and the mean of all the blue.

If we could observe all of these points, then the ATE would be the difference between those two means. However, the outcomes illustrated by the hollow circles are unobserved. IPW estimators view the hollow circles as missing data and use weights to correct the estimates of the treated and untreated sample means for the missing data. If we calculate the mean nonsmoking birthweight using just the solid green points, that mean is biased downward because we are ignoring the hollow green points, which correspond to higher birthweights.

In IPW, we apply more weight to the solid green points corresponding to older mothers and less weight to those corresponding to younger mothers. Using this weighting scheme will pull up the estimated mean birthweight of babies born to nonsmoking mothers to estimate the true mean of all nonsmoking outcomes. The method for obtaining the mean smoking birthweight is virtually the same: we need to apply more weight to the younger smoking mothers than to the older smoking mothers to better approximate the true mean of all smoking outcomes.

Where do these weights for the weighted means come from? As the name implies, IPW uses the inverse (reciprocal) of the probability of being in the observed treatment group. These probabilities are obtained by modeling the observed treatment as a function of subject characteristics that determine treatment group. In our exposition of the RA method, we focused solely on the mother’s age and smoking status as determinants of each baby’s birthweight. To make the results comparable, we will use the same model in this example.
We first fit a logistic model of the mother’s smoking status, `mbsmoke`, as a function of the mother’s age (`mage`):

```
use http://www.stata-press.com/data/r15/bweightex
(Hypothetical birthweight data)
logistic mbsmoke mage
```

|              | Odds Ratio | Std. Err. | z     | P>|z|     | [95% Conf. Interval] |
|--------------|------------|-----------|-------|---------|---------------------|
| `mbsmoke`    | 1.631606   | .21316    | 3.75  | 0.000   | 1.263022 2.107754   |
| `_cons`      | 7.76e-06   | .0000243  | -3.76 | 0.000   | 1.69e-08 .0035718  |

Note: `_cons` estimates baseline odds.

Next, we compute the inverse-probability weights, which we will store in a variable called `ps`. In the IPW method, for subjects who did receive treatment, the weight is equal to the reciprocal of the predicted probability of treatment. For subjects who did not receive treatment, the weight is equal to the reciprocal of the predicted probability of not receiving treatment; the probability of not receiving treatment is just one minus the probability of receiving treatment:

```
predict ps
(option `pr` assumed; Pr(mbsmoke))
replace ps = 1/ps if mbsmoke==1
(30 real changes made)
replace ps = 1/(1-ps) if mbsmoke==0
(30 real changes made)
```

Figure 6 replicates figure 3 with one twist. Rather than making all the points the same size, we have made the size of the points proportional to the IPW variable `ps`. Notice that the largest blue points correspond to the youngest smoking mothers in our sample, so they will receive the most weight when we compute the weighted mean birthweight of babies born to smoking mothers, just as we explained we wanted to do. Similarly, the green points corresponding to older nonsmoking mothers are larger, representing larger weights.
There is a caveat to the IPW estimator. When we fit our logistic or probit model to obtain the predicted probabilities, it is possible that some of the predictions will be close to zero. Because the IPW is the reciprocal of that probability, the weight becomes arbitrarily large as the probability goes to zero. In those cases, the IPW can become unstable. We can improve the estimated IPW by developing a more accurate treatment model. For example, in our dataset, we have other variables such as marital status and the education level of the baby’s father that may also help predict whether the mother smoked during pregnancy. We excluded these variables for simplicity, but in a real analysis, we would want to use all relevant data.

This phenomenon of unstable IPWs is related to the concept of overlap, which means that every subject must have a strictly positive probability of obtaining treatment. We remarked that in our sample, we had few young mothers who smoked. As should be clear from figure 6, the overlap assumption is likely to be violated—young mothers do not appear to have a positive probability of being smokers. We would want to check this assumption before proceeding with an IPW analysis. See [TE] teffects overlap and [TE] teffects intro advanced for more information about overlap.

Another limitation of the IPW estimator is that we are using weighted means to estimate the POMs and ATE. Thus, unlike the RA estimator, we cannot obtain subject-level predictions of the treatment effects or potential outcomes, because we do not have the two regression lines that we can use to predict outcomes for each subject.

**Doubly robust combinations of RA and IPW**

You may have noticed a clear distinction between the RA and IPW estimators. In the case of RA, we built linear regression models to predict the outcomes (birthweights) of each subject but said nothing about how treatment (smoking) arises. In the case of IPW, we built a logistic regression model to predict treatment status but did not build a formal model of the outcome. Doubly robust estimators combine the outcome modeling strategy of RA and the treatment modeling strategy of IPW. These estimators have a remarkable property: although they require us to build two models, we only need to specify one of the two models correctly. If we misspecify the treatment model but correctly specify the outcome model, we still obtain correct estimates of the treatment effect. If we correctly specify the treatment model but misspecify the outcome model, we again will obtain correct estimates of the treatment effect.

Stata’s teffects command implements two doubly robust estimators, the augmented inverse-probability-weighted (AIPW) estimator and the inverse-probability-weighted regression-adjustment (IPWRA) estimator. These estimators combine elements of RA and IPW to be more robust to misspecification.

The AIPW estimator is an IPW estimator that includes an augmentation term that corrects the estimator when the treatment model is misspecified. When the treatment model is correctly specified, the augmentation term vanishes as the sample size becomes large. Like the IPW, the AIPW does not perform well when the predicted treatment probabilities are too close to zero or one.

The IPWRA estimator is an RA estimator that uses estimated inverse-probability weights to correct the estimator when the regression function is misspecified. When the regression function is correctly specified, the weights do not affect the consistency of the estimator.
Matching

Matching estimators are based on the idea of comparing the outcomes of subjects that are as similar as possible with the sole exception of their treatment status. In our birthweight and smoking example, we could select a mother who smokes and select a mother of the same age who does not smoke and compare the birthweights of their infants. The data of each mother serve as the potential outcome for the other mother.

For a single covariate such as age, identifying a pair of comparable mothers is not difficult. If we have a second covariate that is categorical, such as race, we might still be able to identify pairs of mothers who are the same age and of the same race assuming our dataset is large enough. However, once we consider covariates that are measured on continuous scales or allow for more than a few discrete ones, then finding identical matches is a challenge. The solution is to use what is called a similarity measure, which is a statistic that measures how “close” two observations are. `teffects` offers two methods to find comparable observations based on similarity measures: nearest-neighbor matching and propensity-score matching.

Nearest-neighbor matching (NNM) is accomplished by calculating the “distance” between pairs of observations with regard to a set of covariates and then “matching” each subject to comparable observations that are closest to it. For example, suppose we have a variable that records each subject’s annual income to the penny. Say one subject who received treatment had an income of $69,234.21. The likelihood that our dataset has an untreated subject who also earned $69,234.21 is nil. However, we can determine the difference between each untreated subject’s income and our treated subject’s income, then match our treated subject with the untreated subjects whose income differences are smallest. Measuring the distance between subjects when we have multiple covariates is no challenge. By default, `teffects` uses what is known as the Mahalanobis distance, which is really nothing more than the Pythagorean theorem adapted to handle the fact that covariates may be correlated and measured on different scales.

NNM does not use a formal model for either the outcome or the treatment status, but this flexibility comes at a price. When matching on more than one continuous covariate, the NNM estimator must be augmented with a bias-correction term. `teffects nnmatch` uses a linear function of the covariates specified in the `biasadj()` option to remove the large-sample bias.

Propensity-score matching (PSM) is an alternative to NNM. PSM matches on the estimated predicted probabilities of treatment, known as the propensity scores. PSM does not require bias correction, because it uses a model for the treatment. If the treatment model is reasonably well specified, PSM will perform at least as well as NNM; see [TE] `teffects intro advanced`.

Caveats and assumptions

To use the estimators implemented in `teffects`, we must make several assumptions about the process that generated our data. Different estimators and statistics may require slightly more or slightly less restrictive assumptions and may exhibit varying degrees of robustness to departures from these assumptions, but in general, all the estimators require some form of the following three assumptions.

The independent and identically distributed (i.i.d.) sampling assumption ensures that the outcome and treatment status of each individual are unrelated to the outcome and treatment status of all the other individuals in the population. Correlated data arising from hierarchical or longitudinal study designs do not meet this assumption.

The conditional-independence (CI) assumption means once we control for all observable variables, the potential outcomes are independent of treatment assignment. The easiest way to understand the CI assumption is to understand when it is violated. In our birthweight example, suppose mothers
who did not smoke were more health conscious and consumed better prenatal diets than those who
did smoke. Unless we explicitly controlled for health awareness or diet, our model would violate
the CI assumption: the mother’s decision to smoke or not smoke would not be independent of the
baby’s birthweight. If we did not control for health awareness, we would overstate the negative
impact of smoking on birthweight. Babies born to mothers who smoke weigh less than babies born to
nonsmoking mothers not just because of the effects of cigarettes but also because of poorer prenatal
diets.

In a study examining the effect of a job-training program, the CI assumption requires that there
not be any unobserved factors such as ambition or work ethic that influence both whether a person
enrolls in the program and the wage received upon completion. To use the methods implemented by
the teffects estimators, we must have variables in our dataset that allow us to control for those
types of factors.

We mentioned the third assumption, overlap, in our discussions of IPW. More formally, the
overlap assumption states that each individual have a positive probability of receiving treatment. In
our birthweight example, we noted that there were no observations on young smokers and older
nonsmokers. Perhaps we just have an unlucky sample, but to accurately assess the impact of treatment
using these methods, we must have overlap to accurately estimate the counterfactual birthweights.
In the context of matching estimators, overlap essentially means that we can actually match treated
subjects with similar nontreated subjects.

**A quick tour of the estimators**

The teffects command implements six estimators of treatment effects. We introduce each one
by showing the basic syntax one would use to apply them to our birthweight example. See each
command’s entry for more information.

**RA**

teffects ra implements the RA estimator. We estimate the effect of a mother’s smoking behavior
(mbsmoke) on the birthweight of her child (bweight), controlling for marital status (mmarried), the
mother’s age (mage), whether the mother had a prenatal doctor’s visit in the baby’s first trimester
(prenatal1), and whether this baby is the mother’s first child (fbaby). We use linear regression
(the default) to model bweight:

. use http://www.stata-press.com/data/r15/cattaneo2
. teffects ra (bweight mmarried mage prenatal1 fbaby) (mbsmoke)

**IPW**

teffects ipw implements the IPW estimator. Here we estimate the effect of smoking by using a
probit model to predict the mother’s smoking behavior as a function of marital status, the mother’s
age, and indicators for first-trimester doctor’s visits and firstborn status:

. teffects ipw (bweight) (mbsmoke mmarried mage prenatal1 fbaby, probit)
teffects intro — Introduction to treatment effects for observational data

**IPWRA**

`teffects ipwra` implements the IPWRA estimator. We model the outcome, birthweight, as a linear function of marital status, the mother’s age, and indicators for first-trimester doctor’s visits and firstborn status. We use a logistic model (the default) to predict the mother’s smoking behavior, using the same covariates as explanatory variables:

```
. teffects ipwra (bweight mmarried mage prenatal1 fbaby) ///
   (mbsmoke mmarried mage prenatal1 fbaby)
```

**AIPW**

`teffects aipw` implements the AIPW estimator. Here we use the same outcome- and treatment-model specifications as we did with the IPWRA estimator:

```
. teffects aipw (bweight mmarried mage prenatal1 fbaby) ///
   (mbsmoke mmarried mage prenatal1 fbaby)
```

**Nearest-neighbor matching**

`teffects nnmatch` implements the NNM estimator. In this example, we match treated and untreated subjects based on marital status, the mother’s age, the father’s age, and indicators for first-trimester doctor’s visits and firstborn status. We use the Mahalanobis distance based on the mother’s and father’s ages to find matches. We use exact matching on the other three variables to enforce the requirement that treated subjects are matched with untreated subjects who have the same marital status and indicators for first-trimester doctor’s visits and firstborn statuses. Because we are matching on two continuous covariates, we request that `teffects nnmatch` include a bias-correction term based on those two covariates:

```
. teffects nnmatch (bweight mage fage) (mbsmoke), ///
   ematch(prenatal1 mmarried fbaby) biasadj(mage fage)
```

**Propensity-score matching**

`teffects psmatch` implements the PSM estimator. Here we model the propensity score using a probit model, incorporating marital status, the mother’s age, and indicators for first-trimester doctor’s visits and firstborn status as covariates:

```
. teffects psmatch (bweight) (mbsmoke mmarried mage prenatal1 fbaby, probit)
```

**Video examples**

- Introduction to treatment effects in Stata, part 1
- Introduction to treatment effects in Stata, part 2
References


Also see

[TE] **teffects** — Treatment-effects estimation for observational data

[TE] **teffects intro advanced** — Advanced introduction to treatment effects for observational data

[TE] **teffects multivalued** — Multivalued treatment effects
Description

This entry provides a technical overview of treatment-effects estimators and their implementation in Stata. Those who are new to treatment-effects estimation may want to instead see [TE] t effects intro.

The t effects command estimates average treatment effects (ATEs), average treatment effects among treated subjects (ATETs), and potential-outcome means (POMs) using observational data.

Treatment effects can be estimated using regression adjustment (RA), inverse-probability weights (IPW), and “doubly robust” methods, including inverse-probability-weighted regression adjustment (IPWRA) and augmented inverse-probability weights (AIPW), and via matching on the propensity score or nearest neighbors.

The outcome can be continuous, binary, count, fractional, or nonnegative. Treatments can be binary or multivalued.

Remarks and examples

This entry presents a technical overview of treatment-effects estimators and their implementation in Stata. Users who are new to treatment-effects estimators for observational data should instead read [TE] t effects intro.

Remarks are presented under the following headings:

- Introduction
- Defining treatment effects
- The potential-outcome model
- Assumptions needed for estimation
  - The CI assumption
  - The overlap assumption
  - The i.i.d. assumption
- Comparing the ATE and ATET
- Overview of treatment-effect estimators
  - RA estimators
  - IPW estimators
  - AIPW estimators
  - IPWRA estimators
  - Nearest-neighbor matching estimators
  - Propensity-score matching estimators
  - Choosing among estimators
  - Model choice
Introduction

The `teffects` commands estimate treatment effects from observed data. A treatment effect is the change in an outcome caused by a subject, often an individual, getting one treatment instead of another. We cannot estimate individual-level treatment effects, because we only observe each individual getting one or another treatment.

Potential-outcome models provide a solution to this missing-data problem and allow us to estimate the distribution of individual-level treatment effects. A potential-outcome model specifies the potential outcomes that each individual would obtain under each treatment level, the treatment assignment process, and the dependence of the potential outcomes on the treatment assignment process.

When the potential outcomes do not depend on the treatment levels, after conditioning on covariates, regression estimators, inverse-probability-weighted estimators, and matching estimators are commonly used.

What we call the potential-outcome model is also known as the Rubin causal model and the counterfactual model. See Rubin (1974); Holland (1986); Robins (1986); Heckman (1997); Heckman and Navarro-Lozano (2004); Imbens (2004); Cameron and Trivedi (2005, chap. 2.7); Imbens and Wooldridge (2009); and Wooldridge (2010, chap. 21) for more detailed discussions.

For a discussion related to health econometrics, see Deb, Norton, and Manning (2017).

Defining treatment effects

Three parameters are often used to measure treatment effects: the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs). In this section, we define each of these terms and introduce the notation and parameters used in the rest of our discussion.

In the binary-treatment case, the two potential outcomes for each individual are $y_{0i}$ and $y_{1i}$; $y_{0i}$ is the outcome that would be obtained if $i$ does not get the treatment, and $y_{1i}$ is the outcome that would be obtained if $i$ gets the treatment. $y_{0i}$ and $y_{1i}$ are realizations of the random variables $y_0$ and $y_1$. Throughout this entry, $i$ subscripts denote realizations of the corresponding un subscripted random variables. We do not discuss multivalued treatments here, because doing so only increases the number of parameters and notation required and detracts from the essential points; see [TE] teffects multivalued for information about multivalued treatments.

The parameters of interest summarize the distribution of the unobservable individual-level treatment effect $y_1 - y_0$. In defining the parameters, $t$ denotes a random treatment, $t_i$ denotes the treatment received by individual $i$, $t = 1$ is the treatment level, and $t = 0$ is the control level. Given this notation, we can now define our parameters of interest.

**ATE** The ATE is the average effect of the treatment in the population:

$$ATE = E(y_1 - y_0)$$

**POM** The POM for treatment level $t$ is the average potential outcome for that treatment level:

$$POM_t = E(y_t)$$

**ATET** The ATET is the average treatment effect among those that receive the treatment:

$$ATET = E(y_1 - y_0 | t = 1)$$

For an illustration of these concepts, see *Defining treatment effects* in [TE] teffects intro.
The potential-outcome model

Next we specify a potential-outcome model that serves as a touchstone for the rest of our discussion. The model described here generates data in which $y_i$ is the observed outcome variable, $t_i$ is the treatment variable, $x_i$ is a vector of covariates that affect the outcome, and $w_i$ is a vector of covariates that affect the treatment assignment. $x_i$ and $w_i$ may have elements in common.

This potential-outcome model specifies that the observed outcome variable $y$ is $y_0$ when $t = 0$ and that $y$ is $y_1$ when $t = 1$. Algebraically, we say that

$$y = (1 - t)y_0 + ty_1$$

The functional forms for $y_0$ and $y_1$ are

$$y_0 = x'\beta_0 + \epsilon_0$$
$$y_1 = x'\beta_1 + \epsilon_1$$

where $\beta_0$ and $\beta_1$ are coefficients to be estimated, and $\epsilon_0$ and $\epsilon_1$ are error terms that are not related to $x$ or $w$. This potential-outcome model separates each potential outcome into a predictable component, $x\beta_t$, and an unobservable error term, $\epsilon_t$.

The treatment assignment process is

$$t = \begin{cases} 
1 & \text{if } w'\gamma + \eta > 0 \\
0 & \text{otherwise}
\end{cases}$$

where $\gamma$ is a coefficient vector, and $\eta$ is an unobservable error term that is not related to either $x$ or $w$. The treatment assignment process is also separated into a predictable component, $w'\gamma$, and an unobservable error term, $\eta$.

We emphasize six points about this model:

1. The observed data from this model contain $y_i$, $t_i$, $w_i$, and $x_i$. The data do not reveal both $y_{0i}$ and $y_{1i}$ for any given $i$.
2. The model for $t$ determines how the data on $y_0$ and $y_1$ are missing.
3. The model separates the potential outcomes and treatment assignment into observable and unobservable components.
4. Whether $\eta$ is independent of the vector $(\epsilon_0, \epsilon_1)$ is essential in specifying the set of available estimators.
5. The coefficient vectors $\beta_0$, $\beta_1$, and $\gamma$ are auxiliary parameters. We use estimates of these coefficient vectors to estimate the ATE, the POMs, and the ATET.
6. For notational simplicity, we represented $y_0$ and $y_1$ as linear functions. In practice, we can use other functional forms.

In specifying this potential-outcome model, we explicitly showed the functional forms for the potential outcomes and the treatment assignment process. To ease subsequent discussions, we refer to the set of functional forms for the potential outcomes as the “outcome model”, and we refer to the treatment assignment process as the “treatment model”.
Assumptions needed for estimation

As with any type of estimator, we must make some assumptions to use treatment-effects estimators. The particular assumptions we need for each estimator implemented by \texttt{teffects} and for each effect parameter vary, but some version of each of the following is required.

**CI** The conditional-independence CI assumption restricts the dependence between the treatment model and the potential outcomes.

**Overlap** The overlap assumption ensures that each individual could receive any treatment level.

**i.i.d.** The independent and identically distributed (i.i.d.) sampling assumption ensures that the potential outcomes and the treatment status of each individual are unrelated to the potential outcomes and treatment statuses of all other individuals in the population.

We now discuss each assumption in detail.

The CI assumption

After conditioning on covariates, when no unobservable variable affects both treatment assignment and the potential outcomes, the potential outcomes are conditionally independent of the treatment. In epidemiological jargon, there are no unmeasured confounders. In econometric jargon, we have selection on observables. If we observe enough covariates, the potential outcomes may indeed be conditionally independent of the treatment.

Intuitively, the CI assumption says that only the covariates \( x \) affect both the treatment and the potential outcomes. Any other factors that affect the treatment must be independent of the potential outcomes, and any other factors that affect the potential outcomes must be independent of the treatment. Formally, the CI assumption states that, conditional on covariates \( x \), the treatment \( t \) is independent of the vector of potential outcomes \((y_0, y_1)'\).

The CI assumption allows us to estimate the effects by regression-adjustment (RA) methods, inverse-probability-weighting (IPW) methods, methods that combine RA and IPW concepts, and matching methods. The data only reveal information about \( E(y_0|x, w, t=0) \) and \( E(y_1|x, w, t=1) \), but we are interested in an average of \( E(y_0|x, w) \) and \( E(y_1|x, w) \), where \( x \) represents the outcome covariates and \( w \) the treatment-assignment covariates. The CI assumption allows us to estimate \( E(y_0|x, w) \) and \( E(y_1|x, w) \) directly from the observations for which \( E(y_0|x, w, t=0) \) and \( E(y_1|x, w, t=1) \), respectively.

For our potential-outcome model presented in (1) through (3), the CI assumption can be viewed as a set of restrictions on the covariance matrix of the error terms. Suppose that the vector of unobservables \((\epsilon_0, \epsilon_1, \eta)\) is normally distributed

\[
\begin{pmatrix}
\epsilon_0 \\
\epsilon_1 \\
\eta
\end{pmatrix}
\sim N \left( \begin{pmatrix}
0 \\
0 \\
0
\end{pmatrix}, \begin{pmatrix}
\sigma_0^2 & \rho_{01}\sigma_0\sigma_1 & \rho_{0\eta}\sigma_0 \rho_{01} \\
\rho_{01}\sigma_0\sigma_1 & \sigma_1^2 & \rho_{1\eta}\sigma_1 \rho_{01} \\
\rho_{0\eta}\sigma_0 & \rho_{1\eta}\sigma_1 & 1
\end{pmatrix} \right)
\]

where \( \sigma_0 \) is the standard deviation of \( \epsilon_0 \), \( \rho_{01} \) is the correlation between \( \epsilon_0 \) and \( \epsilon_1 \), \( \sigma_1 \) is the standard deviation of \( \epsilon_1 \), \( \rho_{\eta 0} \) is the correlation between \( \epsilon_\eta \) and \( \epsilon_0 \), and \( \rho_{\eta 1} \) is the correlation between \( \epsilon_\eta \) and \( \epsilon_1 \). As is standard in the normally distributed latent-variable specification of a binary-dependent variable, we normalize the variance of \( \epsilon_\eta \) to 1.
CI specifies that \( \rho_{\eta 0} = \rho_{\eta 1} = 0 \) so that we can write (4) as

\[
\begin{pmatrix}
\epsilon_0 \\
\epsilon_1 \\
\eta
\end{pmatrix}
\sim N
\begin{pmatrix}
0 \\
0 \\
0
\end{pmatrix},
\begin{pmatrix}
\sigma_0^2 & \rho_{01}\sigma_0\sigma_1 & 0 \\
\rho_{01}\sigma_0\sigma_1 & \sigma_1^2 & 0 \\
0 & 0 & 1
\end{pmatrix}
\]

Writing the covariance matrix this way makes clear what we mean by conditional independence: unobserved shocks that affect whether a subject is treated do not have any effect on the potential outcomes, and unobserved shocks that affect a potential outcome do not affect treatment.

The command `teffects` implements estimators that require the CI assumption. See `TE etregress` and `TE etpoisson` for commands that handle two cases in which the CI assumption is replaced by precise specifications of the joint dependence among the unobservables. Brown and Mergoupis (2011) discuss the `itreatreg` command that extends `TE etregress`.

The CI assumption is also known as unconfoundedness and selection-on-observables in the literature. See Rosenbaum and Rubin (1983); Heckman (1997); Heckman and Navarro-Lozano (2004); Cameron and Trivedi (2005, sec. 25.2.1); Tsiatis (2006, sec. 13.3); Angrist and Pischke (2009, chap. 3); Imbens and Wooldridge (2009); and Wooldridge (2010, sec. 21.3). Some discussions with Stata commands can be found in Becker and Caliendo (2007), Nichols (2007), and Daniel, De Stavola, and Cousens (2011).

### Technical note

In fact, full CI is stronger than what we need to estimate the ATE, the ATET, or the POMs. For the estimators implemented in `teffects`, we only need a conditional mean independence (CMI) assumption. Intuitively, the CMI assumption says that after accounting for the covariates \( x_i \), the treatment does not affect the conditional mean of each potential outcome. Formally, the CMI requires that \( E(y_0|x, t) = E(y_0|x) \) and that \( E(y_1|x, t) = E(y_1|x) \). The CMI assumption allows the conditional variance to depend on the treatment, while the CI assumption does not.

The CI assumption implies the CMI assumption, but not vice versa.

See Wooldridge (2010, sec. 21.2 and 21.3) for an excellent introduction to this topic, and see Cattaneo, Drukker, and Holland (2013) for some discussion of the multiple treatment case.

### The overlap assumption

The overlap assumption requires that each individual have a positive probability of receiving each treatment level. Formally, the overlap assumption requires that for each possible \( x \) in the population and each treatment level \( \tilde{t} \), \( 0 < \Pr(t = \tilde{t}|x) < 1 \). Rosenbaum and Rubin (1983) call the combination of the CI and overlap assumptions strong ignorability; see also Abadie and Imbens (2006, 237–238) and Imbens and Wooldridge (2009).

### The i.i.d. assumption

The third of the three assumptions listed above is the i.i.d. assumption; it is the standard assumption of having an i.i.d. sample from the population. In potential-outcome models, i.i.d. sampling implies that the potential outcomes and treatment status of each individual are unrelated to the potential outcomes and treatment statuses of all the other individuals in the population. I.i.d. sampling rules out interactions among the individuals. For instance, models of vaccinations in epidemiology and general
equilibrium effects in economics violate the independence assumption. This third assumption is a part of what is known as the stable unit treatment value assumption (SUTVA); see Wooldridge (2010, 905) and Imbens and Wooldridge (2009).

Comparing the ATE and ATET

When comparing the ATE and the ATET, two points should be mentioned.

First, the assumptions required to estimate the ATET are less restrictive than the assumptions required to estimate the ATE. Estimating the ATET requires a weaker form of the CI assumption and a weaker version of the overlap assumption.

To estimate the ATE under CI, we require that the unobservables in the treatment model be conditionally independent of the unobservables in both potential outcomes. In contrast, we can estimate the ATET under CI when the unobservables in the treatment model are conditionally independent of just the control-level potential outcome; see Wooldridge (2010, 906–912).

Although the ATE version of overlap requires that all covariate patterns have a positive probability of being allocated to each treatment state, we can estimate the ATET when only the covariate patterns for which someone is treated have a positive probability of being allocated to each treatment state. This weaker overlap assumption can be important in some studies. For example, Heckman (1997) discusses how the ATET makes sense in job-training programs for which many types of individuals have zero chance of signing up. See also Wooldridge (2010, 911–913).

Second, the ATET reduces to the ATE when the mean of the covariates among the treated is the same as the mean of the covariates in the population and when the average contribution from the unobservables for the participants is zero.

Overview of treatment-effect estimators

We can classify the estimators implemented by teffects into five categories: 1) estimators based on a model for the outcome variable; 2) estimators based on a model for treatment assignment; 3) estimators based on models for both the outcome variable and the treatment assignment; 4) estimators that match on covariates; and 5) estimators that match on predicted probabilities of treatment. Within each category of estimator, there is a variety of choices about the functional forms for the models.

Because there are several categories of estimators, the user must decide whether to model the outcomes, the probability of treatment, or both. Under correct model specification, using an outcome model and a model for the probability of treatment will produce more efficient estimates. Surprisingly, some of the estimators that use both models only require that one of the two be correctly specified to consistently estimate the effects of interest, a property known as the double-robust property.

With the exception of using a matching estimator with a single continuous covariate, some choice of functional forms is required. There are two aspects one must consider when choosing the functional form for the outcome or treatment assignment. First, one must select the functional form for the conditional mean or conditional probability; depending on the variable being modeled, a linear, a binary choice, or an exponential model may be appropriate. Second, one must determine the appropriate polynomials of the covariates to include in the model. teffects offers a wide variety of options to specify different functional form choices for the conditional mean and conditional probability models. The factor variable notation in Stata allows us to easily specify the desired polynomial in the covariates.

We now provide some intuition behind each type of estimator.
RA estimators

RA estimators use means of predicted outcomes for each treatment level to estimate each POM. ATEs and ATETs are differences in estimated POMs.

The CI assumption implies that we can estimate $E(y_0|x)$ and $E(y_1|x)$ directly from the observations for which $t = 0$ and $t = 1$, respectively. Regression adjustment fits separate regressions for each treatment level and uses averages of the predicted outcomes over all the data to estimate the POMs. The estimated ATEs are differences in the estimated POMs. The estimated ATETs are averages of the predicted outcomes over the treated observations.

RA is a venerable estimator. See Lane and Nelder (1982); Cameron and Trivedi (2005, chap. 25); Wooldridge (2010, chap. 21); and Vittinghoff et al. (2012, chap. 9). The usefulness of RA has been periodically questioned in the literature because it relies on specifying functional forms for the conditional means and because it requires having sufficient observations of each covariate pattern in each treatment level; see Rubin (1973) for an early salvo. Our experience is that RA is an exceptionally useful base-case estimator. We describe its relative advantages and disadvantages in the course of covering other estimators.

IPW estimators

IPW estimators use weighted averages of the observed outcome variable to estimate means of the potential outcomes. The weights account for the missing data inherent in the potential-outcome framework. Each weight is the inverse of the estimated probability that an individual receives a treatment level. Outcomes of individuals who receive a likely treatment get a weight close to one. Outcomes of individuals who receive an unlikely treatment get a weight larger than one, potentially much larger.

IPW estimators model the probability of treatment without any assumptions about the functional form for the outcome model. In contrast, RA estimators model the outcome without any assumptions about the functional form for the probability of treatment model.

IPW estimators become extremely unstable as the overlap assumption gets close to being violated. When the overlap assumption is nearly violated, some of the inverse-probability weights become very large, IPW estimators produce erratic estimates, and the large-sample distribution provides a poor approximation to the finite-sample distribution of IPW estimators. This instability occurs even though the functional form for the treatment model is correctly specified.

In contrast, when the overlap assumption is nearly violated, there are very few observations in a treatment level for some covariate patterns, so RA estimators use the model to predict in regions in which there are very little data. If the model is well specified and there are “enough” observations, an RA estimator will not become unstable as quickly as an IPW estimator, and the large-sample distribution of the RA estimator still provides a good approximation to the finite-sample distribution. However, in real situations in which “all models are approximate”, relying on a correctly specified outcome model with little data is extremely risky.

IPW estimators are a general approach to missing-data problems that obey some CI assumptions. While IPW is an old idea in statistics that dates back to Horvitz and Thompson (1952), biostatisticians and econometricians have been actively working on extending it to handle modern problems and estimation methods. See Robins and Rotnitzky (1995); Robins, Rotnitzky, and Zhao (1994, 1995); and Wooldridge (2002, 2007). IPW has been used extensively in the modern treatment-effect estimation literature. See Imbens (2000); Hirano, Imbens, and Ridder (2003); Tan (2010); Wooldridge (2010, chap. 19); van der Laan and Robins (2003); and Tsiatis (2006, chap. 6).

To see the intuition behind IPW, consider a study with observed outcome variable $y$, treatment variable $t \in \{0, 1\}$, and potential outcomes $y_0$ and $y_1$. As part of this process, we need to estimate
the POM for treatment \( t = 1 \), \( E(y_1) \). Using the observed data, \( y_{i1} t_i \) is \( y_{i1} \) when \( t = 1 \), but \( y_{i1} \) is unobserved when \( t = 0 \). An IPW estimator for \( E(y_1) \) is \( 1/N \sum_{i=1}^{N} y_{i1} t_i / p(x_i) \), where \( p(x_i) \) is the probability that \( t_i = 1 \) and is a function of the covariates \( x_i \). If \( y_{i1} \) were always observed, the weights would all equal 1. This IPW estimator places a larger weight on those observations for which \( y_{i1} \) is observed even though its observation was not likely.

### AIPW estimators

Instead of modeling either the outcome, like RA, or the treatment probability, like IPW, augmented inverse-probability-weighted (AIPW) estimators model both the outcome and the treatment probability. A surprising fact is that only one of the two models must be correctly specified to consistently estimate the treatment effects, a property of the AIPW estimators known as being “doubly robust”. Given that two models instead of one are used, it is less surprising that the AIPW estimators can be more efficient than either the RA or the IPW estimators, though deriving this result is rather technical and relies on the theory of semiparametric estimators.

Intuitively, the AIPW estimator is an IPW that includes an augmentation term that corrects the estimator when the treatment model is misspecified. When the treatment is correctly specified, the augmentation term vanishes as the sample size becomes large. Like the IPW, the AIPW does not perform well when the predicted treatment probabilities are too close to zero or one.

AIPW estimators emerge naturally from a technique of producing more efficient estimators from estimators that have a few main parameters of interest and some auxiliary, or nuisance, parameters used in estimating the few main parameters. This method constructs efficient estimating equations for the main parameters that are orthogonal to the auxiliary parameters. The estimators produced by this method are known as efficient-influence function (EIF) estimators.

To gain some intuition, consider finding an EIF estimator from an IPW estimator for two POMs. Note that we only care about the two POM parameters and not about the many auxiliary parameters used to estimate the treatment probabilities. EIF estimators project the equations that yield the POM parameters onto the equations that yield the auxiliary treatment-model parameters and then use the residuals from this projection to estimate the POM parameters.

We refer to these estimators as “AIPW estimators” instead of “EIF estimators” because the former is commonly used in the biostatistical literature for some of the binary-treatment estimators and because the term “augmented inverse-probability-weighted” tells more about how these estimators relate to the other implemented estimators; see Tsiatis (2006) and Tan (2010). The estimators implemented in `teffects aipw` with the `wnls` option are based on those of Rubin and van der Laan (2008), which did well in simulations reported by Tan (2010), and denoted as \( \hat{\alpha}_{RV}(\pi) \) in Tan (2010, 663).

When either the outcome model or the treatment model is well specified, the AIPW estimators implemented in `teffects aipw` are more robust than either the RA or the IPW estimators because the AIPW estimators are doubly robust but the RA and IPW estimators are not. When both the outcome and the treatment model are misspecified, which estimator is more robust is a matter of debate in the literature; see Kang and Schafer (2007) and Robins et al. (2007) for some debate, and see Tan (2010) for a more recent discussion.

To the best of our knowledge, there is no general solution to the question of which estimator performs best when both the outcome and the treatment models are misspecified. We suspect that the answer depends on the true models, the implemented specifications, and the polynomials in the covariates used. To help users through this process, the estimators implemented in `teffects` offer many functional forms to approximate either the outcome process or the treatment process. In addition, Stata’s factor-variable notation makes it easy to include polynomials in the covariates. Both of these approximation methods rely on having enough data. `teffects` also makes it easy to compare the results produced by different estimators.
The literature on these methods is vast and growing. For double-robust results and explanations, see Robins and Rotnitzky (1995); Robins, Rotnitzky, and Zhao (1995); van der Laan and Robins (2003, chap. 6); Bang and Robins (2005); Tsiatis (2006, chap. 13); Wooldridge (2007; 2010, sec. 21.3.4); and Tan (2010).

**IPWRA estimators**

Like AIPW estimators, inverse-probability-weighted regression-adjustment (IPWRA) estimators combine models for the outcome and treatment status; also like AIPW estimators, IPWRA estimators are doubly robust. IPWRA estimators emerge naturally from a robust approach to missing-data methods. IPWRA estimators use the inverse of the estimated treatment-probability weights to estimate missing-data-corrected regression coefficients that are subsequently used to compute the POMs.

As far as we know, there is no literature that compares the relative efficiency of AIPW estimators, which emerge from a general approach to creating efficient estimators, and the IPWRA estimators, which emerge from a robust-correction approach to missing-data analysis.

The IPWRA estimators are also know as “Wooldridge’s double-robust” estimators because they were derived in Wooldridge (2007) and discussed at length in Wooldridge (2010, section 21.3.4).

**Nearest-neighbor matching estimators**

Matching estimators use an average of the outcomes of the nearest individuals to impute the missing potential outcome for each sampled individual. The difference between the observed outcome and the imputed potential outcome is an estimate of the individual-level treatment effect. These estimated individual-level treatment effects are averaged to estimate the ATE or the ATET.

tffects nnmatch determines the “nearest” by using a weighted function of the covariates for each observation. This type of matching is known as nearest-neighbor matching (NNM). teffects psmatch determines the “nearest” by using the estimated treatment probabilities, which are known as the propensity scores. This second type of matching is known as propensity-score matching (PSM).

NNM is nonparametric in that no explicit functional form for either the outcome model or the treatment model is specified. This flexibility comes at a price; the estimator needs more data to get to the true value than an estimator that imposes a functional form. More formally, the NNM estimator converges to the true value at a rate slower than the parametric rate, which is the square root of the sample size, when matching on more than one continuous covariate. teffects nnmatch uses bias correction to fix this problem. PSM provides an alternative to bias correction because it matches on a single continuous covariate, the estimated treatment probabilities.

Abadie and Imbens (2006, 2011) derived the rate of convergence of the NNM estimator and the bias-corrected NNM estimator and the large-sample distributions of the NNM and the bias-corrected NNM estimators. These articles provided the formal results that built on methods suggested in Rubin (1973, 1977).

tffects nnmatch is based on the results in Abadie and Imbens (2006, 2011) and a previous implementation in Abadie et al. (2004).
Propensity-score matching estimators

Instead of performing bias correction to handle the case of more than one continuous covariate, a common solution is to combine all the covariate information into estimated treatment probabilities, known as propensity scores, and use this single continuous covariate as the matching variable.

The term “propensity score” is widely used, but we continue to refer to it as the “treatment probability” to be consistent with the other estimators. We call the estimator that matches on the estimated treatment probabilities the “propensity-score matching (PSM) estimator” because the latter term is ubiquitous.

In effect, the PSM estimator parameterizes the bias-correction term in the treatment probability model. One advantage of matching on the estimated treatment probabilities over the bias-correction method is that one can explore the fit of different treatment probability models using standard methods before performing the nonparametric matching. For example, one can select the treatment model by minimizing an information criterion under i.i.d. sampling. We know of no counterpart for selecting the proper order of the bias-correction term for the NNM estimator.

Matching on estimated treatment probability models has been very popular since Rosenbaum and Rubin (1983) showed that if adjusting for covariates \(x_i\) is sufficient to estimate the effects, then one can use the probability of treatment to perform the adjustment. Abadie and Imbens (2012) derived a method to estimate the standard errors of the estimator that matches on estimated treatment probabilities, and this method is implemented in `teffects psmatch`.

Choosing among estimators

There is no definitive way to select one of the estimators implemented in `teffects` over the others. We offer three observations about the tradeoffs among the estimators.

First, if the outcome model is correctly specified, the RA estimator will break down more slowly than the IPW, AIPW, IPWRA, or PSM estimators as the overlap assumption begins to fail. This result depends critically on the ability of the RA estimator to predict into regions in which there are little data.

Second, if the overlap assumption holds, the AIPW and IPWRA estimators have the double-robust property for some functional form combinations. The double-robust property says that if either the outcome model or the treatment model is correctly specified, we can consistently estimate the effects. The properties of double-robust estimators when both models are misspecified are not known, although there is some discussion in the literature about the properties of the AIPW estimators; see Kang and Schafer (2007), Robins et al. (2007), and Tan (2010).

Third, all the estimators require the same assumptions, so if each is correctly specified, they should all produce similar results. Of course, just because they produce similar results does not mean that they are correctly specified; it is possible that they are just behaving similarly in response to some underlying problem.

Model choice

`teffects` offers a broad selection of functional form combinations so that you can choose a combination that fits your data. Picking a functional form that respects the values of the observed outcomes is usually best. Select `linear` for continuous outcomes over the real line; `logit`, `probit`, or `hetprobit` for binary outcomes; and `poisson` for counts or nonnegative outcomes.

For binary treatments, you can select among `logit`, `probit`, or `hetprobit` models. For multivalued treatments, only the multinomial logit is available to model the treatment probabilities.
Selecting a functional form of a given set of covariates is a famously difficult problem in statistics. In the treatment-effects context, Cattaneo, Drukker, and Holland (2013) found that model selection by minimizing an information criterion worked well. Cattaneo, Drukker, and Holland (2013) discuss a method and a community-contributed command to facilitate the process.

Acknowledgments

We thank Matias Cattaneo, Guido Imbens, Barbara Sianesi, and Jeffrey Wooldridge for helpful conversations and advice about the literature and methods in this area.

We thank Matias Cattaneo for his collaboration on Cattaneo, Drukker, and Holland (2013) and for many conversations about Cattaneo (2010), a part of which is implemented in teffects ipw and teffects aipw.

We thank Ashley Holland for her collaboration on Cattaneo, Drukker, and Holland (2013).

We thank Alberto Abadie, Jane Herr, and Guido Imbens for collaboration on Abadie et al. (2004), which is implemented in teffects nmnm.

We thank Matias Cattaneo for sharing an extract of the dataset that he used in Cattaneo (2010) and Barbara Sianesi for sharing an extract of the dataset that she used in Blundell, Dearden, and Sianesi (2005).

We thank Alberto Abadie, Guido Imbens, and László Sándor for sharing their implementation of the standard errors derived in Abadie and Imbens (2012), which are implemented in teffects psmatch.

We thank Edwin Leuven and Barbara Sianesi for their inspirational command psmatch2, which computes several of the estimators in teffects and a few more that we have not implemented.

References


**Also see**

[TE] **teffects** — Treatment-effects estimation for observational data

[TE] **teffects intro** — Introduction to treatment effects for observational data

[TE] **teffects multivalued** — Multivalued treatment effects
teffects aipw — Augmented inverse-probability weighting

Description

tffects aipw estimates the average treatment effect (ATE) and the potential-outcome means (POMs) from observational data by augmented inverse-probability weighting (AIPW). AIPW estimators combine aspects of regression-adjustment and inverse-probability-weighted methods. AIPW estimators have the double-robust property. teffects aipw accepts a continuous, binary, count, fractional, or nonnegative outcome and allows a multivalued treatment.

See [TE] teffects intro or [TE] teffects intro advanced for more information about estimating treatment effects from observational data.

Quick start

ATE of binary treatment treat2 by AIPW using a linear model for outcome y1 on x1 and x2 and a logistic model for treat2 on x1 and w

tffects aipw (y1 x1 x2) (treat2 x1 w)

As above, but use a fractional logistic model for fractional outcome y2

tffects aipw (y2 x1 x2, flogit) (treat2 x1 w)

As above, but use a heteroskedastic probit model for binary outcome y3 and a probit model for treat2

tffects aipw (y3 x1 x2, hetprobit(x1 x2)) (treat2 x1 w, probit)

ATE for each level of three-valued treatment treat3 on y1

tffects aipw (y1 x1 x2) (treat3 x1 w)

As above, and specify that treat3 = 3 is the control level

tffects aipw (y1 x1 x2) (treat3 x1 w), control(3)

Same as above, specified using the label “MyControl” corresponding to treat3 = 3

tffects aipw (y1 x1 x2) (treat3 x1 w), control(MyControl)

Menu

Statistics > Treatment effects > Continuous outcomes > Augmented inverse-probability weighting
Statistics > Treatment effects > Binary outcomes > Augmented inverse-probability weighting
Statistics > Treatment effects > Count outcomes > Augmented inverse-probability weighting
Statistics > Treatment effects > Fractional outcomes > Augmented inverse-probability weighting
Statistics > Treatment effects > Nonnegative outcomes > Augmented inverse-probability weighting
teffects aipw (ovar omvarlist [, omodel noconstant])
  (tvar tmvarlist [, tmodel noconstant]) [if] [in] [weight]
  [, stat options]

.ovar is a binary, count, continuous, fractional, or nonnegative outcome of interest.
.omvarlist specifies the covariates in the outcome model.
.tvar must contain integer values representing the treatment levels.
.tmvarlist specifies the covariates in the treatment-assignment model.

<table>
<thead>
<tr>
<th>omodel</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>linear</td>
<td>linear outcome model; the default</td>
</tr>
<tr>
<td>logit</td>
<td>logistic outcome model</td>
</tr>
<tr>
<td>probit</td>
<td>probit outcome model</td>
</tr>
<tr>
<td>hetprobit(varlist)</td>
<td>heteroskedastic probit outcome model</td>
</tr>
<tr>
<td>poisson</td>
<td>exponential outcome model</td>
</tr>
<tr>
<td>flogit</td>
<td>fractional logistic outcome model</td>
</tr>
<tr>
<td>fprobit</td>
<td>fractional probit outcome model</td>
</tr>
<tr>
<td>fhetprobit(varlist)</td>
<td>fractional heteroskedastic probit outcome model</td>
</tr>
</tbody>
</table>

.ovar specifies the model for the outcome variable.

<table>
<thead>
<tr>
<th>tmodel</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>logit</td>
<td>logistic treatment model; the default</td>
</tr>
<tr>
<td>probit</td>
<td>probit treatment model</td>
</tr>
<tr>
<td>hetprobit(varlist)</td>
<td>heteroskedastic probit treatment model</td>
</tr>
</tbody>
</table>

.tmodel specifies the model for the treatment variable.
For multivalued treatments, only logit is available and multinomial logit is used.

<table>
<thead>
<tr>
<th>stat</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>ate</td>
<td>estimate average treatment effect in population; the default</td>
</tr>
<tr>
<td>pomeans</td>
<td>estimate potential-outcome means</td>
</tr>
</tbody>
</table>
**Options**

---

### Model

<table>
<thead>
<tr>
<th>Option</th>
<th>Description</th>
</tr>
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<tbody>
<tr>
<td><strong>noconstant</strong></td>
<td>see [R] estimation options.</td>
</tr>
</tbody>
</table>

**nls** specifies that the parameters of the outcome model be estimated by nonlinear least squares instead of the default maximum likelihood.

**wnls** specifies that the parameters of the outcome model be estimated by weighted nonlinear least squares instead of the default maximum likelihood. The weights make the estimator of the effect parameters more robust to a misspecified outcome model.

### Stat

**stat** is one of two statistics: **ate** or **pomeans**. **ate** is the default.

- **ate** specifies that the average treatment effect be estimated.
- **pomeans** specifies that the potential-outcome means for each treatment level be estimated.
vce(vcetype) specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (robust), that allow for intragroup correlation (cluster clustvar), and that use bootstrap or jackknife methods (bootstrap, jackknife); see [R] vce_option.

Reporting

Reporting level(#); see [R] estimation options.

eaequations specifies that the results for the outcome-model or the treatment-model parameters be displayed. By default, the results for these auxiliary parameters are not displayed.

display_options: noci, nopvalues, noomitted, vsquish, noemptycells, baselevels, allbaselevels, nolabel, fvwrap(#), fvwrapon(style), cformat(%,fmt), pformat(%,fmt), sformat(%,fmt), and nolstretch; see [R] estimation options.

Maximization

maximize_options: iterate(#), [no]log, and from(init_specs); see [R] maximize. These options are seldom used.

init_specs is one of

matname [, skip copy]
# [, # ..., copy]

Advanced

pstolerance(#) specifies the tolerance used to check the overlap assumption. The default value is pstolerance(1e-5). teffects will exit with an error if an observation has an estimated propensity score smaller than that specified by pstolerance().

osample(newvar) specifies that indicator variable newvar be created to identify observations that violate the overlap assumption.

ccontrol(#) | label) specifies the level of tvar that is the control. The default is the first treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. control() may not be specified with statistic pomeans.

The following option is available with teffects aipw but is not shown in the dialog box:

coefflegend; see [R] estimation options.

Remarks and examples

Remarks are presented under the following headings:

Overview
Video example

Overview

AIPW estimators use inverse-probability weights to correct for the missing-data problem arising from the fact that each subject is observed in only one of the potential outcomes; these estimators also use an augmentation term in the outcome model to correct the estimator in case the treatment model is misspecified. If the treatment model is correctly specified, the augmentation term goes to zero in large samples.
AIPW estimators compute averages of the augmented inverse-probability-weighted outcomes for each treatment level. Contrasts of these averages provide estimates of the treatment effects.

AIPW estimators use a model to predict treatment status, and they use another model to predict outcomes. Because of the double-robust property, only one of these two models must be correctly specified for the AIPW estimator to be consistent.

AIPW estimators use a three-step approach to estimating treatment effects:

1. They estimate the parameters of the treatment model and compute inverse-probability weights.
2. They estimate separate regression models of the outcome for each treatment level and obtain the treatment-specific predicted outcomes for each subject.
3. They compute the weighted means of the treatment-specific predicted outcomes, where the weights are the inverse-probability weights computed in step 1. The contrasts of these weighted averages provide the estimates of the ATEs.

These steps produce consistent estimates of the effect parameters because the treatment is assumed to be independent of the potential outcomes after conditioning on the covariates. The overlap assumption ensures that predicted inverse-probability weights do not get too large. The standard errors reported by `teffects aipw` correct for the three-step process. See [TE] `teffects intro` or [TE] `teffects intro advanced` for more information about this estimator.

We will illustrate the use of `teffects aipw` by using data from a study of the effect of a mother’s smoking status during pregnancy (`mbsmoke`) on infant birthweight (`bweight`) as reported by Cattaneo (2010). This dataset also contains information about each mother’s age (`mage`), education level (`medu`), marital status (`mmarried`), whether the first prenatal exam occurred in the first trimester (`prenatal1`), and whether this baby was the mother’s first birth (`fbaby`).

#### Example 1: Estimating the ATE

We begin by using `teffects aipw` to estimate the average treatment effect of `mbsmoke` on `bweight`. We use a probit model to predict treatment status as a function of `mmarried`, `mage`, and `fbaby`; to maximize the predictive power of this model, we use factor-variable notation to incorporate quadratic effects of the mother’s age, the only continuous covariate in our model. We use linear regression to model birthweight, using `prenatal1`, `mmarried`, `mage`, and `fbaby` as explanatory variables. We type
. use http://www.stata-press.com/data/r15/cattaneo2
. teffects aipw (bweight prenatal1 mmarried mage fbaby)
> (mbsmoke mmarried c.mage##c.mage fbaby medu, probit)
Iteration 0:   EE criterion =  4.629e-21
Iteration 1:   EE criterion =  1.944e-25

Treatment-effects estimation                  Number of obs  =    4,642
Estimator    : augmented IPW
Outcome model : linear by ML
Treatment model: probit

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
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<tbody>
<tr>
<td></td>
<td>Coef.  Std. Err.  z  P&gt;</td>
</tr>
<tr>
<td><strong>ATE</strong></td>
<td></td>
</tr>
<tr>
<td>mbsmoke (smoker vs</td>
<td></td>
</tr>
<tr>
<td>nonsmoker)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>-230.9892</td>
</tr>
<tr>
<td></td>
<td>26.21056</td>
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<tr>
<td></td>
<td>-8.81</td>
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<td>0.000</td>
</tr>
<tr>
<td></td>
<td>-282.361 -179.6174</td>
</tr>
<tr>
<td><strong>POmean</strong></td>
<td></td>
</tr>
<tr>
<td>mbsmoke nonsmoker</td>
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<td></td>
<td>3403.355</td>
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<td></td>
<td>9.568472</td>
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<td></td>
<td>355.68</td>
</tr>
<tr>
<td></td>
<td>0.000</td>
</tr>
<tr>
<td></td>
<td>3384.601 3422.109</td>
</tr>
</tbody>
</table>

The average birthweight if all mothers were to smoke would be 231 grams less than the average of 3,403 grams that would occur if none of the mothers had smoked.

By default, `teffects aipw` reports the ATE and the POM for the base (untreated) subjects. The `pomeans` option allows us to view the treated subjects’ POM as well; the `aequations` option displays the regression model coefficients used to predict the POMs as well as the coefficients from the model used to predict treatment.

➤ Example 2: Displaying the POMs and equations

Here we use the `pomeans` and `aequations` options to obtain estimates of both POMs and view all the fitted equations underlying our estimates:
The coefficient table indicates that the treated PM is 3,172 grams, 231 grams less than the untreated PM. The sections of the table labeled OME0 and OME1 represent the linear regression coefficients for the untreated and treated potential-outcome equations, respectively. The coefficients of the TME1 equation are used in the probit model to predict treatment status.

As is well known, the standard probit model assumes that the error terms in the latent-utility framework are homoskedastic; the model is not robust to departures from this assumption. An alternative is to use the heteroskedastic probit model, which explicitly models the error variance as a function of a set of variables.

Example 3: Heteroskedastic probit treatment model

Here we refit our model as in the previous examples, but we instead use heteroskedastic probit to model the treatment variable. We posit that the heteroskedasticity is a function of the mother’s age. We type
. teffects aipw (bweight prenatal1 mmarried fbaby) > (mbsmoke mmarried c.mage##c.mage fbaby medu, hetprobit(c.mage)), aequations

|             | Coef. | Std. Err. | z      | P>|z|   | [95% Conf. Interval] |
|-------------|-------|-----------|--------|-------|----------------------|
| **ATE**     |       |           |        |       |                      |
| mbsmoke     | -230.2699 | 27.49461  | -8.38  | 0.000 | -284.1584 to -176.3815 |
| (smoker vs nonsmoker) |
| **POmean**  |       |           |        |       |                      |
| mbsmoke     | 3403.657  | 9.540713  | 356.75 | 0.000 | 3384.957 to 3422.356  |
| nonsmoker   |       |           |        |       |                      |
| **OME0**    |       |           |        |       |                      |
| prenatal1   | 69.5048  | 27.04642  | 2.57   | 0.010 | 16.49479 to 122.5148  |
| mmarried    | 173.74   | 24.63865  | 7.05   | 0.000 | 125.4491 to 222.0308  |
| fbaby       | -79.19473| 18.62584  | -4.25  | 0.000 | -115.7007 to -42.68875|
| _cons       | 3260.768 | 28.29282  | 115.25 | 0.000 | 3205.315 to 3316.221  |
| **OME1**    |       |           |        |       |                      |
| prenatal1   | 12.86437 | 39.83916  | 0.32   | 0.747 | -65.21894 to 90.94768 |
| mmarried    | 113.3491 | 39.47422  | 2.87   | 0.004 | 35.9811 to 190.7172   |
| fbaby       | 64.22326 | 38.42042  | 1.67   | 0.095 | -11.07939 to 139.5259 |
| _cons       | 3051.268 | 37.30413  | 81.79  | 0.000 | 2978.153 to 3124.383  |
| **TME1**    |       |           |        |       |                      |
| mmarried    | -.3551755| .1044199  | -3.40  | 0.001 | -.5598347 to -.1505162|
| mage        | .0831898 | .0349088  | 2.38   | 0.017 | .0147699 to .1516097  |
| c.mage#c.mage | -.0013458| .0006659  | -2.02  | 0.043 | -.002651 to -.0000406 |
| fbaby       | -.1170697| .044998   | -2.60  | 0.009 | -.2052643 to -.0288752|
| medu        | -.0435067| .0147852  | -2.94  | 0.003 | -.0724842 to -.015272  |
| _cons       | -.8757021| .347814   | -2.52  | 0.012 | -1.557405 to -.1939993|
| **TME1_lnsigma** |       |           |        |       |                      |
| mage        | -.0236336| .0107134  | -2.21  | 0.027 | -.0446315 to -.0026357|

The equation labeled TME1_lnsigma represents the heteroskedasticity function used to model the logarithm of the variance. Because the coefficient on the single variable we specified is significant below the 5% level, we conclude that allowing for heteroskedasticity was indeed necessary.

Rather than using maximum likelihood to fit the outcome model, you can instruct `teffects aipw` to use a weighted nonlinear least-squares (WNLS) estimator instead. The WNLS estimator may be more robust to outcome model misspecification.
Example 4: Using the WNLS estimator

Here we use WNLS to fit our outcome model:

```stata
use http://www.stata-press.com/data/r15/cattaneo2
> (mbsmoke mmarried c.mage##c.mage fbaby medu, probit), wnls
Iteration 0:  EE criterion = 2.742e-20
Iteration 1:  EE criterion = 3.436e-24
Treatment-effects estimation  Number of obs = 4,642
Estimator : augmented IPW
Outcome model : linear by WNLS
Treatment model: probit
```

| bweight | Coef. | Std. Err. | z    | P>|z| | 95% Conf. Interval |
|---------|-------|-----------|------|-----|-------------------|
| ATE     |       |           |      |     |                   |
| mbsmoke (smoker vs nonsmoker) | -227.1956 | 27.34794 | -8.31 | 0.000 | -280.7966 -173.5946 |
| POmean  |       |           |      |     |                   |
| mbsmoke nonsmoker | 3403.251 | 9.596622 | 354.63 | 0.000 | 3384.442 3422.06 |

The ATE of $-227$ is slightly greater than the ATE of $-231$ estimated in example 1. The estimated POMs are nearly indistinguishable.

Video example

Treatment effects: Augmented inverse-probability weighting
Stored results

teffects aipw stores the following in e():

Scalars
- e(N) number of observations
- e(nj) number of observations for treatment level j
- e(N_clust) number of clusters
- e(k_eq) number of equations in e(b)
- e(k_levels) number of levels in treatment variable
- e(treated) level of treatment variable defined as treated
- e(control) level of treatment variable defined as control
- e(converged) 1 if converged, 0 otherwise

Macros
- e(cmd) teffects
- e(cmdline) command as typed
- e(depvar) name of outcome variable
- e(tvar) name of treatment variable
- e(subcmd) aipw
- e(tmodel) logit, probit, or hetprobit
- e(omodel) linear, logit, probit, hetprobit, poisson, flogit, fprobit, or fhetprobit
- e(stat) statistic estimated, ate or pomeans
- e(vtype) weight type
- e(wexp) weight expression
- e(title) title in estimation output
- e(clustvar) name of cluster variable
- e(tlevels) levels of treatment variable
- e(cme) ml, nls, or wnls
- e(vce) vcetype specified in vce()
- e(vcetype) title used to label Std. Err.
- e(properties) b V
- e(estat_cmd) program used to implement estat
- e(predict) program used to implement predict
- e(marginsnotok) predictions disallowed by margins
- e(asbalanced) factor variables fvset as asbalanced
- e(asobserved) factor variables fvset as asobserved

Matrices
- e(b) coefficient vector
- e(V) variance–covariance matrix of the estimators

Functions
- e(sample) marks estimation sample

Methods and formulas

The methods and formulas presented here provide the technical details underlying the estimators implemented in teffects ra, teffects ipw, teffects aipw, and teffects ipwra. See Methods and formulas of [TE] teffects nnmatch for the methods and formulas used by teffects nnmatch and teffects psmatch.

Methods and formulas are presented under the following headings:
- Parameters and notation
- Overview of EE estimators
- VCE for EE estimators
- TM and OM estimating functions
  - TM estimating functions
  - OM estimating functions
- Effect estimating functions
- RA estimators
Parameters and notation

We begin by reviewing the effect parameters estimated by teffects and some essential notation.

The potential outcome that an individual would obtain if given treatment level \( t \in \{0, 1, \ldots, q\} \) is \( y_t \). Each \( y_t \) is a random variable, the realizations of which are \( y_{ti} \). Throughout this document, \( i \) subscripts denote realizations of the corresponding, unsubscripted random variables.

The three parameters of interest are

1. the potential-outcome mean (POM) \( \alpha_t = E(y_t) \);
2. the average treatment effect (ATE) \( \tau_t = E(y_t - y_0) \); and
3. the average treatment effect on the treated (ATET) \( \delta_t = E(y_t - y_0 | t = \tilde{t}) \).

The no-treatment level is 0.

The estimators implemented in teffects use three assumptions to justify the equations used for estimation and inference about the effect parameters of interest:

1. Conditional mean independence (CMI) allows us to estimate potential-outcome means from the observed outcomes in the sample.
2. Overlap ensures that we have data on each type of individual in each treatment level.
3. Independent observations ensure that the outcome and treatment for one individual has no effect on the outcome or treatment for any other individual.

teffects ra implements some regression-adjustment (RA) estimators; teffects ipw implements some inverse-probability-weighted (IPW) estimators; teffects ipwra implements some inverse-probability-weighted regression-adjustment (IPWRA) estimators; and teffects aipw implements some augmented inverse-probability-weighted (AIPW) estimators. All are implemented as estimating-equation (EE) estimators. The estimators are consistent and asymptotically normally distributed under the CMI, overlap, and independence assumptions.

Overview of EE estimators

EE estimators compute estimates by solving sample estimating equations. The sample estimating equations are the sample equivalents of population expectation equations.

Each EE estimator specifies a set of estimating equations for the effect parameters of interest and a set of estimating equations for the auxiliary parameters in the outcome model (OM) or the treatment model (TM). The next few sections provide tremendous detail about the estimating equations that define the RA, IPW, AIPW, and IPWRA estimators.

Ignoring the details for a moment, EE estimators solve systems of equations to compute estimates. A standard robust estimator is consistent for the variance of the estimator (VCE). All the details involve the equations specified by choices of estimator and functional forms for the OM or TM.

When used, the OM is a model for the conditional mean of the outcome variable. We let \( \mu(x, t, \beta_t) \) denote a conditional-mean model for the outcome \( y \) conditional on covariates \( x \) and treatment level \( t \). Mathematically, \( E(y|x, t) = \mu(x, t, \beta_t) \), where \( \beta_t \) are the parameters of the conditional-mean model given treatment level \( t \). The table below provides details about the available functional forms.
Outcome model | Functional form for $\mu(x, t, \beta_t)$
--- | ---
linear | $x\beta_t$
logit, flogit | $\exp(x\beta_t)/(1 + \exp(x\beta_t))$
probit, fprobit | $\Phi(x\beta_t)$
poisson | $\exp(x\beta_t)$
hetprobit, fhetprobit | $\Phi\{\hat{x}\beta_t/\exp(\hat{x}\beta_t)\}$

In the cases of hetprobit and fhetprobit, we use $\hat{x}$ and $\hat{\beta}_t$ to denote the variables and parameters in the index function, and we use $\hat{x}$ and $\hat{\beta}_t$ to denote the variables and parameters in the variance equation. We define $x' = (\hat{x}', \hat{x}')$ and $\beta'_t = (\hat{\beta}'_t, \hat{\beta}'_t)$.

We write the vector of parameters for the outcome model over all treatment levels as $\beta' = (\beta'_0, \beta'_1, \ldots, \beta'_q)$.

Next we provide details about the estimating equations implied by each functional form choice.

When used, the TM is a model for the conditional probability of treatment. We let $p(z, t, \gamma)$ denote the conditional probability model for the probability that a person receives treatment $t$, conditional on covariates $z$. The table below provides details about the functional form options allowed in the case of a binary treatment.

| Treatment model | Functional form for $p(z, t, \gamma)$
--- | ---
logit | $\exp(z\gamma)/(1 + \exp(z\gamma))$
probit | $\Phi(z\gamma)$
hetprobit | $\Phi\{\hat{z}\gamma/\exp(\hat{z}\gamma)\}$

In the case of hetprobit, we use $\hat{z}$ and $\hat{\gamma}$ to denote the variables and parameters in the index function, and we use $\hat{z}$ and $\hat{\gamma}$ to represent the variables and parameters in the variance equation. We define $z' = (\hat{z}', \hat{z}')$, and $\gamma' = (\hat{\gamma}', \hat{\gamma}')$.

In the multivalued-treatment case, $p(z, t, \gamma)$ is specified as a multinomial logit with $p(z, t, \gamma) = \exp(z\gamma)/\{1 + \sum_{k=1}^q \exp(z\gamma_k)\}$ and $\gamma' = (\gamma'_1, \gamma'_2, \ldots, \gamma'_q)$. (We present formulas for the case with treatment level 0 as the base with $\gamma'_0 = 0'$; see [R] mlogit for background.) In teffects, the logit option in the treatment-model specification means binary logit for the binary-treatment case and multinomial logit for the multivalued-treatment case: this simplifies the use of the command and makes statistical sense.

Below we provide details about the estimating equations implied by each functional form choice. The effect parameters of interest are

1. the POMs denoted by $\alpha' = (\alpha_0, \alpha_1, \ldots, \alpha_q)$;
2. the ATEs denoted by $\tau' = (\tau_1, \tau_2, \ldots, \tau_q)$; and
3. the ATETs denoted by $\delta' = (\delta_1, \delta_2, \ldots, \delta_q)$.

We denote the effect parameters by $\vartheta$ and all the parameters in any particular case by $\theta$. More formally, $\theta$ is the concatenation of the effect parameters, the OM parameters, and the TM parameters; $\theta' = (\vartheta', \beta', \gamma')$, where $\vartheta$ is $\alpha$, $\tau$, or $\delta$, and $\beta$ or $\gamma$ may not be present, depending on the case at hand.
In the subsections below, we discuss estimators for the elements in $\theta$ in detail and note how these elements change over the cases defined by effect parameters and estimators. The parameter vector $\theta$ denotes all the parameters, no matter which particular case is under consideration.

The EE estimators described in this section are defined by a set of equations,

$$E\{s(x, z, \theta)\} = 0$$

where $s(x, z, \theta)$ is a vector of estimating functions. Note the notation: estimating equations equate the expected value of a vector of estimating functions to zero.

Because each of the estimating functions has mean zero, we can construct estimators that find the estimates $\hat{\theta}$ by solving a system of equations,

$$\frac{1}{N} \sum_{i} s_i(x_i, z_i, \hat{\theta}) = 0$$

where $s_i(x_i, z_i, \hat{\theta})$ are the sample realizations of the estimating functions. In other words, the parameter estimates set the average of the realizations of each estimating function to zero. Almost all the details below involve specifying the sample realizations $s_i(x_i, z_i, \hat{\theta})$.

Estimators that set the expected value of estimating functions to zero are known as estimating-equations (EE) estimators, $M$ estimators, or $Z$ estimators in the statistics literature and as generalized method of moments (GMM) estimators in the econometrics literature. See van der Vaart (1998, 41), Stefanski and Boos (2002), and Tsiatis (2006, sec. 3.2) for statistics; and see Wooldridge (2010, chap. 14), Cameron and Trivedi (2005, chap. 6), and Newey and McFadden (1994) for econometrics.

We refer to them as EE estimators because this name is closest to being independent of any discipline. The estimators are implemented using gmm because they are exactly identified generalized method-of-moments (GMM) estimators. When weights are specified by the user, they are applied to the estimating equations just as gmm applies user-specified weights.

Each estimator has a set of estimating equations for the effect parameters and either an OM or a TM, or both. The OM parameters or the TM parameters are auxiliary parameters used to estimate the effect parameters of interest.

Each set of parameters has its own set of sample estimating equations:

$$\frac{1}{N} \sum_{i} s_{e,i}(x_i, z_i, \hat{\theta}) = 0$$

are the sample estimating equations for the effect parameters. These equations determine the effect parameter estimates $\hat{\vartheta}$ as functions of the data and the other estimated parameters.

$$\frac{1}{N} \sum_{i} s_{om,i}(x_i, w_i, \hat{\beta}) = 0$$

are the sample estimating equations for OM parameters that use the weights $w_i$, which are functions of the TM parameters.

$$\frac{1}{N} \sum_{i} s_{tm,i}(z_i, \hat{\gamma}) = 0$$

are the sample estimating equations for TM parameters.

The whole set of sample estimating functions is $s_i(x_i, z_i, \hat{\theta})$ with

$$s_i(x_i, z_i, \hat{\theta})' = (s_{e,i}(x_i, z_i, \hat{\theta})', s_{om,i}(x_i, w_i(t), \hat{\beta})', s_{tm,i}(z_i, \hat{\gamma})')$$

although not all the estimators have each of three components.
VCE for EE estimators

The Huber/White/robust sandwich estimator is consistent for the variance–covariance of the estimator (VCE). See van der Vaart (1998, 41), Stefanski and Boos (2002), and Tsiatis (2006, sec. 3.2) for statistics; and see Wooldridge (2010, chap. 14), Cameron and Trivedi (2005, chap. 6), and Newey and McFadden (1994) for econometrics.

The formula is

$$\hat{V} = (1/N)\overline{G} \overline{S} \overline{G}'$$

where

$$\overline{G} = \left\{ (1/N) \sum_i \frac{\partial s_i(x_i, z_i, \theta)}{\partial \theta} \right\}^{-1}$$

and

$$\overline{S} = (1/N) \sum_i s_i(x_i, z_i, \theta) s_i(x_i, z_i, \theta)'$$

The matrix $\overline{G}$ is not symmetric because our EE estimators come from stacking moment conditions instead of optimizing a single objective function. The implication is that the robust formula should always be used because, even under correct specification, the nonsymmetric $\overline{G}$ and the symmetric $\overline{S}$ converge to different matrices.

TM and OM estimating functions

Although the sample estimating functions for the effect parameters, the $s_{e,i}(x_i, z_i, \hat{\theta})$, are estimator specific, the sample estimating functions for the TM parameters, the $s_{tm,i}(z_i, \hat{\gamma})$, and the sample estimating functions for the OM parameters, the $s_{om,i}(x_i, w_i(t), \hat{\beta})'$, are used in multiple estimators. We provide details about the TM and the OM sample estimating functions here.

TM estimating functions

The sample estimating functions used to estimate the parameters of the TM $p(z, t, \gamma)$ are the sample score equations from the quasimaximum likelihood (QML) estimator.

In the binary-treatment case, $p(z, t, \gamma)$ may be logit, probit, or heteroskedastic probit. In the multivalued-treatment case, $p(z, t, \gamma)$ is a multinomial logit. We now give formulas for the $s_{tm,i}(z_i, \hat{\gamma})$ for each case.

logit and probit

In the logit and probit cases,

$$s_{tm,i}(z_i, \hat{\gamma}) = \left[ g(z_i \hat{\gamma}) \left\{ t_i - G(z_i \hat{\gamma}) \right\} \right] z_i$$

where $G(z)$ is the logistic cumulative distribution function for the logit, $G(z)$ is the normal cumulative distribution function for the probit, and $g(\cdot) = \{\partial G(\cdot)\}/(\partial z)$ is the corresponding density function.
hetprobit

In the hetprobit case, there are two sets of sample score equations, $s_{tm,1,i}(z_i, \gamma)$ and $s_{tm,2,i}(z_i, \gamma)$:

$$s_{tm,1,i}(z_i, \gamma) = \left( \frac{\phi \{ q(z_i, \gamma) \} [t_i - \Phi \{ q(z_i, \gamma) \}] - \Phi \{ q(z_i, \gamma) \} [1 - \Phi \{ q(z_i, \gamma) \}] \exp(\hat{z}_i \hat{\gamma})}{\Phi \{ q(z_i, \gamma) \} [1 - \Phi \{ q(z_i, \gamma) \}] \exp(\hat{z}_i \hat{\gamma})} \right) \hat{z}_i'$$

and

$$s_{tm,2,i}(z_i, \gamma) = \left( \frac{\phi \{ q(z_i, \gamma) \} \hat{z}_i \hat{\gamma} \Phi \{ q(z_i, \gamma) \} - \Phi \{ q(z_i, \gamma) \} [1 - \Phi \{ q(z_i, \gamma) \}] \exp(\hat{z}_i \hat{\gamma})}{\Phi \{ q(z_i, \gamma) \} [1 - \Phi \{ q(z_i, \gamma) \}] \exp(\hat{z}_i \hat{\gamma})} \right) \hat{z}_i'$$

where $\phi(\cdot)$ is the standard normal density function, and $q(z_i, \gamma) = \left( \hat{z}_i \hat{\gamma} / \exp(\hat{z}_i \hat{\gamma}) \right)$.

mlogit

In the mlogit case, $p(z, t, \gamma) = \exp(z \gamma_t) / \{1 + \sum_{k=1}^q \exp(z \gamma_k)\}$. We present formulas for the case with treatment level 0 as the base with $\gamma_0 = 0'$; see [R] mlogit for background.

There are $q$ vectors of sample estimating functions for the mlogit case, $s_{tm,k,i}(z_i, \gamma)$ for each $k \in \{1, \ldots, q\}$, 1 for each vector $\gamma_k$, $k \in \{1, \ldots, q\}$. These sample estimating functions are

$$s_{tm,k,i}(z_i, \gamma) = \begin{cases} \{1 - p(z_i, k, \gamma)\} z_i' & t_i = k \\ -p(z_i, k, \gamma) z_i' & \text{otherwise} \end{cases}$$

OM estimating functions

The parameters of the OM $\mu(x, t, \beta_t)$ are estimated either by weighted QML or by weighted nonlinear least squares. The estimating functions used to estimate the parameters of the OM are either the score equations from the weighted QML estimator or the moment conditions for the weighted nonlinear least-squares estimator.

The estimating functions for the OM parameters in $\mu(x, t, \beta_t)$ vary over the models for the conditional mean because $\mu(x, t, \beta_t)$ may be linear, logit, probit, heteroskedastic probit, or poisson.

Let $N_t$ be the number of observations in treatment level $t$, and let $t_i(t) = 1$ if $t_i = t$, with $t_i(t) = 0$ if $t_i \neq t$.

There are two sets of sample estimating functions for the OM parameters with weights $w_i(t)$:

1. $s_{ml,om,i} \{x_i, w_i(t), \hat{\beta}_t\}$ are the sample estimating functions for the weighted QML estimator.
2. $s_{nls,om,i} \{x_i, w_i(t), \hat{\beta}_t\}$ are the sample estimating functions for the weighted nonlinear least-squares estimator.

OM QML

Here are the formulas for the $s_{ml,om,i} \{x_i, w_i(t), \hat{\beta}_t\}$ for each functional form choice.
linear

In the linear case,

\[ s_{\text{ml, om}, i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \} = w_i(t)t_i(t)(y_i - x_i\hat{\beta}_t)x'_i \]

logit, flogit, probit, and fprobit

In the logit, flogit, probit, and fprobit cases,

\[ s_{\text{ml, om}, i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \} = w_i(t)t_i(t) \left[ g(\mathbf{x}_i\hat{\beta}'_t) \left\{ y_i - G(\mathbf{x}_i\hat{\beta}_t) \right\} \right. \\
\left. \frac{G(\mathbf{x}_i\hat{\beta}_t)}{G(\mathbf{x}_i\hat{\beta}_t) \{ 1 - G(\mathbf{x}_i\hat{\beta}_t) \}} \right] x_i \]

where \( G(z) \) is the logistic cumulative distribution function for the logit and flogit, \( G(z) \) is the normal cumulative distribution function for the probit and fprobit, and \( g(\cdot) = \{ \partial G(z) \}/(\partial z) \) is the corresponding density function.

hetprobit and fhethetprobit

In the hetprobit and fhethetprobit cases, there are two sets of sample score equations, \( s_{\text{ml, om}, 1,i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \} \) and \( s_{\text{ml, om}, 2,i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \} \):

\[ s_{\text{ml, om}, 1,i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \} = w_i(t)t_i(t) \left\{ \frac{\phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \} \left[ y_i - \Phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \right\} \right]}{\Phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \} \left[ 1 - \Phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \right\} \right]} \exp(\mathbf{x}_i\hat{\beta}_t) \right\} x'_i \]

and

\[ s_{\text{ml, om}, 2,i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \} = w_i(t)t_i(t) \left\{ \frac{\phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \} \left[ x_i\hat{\beta}'_t \left( \Phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \right\} - y_i \right] \right]}{\Phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \} \left[ 1 - \Phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \right\} \right]} \exp(\mathbf{x}_i\hat{\beta}_t) \right\} x'_i \]

where \( \phi(\cdot) \) is the standard normal density function, \( s_{\text{ml, om}, i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \}' = \left[ s_{\text{ml, om}, 1,i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \}', s_{\text{ml, om}, 2,i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \}' \right] \), and \( q \left( \mathbf{x}_i, \hat{\beta}_t \right) = \left( \hat{x}_i\hat{\beta}'_t / \exp(\mathbf{x}_i\hat{\beta}_t) \right) \).

poisson

In the poisson case,

\[ s_{\text{ml, om}, i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \} = w_i(t)t_i(t)(y_i - \exp(\mathbf{x}_i\hat{\beta}_t))x'_i \]

OM WNL

Here are the formulas for the \( s_{\text{nls, om}, i} \{ \mathbf{x}_i, w_i(t), \hat{\beta}_t \} \) for each functional form choice.
linear

In the linear case,
\[
\mathbf{s}_{\text{nls,om},i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\} = w_i(t) t_i(t) (y_i - \mathbf{x}_i \hat{\beta}_t)^\prime \mathbf{x}_i
\]

logit, flogit, probit, and fprobit

In the logit, flogit, probit, and fprobit cases,
\[
\mathbf{s}_{\text{nls,om},i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\} = w_i(t) t_i(t) \left[ g(\mathbf{x}_i \hat{\beta}_t) \left\{ y_i - G(\mathbf{x}_i \hat{\beta}_t) \right\} \right] \mathbf{x}_i
\]

where \( G(z) \) is the logistic cumulative distribution function for the logit and flogit, \( G(z) \) is the normal cumulative distribution function for the probit and fprobit, and \( g(\cdot) = \partial G(z) / \partial z \) is the corresponding density function.

hetprobit and fhetprobit

In the hetprobit and fhetprobit cases, there are two sets of sample score equations, \( \mathbf{s}_{\text{nls,om},1,i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\} \) and \( \mathbf{s}_{\text{nls,om},2,i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\} \):
\[
\mathbf{s}_{\text{nls,om},1,i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\} = w_i(t) t_i(t) \left( \frac{\phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \right\}}{\exp(\dot{\mathbf{x}}_i \hat{\beta}_t)} \right) \mathbf{y}_i - \Phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \right\} \right] \mathbf{x}_i
\]

and
\[
\mathbf{s}_{\text{nls,om},2,i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\} = w_i(t) t_i(t) \left( \frac{\phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \right\}}{\exp(\dot{\mathbf{x}}_i \hat{\beta}_t)} \right) \mathbf{x}_i \hat{\beta}_t \left[ \Phi \left\{ q \left( \mathbf{x}_i, \hat{\beta}_t \right) \right\} - y_i \right] \mathbf{x}_i
\]

where \( \phi(\cdot) \) is the standard normal density function. \( \mathbf{s}_{\text{nls,om},i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\}' = \left[ \mathbf{s}_{\text{nls,om},1,i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\}', \mathbf{s}_{\text{nls,om},2,i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\}' \right] \), and \( q \left( \mathbf{x}_i, \hat{\beta}_t \right) = \left( \mathbf{x}_i \hat{\beta}_t / \exp(\dot{\mathbf{x}}_i \hat{\beta}_t) \right) \).

poisson

In the poisson case,
\[
\mathbf{s}_{\text{nls,om},i}\{\mathbf{x}_i, w_i(t), \hat{\beta}_t\} = w_i(t) t_i(t) \{y_i - \exp(\mathbf{x}_i \hat{\beta}_t)\} \exp(\mathbf{x}_i \hat{\beta}_t) \mathbf{x}_i
\]

Effect estimating functions

We now describe the sample estimating functions for the effect parameters, which vary over estimator and effect parameter.
RA estimators

RA estimators estimate the effect parameters using means of the observation-level predictions of the conditional means of the outcomes. There is no model for the conditional probability of treatment.

The RA estimators use unweighted QML estimators to estimate the parameters of the conditional mean model. In other words, the RA estimators use the sample estimating functions $s_{ml,om,i}(x_i, 1, \hat{\beta})$, given above.

For the RA estimators, the vector of sample estimating functions is the concatenation of the sample estimating functions for the effect parameters with the sample estimating functions for the OM parameters. Algebraically,

$$s_{ra,i}(x_i, \hat{\theta})' = s_{ra,e,i}(x_i, \hat{\theta}, \hat{\beta})', s_{ml,om,i}(x_i, 1, \hat{\beta})'$$

The estimating functions $s_{ra,e,i}(x_i, \hat{\theta}, \hat{\beta})'$ vary over the effect parameter.

RA for POM

The RA estimators for the POM parameters estimate $\theta' = (\alpha', \beta')$ using two types of estimating equations: 1) those for the POM parameters $\alpha$, and 2) those for the conditional-mean model parameters $\beta_t$ in $\mu(x, t, \beta_t)$.

The sample estimating functions for the $\hat{\beta}_t$ are given in OM estimating functions above.

The elements of $s_{ra,e,i}(x_i, \hat{\alpha}, \hat{\beta})$ for the POM parameters $\alpha$ are given by

$$\mu(x_i, t, \hat{\beta}_t) - \hat{\alpha}_t \quad (RAPOM)$$

RA for ATE

The RA estimators for the ATE parameters estimate $\theta' = (\tau', \beta')$ using two types of estimating equations: 1) those for the ATE parameters $\tau$, and 2) those for the OM parameters $\beta_t$ in $\mu(x, t, \beta_t)$.

The sample estimating functions that determine the $\hat{\beta}_t$ are given in OM estimating functions with $w_i(t) = 1$.

The elements of $s_{ra,e,i}(x_i, \hat{\tau}, \hat{\beta})$ for the ATE parameters $\tau$ are given by

$$\mu(x_i, t, \hat{\beta}_t) - \mu(x_i, 0, \hat{\beta}_t) - \hat{\tau}_t \quad (RAATE)$$

RA for ATET

The RA estimators for the ATET parameters estimate $\theta' = (\delta', \beta')$ using two types of estimating equations: 1) those for the ATET parameters $\delta$, and 2) those for the OM parameters $\beta_t$ in $\mu(x, t, \beta_t)$.

The sample estimating functions that determine the $\hat{\beta}_t$ are given in OM estimating functions above with $w_i(t) = 1$.

The elements of $s_{ra,e,i}(x_i, \hat{\delta}, \hat{\beta})$ for the ATET parameters $\delta$ are given by

$$Nt_i(\hat{t})/N\tilde{t} \left\{ \mu(x_i, t, \hat{\beta}_t) - \mu(x_i, 0, \hat{\beta}_t) - \hat{\delta}_t \right\} \quad (RAATET)$$
IPW estimators

IPW estimators estimate the effect parameters using means of the observed outcomes weighted by the inverse probability of treatment. There is no outcome model. The IPW estimators use QML estimators to estimate the parameters of the conditional probability model.

The vector of estimating functions is the concatenation of the estimating functions for the effect parameters with the estimating functions for the conditional-probability parameters. The sample estimating functions used by the IPW estimators are

\[ s_{ipw, i}(\mathbf{x}_i, \hat{\theta})' = s_{ipw, e, i}(\mathbf{x}_i, \hat{\theta}, \hat{\gamma})', s_{tm, i}(\mathbf{z}_i, 1, \hat{\gamma})' \]

The estimating functions \( s_{ipw, e, i}(\mathbf{z}_i, \hat{\theta}, \hat{\gamma})' \) vary over the effect parameter.

All the IPW estimators use normalized inverse-probability weights. These weights are not related to the weights \( w_i(t) \) that were used in the OM equations. The functional form for the normalized inverse-probability weights varies over the effect parameters POM, ATE, and ATET.

The POM and ATE estimators use normalized inverse-probability weights. The unnormalized weights for individual \( i \) and treatment level \( t \) are \( d_i(t) = t_i(t)/p(z_i, t, \hat{\gamma}) \), and the normalized weights are \( \bar{d}_i(t) = N_i d_i(t)/\sum_i N d_i(t) \).

The ATET estimator uses normalized treatment-adjusted inverse-probability weights. The treatment-adjusted inverse-probability weights adjust the inverse-probability weights for the probability of getting the conditional treatment \( \tilde{t} \). The unnormalized weights are \( f_i = p(z_i, \tilde{t}, \hat{\gamma})/p(z_i, t_i, \hat{\gamma}) \), and the normalized weights are \( \bar{f}_i = N f_i/\sum_i N f_i \).

The IPW effect estimators are weighted cell averages. The sample estimating functions used in POM estimators are the sample estimating functions from weighted OLS regression on treatment-cell indicators. The POM estimators use a full set of \( q + 1 \) of treatment indicator variables \( a_i \). (The \( i \)th observation on the \( k \)th variable in \( a_i \) is 1 if \( i \) received treatment \( k - 1 \) and 0 otherwise, for \( k \in \{1, 2, \ldots, q + 1\} \).)

The sample estimating functions used in the ATE and the ATET estimators are the sample estimating functions from weighted OLS regression on treatment-cell indicators, excluding the indicator for the control level and including a constant term. The variables \( \tilde{a}_i \) used in the ATE and ATET sample estimating functions include \( q \) of treatment indicator variables and a variable that is always 1. (The first \( q \) variables in \( \tilde{a}_i \) are treatment indicators: the \( i \)th observation on the \( k \)th variable in \( \tilde{a}_i \) is 1 if \( i \) received treatment \( k \) and 0 otherwise, for \( k \in \{1, 2, \ldots, q\} \). The \((q + 1)\)th variable is always 1.) This definition of \( \tilde{a}_i \) sets the last treatment level to be the control; renaming the treatments handles the more general case allowed for by teffects.

Having defined \( a_i \) and \( \tilde{a}_i \), we can give the sample estimating functions that the IPW estimators use for the effects parameters.

IPW for POM

We begin with the IPW estimators for the potential-outcome means. In this case, \( \theta' = (\alpha', \gamma') \).

The sample estimating functions for the \( \hat{\gamma} \) are given in TM estimating functions above.

The sample estimating functions for \( \hat{\alpha} \) are given by

\[ s_{ipw, e, i, t}(\mathbf{z}_i, \hat{\alpha}, \hat{\gamma})' = \bar{d}_i(t)(y_i - a_i \hat{\alpha})\tilde{a}_i' \]  

(IPWPOM)
IPW for ATE

The full parameter vector for the IPW estimators for the ATE is $\theta' = (\tau', \gamma')$.
The sample estimating functions for the $\widehat{\gamma}$ are given in *TM estimating functions* above.
The sample estimating functions for $\widehat{\tau}$ are given by

$$s_{ipw,e,i,t}(z_i, \widehat{\gamma}, \widehat{\tau})' = \overline{d}_i(t)(y_i - \tilde{a}_i\widehat{\tau})\tilde{a}_i' \tag{IPWATE}$$

IPW for ATET

The full parameter vector for the IPW estimators for the ATET is $\theta' = (\delta', \gamma')$.
The sample estimating functions for the $\widehat{\gamma}$ are given in *TM estimating functions* above.
The sample estimating functions for $\widehat{\delta}$ are given by

$$s_{ipw,e,i,t}(z_i, \widehat{\delta}, \widehat{\gamma})' = \overline{f}_i(t)(y_i - \tilde{a}_i\widehat{\delta})\tilde{a}_i' \tag{IPWATET}$$

AIPW estimators

This section documents the sample estimating functions used by the augmented inverse-probability-weighted (AIPW) estimators implemented in *teffects*. These AIPW estimators are efficient-influence-function estimators as discussed in [TE] *teffects intro* and [TE] *teffects intro advanced*. The *teffects* implementation was primarily inspired by Cattaneo, Drukker, and Holland (2013), which was based on Cattaneo (2010). Tan (2010) was influential by identifying the implemented weighted nonlinear least-squares estimator as having relatively good properties when both the conditional mean function and the conditional probability function are misspecified.

The AIPW estimating functions for the treatment parameters include terms from a conditional probability model and from a conditional mean model for the outcome.

The sample-estimation-equations vector has three parts for the AIPW estimators:

$$s_{aipw,i}(x_i, z_i, \widehat{\theta})' = [s_{aipw,e,i}(x_i, z_i, \widehat{\theta})', s_{aipw,tm,i}(z_i, \widehat{\gamma})', s_{aipw,om,i}(x_i, w_i(t), \widehat{\beta})']$$

The sample estimating functions for the parameters of the TM are the $s_{tm,i}(z_i, \widehat{\gamma})$ given in *TM estimating functions* above.

*teffects aipw* implements three AIPW estimators: the standard AIPW estimator, the NLS AIPW estimator, and the WNLS AIPW estimator.

The three AIPW estimators differ in how they estimate the parameters of the OM.

By default, *teffects aipw* uses the standard AIPW estimator that estimates the parameters of the OM by the unweighted ML estimator, whose sample estimating functions, $s_{ml,om,i}(x_i, 1, \widehat{\beta})$, are given in *OM estimating functions* above.

When the *nls* option is specified, *teffects aipw* uses the NLS AIPW estimator that estimates the parameters of the OM by the unweighted NLS estimator, whose sample estimating functions, $s_{nls,om,i}(x_i, 1, \widehat{\beta})$, are given in *OM estimating functions* above.
When the \texttt{wnls} option is specified, \texttt{teffects aipw} uses the WNLS AIPW estimator that estimates the parameters of the OM by the WNLS estimator, whose sample estimating functions, \( s_{\text{nl}, \text{om}, i}(x_i, \hat{w}_i(t), \hat{\beta}) \), are given in \textit{OM estimating functions} above. The weights for person \( i \) in treatment level \( t \) are

\[
\hat{w}_i(t) = \frac{t_i(t)}{p(z_i, t, \gamma)} \left\{ \frac{t_i(t)}{p(z_i, t, \gamma)} - 1 \right\} \quad \text{(WNLSW)}
\]

Now we discuss the sample estimating functions for the effect parameters, the \( s_{e, i}(x_i, z_i, \hat{\theta}) \).

\textbf{AIPW for POM}

We begin with the AIPW estimators for the potential-outcome means. In this case, \( \theta' = (\alpha', \gamma', \beta') \), and the elements of \( s_{\text{aipw}, e, i}(x_i, z_i, \hat{\theta}) \) are given by

\[
\frac{t_i(t)}{p(z_i, t, \gamma)} y_i - \mu(x_i, \hat{\beta}_t) \left\{ \frac{t_i(t)}{p(z_i, t, \gamma)} - 1 \right\} - \alpha_t
\]

\textbf{AIPW for ATE}

The AIPW estimators for the ATE estimate \( \theta' = (\tau', \gamma', \beta') \), and the elements of \( s_{\text{aipw}, e, i}(x_i, z_i, \hat{\theta}) \) are given by

\[
\begin{align*}
\frac{t_i(t)}{p(z_i, t, \gamma)} y_i - \mu(x_i, \hat{\beta}_t) \left\{ \frac{t_i(t)}{p(z_i, t, \gamma)} - 1 \right\} - \tau_0 & \text{ if } t = 0 \\
\frac{t_i(t)}{p(z_i, t, \gamma)} y_i - \mu(x_i, \hat{\beta}_t) \left\{ \frac{t_i(t)}{p(z_i, t, \gamma)} - 1 \right\} - \tau_t - \tau_0 & \text{ if } t > 0
\end{align*}
\]

\textbf{IPWRA estimators}

The IPWRA estimators combine inverse-probability weighting (IPW) with regression-adjustment (RA) methods. The sample estimating functions for IPWRA include sample estimating functions from both RA and IPW. The vector of sample estimating functions is

\[
s_{\text{ipwra}, i}(x_i, \hat{\theta})' = s_{\text{ra}, e, i}(x_i, \hat{\vartheta}, \hat{\beta})', s_{\text{ml}, \text{om}, i}(x_i, w_i(j), \hat{\beta})', s_{\text{tm}, i}(z_i, \hat{\gamma})'
\]

where \( \hat{\theta}' = (\hat{\vartheta}', \hat{\beta}', \hat{\gamma}') \), \( \hat{\vartheta} = \hat{\alpha} \) for POM, \( \hat{\vartheta} = \hat{\tau} \) for ATE, and \( \hat{\vartheta} = \hat{\delta}_t \) for ATET. The sample estimating functions, \( s_{\text{ra}, e, i}(x_i, \hat{\vartheta}, \hat{\beta}) \), for POM, ATE, and ATET are given in equations (RAPOM), (RAATE), and (RAATET). For the OM sample estimating functions, \( s_{\text{ml}, \text{om}, i}(\cdot) \), we replace the RA unity weights, \( w_i(t) = 1 \), with \( d_i(j) \) for POM or ATE and \( \bar{f}_i \) for ATET, the normalized inverse-probability weights described in \textit{IPW estimators} above. The specific form of the OM sample estimating functions are given in \textit{OM estimating functions} above. The TM sample estimating functions are given in \textit{TM estimating functions} above.
References


Also see

[TE] tefects postestimation — Postestimation tools for tefects

[TE] tefects — Treatment-effects estimation for observational data

[TE] tefects ipwra — Inverse-probability-weighted regression adjustment

[U] 20 Estimation and postestimation commands


teffects ipw — Inverse-probability weighting

**Description**

teffects ipw estimates the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs) from observational data by inverse-probability weighting (IPW). IPW estimators use estimated probability weights to correct for missing data on the potential outcomes. teffects ipw accepts a continuous, binary, count, fractional, or nonnegative outcome and allows a multivalued treatment.

See [TE] teffects intro or [TE] teffects intro advanced for more information about estimating treatment effects from observational data.

**Quick start**

ATE of binary treat2 on y by IPW using a logistic model of treat2 on x and w

teffects ipw (y) (treat2 x w)

As above, but estimate ATET

teffects ipw (y) (treat2 x w), atet

As above, but estimate potential-outcome means

teffects ipw (y) (treat2 x w), pomeans

ATE of treat2 on y using heteroskedastic probit for treat2 as a function of x and w

teffects ipw (y) (treat2 x w, hetprobit(x w))

ATE for treatment levels 2 and 3 of three-valued treatment treat3

teffects ipw (y) (treat3 x w)

As above, and specify that treat3 = 3 is the control level

teffects ipw (y) (treat3 x w), control(3)

Same as above, specified using the label “MyControl” corresponding to treat3 = 3

teffects ipw (y) (treat3 x w), control(MyControl)

**Menu**

Statistics > Treatment effects > Continuous outcomes > Inverse-probability weighting (IPW)

Statistics > Treatment effects > Binary outcomes > Inverse-probability weighting (IPW)

Statistics > Treatment effects > Count outcomes > Inverse-probability weighting (IPW)

Statistics > Treatment effects > Fractional outcomes > Inverse-probability weighting (IPW)

Statistics > Treatment effects > Nonnegative outcomes > Inverse-probability weighting (IPW)
**Syntax**

```
teffects ipw (ovar) (tvar tmvarlist [, tmodel noconstant]) [if] [in] [weight] 
    [, stat options]
```

`ovar` is a binary, count, continuous, fractional, or nonnegative outcome of interest.

`tvar` must contain integer values representing the treatment levels.

`tmvarlist` specifies the variables that predict treatment assignment in the treatment model.

**tmodel**

<table>
<thead>
<tr>
<th>Model</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>logit</td>
<td>logistic treatment model; the default</td>
</tr>
<tr>
<td>probit</td>
<td>probit treatment model</td>
</tr>
<tr>
<td>hetprobit(varlist)</td>
<td>heteroskedastic probit treatment model</td>
</tr>
</tbody>
</table>

`tmodel` specifies the model for the treatment variable.
For multivalued treatments, only `logit` is available and multinomial logit is used.

**stat**

<table>
<thead>
<tr>
<th>Stat</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>ate</td>
<td>estimate average treatment effect in population; the default</td>
</tr>
<tr>
<td>atet</td>
<td>estimate average treatment effect on the treated</td>
</tr>
<tr>
<td>pomeans</td>
<td>estimate potential-outcome means</td>
</tr>
</tbody>
</table>

**options**

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
</table>

**SE/Robust**

```
vce(vcetype) vcetype may be robust, cluster clustvar, bootstrap, or jackknife
```

**Reporting**

```
level(#) set confidence level; default is level(95)
aequations display auxiliary-equation results
display_options control columns and column formats, row spacing, line width,
                   display of omitted variables and base and empty cells, and
                   factor-variable labeling
```

**Maximization**

```
maximize_options control the maximization process; seldom used
```

**Advanced**

```
pstolerance(#) set tolerance for overlap assumption
osample(newvar) newvar identifies observations that violate the overlap assumption
control(# | label) specify the level of `tvar` that is the control
tlevel(# | label) specify the level of `tvar` that is the treatment
coeflegend display legend instead of statistics
```
tmvarlist may contain factor variables; see [U] 11.4.3 Factor variables.

bootstrap, by, jackknife, and statsby are allowed; see [U] 11.1.10 Prefix commands.

Weights are not allowed with the bootstrap prefix; see [R] bootstrap.

fweights, iweights, and pweights are allowed; see [U] 11.1.6 weight.

coefflegend does not appear in the dialog box.

See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.

Options

Model

tnoconstant; see [R] estimation options.

Stat

stat is one of three statistics: ate, atet, or pomeans. ate is the default.

ate specifies that the average treatment effect be estimated.

atet specifies that the average treatment effect on the treated be estimated.

pomeans specifies that the potential-outcome means for each treatment level be estimated.

SE/Robust

vce(vcetype) specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (robust), that allow for intragroup correlation (cluster clustvar), and that use bootstrap or jackknife methods (bootstrap, jackknife); see [R] vce_option.

Reporting

level(#); see [R] estimation options.

eaequations specifies that the results for the outcome-model or the treatment-model parameters be displayed. By default, the results for these auxiliary parameters are not displayed.

display_options: noci, nopvalues, noomitted, vsquish, noemptycells, baselevels, allbaselevels, nolabel, fvwrap(#), fvwrapon(style), cformat(%fmt), pformat(%fmt), sformat(%fmt), and nolstretch; see [R] estimation options.

Maximization

maximize_options: iterate(#), [no]log, and from(init specs); see [R] maximize. These options are seldom used.

init Specs is one of

matname [, skip copy]

# [, # ...], copy

Advanced

pstolerance(#) specifies the tolerance used to check the overlap assumption. The default value is pstolerance(1e-5). teffects will exit with an error if an observation has an estimated propensity score smaller than that specified by pstolerance().

osample(newvar) specifies that indicator variable newvar be created to identify observations that violate the overlap assumption.
control(# | label) specifies the level of tvar that is the control. The default is the first treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. control() may not be specified with statistic pomeans. control() and tlevel() may not specify the same treatment level.

tlevel(# | label) specifies the level of tvar that is the treatment for the statistic atet. The default is the second treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. tlevel() may only be specified with statistic atet. tlevel() and control() may not specify the same treatment level.

The following option is available with teffects ipw but is not shown in the dialog box: coeflegend; see [R] estimation options.

Remarks and examples

Remarks are presented under the following headings:

Overview
Video example

Overview

IPW estimators use estimated probability weights to correct for the missing-data problem arising from the fact that each subject is observed in only one of the potential outcomes. IPW estimators use a two-step approach to estimating treatment effects:

1. They estimate the parameters of the treatment model and compute the estimated inverse-probability weights.
2. They use the estimated inverse-probability weights to compute weighted averages of the outcomes for each treatment level. The contrasts of these weighted averages provide the estimates of the ATEs. Using this weighting scheme corrects for the missing potential outcomes.

These steps produce consistent estimates of the effect parameters because the treatment is assumed to be independent of the potential outcomes after conditioning on the covariates. The overlap assumption ensures that predicted inverse-probability weights do not get too large. In fact, teffects ipw uses an estimation technique that implements both steps at once so that we do not need to correct the standard errors in the second step to reflect the uncertainty associated with the predicted treatment probabilities.

We will illustrate the use of teffects ipw by using data from a study of the effect of a mother’s smoking status during pregnancy (mbsmoke) on infant birthweight (bweight) as reported by Cattaneo (2010). This dataset also contains information about each mother’s age (mage), education level (medu), marital status (mmarried), whether the first prenatal exam occurred in the first trimester (prenatal1), and whether this baby was the mother’s first birth (fbaby).

Example 1: Estimating the ATE

We begin by using teffects ipw to estimate the average treatment effect of smoking on birthweight. We will use a probit model to predict treatment status, using prenatal1, mmarried, mage, the square of mage, and fbaby as explanatory variables:
The average birthweight if all mothers were to smoke would be 231 grams less than the average of 3,403 grams that would occur if none of the mothers had smoked.

Sometimes, we are mainly concerned about those subjects that did in fact receive treatment, and we want to know how much the outcome changes as a result of treatment for that subpopulation. The ATET provides us with the answer. Moreover, the ATET can be estimated using weaker assumptions than are required to estimate the ATE; see [TE] `teffects intro advanced`.

Example 2: Estimating the ATET

```
. teffects ipw (bweight) (mbsmoke mmarried c.mage##c.mage fbaby medu, probit), > atet
Iteration 0: EE criterion = 4.636e-21
Iteration 1: EE criterion = 6.467e-27
Treatment-effects estimation Number of obs = 4,642
Estimator : inverse-probability weights
Outcome model : weighted mean
Treatment model: probit

<table>
<thead>
<tr>
<th>bweight</th>
<th>Robust</th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coef. Std. Err.</td>
<td>z</td>
<td>P&gt;</td>
<td>z</td>
</tr>
<tr>
<td>ATE</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mbsmoke</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>smoker</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>vs nonsmoker</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>-225.1773</td>
<td>23.66458</td>
<td>-9.52</td>
<td>0.000</td>
<td>-271.559</td>
</tr>
<tr>
<td>P0mean</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mbsmoke</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>nonsmoker</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3362.837</td>
<td>14.20149</td>
<td>236.79</td>
<td>0.000</td>
<td>3335.003</td>
</tr>
</tbody>
</table>
```

```
The average birthweight is 225 grams less when all the mothers who smoke do so than the average of 3,363 grams that would have occurred if none of these mothers had smoked.

We often express statistics as percentages to alleviate scaling issues and aid interpretation. In the present context, we may wish to express an ATE as a percentage of the untreated POM to gain a more intuitive measure of the effect of treatment.

Example 3: Reporting the ATE as a percentage

Here we use the same model as in example 1, but we report the ATE as a percentage of the mean birthweight that would occur if no mothers smoke. First, we use `teffects ipw` to fit the model. We use the `coeflegend` option so that `teffects ipw` reports the names of the parameters. Then we use `nlcom` to obtain the statistic we want along with its delta-method-based standard error. We type

```
. teffects ipw (bweight) (mbsmoke mmarried c.mage##c.mage fbaby medu, probit), 
>    > coeflegend
```

```
  Iteration 0: EE criterion = 4.621e-21
  Iteration 1: EE criterion = 7.358e-26

Treatment-effects estimation Number of obs = 4,642
Estimator : inverse-probability weights
Outcome model : weighted mean
Treatment model: probit

<table>
<thead>
<tr>
<th>bweight</th>
<th>Coef. Legend</th>
</tr>
</thead>
<tbody>
<tr>
<td>ATE</td>
<td></td>
</tr>
<tr>
<td>mbsmoke</td>
<td></td>
</tr>
<tr>
<td>(smoker vs</td>
<td></td>
</tr>
<tr>
<td>nonsmoker)</td>
<td>-230.6886 _b[ATE:r1vs0.mbsmoke]</td>
</tr>
<tr>
<td>POmean</td>
<td></td>
</tr>
<tr>
<td>mbsmoke</td>
<td></td>
</tr>
<tr>
<td>nonsmoker</td>
<td>3403.463 _b[POmean:0.mbsmoke]</td>
</tr>
</tbody>
</table>
```

```
. nlcom _b[ATE:r1vs0.mbsmoke] / _b[POmean:0.mbsmoke]
  _nl_1: _b[ATE:r1vs0.mbsmoke] / _b[POmean:0.mbsmoke]

| bweight | Coef.   | Std. Err. | z     | P>|z|  | [95% Conf. Interval] |
|---------|---------|-----------|-------|------|---------------------|
| _nl_1   | -.0677806 | .0075169  | -9.02 | 0.000 | -.0825133 -.0530478  |
```

The average birthweight falls by an estimated 6.8% when every mother smokes relative to the case when no mothers smoke. We also obtain a 95% confidence interval of a 5.3% to 8.3% reduction.

Video example

Treatment effects: Inverse-probability weighting
t`effe`cts ipw — Inverse-probability weighting

Stored results

t`effe`cts ipw stores the following in `e`():

Scalars
- `e(N)`: number of observations
- `e(n_j)`: number of observations for treatment level `j`
- `e(N_clust)`: number of clusters
- `e(k_eq)`: number of equations in `e(b)`
- `e(k_levels)`: number of levels in treatment variable
- `e(treated)`: level of treatment variable defined as treated
- `e(control)`: level of treatment variable defined as control
- `e(converged)`: 1 if converged, 0 otherwise

Macros
- `e(cmd)`: `teffects`
- `e(cmdline)`: command as typed
- `e(depvar)`: name of outcome variable
- `e(tvar)`: name of treatment variable
- `e(subcmd)`: `ipw`
- `e(tmodel)`: `logit`, `probit`, or `hetprobit`
- `e(stat)`: statistic estimated, `ate`, `atet`, or `pomeans`
- `e(vtype)`: weight type
- `e(wexp)`: weight expression
- `e(title)`: title in estimation output
- `e(clustvar)`: name of cluster variable
- `e(tlevels)`: levels of treatment variable
- `e(vce)`: `vcetype` specified in `vce()`
- `e(vcetype)`: title used to label Std. Err.
- `e(properties)`: `b V`
- `e(estat_cmd)`: program used to implement `estat`
- `e(predict)`: program used to implement `predict`
- `e(marginsnotok)`: predictions disallowed by `margins`
- `e(asbalanced)`: factor variables `fvset` as `asbalanced`
- `e(asobserved)`: factor variables `fvset` as `asobserved`

Matrices
- `e(b)`: coefficient vector
- `e(V)`: variance–covariance matrix of the estimators

Functions
- `e(sample)`: marks estimation sample

Methods and formulas

t`effe`cts ipw implements a smooth treatment-effects estimator. All smooth treatment-effects estimators are documented in Methods and formulas of [TE] t`effe`cts aipw.

References


Also see

[TE] teffects postestimation — Postestimation tools for teffects

[TE] teffects — Treatment-effects estimation for observational data

[U] 20 Estimation and postestimation commands
t effects ipwra — Inverse-probability-weighted regression adjustment

Description

t effects ipwra estimates the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs) from observational data by inverse-probability-weighted regression adjustment (IPWRA). IPWRA estimators use weighted regression coefficients to compute averages of treatment-level predicted outcomes, where the weights are the estimated inverse probabilities of treatment. The contrasts of these averages estimate the treatment effects. IPWRA estimators have the double-robust property. t effects ipwra accepts a continuous, binary, count, fractional, or nonnegative outcome and allows a multivalued treatment.

See [TE] t effects intro or [TE] t effects intro advanced for more information about estimating treatment effects from observational data.

Quick start

ATE of binary treatment treat2 estimated by IPWRA using a linear model for outcome y1 on x1 and x2 and a logistic model for treat2 on x1 and w

t effects ipwra (y1 x1 x2) (treat2 x1 w)

As above, but estimate the ATET

t effects ipwra (y1 x1 x2) (treat2 x1 w), atet

Probit model for binary outcome y3

t effects ipwra (y3 x1 x2, probit) (treat2 x1 w)

As above, but use a heteroskedastic probit model for y3 and a probit model for treat2

t effects ipwra (y3 x1 x2, hetprobit(x1 x2)) (treat2 x1 w, probit)

As above, but use a fractional heteroskedastic probit model for y4 and a probit model for treat2

t effects ipwra (y4 x1 x2, fhetprobit(x1 x2)) (treat2 x1 w, probit)

ATE for each level of a three-valued treatment treat3

t effects ipwra (y1 x1 x2) (treat3 x1 w)

As above, and specify that treat3 = 3 is the control level using the value label “MyControl” for 3

t effects ipwra (y1 x1 x2) (treat3 x1 w), control(MyControl)

Menu

Statistics > Treatment effects > Continuous outcomes > Regression adjustment with IPW
Statistics > Treatment effects > Binary outcomes > Regression adjustment with IPW
Statistics > Treatment effects > Count outcomes > Regression adjustment with IPW
Statistics > Treatment effects > Fractional outcomes > Regression adjustment with IPW
Statistics > Treatment effects > Nonnegative outcomes > Regression adjustment with IPW
teffects ipwra — Inverse-probability-weighted regression adjustment

Syntax

```
teffects ipwra (ovar omvarlist [, omodel noconstant])
    (tvar tmvarlist [, tmodel noconstant]) [if] [in] [weight]
    [, stat options]
```

`ovar` is a binary, count, continuous, fractional, or nonnegative outcome of interest.

`omvarlist` specifies the covariates in the outcome model.

`tvar` must contain integer values representing the treatment levels.

`tmvarlist` specifies the covariates in the treatment-assignment model.

**omodel**

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Model</strong></td>
</tr>
<tr>
<td>linear</td>
</tr>
<tr>
<td>logit</td>
</tr>
<tr>
<td>probit</td>
</tr>
<tr>
<td>hetprobit(varlist)</td>
</tr>
<tr>
<td>poisson</td>
</tr>
<tr>
<td>flogit</td>
</tr>
<tr>
<td>fprobit</td>
</tr>
<tr>
<td>fhetprobit(varlist)</td>
</tr>
</tbody>
</table>

`omodel` specifies the model for the outcome variable.

**tmodel**

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Model</strong></td>
</tr>
<tr>
<td>logit</td>
</tr>
<tr>
<td>probit</td>
</tr>
<tr>
<td>hetprobit(varlist)</td>
</tr>
</tbody>
</table>

`tmodel` specifies the model for the treatment variable.

For multivalued treatments, only `logit` is available and multinomial logit is used.

**stat**

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Stat</strong></td>
</tr>
<tr>
<td>ate</td>
</tr>
<tr>
<td>atet</td>
</tr>
<tr>
<td>pomeans</td>
</tr>
</tbody>
</table>

```
### Description

<table>
<thead>
<tr>
<th>Options</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>SE/Robust</td>
<td></td>
</tr>
<tr>
<td><code>vce(vcetype)</code></td>
<td><code>vcetype</code> may be <code>robust</code>, <code>cluster clustvar</code>, <code>bootstrap</code>, or <code>jackknife</code></td>
</tr>
<tr>
<td>Reporting</td>
<td></td>
</tr>
<tr>
<td><code>level(#)</code></td>
<td>set confidence level; default is <code>level(95)</code></td>
</tr>
<tr>
<td><code>aequations</code></td>
<td>display auxiliary-equation results</td>
</tr>
<tr>
<td><code>display_options</code></td>
<td>control columns and column formats, row spacing, line width, display of omitted variables and base and empty cells, and factor-variable labeling</td>
</tr>
<tr>
<td>Maximization</td>
<td></td>
</tr>
<tr>
<td><code>maximize_options</code></td>
<td>control the maximization process; seldom used</td>
</tr>
<tr>
<td>Advanced</td>
<td></td>
</tr>
<tr>
<td><code>pstorencelion(#)</code></td>
<td>set tolerance for overlap assumption</td>
</tr>
<tr>
<td><code>osample(newvar)</code></td>
<td><code>newvar</code> identifies observations that violate the overlap assumption</td>
</tr>
<tr>
<td>`control(#</td>
<td>label)`</td>
</tr>
<tr>
<td>`tlevel(#</td>
<td>label)`</td>
</tr>
<tr>
<td><code>coeflegend</code></td>
<td>display legend instead of statistics</td>
</tr>
</tbody>
</table>

`omvarlist` and `tmvarlist` may contain factor variables; see [U] 11.4.3 Factor variables.

`bootstrap`, `by`, `jackknife`, and `statsby` are allowed; see [U] 11.1.10 Prefix commands.

Weights are not allowed with the `bootstrap` prefix; see [R] bootstrap.

`fweights`, `iweights`, and `pweights` are allowed; see [U] 11.1.6 weight.

`coeflegend` does not appear in the dialog box.

See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.

### Options

**Model**

- `noconstant`; see [R] estimation options.

**Stat**

- `stat` is one of three statistics: `ate`, `atet`, or `pomeans`. `ate` is the default.
  
  - `ate` specifies that the average treatment effect be estimated.
  
  - `atet` specifies that the average treatment effect on the treated be estimated.
  
  - `pomeans` specifies that the potential-outcome means for each treatment level be estimated.

**SE/Robust**

- `vce(vcetype)` specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (`robust`), that allow for intragroup correlation (`cluster clustvar`), and that use bootstrap or jackknife methods (`bootstrap`, `jackknife`); see [R] vce_option.
Reporting
level(#) \[\text{;} \text{see \textit{[R] estimation options}.}\]

equations specifies that the results for the outcome-model or the treatment-model parameters be
displayed. By default, the results for these auxiliary parameters are not displayed.

display_options: noci, nopvalues, noomitted, vsquish, noemptycells, baselevels,
allbaselevels, nofvlabel, fvwrap(#), fvwrapon(style), cformat(\%fmt), pformat(\%fmt),
sformat(\%fmt), and nolstretch; see \textit{[R] estimation options}.\]

Maximization

maximize_options: iterate(#), \[\text{no} \log\text{, and from}\text{\textit{\init\_specs}; see \textit{[R] maximize}. These options}\]

\init\_specs is one of
matname [\text{, skip copy}]
# [\text{, # …}], copy

Advanced

\textit{\texttt{pstands}}(#\text{)} \textit{specifies the tolerance used to check the overlap assumption. The default value is \texttt{pstands}(1e-5). \texttt{teffects} will exit with an error if an observation has an estimated}
\textit{propensity score smaller than that specified by \texttt{pstands}().}\textit{\textit{\osample\texttt{(newvar)} \textit{specifies that indicator variable \texttt{newvar} be created to identify observations that}
\textit{violate the overlap assumption.}\textit{\texttt{\textit{control}(\#|\textit{label}) \textit{specifies the level of \texttt{tvar} that is the control. The default is the first treatment}
level. You may specify the numeric level \# (a nonnegative integer) or the label associated with the}
numeric level. \texttt{control()} may not be specified with statistic \texttt{pomeans}. \texttt{control()} and \texttt{tlevel()} may not specify the same treatment level.}\textit{\texttt{\textit{tlevel}(\#|\textit{label}) \textit{specifies the level of \texttt{tvar} that is the treatment for the statistic \texttt{atet}. The default is the second treatment level. You may specify the numeric level \# (a nonnegative integer) or the label associated with the numeric level. \texttt{tlevel()} may only be specified with statistic \texttt{atet}. \texttt{tlevel()} and \texttt{control()} may not specify the same treatment level.}\textit{\textit{The following option is available with \texttt{teffects ipwra} but is not shown in the dialog box:}
coeflegend; see \textit{[R] estimation options}.\textit{}}\textit{}}

Remarks and examples

Remarks are presented under the following headings:

\textit{\textit{\texttt{Overview}}} \textit{\texttt{Video example}}

Overview

\textit{\texttt{IPWRA} estimators use probability weights to obtain outcome-regression parameters that account}
for the missing-data problem arising from the fact that each subject is observed in only one of the
potential outcomes. The adjusted outcome-regression parameters are used to compute averages of
treatment-level predicted outcomes. The contrasts of these averages provide estimates of the treatment
effects.}
IPWRA estimators use a model to predict treatment status, and they use another model to predict outcomes. Because IPWRA estimators have the double-robust property, only one of the two models must be correctly specified for the IPWRA estimator to be consistent.

IPWRA estimators use a three-step approach to estimating treatment effects:

1. They estimate the parameters of the treatment model and compute inverse-probability weights.
2. Using the estimated inverse-probability weights, they fit weighted regression models of the outcome for each treatment level and obtain the treatment-specific predicted outcomes for each subject.
3. They compute the means of the treatment-specific predicted outcomes. The contrasts of these averages provide the estimates of the ATEs. By restricting the computations of the means to the subset of treated subjects, we can obtain the ATETs.

These steps produce consistent estimates of the effect parameters because the treatment is assumed to be independent of the potential outcomes after conditioning on the covariates. The overlap assumption ensures that predicted inverse-probability weights do not get too large. The standard errors reported by `teffects ipwra` correct for the three-step process. See [TE] `teffects intro` or [TE] `teffects intro advanced` for more information about this estimator.

We will illustrate the use of `teffects ipwra` by using data from a study of the effect of a mother’s smoking status during pregnancy (`mbsmoke`) on infant birthweight (`bweight`) as reported by Cattaneo (2010). This dataset also contains information about each mother’s age (`mage`), education level (`medu`), marital status (`mmarried`), whether the first prenatal exam occurred in the first trimester (`prenatal1`), and whether this baby was the mother’s first birth (`fbaby`).

### Example 1: Estimating the ATE

We begin by using `teffects ipwra` to estimate the average treatment effect of smoking on birthweight. We will use a probit model to predict treatment status as a function of `mmarried`, `mage`, and `fbaby`; to maximize the predictive power of this model, we use factor-variable notation to incorporate quadratic effects of the mother’s age, the only continuous covariate in our model. We will use linear regression (the default) to model birthweight, using `prenatal1`, `mmarried`, `mage`, and `fbaby` as explanatory variables. We type
. use http://www.stata-press.com/data/r15/cattaneo2
. teffects ipwra (bweight prenatal1 mmarried mage fbaby)
> (mbsmoke mmarried c.mage##c.mage fbaby medu, probit)
Iteration 0:  EE criterion =  9.885e-21
Iteration 1:  EE criterion =  7.847e-26
Treatment-effects estimation  Number of obs =  4,642
Estimator : IPW regression adjustment
Outcome model : linear
Treatment model: probit

| bweight     | Robust Coef.  | Std. Err.  | z     | P>|z|    | [95% Conf. Interval] |
|-------------|---------------|------------|-------|--------|----------------------|
| ATE mbsmoke (smoker vs nonsmoker) | -229.9671 | 26.62668 | -8.64 | 0.000 | -282.1544 -177.7798 |
| POmean mbsmoke nonsmoker | 3403.336 | 9.57126 | 355.58 | 0.000 | 3384.576 3422.095 |

The average birthweight if all mothers were to smoke would be 230 grams less than the average of 3,403 grams that would occur if none of the mothers had smoked.

By default, `teffects ipwra` displays the ATE and untreated POM. We can specify the `pomeans` option to display both the treated and untreated POMs, and we can use the `aequations` option to display the regression model coefficients used to predict the POMs as well as the coefficients from the model used to predict treatment.
Example 2: Displaying the POMs and equations

```
use http://www.stata-press.com/data/r15/cattaneo2
.teffects ipwra (bweight prenatal1 mmarried mage fbaby)
> (mbsmoke mmarried c.mage##c.mage fbaby medu, probit), pomeans aequations
```

Iteration 0: EE criterion = 9.885e-21
Iteration 1: EE criterion = 6.922e-26
 Treatment-effects estimation  Number of obs = 4,642
 Estimator : IPW regression adjustment
 Outcome model : linear
 Treatment model: probit

```
<table>
<thead>
<tr>
<th>bweight</th>
<th>Robust</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coef.</td>
</tr>
<tr>
<td>POmeans</td>
<td></td>
</tr>
<tr>
<td>mbsmoke</td>
<td></td>
</tr>
<tr>
<td>nonsmoker</td>
<td>3403.336</td>
</tr>
<tr>
<td>smoker</td>
<td>3173.369</td>
</tr>
<tr>
<td>OME0</td>
<td></td>
</tr>
<tr>
<td>prenatal1</td>
<td>67.98549</td>
</tr>
<tr>
<td>mmarried</td>
<td>155.5893</td>
</tr>
<tr>
<td>mage</td>
<td>2.893051</td>
</tr>
<tr>
<td>fbaby</td>
<td>-71.9215</td>
</tr>
<tr>
<td>_cons</td>
<td>3194.808</td>
</tr>
<tr>
<td>OME1</td>
<td></td>
</tr>
<tr>
<td>prenatal1</td>
<td>34.76923</td>
</tr>
<tr>
<td>mmarried</td>
<td>124.0941</td>
</tr>
<tr>
<td>mage</td>
<td>-5.068833</td>
</tr>
<tr>
<td>fbaby</td>
<td>39.89692</td>
</tr>
<tr>
<td>_cons</td>
<td>3175.551</td>
</tr>
<tr>
<td>TME1</td>
<td></td>
</tr>
<tr>
<td>mmarried</td>
<td>-.6484821</td>
</tr>
<tr>
<td>mage</td>
<td>.1744327</td>
</tr>
<tr>
<td>c.mage#</td>
<td>-.0032559</td>
</tr>
<tr>
<td>c.mage</td>
<td>-.2175962</td>
</tr>
<tr>
<td>fbaby</td>
<td>-.0863631</td>
</tr>
<tr>
<td>_cons</td>
<td>-1.558255</td>
</tr>
</tbody>
</table>
```

As is well known, the standard probit model assumes that the error terms in the latent-utility framework are homoskedastic; the model is not robust to departures from this assumption. An alternative is to use the heteroskedastic probit model, which explicitly models the error variance as a function of a set of variables.

Example 3: Heteroskedastic probit treatment model

Here we use the variables as before, but we use a heteroskedastic probit model to predict treatment status, modeling the heteroskedasticity as a quadratic function of the mother’s age:
. teffects ipwra (bweight prenatal1 mmarried fbaby c.mage) > (mbsmoke mmarried c.mage##c.mage fbaby medu, hetprobit(c.mage##c.mage)), > aequations

Iteration 0:  EE criterion =  4.443e-09
Iteration 1:  EE criterion =  4.325e-18

Treatment-effects estimation

Number of obs  =  4,642

Estimator : IPW regression adjustment
Outcome model : linear
Treatment model: heteroskedastic probit

| bweight | Coef. | Std. Err. | z     | P>|z|   | [95% Conf. Interval] |
|---------|-------|-----------|-------|-------|----------------------|
| **ATE** |       |           |       |       |                      |
| mbsmoke (smoker vs nonsmoker) | -229.6322 | 26.33452 | -8.72 | 0.000 | -281.2469 -178.0175 |
| **POmean** |       |           |       |       |                      |
| mbsmoke nonsmoker | 3403.74 | 9.545798 | 356.57 | 0.000 | 3385.03 3422.449 |
| **OME0** |       |           |       |       |                      |
| prenatal1 mmarried fbaby mage _cons | 64.95125 | 28.62159 | 2.27 | 0.023 | 8.853958 121.0485 |
| mmarried | 154.2297 | 26.45867 | 5.83 | 0.000 | 102.3717 206.0878 |
| fbaby | -71.61131 | 20.33774 | -3.52 | 0.000 | -111.4725 -31.7506 |
| mage | 3.010148 | 2.133812 | 1.41 | 0.158 | -1.172047 7.192343 |
| _cons | 3195.355 | 55.05451 | 58.04 | 0.000 | 3087.45 3303.26 |
| **OME1** |       |           |       |       |                      |
| prenatal1 mmarried fbaby mage _cons | 38.55274 | 43.57024 | -0.88 | 0.376 | -16.22735 93.33283 |
| mmarried | 126.3377 | 40.7398 | 3.10 | 0.002 | 46.48921 206.1863 |
| fbaby | -6.069913 | 5.95251 | -1.02 | 0.308 | -17.73662 5.596792 |
| mage | 3195.795 | 152.3979 | 20.97 | 0.000 | 2897.101 3394.49 |
| **TME1** |       |           |       |       |                      |
| mmarried | -.0295523 | .0238951 | -1.24 | 0.216 | -.0763857 .0172812 |
| mage | .0157893 | .0105486 | 1.50 | 0.134 | -.0048857 .0364643 |
| c.mage#c.mage | -.0002837 | .001901 | -1.49 | 0.136 | -.0006563 .0000888 |
| fbaby | -.0093306 | .0080003 | -1.17 | 0.244 | -.025011 .0063497 |
| medu | -.0036773 | .0030317 | -1.21 | 0.225 | -.0096193 .0022647 |
| _cons | -.1822201 | .1180587 | -1.54 | 0.123 | -.4136109 .0491707 |
| **TME1_lnsigma** |       |           |       |       |                      |
| mage | -.2211492 | .0631504 | -3.50 | 0.000 | -.3449217 -.0973767 |
| c.mage#c.mage | .0037613 | .0012437 | 3.02 | 0.002 | .0013236 .006199 |

The estimated ATE and base-level POM are essentially the same as those produced by the model that used a homoskedastic probit.
Video example

Treatment effects: Inverse-probability-weighted regression adjustment

Stored results

teffects ipwra stores the following in e():

Scalars

- `e(N)`: number of observations
- `e(Nj)`: number of observations for treatment level \( j \)
- `e(N_clust)`: number of clusters
- `e(k_eq)`: number of equations in `e(b)`
- `e(k_levels)`: number of levels in treatment variable
- `e(treated)`: level of treatment variable defined as treated
- `e(control)`: level of treatment variable defined as control
- `e(converged)`: 1 if converged, 0 otherwise

Macros

- `e(cmd)`: teffects
- `e(cmdline)`: command as typed
- `e(depvar)`: name of outcome variable
- `e(tvar)`: name of treatment variable
- `e(subcmd)`: ipwra
- `e(tmodel)`: logit, probit, or hetprobit
- `e(omodel)`: linear, logit, probit, hetprobit, poisson, flogit, fprobit, or fhetprobit
- `e(stat)`: statistic estimated, ate, atet, or pomeans
- `e(vtype)`: weight type
- `e(wexp)`: weight expression
- `e(title)`: title in estimation output
- `e(tlevels)`: levels of treatment variable
- `e(vce)`: vcetype specified in vce()
- `e(vcetype)`: title used to label Std. Err.
- `e(properties)`: b V
- `e(estat_cmd)`: program used to implement estat
- `e(predict)`: program used to implement predict
- `e(marginsnotok)`: predictions disallowed by margins
- `e(asbalanced)`: factor variables fvset as asbalanced
- `e(asobserved)`: factor variables fvset as asobserved

Matrices

- `e(b)`: coefficient vector
- `e(V)`: variance–covariance matrix of the estimators

Functions

- `e(sample)`: marks estimation sample

Methods and formulas

teffects ipwra implements a smooth treatment-effects estimator. All smooth treatment-effects estimators are documented in Methods and formulas of [TE] teffects aipw.

References


Also see

[TE] **teffects postestimation** — Postestimation tools for teffects

[TE] **teffects** — Treatment-effects estimation for observational data

[TE] **teffects aipw** — Augmented inverse-probability weighting

[U] **20 Estimation and postestimation commands**
Description

This entry discusses the use of `teffects` when the treatment is multivalued. This entry presumes you are already familiar with the potential-outcome framework and the use of the `teffects` commands with binary treatments. See [TE] `teffects intro` or [TE] `teffects intro advanced` for more information.

Remarks and examples

Remarks are presented under the following headings:

- Introduction
- Parameters and notation
- Illustrating multivalued treatments
- Examples

Introduction

When the treatment is binary, each subject could either receive the treatment or not receive the treatment. In contrast, multivalued treatments refer to cases in which each subject could receive one of several different treatments or else not receive treatment at all. For example, in testing the efficacy of a drug, a patient could receive a 10 milligram (mg) dose, a 20 mg dose, a 30 mg dose, or no dose at all. We first want to be able to compare a patient receiving the 10 mg dose with a patient receiving no dose, a patient receiving the 20 mg dose with a patient receiving no dose, and a patient receiving the 30 mg dose with a patient receiving no dose. Once we can make those comparisons, we can then, for example, compare the efficacy of a 30 mg dose with that of a 20 mg dose or a 10 mg dose.

To highlight an example in economics, we consider an unemployed person who could participate in a comprehensive skills training program, attend a one-day workshop that helps job seekers write their resumés, or choose not to participate in either. We want to know how effective each of those programs is relative to not participating; once we know that, we can then compare the effectiveness of the comprehensive program with that of the one-day program.

Multivalued treatments increase the number of parameters that must be estimated and complicate the notation. Fortunately, however, using the `teffects` commands is not much more difficult with multivalued treatments than with binary treatments.

You can use `teffects ra`, `teffects ipw`, `teffects ipwra`, and `teffects aipw` to estimate multivalued treatment effects. However, the theory developed in [Abadie and Imbens (2006, 2012)] has not been extended to handle multivalued treatments, so you cannot use `teffects nnmatch` or `teffects psmatch` in these cases.

Parameters and notation

We denote the potential outcome that subject \( i \) would obtain if given treatment-level \( t \) as \( y_{ti} \), where \( y_{ti} \) is the realization of the random variable \( y_t \). Throughout this entry, \( i \) subscripts denote realizations of the corresponding unsubscripted random variables. We again let \( y_0 \) denote the potential outcome of a subject who did not receive any treatment. To handle the case of multivalued treatments, we extend the definition of the unobservable, individual-level treatment effects to be \( y_t - y_0 \) for \( t \in \{1, \ldots, q\} \).

As in the binary-valued case, we again focus on three parameters of interest: the average treatment effect (ATE), the potential-outcome mean (POM), and the average treatment effect on the treated (ATET).

**ATE** The ATE is the average effect of giving each individual treatment \( t \) instead of treatment 0:

\[
\text{ATE}_t = E(y_t - y_0)
\]

**POM** The POM for each treatment level is an average of each potential outcome:

\[
\text{POM}_t = E(y_t)
\]

**ATET** The ATET is the average effect among those subjects that receive treatment level \( \tilde{t} \) of giving each subject treatment \( \tilde{t} \) instead of treatment 0:

\[
\text{ATET}_{\tilde{t}, \tilde{t}} = E \{(y_{\tilde{t}} - y_0) | t = \tilde{t}\}
\]

The extra notation required to define the ATET in this case indicates the difficulties surrounding this parameter.

Defining the ATET in the multivalued treatment case requires three different treatment levels: \( \tilde{t} \) defines the treatment level of the treated potential outcome; 0 is the treatment level of the control potential outcome; and \( t = \tilde{t} \) restricts the expectation to include only those individuals who actually receive treatment level \( \tilde{t} \).

Illustrating multivalued treatments

To illustrate the concept of a potential outcome and the parameters we would like to estimate, we consider the following table:

<table>
<thead>
<tr>
<th>( y )</th>
<th>( t )</th>
<th>( y_0 )</th>
<th>( y_1 )</th>
<th>( y_2 )</th>
</tr>
</thead>
<tbody>
<tr>
<td>( 0.50 )</td>
<td>0</td>
<td>( -0.50 )</td>
<td>1.06</td>
<td>1.93</td>
</tr>
<tr>
<td>2.42</td>
<td>1</td>
<td>2.13</td>
<td>2.42</td>
<td>2.43</td>
</tr>
<tr>
<td>3.15</td>
<td>2</td>
<td>1.26</td>
<td>2.57</td>
<td>3.15</td>
</tr>
<tr>
<td>(-0.39)</td>
<td>0</td>
<td>(-0.39)</td>
<td>(-0.18)</td>
<td>0.52</td>
</tr>
<tr>
<td>2.22</td>
<td>2</td>
<td>(-0.24)</td>
<td>(-0.01)</td>
<td>2.22</td>
</tr>
</tbody>
</table>

We observe the outcome \( y \) as well as the treatment indicator \( t \). There are three levels of treatment: 0, 1, or 2. Ideally, we would observe \( y_0 \), \( y_1 \), and \( y_2 \), but in fact all we have is \( y \). In the first row, the subject received treatment level 0, so \( y = y_0 \) for that subject. In the last row, the subject received treatment 2, so \( y = y_2 \). We reiterate that we do not actually observe \( y_0 \), \( y_1 \), or \( y_2 \).
If we did have data on $y_0$, $y_1$, and $y_2$, then we could define subject-level treatment variables $te_1 = y_1 - y_0$ and $te_2 = y_2 - y_0$. Here we would be following the convention of taking treatment level 0 to be the control level. The following table adds these two variables:

<table>
<thead>
<tr>
<th></th>
<th></th>
<th>$y_0$</th>
<th>$y_1$</th>
<th>$y_2$</th>
<th>$te_1$</th>
<th>$te_2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>-0.50</td>
<td>0</td>
<td>-0.50</td>
<td>1.06</td>
<td>1.93</td>
<td>1.56</td>
<td>2.43</td>
</tr>
<tr>
<td>2.42</td>
<td>1</td>
<td>2.13</td>
<td>2.42</td>
<td>2.43</td>
<td>0.29</td>
<td>0.30</td>
</tr>
<tr>
<td>3.15</td>
<td>2</td>
<td>1.26</td>
<td>2.57</td>
<td>3.15</td>
<td>1.31</td>
<td>1.89</td>
</tr>
<tr>
<td>-0.39</td>
<td>0</td>
<td>-0.39</td>
<td>-0.18</td>
<td>0.52</td>
<td>0.21</td>
<td>0.91</td>
</tr>
<tr>
<td>2.22</td>
<td>2</td>
<td>-0.24</td>
<td>-0.01</td>
<td>2.22</td>
<td>0.23</td>
<td>2.46</td>
</tr>
</tbody>
</table>

Once we have $te_1$ and $te_2$, obtaining the ATEs is straightforward. The ATE of going from treatment 0 to treatment 1 is simply the mean of the five entries in the column labeled $te_1$, which here works out to 0.72. Going from treatment level 0 to treatment level 1 causes the outcome to increase an average of 0.72. Similarly, the ATE of going from treatment 0 to treatment 2 is the mean of the entries in the column labeled $te_2$, which is 1.60. Exposing all subjects to treatment level 2 would cause the outcome to rise by an average of 1.60 relative to the outcome obtained by exposing them to treatment level 0.

The ATET is the average difference in the potential outcomes among those that get a particular treatment level. To compute this, we must specify two treatment levels: the actual treatment level the subjects we are interested in received as well as the treatment level we want to compare them with. For example, suppose we are interested in the ATET of going from treatment 0 to treatment 1 for those who received treatment 0. This ATET is the average of $te_1$ for those subjects for which $t = 0$. Here that ATET is just $(1.56 + 0.21)/2 \approx 0.89$. If we exposed the subjects who received treatment 0 to treatment 1 instead, the outcome would increase an average of 0.89.

The ATET of going from treatment 0 to treatment 2 for those subjects who received treatment 2 is the mean of $te_2$ for those subjects for which $t = 2$, which is $(1.89 + 2.46)/2 \approx 2.18$. Receiving treatment 2 increased the outcome of those who received treatment 2 by an average of 2.18 relative to receiving the control.

**Examples**

In the remainder of this entry, we provide several examples demonstrating how to estimate multivalued treatments using `teffects`.

**Example 1: Potential outcomes with four treatment levels**

*bdsianesi5.dta* contains an extract of data from Blundell, Dearden, and Sianesi (2005). In this dataset on individuals in the United Kingdom, *wages* records hourly wages in pounds; *ed* records the highest educational degree obtained; *paed* records the highest educational level obtained by each individual’s father; *math7* records a score obtained on a standardized math test when the individual was seven; *read7* records a score obtained on a standardized reading test when the individual was seven; and *london* and *eastern* are indicators for whether an individual lives in the expensive area of London or the east. We want to know how the level of education obtained affects a person’s wage.

We begin by using `mean` to report the estimated means of wages over the four education levels. The value labels on `mean` are coded as `none` for no degree, `O` for an O-level degree, `A` for an A-level degree, or `H` for a higher-education degree.
. use http://www.stata-press.com/data/r15/bdsianesi5
(Excerpt from Blundell, Dearden, & Sianesi (2005) JRSSA 168: 473)
. mean wage, over(ed)

<table>
<thead>
<tr>
<th>Over</th>
<th>Mean</th>
<th>Std. Err.</th>
<th>[95% Conf. Interval]</th>
</tr>
</thead>
<tbody>
<tr>
<td>none</td>
<td>6.057816</td>
<td>.154332</td>
<td>5.755114 - 6.360518</td>
</tr>
<tr>
<td>O</td>
<td>7.501648</td>
<td>.1807359</td>
<td>7.147158 - 7.856137</td>
</tr>
<tr>
<td>A</td>
<td>8.220637</td>
<td>.1540359</td>
<td>7.918516 - 8.522758</td>
</tr>
<tr>
<td>H</td>
<td>10.87703</td>
<td>.2257888</td>
<td>10.43417 - 11.31988</td>
</tr>
</tbody>
</table>

The output reveals that the estimated mean wage increases as the education level goes from no degree to an O-level degree, to an A-level degree, and to a higher-education degree, as we would expect. Once we control for other characteristics of each individual, do we still observe a positive effect of education on wage?

We use `teffects ra` (see [TE] `teffects ra`) to estimate the ATEs of the different education levels by regression adjustment (RA), controlling for each person’s location, math score, and father’s education level:

. teffects ra (wage london eastern paed math7, poisson) (ed)
Iteration 0: EE criterion = 1.865e-18
Iteration 1: EE criterion = 4.077e-30

| Treatment | Robust Coef. | Std. Err. | z   | P>|z| | [95% Conf. Interval] |
|-----------|--------------|-----------|-----|------|----------------------|
| ATE ed    |              |           |     |      |                      |
| (O vs none) | 1.181543 | .3520371  | 3.36| 0.001| .4915626 - 1.871523  |
| (A vs none) | 1.743079 | .3282152  | 5.31| 0.000| 1.099789 - 2.386369  |
| (H vs none) | 3.972829 | .3840024  | 10.35| 0.000| 3.220199 - 4.72546   |

POmean ed

| Robust Coef. | Std. Err. | z   | P>|z| | [95% Conf. Interval] |
|--------------|-----------|-----|------|----------------------|
| none         | 6.525873  | .2931933 | 22.26| 0.000| 5.951224 - 7.100521  |

Because wages are necessarily positive, we used the `poisson` option inside the outcome-model specification. The estimated POM of the control level of no degree is 6.53 pounds per hour. The estimated ATE of going from no degree to an O-level degree is 1.18 pounds per hour; the estimated ATE of going from no degree to an A-level degree is 1.74 pounds per hour; and the estimated ATE of going from no degree to a higher-education degree is 3.97 pounds per hour. All of these effects are highly significant.

For comparison purposes, we also use `teffects aipw` (see [TE] `teffects aipw`). We use the same outcome model as before. We use a multinomial logit model to predict education level, using math and reading scores and both the father’s and the mother’s educational attainment levels as predictors:
. teffects aipw (wage london eastern paed math7, poisson)
> (ed math7 read7 maed paed)

Iteration 0: EE criterion = 1.877e-18
Iteration 1: EE criterion = 1.029e-30

Treatment-effects estimation

Estimator : augmented IPW
Outcome model : Poisson by ML
Treatment model: (multinomial) logit

| wage | Coef. | Std. Err. | z | P>|z| | [95% Conf. Interval] |
|------|-------|-----------|---|-----|----------------------|
| ATE  | ed    |           |   |     |                      |
| (O vs none) | 1.748197 | .3911167 | 4.47 | 0.000 | .9816221 | 2.514771 |
| (A vs none) | 2.363228 | .3741584 | 6.32 | 0.000 | 1.629891 | 3.096565 |
| (H vs none) | 4.359777 | .4133059 | 10.55 | 0.000 | 3.549712 | 5.169842 |
| POmean | ed    |           |   |     |                      |
| none | 5.946184 | .3391531 | 17.53 | 0.000 | 5.281456 | 6.610912 |

The results indicate slightly higher treatment effects relative to those indicated by \texttt{teffects ra}. That is largely because the AIPW estimator predicts a lower no-higher-education POM than the RA estimator.

Example 2: Expressing ATEs as percentages

As in the binary-treatment case, expressing the ATEs as percentages of the POM for the control level often aids interpretation. Here we first use the replay facility of \texttt{teffects aipw} along with the \texttt{coeflegend} option to see how the parameters are named.

. teffects, coeflegend

<table>
<thead>
<tr>
<th>wage</th>
<th>Coef.</th>
<th>Legend</th>
</tr>
</thead>
<tbody>
<tr>
<td>ATE</td>
<td>ed</td>
<td></td>
</tr>
<tr>
<td>(O vs none)</td>
<td>1.748197</td>
<td>_b[ATE:r1vs0.ed]</td>
</tr>
<tr>
<td>(A vs none)</td>
<td>2.363228</td>
<td>_b[ATE:r2vs0.ed]</td>
</tr>
<tr>
<td>(H vs none)</td>
<td>4.359777</td>
<td>_b[ATE:r3vs0.ed]</td>
</tr>
<tr>
<td>POmean</td>
<td>ed</td>
<td></td>
</tr>
<tr>
<td>none</td>
<td>5.946184</td>
<td>_b[POmean:0.ed]</td>
</tr>
</tbody>
</table>
Now that we know the names, we can use `nlcom` to obtain the ATEs relative to the base-level POM:

```
. nlcom (_b[ATE:r1vs0.ed] / _b[POmean:0.ed])
  > (_b[ATE:r2vs0.ed] / _b[POmean:0.ed])
  > (_b[ATE:r3vs0.ed] / _b[POmean:0.ed])
```

|      | Coef. Std. Err. | z    | P>|z| [95% Conf. Interval] |
|------|----------------|------|------|------------------------|
| _nl_1 | .2940031       | .0808156 | 3.64 | 0.000 [.1356075 .4523988] |
| _nl_2 | .3974361       | .0840545 | 4.73 | 0.000 [.2326923 .5621799] |
| _nl_3 | .7332059       | .1068848 | 6.86 | 0.000 [.5237156 .9426962] |

Wages are 29% higher when everyone receives an O-level degree than if no one receives a degree. Wages are 40% higher when everyone receives an A-level degree than if no one receives a degree. Wages are 73% higher when everyone receives an H-level degree than if no one receives a degree.

Although impressive, these changes are not presented in the way that is most commonly discussed. (There is a large amount of literature on the treatment effect of getting a higher-education degree.) In particular, we might rather want to know the percentage changes in wages relative to a person with an A-level degree. Next we estimate the ATEs treating an A-level degree as the control level; to do that, we use the `control()` option. We also specify `coeflegend` again because we are more interested in how the parameters are named rather than in their standard errors at this point:

```
. teffects aipw (wage london eastern paed math7, poisson)
  > (ed math7 read7 maed paed), control(A) coeflegend
```

```
Iteration 0: EE criterion = 1.870e-18
Iteration 1: EE criterion = 2.882e-30
```

```
Treatment-effects estimation Number of obs = 1,693
Estimator : augmented IPW
Outcome model : Poisson by ML
Treatment model: (multinomial) logit
```

```
<table>
<thead>
<tr>
<th></th>
<th>Coef. Legend</th>
</tr>
</thead>
<tbody>
<tr>
<td>wage</td>
<td></td>
</tr>
<tr>
<td>ATE</td>
<td></td>
</tr>
<tr>
<td>ed</td>
<td></td>
</tr>
<tr>
<td>(none vs A)</td>
<td>-2.363228 _b[ATE:r0vs2.ed]</td>
</tr>
<tr>
<td>(O vs A)</td>
<td>-.6150312 _b[ATE:r1vs2.ed]</td>
</tr>
<tr>
<td>(H vs A)</td>
<td>1.996549 _b[ATE:r3vs2.ed]</td>
</tr>
<tr>
<td>POmean</td>
<td></td>
</tr>
<tr>
<td>ed</td>
<td></td>
</tr>
<tr>
<td>A</td>
<td>8.309412 _b[POmean:2.ed]</td>
</tr>
</tbody>
</table>
```

Now we use `nlcom` to obtain the ATE of obtaining a higher-education degree as a percentage of the expected A-level wage:

```
. nlcom _b[ATE:r3vs2.ed] / _b[POmean:2.ed], noheader
```

|      | Coef. Std. Err. | z    | P>|z| [95% Conf. Interval] |
|------|----------------|------|------|------------------------|
| _nl_1 | .2402756       | .0355404 | 6.76 | 0.000 [.1706177 .3099335] |
The average wage increases by 24% when everyone receives an H-level degree relative to when everyone receives an A-level degree.

Example 3: Obtaining ATETs

In the previous example, we showed that on average, a higher-education degree increases a person’s wage by 24% relative to someone with only an A-level degree. Sometimes, though, we would rather know how much the higher-education degree increases wages among the people who actually have a higher-education degree. To answer that question, we want to examine the ATET rather than the ATE.

Here we use the IPWRA estimator to obtain our answer. We specify the control(A) option so that an A-level education is treated as the basis for comparisons. We specify the atet option to obtain ATETs rather than ATEs, and we specify the tlevel(H) option to indicate that we want the ATETs to be calculated for the subset of people who actually receive higher-education degrees.

```
. teffects ipwra (wage london eastern paed math7, poisson)>
  > (ed math7 read7 maed paed), atet control(A) tlevel(H)
```

```
Iteration 0:  EE criterion = 2.731e-18
Iteration 1:  EE criterion = 5.636e-31
```

```
Treatment-effects estimation
Number of obs = 1,693
Estimator : IPW regression adjustment
Outcome model : Poisson
Treatment model: (multinomial) logit

Robust

|      | Coef.  | Std. Err. |    z   |   P>|z|  | [95% Conf. Interval] |
|------|--------|-----------|--------|--------|----------------------|
| wage |        |           |        |        |                      |
| ATET | ed     |           |        |        |                      |
| (none vs A) | -2.87423 |  .361093  | -7.96  | 0.000  | -3.58196 to -2.166501 |
| (O vs A)   | -.8246604 |  .3609131 | -2.28  | 0.022  | -1.532037 to -.1172837 |
| (H vs A)   | 1.866757  |  .3277701 |  5.70  | 0.000  | 1.224339 to 2.509174  |
| POmean ed  | 9.010271  |  .2503971 | 35.98  | 0.000  | 8.519501 to 9.50104   |
| A          |         |           |        |        |                      |

The point estimates are similar to the ATEs we obtained above, suggesting that the means of the covariates among those with a higher-education degree are similar to the means for the entire population.

In output not shown to save space, we replayed the previous results with the coeflegend option to determine how the parameters are named. Armed with that information, we call nlcom:

```
. nlcom _b[ATET:r3vs2.ed] / _b[POmean:2.ed], noheader
```

```
|      | Coef.  | Std. Err. |    z   |   P>|z|  | [95% Conf. Interval] |
|------|--------|-----------|--------|--------|----------------------|
| _nl_1 |  .207181 |  .0407528 |  5.08  | 0.000  |  .127307 to  .287055  |
```

Our estimate of the percentage increase is now noticeably smaller once we restrict ourselves to only those people who actually received a higher-education degree. However, because of the width of the confidence intervals, there is no evidence to suggest that the difference between the estimates is statistically significant.
Example 4: ATEs comparing adjacent treatments

In the first example, we obtained the three ATEs, and they were all expressed relative to the base level of no degree. Now we show how we can express the gains to an O-level degree relative to no degree, the gains to an A-level degree relative to an O-level degree, and the gains to a higher-education degree relative to an A-level degree.

First, we use an AIPW estimator to obtain all the POMs for our example dataset:

```
. teffects aipw (wage london eastern paed math7, poisson)
> (ed math7 read7 maed paed), pom
Iteration 0: EE criterion = 1.877e-18
Iteration 1: EE criterion = 1.542e-30
```

| POmeans | Robust Coef. | Std. Err. | z | P>|z| | [95% Conf. Interval] |
|---------|--------------|-----------|---|-----|-----------------------|
|         | wage         |           |   |     |                       |
| none    | 5.946184     | .3391531  | 17.53 | 0.000 | 5.281456 | 6.610912 |
| O       | 7.694381     | .1915192  | 40.18 | 0.000 | 7.31901  | 8.069752 |
| A       | 8.309412     | .1563348  | 53.15 | 0.000 | 8.003001 | 8.615823 |
| H       | 10.30596     | .2285837  | 45.09 | 0.000 | 9.857945 | 10.75398 |

ATEs are contrasts of POMs, and here we show how to use contrast to obtain the estimated ATEs:

```
. contrast r.ed, nowald
```

Now that we know how to use contrast to obtain the ATEs based on the POMs, we can take advantage of contrast's ability to obtain “reverse adjacent” contrasts, which compare each level with the previous level. We use the ar. operator with contrast to accomplish this:

```
. contrast r.ar.ed, nowald
```
teffects multivalued — Multivalued treatment effects

. contrast ar.ed, nowald
Warning: cannot perform check for estimable functions.
Contrasts of marginal linear predictions
Margins : asbalanced

<table>
<thead>
<tr>
<th>Contrast</th>
<th>Std. Err.</th>
<th>[95% Conf. Interval]</th>
</tr>
</thead>
<tbody>
<tr>
<td>ed (O vs none)</td>
<td>1.748197</td>
<td>.3911167 .9816221 2.514771</td>
</tr>
<tr>
<td>ed (A vs O)</td>
<td>.6150312</td>
<td>.2432806 .13821 1.091852</td>
</tr>
<tr>
<td>ed (H vs A)</td>
<td>1.996549</td>
<td>.2730712 1.461339 2.531759</td>
</tr>
</tbody>
</table>

These ATEs are for incremental increases. In contrast, the ATEs considered above had a common base.

Technical note

The multivalued treatment AIPW estimators implemented in teffects aipw are EIF estimators based on the results of Cattaneo (2010). The results in Cattaneo (2010) are for semiparametric estimators, and we implement parametric versions. Of more practical importance, Cattaneo (2010) contains results for quantile treatment effects that are not implemented in teffects but implemented in the community-contributed poparms command discussed in Cattaneo, Drukker, and Holland (2013). See Emsley et al. (2008) for another implementation of the AIPW estimator, and see Frölich and Melly (2010) for other commands that estimate quantile treatment effects.

References


Also see

[TE] teffects — Treatment-effects estimation for observational data

[TE] teffects intro — Introduction to treatment effects for observational data

[TE] teffects intro advanced — Advanced introduction to treatment effects for observational data
Description

teffects nnmatch estimates the average treatment effect (ATE) and average treatment effect on the treated (ATET) from observational data by nearest-neighbor matching (NNM). NNM estimators impute the missing potential outcome for each subject by using an average of the outcomes of similar subjects that receive the other treatment level. Similarity between subjects is based on a weighted function of the covariates for each observation. The treatment effect is computed by taking the average of the difference between the observed and imputed potential outcomes for each subject. teffects nnmatch accepts a continuous, binary, count, fractional, or nonnegative outcome.

See [TE] teffects intro or [TE] teffects intro advanced for more information about estimating treatment effects from observational data.

Quick start

ATE of treat on y estimated by NNM on x1 and indicators for levels of categorical variable a

teffects nnmatch (y x1 i.a) (treat)

As above, but estimate the ATET

teffects nnmatch (y x1 i.a) (treat), atet

Add continuous covariate x2 and perform bias correction

teffects nnmatch (y x1 x2 i.a) (treat), biasadj(x1 x2)

As above, and match exactly on values of a

teffects nnmatch (y x1 x2 i.a) (treat), biasadj(x1 x2) ematch(i.a)

With robust standard errors

teffects nnmatch (y x1 x2 i.a) (treat), vce(robust)

With four matches per observation

teffects nnmatch (y x1 x2 i.a) (treat), nneighbor(4)

Menu

Statistics > Treatment effects > Continuous outcomes > Nearest-neighbor matching
Statistics > Treatment effects > Binary outcomes > Nearest-neighbor matching
Statistics > Treatment effects > Count outcomes > Nearest-neighbor matching
Statistics > Treatment effects > Fractional outcomes > Nearest-neighbor matching
Statistics > Treatment effects > Nonnegative outcomes > Nearest-neighbor matching
**Syntax**

```plaintext
teffects nnmatch (ovar omvarlist) (tvar) [if] [in] [weight]
    [ , stat options ]
```

`ovar` is a binary, count, continuous, fractional, or nonnegative outcome of interest.

`omvarlist` specifies the covariates in the outcome model.

`tvar` must contain integer values representing the treatment levels. Only two treatment levels are allowed.

**stat**

<table>
<thead>
<tr>
<th>Stat</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>ate</td>
<td>estimate average treatment effect in population; the default</td>
</tr>
<tr>
<td>atet</td>
<td>estimate average treatment effect on the treated</td>
</tr>
</tbody>
</table>

**options**

<table>
<thead>
<tr>
<th>Model</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>nneighbor(#)</code></td>
<td>specify number of matches per observation; default is <code>nneighbor(1)</code></td>
</tr>
<tr>
<td><code>biasadj(varlist)</code></td>
<td>correct for large-sample bias using specified variables</td>
</tr>
<tr>
<td><code>ematch(varlist)</code></td>
<td>match exactly on specified variables</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>SE/Robust</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>vce(vcetype)</code></td>
<td><code>vcetype</code> may be</td>
</tr>
<tr>
<td><code>vce(robust [ , nn(#)]);</code></td>
<td>use robust Abadie–Imbens standard errors with # matches</td>
</tr>
<tr>
<td><code>vce(iid);</code></td>
<td>use default Abadie–Imbens standard errors</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Reporting</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>level(#)</code></td>
<td>set confidence level; default is <code>level(95)</code></td>
</tr>
<tr>
<td><code>dmvariables</code></td>
<td>display names of matching variables</td>
</tr>
<tr>
<td><code>display_options</code></td>
<td>control columns and column formats, row spacing, line width, display of omitted variables and base and empty cells, and factor-variable labeling</td>
</tr>
</tbody>
</table>

**Advanced**

<table>
<thead>
<tr>
<th>Advanced</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>caliper(#)</code></td>
<td>specify the maximum distance for which two observations are potential neighbors</td>
</tr>
<tr>
<td><code>dtolerance(#)</code></td>
<td>set maximum distance between individuals considered equal</td>
</tr>
<tr>
<td><code>osample(newvar)</code></td>
<td><code>newvar</code> identifies observations that violate the overlap assumption</td>
</tr>
<tr>
<td>`control(#</td>
<td>label)`</td>
</tr>
<tr>
<td>`tlevel(#</td>
<td>label)`</td>
</tr>
<tr>
<td><code>generate(stub)</code></td>
<td>generate variables containing the observation numbers of the nearest neighbors</td>
</tr>
<tr>
<td><code>metric(metric)</code></td>
<td>select distance metric for covariates</td>
</tr>
<tr>
<td><code>coeflegend</code></td>
<td>display legend instead of statistics</td>
</tr>
</tbody>
</table>
**metric** | Description
---|---
mahalanobis | inverse sample covariate covariance; the default
ivariance | inverse diagonal sample covariate covariance
euclidean | identity
matrix matname | user-supplied scaling matrix

*omvarlist* may contain factor variables; see [U] 11.4.3 Factor variables.

by and stataby are allowed; see [U] 11.1.10 Prefix commands.

fweights are allowed; see [U] 11.1.6 weight.

coeflegend does not appear in the dialog box.

See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.

### Options

**Model**

**nneighbor(#)** specifies the number of matches per observation. The default is **nneighbor(1)**. Each observation is matched with at least the specified number of observations from the other treatment level. **nneighbor()** must specify an integer greater than or equal to 1 but no larger than the number of observations in the smallest treatment group.

**biasadj(varlist)** specifies that a linear function of the specified covariates be used to correct for a large-sample bias that exists when matching on more than one continuous covariate. By default, no correction is performed.

Abadie and Imbens (2006, 2011) show that nearest-neighbor matching estimators are not consistent when matching on two or more continuous covariates and propose a bias-corrected estimator that is consistent. The correction term uses a linear function of variables specified in **biasadj()**; see example 3.

**ematch(varlist)** specifies that the variables in **varlist** match exactly. All variables in **varlist** must be numeric and may be specified as factors. **teffects nnmatch** exits with an error if any observations do not have the requested exact match.

**Stat**

**stat** is one of two statistics: **ate** or **atet**. **ate** is the default.

**ate** specifies that the average treatment effect be estimated.

**atet** specifies that the average treatment effect on the treated be estimated.

**SE/Robust**

**vce(vcetype)** specifies the standard errors that are reported. By default, **teffects nnmatch** uses two matches in estimating the robust standard errors.

**vce(robust [ , nn(#)]** specifies that robust standard errors be reported and that the requested number of matches be used optionally.

**vce(iid)** specifies that standard errors for independently and identically distributed data be reported.

The standard derivative-based standard-error estimators cannot be used by **teffects nnmatch**, because these matching estimators are not differentiable. The implemented methods were derived by Abadie and Imbens (2006, 2011, 2012); see Methods and formulas.
As discussed in Abadie and Imbens (2008), bootstrap estimators do not provide reliable standard errors for the estimator implemented by `teffects nnmatch`.

Reporting

```plaintext
level(#) ; see [R] estimation options.
```

dmvariables specifies that the matching variables be displayed.

```plaintext
display_options: noci, nopvalues, noomitted, vsquish, noemptycells, baselevels,
allbaselevels, nofvlabel, fvwrap(#), fvwrapon(style), cformat(%fmt), pformat(%fmt),
sformat(%fmt), and nolstretch ; see [R] estimation options.
```

Advanced

```plaintext
caliper(#) specifies the maximum distance at which two observations are a potential match. By default, all observations are potential matches regardless of how dissimilar they are.

The distance is based on `omvlist`. If an observation does not have at least nneighbor(#) matches, `teffects nnmatch` exits with an error message. Use option osample(newvar) to identify all observations that are deficient in matches.

dtolerance(#) specifies the tolerance used to determine exact matches. The default value is dtolerance(sqrt(c(epsdouble))).

Integer-valued variables are usually used for exact matching. The dtolerance() option is useful when continuous variables are used for exact matching.

osample(newvar) specifies that indicator variable newvar be created to identify observations that violate the overlap assumption. This variable will identify all observations that do not have at least nneighbor(#) matches in the opposite treatment group within caliper(#) (for metric() distance matching) or dtolerance(#) (for ematch(varlist) exact matches).

The vce(robust, nn(#)) option also requires at least # matches in the same treatment group within the distance specified by caliper(#) or within the exact matches specified by dtolerance(#).

The average treatment effect on the treated, option atet, using vce(iid) requires only nneighbor(#) control group matches for the treated group.

control(# | label) specifies the level of tvar that is the control. The default is the first treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. control() and tlevel() may not specify the same treatment level.

tlevel(# | label) specifies the level of tvar that is the treatment for the statistic atet. The default is the second treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. tlevel() may only be specified with statistic atet. tlevel() and control() may not specify the same treatment level.

generate(stub) specifies that the observation numbers of the nearest neighbors be stored in the new variables stub1, stub2, ... . This option is required if you wish to perform postestimation based on the matching results. The number of variables generated may be more than nneighbors(#) because of tied distances. These variables may not already exist.

metric(metric) specifies the distance matrix used as the weight matrix in a quadratic form that transforms the multiple distances into a single distance measure; see Nearest-neighbor matching estimator in Methods and formulas for details.

The following option is available with `teffects nnmatch` but is not shown in the dialog box:

coflegend; see [R] estimation options.
Remarks and examples

The NNM method of treatment-effect estimation imputes the missing potential outcome for each individual by using an average of the outcomes of similar subjects that receive the other treatment level. Similarity between subjects is based on a weighted function of the covariates for each observation. The average treatment effect (ATE) is computed by taking the average of the difference between the observed and potential outcomes for each subject.

t/effects nnmatch determines the “nearest” by using a weighted function of the covariates for each observation. By default, the Mahalanobis distance is used, in which the weights are based on the inverse of the covariates’ variance-covariance matrix. t/effects nnmatch also allows you to request exact matching for categorical covariates. For example, you may want to force all matches to be of the same gender or race.

NNM is nonparametric in that no explicit functional form for either the outcome model or the treatment model is specified. This flexibility comes at a price; the estimator needs more data to get to the true value than an estimator that imposes a functional form. More formally, the NNM estimator converges to the true value at a rate slower than the parametric rate, which is the square root of the sample size, when matching on more than one continuous covariate. t/effects nnmatch uses bias correction to fix this problem. t/effects psmatch implements an alternative to bias correction; the method matches on a single continuous covariate, the estimated treatment probabilities. See [TE] t/effects intro or [TE] t/effects intro advanced for more information about this estimator.

We will illustrate the use of t/effects nnmatch by using data from a study of the effect of a mother’s smoking status during pregnancy (mbsmoke) on infant birthweight (bweight) as reported by Cattaneo (2010). This dataset also contains information about each mother’s age (mage), education level (medu), marital status (mmarried), whether the first prenatal exam occurred in the first trimester (prenatal1), whether this baby was the mother’s first birth (fbaby), and the father’s age (fage).

Example 1: Estimating the ATE

We begin by using t/effects nnmatch to estimate the average treatment effect of mbsmoke on bweight. Subjects are matched using the Mahalanobis distance defined by covariates mage, prenatal1, mmarried, and fbaby.

```
. use http://www.stata-press.com/data/r15/cattaneo3
. teffects nnmatch (bweight mage prenatal1 mmarried fbaby) (mbsmoke)
```

<table>
<thead>
<tr>
<th>Treatment-effects estimation</th>
<th>Number of obs  =  4,642</th>
</tr>
</thead>
<tbody>
<tr>
<td>Estimator</td>
<td>nearest-neighbor matching</td>
</tr>
<tr>
<td>Matches: requested =</td>
<td>1</td>
</tr>
<tr>
<td>Outcome model</td>
<td>matching</td>
</tr>
<tr>
<td>Distance metric: Mahalanobis</td>
<td>max = 139</td>
</tr>
</tbody>
</table>

| bweight | AI Robust Coef.  | Std. Err.  | z   | P>|z|   | [95% Conf. Interval] |
|---------|------------------|------------|-----|--------|---------------------|
| ATE     |                  |            |     |        |                     |
| mbsmoke | (smoker vs       | -240.3306  | 28.43006 | -8.45 | 0.000  | -296.0525 | -184.6087 |
|         | nonsmoker)       |            |     |        |                     |

The average birthweight if all mothers were to smoke would be 240 grams less than the average that would occur if none of the mothers had smoked.
When the model includes indicator and categorical variables, you may want to restrict matches to only those subjects who are in the same category. The `ematch()` option of \texttt{teffects nnmatch} allows you to specify such variables that must match exactly.

\textbf{Example 2: Exact matching}

Here we use the `ematch()` option to require exact matches on the binary variables `prenatal1`, `mmarried`, and `fbaby`. We also use Euclidean distance, rather than the default Mahalanobis distance, to match on the continuous variable `mage`, which uses Euclidean distance.

\begin{verbatim}
. teffects nnmatch (bweight mage) (mbsmoke),
    > ematch(prenatal1 mmarried fbaby) metric(euclidean)
\end{verbatim}

\begin{verbatim}
Treatment-effects estimation
 Number of obs = 4,642
Estimator : nearest-neighbor matching
 Matches: requested = 1
Outcome model : matching
 min = 1
Distance metric: Euclidean
 max = 139
\end{verbatim}

\begin{verbatim}
AI Robust
 bweight       Coef.  Std. Err.     z    P>|z|     [95% Conf. Interval]
ATE
mbsmoke
(smoker vs nonsmoker)
 -240.3306  28.43006  -8.45  0.000  -296.0525  -184.6087
\end{verbatim}

\textit{Abadie and Imbens} (2006, 2011) have shown that nearest-neighbor matching estimators are not consistent when matching on two or more continuous covariates. A bias-corrected estimator that uses a linear function of variables can be specified with `biasadj()`.

\textbf{Example 3: Bias adjustment}

Here we match on two continuous variables, `mage` and `fage`, and we use the bias-adjusted estimator:

\begin{verbatim}
. teffects nnmatch (bweight mage fage) (mbsmoke),
    > ematch(prenatal1 mmarried fbaby) biasadj(mage fage)
\end{verbatim}

\begin{verbatim}
Treatment-effects estimation
 Number of obs = 4,642
Estimator : nearest-neighbor matching
 Matches: requested = 1
Outcome model : matching
 min = 1
Distance metric: Mahalanobis
 max = 25
\end{verbatim}

\begin{verbatim}
AI Robust
 bweight       Coef.  Std. Err.     z    P>|z|     [95% Conf. Interval]
ATE
mbsmoke
(smoker vs nonsmoker)
 -223.8389  26.19973  -8.54  0.000  -275.1894  -172.4883
\end{verbatim}

These results are similar to those reported in example 1.
Example 4: NNM can reduce to RA

NNM reduces to RA when matching exactly and all the covariates are discrete. We begin our illustration of this point by estimating the ATE by NNM using exact matching on \texttt{mmarried} and the mother’s age-categories \texttt{magecat}.

```
. teffects nnmatch (bweight) (mbsmoke), ematch(i.mmarried i.magecat)
```

```
Treatment-effects estimation
Estimator : nearest-neighbor matching
Outcome model : matching
Distance metric: Mahalanobis

|                  | Coef. | Std. Err. | z     | P>|z|  | [95% Conf. Interval] |
|------------------|-------|-----------|-------|------|---------------------|
| bweight          | -241.5264 | 24.39661  | -9.90 | 0.000 | -289.3429 to -193.71 |
| ATE              |        |           |       |      |                     |
| mbsmoke          |       |           |       |      |                     |
| smoker vs nonsmoker | -241.5264 | 24.39661  | -9.90 | 0.000 | -289.3429 to -193.71 |
```

The RA estimator that includes the interactions among the discrete covariates produces the same point estimate.

```
. teffects ra (bweight i.mmarried##i.magecat) (mbsmoke)
```

```
Iteration 0: EE criterion = 1.523e-23
Iteration 1: EE criterion = 7.899e-26
Treatment-effects estimation
Estimator : regression adjustment
Outcome model : linear
Treatment model: none

|                  | Coef. | Std. Err. | z     | P>|z|  | [95% Conf. Interval] |
|------------------|-------|-----------|-------|------|---------------------|
| bweight          | -241.5264 | 24.39661  | -9.90 | 0.000 | -289.3429 to -193.71 |
| ATE              |        |           |       |      |                     |
| mbsmoke          |       |           |       |      |                     |
| smoker vs nonsmoker | -241.5264 | 24.39661  | -9.90 | 0.000 | -289.0797 to -193.9732 |
```

The two estimates of the ATE are the same. The standard errors differ in finite samples because the RA and NNM estimators use different robust estimators of the variance of the estimator.

With exact matching on discrete covariates, the NNM estimator reduces to an average of differences in cell means. With fully interacted discrete covariates, the RA estimator reduces to the same average of difference in cell means.
**Video example**

Treatment effects in Stata: Nearest-neighbor matching

**Stored results**

t/effects nnmatch stores the following in e():

Scalars
- `e(N)` number of observations
- `e(nj)` number of observations for treatment level j
- `e(k_levels)` number of levels in treatment variable
- `e(treated)` level of treatment variable defined as treated
- `e(control)` level of treatment variable defined as control
- `e(k_neighbor)` requested number of matches
- `e(k_min)` minimum number of matches
- `e(k_max)` maximum number of matches
- `e(k_robust)` matches for robust VCE

Macros
- `e(cmd)` `teffects`
- `e(cmdline)` command as typed
- `e(depvar)` name of outcome variable
- `e(tvar)` name of treatment variable
- `e(emvarlist)` exact match variables
- `e(bavarlist)` variables used in bias adjustment
- `e(mvarlist)` match variables
- `e(subcmd)` `nnmatch`
- `e(metric)` `mahalanobis, ivariance, euclidean`, or `matrix matname`
- `e(stat)` statistic estimated, `ate` or `atet`
- `e(wtype)` weight type
- `e(wexp)` weight expression
- `e(title)` title in estimation output
- `e(tlevels)` levels of treatment variable
- `e(vce)` `vcetype` specified in `vce()`
- `e(vcetype)` title used to label Std. Err.
- `e(datasignature)` the checksum
- `e(datasignaturevars)` variables used in calculation of checksum
- `e(properties)` `b V`
- `e(estat_cmd)` program used to implement `estat`
- `e(predict)` program used to implement `predict`
- `e(marginsnotok)` predictions disallowed by `margins`

Matrices
- `e(b)` coefficient vector
- `e(V)` variance–covariance matrix of the estimators

Functions
- `e(sample)` marks estimation sample

**Methods and formulas**

The methods and formulas presented here provide the technical details underlying the estimators implemented in `teffects nnmatch` and `teffects psmatch`. See Methods and formulas of [TE] teffects aipw for the methods and formulas used by `teffects aipw`, `teffects ipw`, `teffects ipwra`, and `teffects ra`.

Methods and formulas are presented under the following headings:

- Nearest-neighbor matching estimator
- Bias-corrected matching estimator
- Propensity-score matching estimator
- PSM, ATE, and ATET variance adjustment
Nearest-neighbor matching estimator

`teffects nnmatch` implements the nearest-neighbor matching (NNM) estimator for the average treatment effect (ATE) and the average treatment effect on the treated (ATET). This estimator was derived by Abadie and Imbens (2006, 2011) and was previously implemented in Stata as discussed in Abadie et al. (2004).

`teffects psmatch` implements nearest-neighbor matching on an estimated propensity score. A propensity score is a conditional probability of treatment. The standard errors implemented in `teffects psmatch` were derived by Abadie and Imbens (2012).

`teffects nnmatch` and `teffects psmatch` permit two treatment levels: the treatment group with $t = 1$ and a control group with $t = 0$.

Matching estimators are based on the potential-outcome model, in which each individual has a well-defined outcome for each treatment level; see [TE] `teffects intro`. In the binary-treatment potential-outcome model, $y_{1i}$ is the potential outcome obtained by an individual if given treatment-level 1 and $y_{0i}$ is the potential outcome obtained by each individual $i$ if given treatment-level 0. The problem posed by the potential-outcome model is that only $y_{1i}$ or $y_{0i}$ is observed, never both. $y_{0i}$ and $y_{1i}$ are realizations of the random variables $y_0$ and $y_1$. Throughout this document, $i$ subscripts denote realizations of the corresponding, unsubscripted random variables.

Formally, the ATE is

$$\tau_1 = E(y_1 - y_0)$$

and the ATET is

$$\delta_1 = E(y_1 - y_0 | t = 1)$$

These expressions imply that we must have some solution to the missing-data problem that arises because we only observe either $y_{1i}$ or $y_{0i}$, not both.

For each individual, NNM uses an average of the individuals that are most similar, but get the other treatment level, to predict the unobserved potential outcome. NNM uses the covariates $\{x_1, x_2, \ldots, x_p\}$ to find the most similar individuals that get the other treatment level.

More formally, consider the vector of covariates $x_i = \{x_{i,1}, x_{i,2}, \ldots, x_{i,p}\}$ and frequency weight $w_i$ for observation $i$. The distance between $x_i$ and $x_j$ is parameterized by the vector norm

$$\|x_i - x_j\|_S = \{(x_i - x_j)' S^{-1} (x_i - x_j)\}^{1/2}$$

where $S$ is a given symmetric, positive-definite matrix.

Using this distance definition, we find that the set of nearest-neighbor indices for observation $i$ is

$$\Omega^x_{m}(i) = \{j_1, j_2, \ldots, j_{m_i} \mid t_{jk} = 1 - t_i, \quad \|x_i - x_{jk}\|_S < \|x_i - x_l\|_S, \quad t_l = 1 - t_i, \quad l \neq j_k\}$$

Here $m_i$ is the smallest number such that the number of elements in each set, $m_i = |\Omega^x_{m}(i)| = \sum_{j \in \Omega^x_{m}(i)} w_j$, is at least $m$, the desired number of matches. You set the size of $m$ using the `nneighbors(#)` option. The number of matches for the $i$th observation may not equal $m$ because of ties or if there are not enough observations with a distance from observation $i$ within the caliper limit, $c$. $\|x_i - x_j\|_S \leq c$. You may set the caliper limit by using the `caliper(#)` option. For ease of notation, we will use the abbreviation $\Omega(i) = \Omega^x_{m}(i)$.

With the `metric(string)` option, you have three choices for the scaling matrix $S$: Mahalanobis, inverse variance, or Euclidean.
\[
S = \begin{cases} 
(X - \bar{x}'1_n)'W(X - \bar{x}'1_n) \\
\text{diag} \left( (X - \bar{x}'1_n)'W(X - \bar{x}'1_n) \right) \\
I_p 
\end{cases}
\] if metric = mahalanobis
\[
\text{diag} \left( (X - \bar{x}'1_n)'W(X - \bar{x}'1_n) \right) 
\] if metric = ivariance
\[
I_p 
\] if metric = euclidean

where \(1_n\) is an \(n \times 1\) vector of ones, \(I_p\) is the identity matrix of order \(p\), \(\bar{x} = (\sum^nw_ix_i)/\sum^nw_i\), and \(W\) is an \(n \times n\) diagonal matrix containing frequency weights.

The NNM method predicting the potential outcome for the \(i\)th observation as a function of the observed \(y_i\) is

\[
\hat{y}_{ti} = \begin{cases} 
y_i & \text{if } t_i = t \\
\sum_{j \in \Omega(i)} w_j y_j / \sum_{j \in \Omega(i)} w_j & \text{otherwise} 
\end{cases}
\]

for \(t \in \{0, 1\}\).

We are now set to provide formulas for estimates \(\hat{\tau}_1\), the ATE, and \(\hat{\delta}_1\), the ATET,

\[
\hat{\tau}_1 = \frac{\sum^nw_i(\hat{y}_{1i} - \hat{y}_{0i})}{\sum^nw_i} = \frac{\sum^nw_i(2t_i - 1)\{1 + K_m(i)\}y_i}{\sum^nw_i}
\]

\[
\hat{\delta}_1 = \frac{\sum^nt_iw_i(\hat{y}_{1i} - \hat{y}_{0i})}{\sum^nt_iw_i} = \frac{\sum^nt_i\{t_i - (1 - t_i)K_m(i)\}y_i}{\sum^nt_iw_i}
\]

where

\[
K_m(i) = \sum_{j=1}^{n} I\{i \in \Omega(j)\} \frac{w_j}{\sum_{k \in \Omega(j)} w_k}
\]

The estimated variance of \(\hat{\tau}_1\) and \(\hat{\delta}_1\) are computed as

\[
\hat{\sigma}^2_{\tau} = \frac{\sum^nw_i \left[ (\hat{y}_{1i} - \hat{y}_{0i} - \hat{\tau}_1)^2 + \hat{\xi}^2_i \{K_m^2(i) + 2K_m(i) - K_m'(i)\} \right]}{\left(\sum^nw_i\right)^2}
\]

\[
\hat{\sigma}^2_{\delta} = \frac{\sum^nt_iw_i \left[ (\hat{y}_{1i} - \hat{y}_{0i} - \hat{\delta}_1)^2 + \hat{\xi}^2_i \{K_m^2(i) - K_m'(i)\} \right]}{\left(\sum^nt_iw_i\right)^2}
\]

where

\[
K_m'(i) = \sum_{j=1}^{n} I\{i \in \Omega(j)\} \frac{w_j}{\left(\sum_{k \in \Omega(j)} w_k\right)^2}
\]

and \(\hat{\xi}^2_i = \text{var}(y_{ti}|x_i)\) is the conditional outcome variance. If we can assume that \(\hat{\xi}^2_i\) does not vary with the covariates or treatment (homoskedastic), then we can compute an ATE estimate of \(\xi^2_{\tau}\) as
Bias-corrected matching estimator

When matching on more than one continuous covariate, the matching estimator described above is biased, even in infinitely large samples; in other words, it is not $\sqrt{n}$-consistent; see Abadie and Imbens (2006, 2011). Following Abadie and Imbens (2011) and Abadie et al. (2004), \texttt{teffects nnnmatch} makes an adjustment based on the regression functions $\mu_t(\tilde{x}_i) = E(y_t \mid \tilde{x} = \tilde{x}_i)$, for $t = 0, 1$ and the set of covariates $\tilde{x}_i = (\tilde{x}_{i,1}, \ldots, \tilde{x}_{i,q})$. The bias-correction covariates may be the same as the NNM covariates $x_i$. We denote the least-squares estimates as $\hat{\mu}_t(\tilde{x}_i) = \hat{\nu}_t + \hat{\beta}_t \tilde{x}_i$, where we regress $\{y_i \mid t_i = t\}$ onto $\{\tilde{x}_i \mid t_i = t\}$ with weights $w_i K_m(i)$, for $t = 0, 1$.

Given the estimated regression functions, the bias-corrected predictions for the potential outcomes are computed as

$$\hat{y}_{ti} = \begin{cases} y_i & \text{if } t_i = t \\ \frac{\sum_{j \in \Omega_{m}(i)} w_j \{y_j + \hat{\mu}_t(\tilde{x}_i) - \hat{\mu}_t(\tilde{x}_j)\}}{\sum_{j \in \Omega_{m}(i)} w_j} & \text{otherwise} \end{cases}$$

The \texttt{biasadj(varlist)} option specifies the bias-adjustment covariates $\tilde{x}_i$. 

$$\hat{\xi}^2_{\tau} = \frac{1}{2\sum_i^n w_i} \sum_{i=1}^n w_i \left[ \frac{\sum_{j \in \Omega(i)} w_j \{y_i - y_j(1 - t_i) - \hat{\tau}_1\}^2}{\sum_{j \in \Omega(i)} w_j} \right]$$

and an ATET estimate of $\xi^2_\delta$ as

$$\hat{\xi}^2_\delta = \frac{1}{2\sum_i^n t_i w_i} \sum_{i=1}^n t_i w_i \left[ \frac{\sum_{j \in \Omega(i)} t_j w_j \{y_i - y_j(1 - t_i) - \hat{\delta}_1\}^2}{\sum_{j \in \Omega(i)} t_j w_j} \right]$$

If the conditional outcome variance is dependent on the covariates or treatment, we require an estimate for $\xi^2_t$ at each observation. In this case, we require a second matching procedure, where we match on observations within the same treatment group.

Define the within-treatment matching set

$$\Psi^x_h(i) = \{j_1, j_2, \ldots, j_{h_i} \mid t_{j_k} = t_i, \|x_i - x_{j_k}\|_S < \|x_i - x_l\|_S, t_l = t_i, l \neq j_k\}$$

where $h$ is the desired set size. As before, the number of elements in each set, $h_i = |\Psi^x_h(i)|$, may vary depending on ties and the value of the caliper. You set $h$ using the \texttt{vce(robust, nn(#))} option. As before, we will use the abbreviation $\Psi(i) = \Psi^x_h(i)$ where convenient.

We estimate $\xi^2_t$ by

$$\hat{\xi}^2_{t_i}(x_i) = \frac{\sum_{j \in \Psi(i)} w_j (y_j - \bar{y}_{\Psi})^2}{\sum_{j \in \Psi(i)} w_j - 1}$$

where $\bar{y}_{\Psi} = \frac{\sum_{j \in \Psi(i)} w_j y_j}{\sum_{j \in \Psi(i)} w_j - 1}$
Propensity-score matching estimator

The propensity-score matching (PSM) estimator uses a treatment model (TM), \( p(z_i, t, \gamma) \), to model the conditional probability that observation \( i \) receives treatment \( t \) given covariates \( z_i \). The literature calls \( p(z_i, t, \gamma) \) a propensity score, and PSM matches on the estimated propensity scores.

When matching on the estimated propensity score, the set of nearest-neighbor indices for observation \( i \), \( i = 1, \ldots, n \), is

\[
\Omega_m(i) = \{j_1, j_2, \ldots, j_m \mid t_{jk} = 1 - t_i, |\hat{p}_i(t) - \hat{p}_{jk}(t)| < |\hat{p}_i(t) - \hat{p}_l(t)|, t_l = 1 - t_i, l \neq jk\}
\]

where \( \hat{p}_i(t) = p(z_i, t, \hat{\gamma}) \). As was the case with the NNM estimator, \( m_i \) is the smallest number such that the number of elements in each set, \( m_i = |\Omega_m(i)| = \sum_{j \in \Omega_m(i)} w_j \), is at least \( m \), the desired number of matches, set by the `nneighbors(#)` option.

Once a matching set is computed for each observation, the potential-outcome mean, \( \tau \) and \( \delta \), have the following forms, respectively:

\[
\hat{\tau}_i = \frac{\sum_{j \in \Omega_1(i)} \hat{y}_j - \sum_{j \in \Omega_0(i)} \hat{y}_j}{m_i}, \quad \hat{\delta}_i = \frac{\sum_{j \in \Omega_1(i)} \hat{y}_j - \sum_{j \in \Omega_0(i)} \hat{y}_j}{m_i}
\]

where \( \Omega_1(i) \) and \( \Omega_0(i) \) are the sets of treated and untreated observations, respectively.

The adjustment term for \( \text{ATE} \) can be expressed as

\[
\hat{c}_c = \frac{1}{\sum_{i=1}^{n} w_i} \sum_{i=1}^{n} w_i f(z_i', \hat{\gamma}) \left( \frac{\text{cov}(z_i, \hat{y}_{i1})}{\hat{p}_i(1)} + \frac{\text{cov}(z_i, \hat{y}_{i0})}{\hat{p}_i(0)} \right)
\]

where

\[
f(z_i', \hat{\gamma}) = \frac{d p(z_i, 1, \hat{\gamma})}{d (z_i')}
\]

is the derivative of \( p(z_i, 1, \hat{\gamma}) \) with respect to \( z_i' \), and

\[
\text{cov}(z_i, \hat{y}_{i1}) = \begin{cases} \sum_{j \in \Omega_1(i)} w_j (z_j - \overline{z}_{\Omega_1(i)}) (y_j - \overline{y}_{\Omega_1(i)}) & \text{if } t_i = t \\ \sum_{j \in \Omega_1(i)} w_j - 1 & \text{otherwise} \end{cases}
\]

\[
\text{cov}(z_i, \hat{y}_{i0}) = \begin{cases} \sum_{j \in \Omega_0(i)} w_j (z_j - \overline{z}_{\Omega_0(i)}) (y_j - \overline{y}_{\Omega_0(i)}) & \text{if } t_i = 0 \\ \sum_{j \in \Omega_0(i)} w_j - 1 & \text{otherwise} \end{cases}
\]

PSM, ATE, and ATET variance adjustment

The variances for \( \hat{\tau}_1 \) and \( \hat{\delta}_1 \) must be adjusted because we use \( \hat{\gamma} \) instead of \( \gamma \). The adjusted variances for \( \hat{\tau}_1 \) and \( \hat{\delta}_1 \) have the following forms, respectively:

\[
\hat{\sigma}_{\tau, \text{adj}}^2 = \hat{\sigma}_{\tau}^2 + \hat{c}' \hat{V}_{\gamma} \hat{c}_c
\]

\[
\hat{\sigma}_{\delta, \text{adj}}^2 = \hat{\sigma}_{\delta}^2 - \hat{c}' \hat{V}_{\gamma} \hat{c}_c + \frac{\partial \hat{\delta}_1}{\partial \gamma} \hat{V}_{\gamma} \frac{\partial \hat{\delta}_1}{\partial \gamma}
\]

In both equations, the matrix \( \hat{V}_{\gamma} \) is the TM coefficient variance–covariance matrix.

The adjustment term for ATE can be expressed as

\[
\hat{c}_c = \frac{1}{\sum_{i=1}^{n} w_i} \sum_{i=1}^{n} w_i f(z_i', \hat{\gamma}) \left( \frac{\text{cov}(z_i, \hat{y}_{i1})}{\hat{p}_i(1)} + \frac{\text{cov}(z_i, \hat{y}_{i0})}{\hat{p}_i(0)} \right)
\]

where

\[
f(z_i', \hat{\gamma}) = \frac{d p(z_i, 1, \hat{\gamma})}{d (z_i')}
\]

is the derivative of \( p(z_i, 1, \hat{\gamma}) \) with respect to \( z_i' \), and

\[
\text{cov}(z_i, \hat{y}_{i1}) = \begin{cases} \sum_{j \in \Omega_1(i)} w_j (z_j - \overline{z}_{\Omega_1(i)}) (y_j - \overline{y}_{\Omega_1(i)}) & \text{if } t_i = t \\ \sum_{j \in \Omega_1(i)} w_j - 1 & \text{otherwise} \end{cases}
\]

\[
\text{cov}(z_i, \hat{y}_{i0}) = \begin{cases} \sum_{j \in \Omega_0(i)} w_j (z_j - \overline{z}_{\Omega_0(i)}) (y_j - \overline{y}_{\Omega_0(i)}) & \text{if } t_i = 0 \\ \sum_{j \in \Omega_0(i)} w_j - 1 & \text{otherwise} \end{cases}
\]
is a $p \times 1$ vector with

$$
\mathbf{z}_{\Psi_i} = \frac{\sum_{j \in \Psi_h(i)} w_j \mathbf{z}_j}{\sum_{j \in \Psi_h(i)} w_j}
$$

$$
\mathbf{z}_{\Omega_i} = \frac{\sum_{j \in \Omega_h(i)} w_j \mathbf{z}_j}{\sum_{j \in \Omega_h(i)} w_j}
$$

and

$$
\mathbf{y}_{\Omega_i} = \frac{\sum_{j \in \Omega_h(i)} w_j y_j}{\sum_{j \in \Omega_h(i)} w_j}
$$

Here we have used the notation $\Psi_h(i) = \Psi^p_h(i)$ and $\Omega_h(i) = \Omega^p_h(i)$ to stress that the within-treatment and opposite-treatment clusters used in computing $\hat{\sigma}^2_{\tau, \text{adj}}$ and $\hat{\sigma}^2_{\tau, \text{adj}}$ are based on $h$ instead of the cluster $\Omega^m(i)$ based on $m$ used to compute $\hat{\tau}_1$ and $\hat{\delta}_1$, although you may desire to have $h = m$.

The adjustment term $\mathbf{c}_\delta$ for the ATET estimate has two components, $\mathbf{c}_\delta = \mathbf{c}_{\delta,1} + \mathbf{c}_{\delta,2}$, defined as

$$
\mathbf{c}_{\delta,1} = \frac{1}{\sum_{i=1}^n t_i w_i} \sum_{i=1}^n w_i \mathbf{z}_i f(\mathbf{z}_i') \left( \hat{y}_{1i} - \hat{y}_{0i} - \hat{\delta}_1 \right)
$$

$$
\mathbf{c}_{\delta,2} = \frac{1}{\sum_{i=1}^n t_i w_i} \sum_{i=1}^n w_i f(\mathbf{z}_i') \left\{ \text{cov}(\mathbf{z}_i, \hat{y}_{1i}) + \frac{\hat{p}_t(1) - \hat{p}_t(0)}{\hat{p}_t(0)} \text{cov}(\mathbf{z}_i, \hat{y}_{0i}) \right\}
$$

where

$$
\hat{y}_{ti} = \begin{cases} 
\frac{\sum_{j \in \Psi_h(t-i)} w_j y_j}{\sum_{j \in \Psi_h(t-i)} w_j} & \text{if } t = t_i \\
\frac{\sum_{j \in \Omega_h} w_j y_j}{\sum_{j \in \Omega_h} w_j} & \text{otherwise}
\end{cases}
$$

and the within-treatment matching sets $\Psi_h(t-i) = \Psi^p_h(t-i)$ are similar to $\Psi^p_h(i)$ but exclude observation $i$:

$$
\Psi^p_h(t-i) = \{ j_1, j_2, \ldots, j_n \mid j_k \neq i, \ t_{jk} = t_i, \ |\hat{p}_i - \hat{p}_{jk}| < |\hat{p}_i - \hat{p}_l|, \ t_l = t_i, \ l \notin \{i, j_k\} \}
$$

Finally, we cover the computation of $\hat{\delta}_{1}'$ in the third term on the right-hand side of $\hat{\sigma}^2_{\delta, \text{adj}}$. Here we require yet another clustering, but we match on the opposite treatment by using the covariates $\mathbf{z}_i = (z_{i,1}, \ldots, z_{i,p})'$. We will denote these cluster sets as $\Omega^m_z(i)$, for $i = 1, \ldots, n$.

The estimator of the $p \times 1$ vector $(\partial \delta_1')/\partial \gamma'$ is computed as

$$
\hat{\delta}_{1}' = \frac{1}{\sum_i t_i w_i} \sum_{i=1}^n \mathbf{z}_i f(\mathbf{z}_i') \left\{ (2t_i - 1)(y_i - \bar{y}_{\Omega^m_z(i)}) - \hat{\delta}_1 \right\}
$$

where

$$
\bar{y}_{\Omega^m_z(i)} = \frac{\sum_{j \in \Omega^m_z(i)} w_j y_j}{\sum_{j \in \Omega^m_z(i)} w_j}
$$
References


Also see

[TE] *teffects postestimation* — Postestimation tools for *teffects*

[TE] *teffects* — Treatment-effects estimation for observational data

[TE] *teffects psmatch* — Propensity-score matching

[U] 20 Estimation and postestimation commands
Description

One of the assumptions required to use the teffects and stteffects estimators is the overlap assumption, which states that each individual has a positive probability of receiving each treatment level. teffects overlap, a postestimation command, plots the estimated densities of the probability of getting each treatment level. These plots can be used to check whether the overlap assumption is violated.

Quick start

Visually check whether the overlap assumption is violated
   teffects overlap

As above, but use the Epanechnikov kernel function
   teffects overlap, kernel(epanechnikov)

Plot probability of getting treatment 3 for subjects receiving treatments 2 or 3 of a multivalued treatment
   teffects overlap, ptlevel(3) tlevels(2 3)

As above, and change legend labels to “Treated 2” and “Treated 3”
   teffects overlap, ptlevel(3) tlevels(2 3) ///
       legend(label(1 "Treated 2") label(2 "Treated 3"))

Menu

Statistics > Treatment effects > Overlap plots
## Syntax

```bash
teffects overlap [, treat_options kden_options]
```

### `treat_options`

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>ptlevel(treat_level)</code></td>
</tr>
<tr>
<td><code>tlevels(treatments)</code></td>
</tr>
<tr>
<td><code>nolabel</code></td>
</tr>
</tbody>
</table>

### `kden_options`

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>kernel(kernel)</code></td>
</tr>
<tr>
<td><code>n(#)</code></td>
</tr>
<tr>
<td><code>bwidth(#)</code></td>
</tr>
<tr>
<td><code>at(var_x)</code></td>
</tr>
</tbody>
</table>

**Kernel plots**

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>line#opts(cline_options)</code></td>
</tr>
</tbody>
</table>

**Add plots**

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>addplot(plot)</code></td>
</tr>
</tbody>
</table>

**Y axis, X axis, Titles, Legend, Overall**

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>twoway_options</code></td>
</tr>
</tbody>
</table>

### `kernel_options`

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>triangle</code></td>
</tr>
<tr>
<td><code>epanechnikov</code></td>
</tr>
<tr>
<td><code>epan2</code></td>
</tr>
<tr>
<td><code>biweight</code></td>
</tr>
<tr>
<td><code>cosine</code></td>
</tr>
<tr>
<td><code>gaussian</code></td>
</tr>
<tr>
<td><code>parzen</code></td>
</tr>
<tr>
<td><code>rectangle</code></td>
</tr>
</tbody>
</table>

## Options

**Main**

- `ptlevel(treat_level)` specifies that predicted probabilities be calculated for treatment level `treat_level`. The default is `ptlevel(first)`, where `first` is the first treatment level.
\texttt{tlevels}(*treatments*) specifies the observations for which to obtain predicted probabilities. By default, all treatment levels are used. Specify *treatments* as a space-delimited list.

For instance,

\begin{verbatim}
   . teffects overlap, ptlevel(1) tlevels(1 2)
\end{verbatim}

says to predict the probability of getting treatment level 1 for those subjects who actually obtained treatment levels 1 or 2.

\texttt{nolabel} specifies that treatment level values and not value labels be used in legend and axis titles.

\texttt{kernel(*kernel*)} specifies the kernel function for use in calculating the kernel density estimates. The default kernel is the triangle kernel (*triangle*).

\texttt{n(*#*)} specifies the number of points at which the density estimate is to be evaluated. The default is \texttt{e(N)}, the estimation sample size.

\texttt{bwidth(*#*)} specifies the half-width of the kernel, the width of the density window around each point. If \texttt{bwidth()} is not specified, the “optimal” width is calculated and used; see \([R]\) \texttt{kdensity}. The optimal width is the width that would minimize the mean integrated squared error if the data were Gaussian and a Gaussian kernel were used, so it is not optimal in any global sense. In fact, for multimodal and highly skewed densities, this width is usually too wide and oversmooths the density (Silverman 1986).

\texttt{at(*var_x*)} specifies a variable that contains the values at which the density should be estimated. This option allows you to more easily obtain density estimates for different variables or different subsamples of a variable and then overlay the estimated densities for comparison.

\underline{Kernel plots}

\texttt{line#opts(*cline_options*)} affect the rendition of the plotted kernel density estimates. See \([G-3]\) \texttt{cline_options}.

\underline{Add plots}

\texttt{addplot(*plot*)} provides a way to add other plots to the generated graph. See \([G-3]\) \texttt{addplot_option}.

\underline{Y axis, X axis, Titles, Legend, Overall}

\texttt{twoway_options} are any of the options documented in \([G-3]\) \texttt{twoway_options}, excluding \texttt{by()}. These include options for titling the graph (see \([G-3]\) \texttt{title_options}) and for saving the graph to disk (see \([G-3]\) \texttt{saving_option}).

\section*{Remarks and examples}

\texttt{teffects overlap} plots the estimated densities of the probability of getting each treatment level after \texttt{teffects}.

These plots can be used to check whether the overlap assumption is violated. The overlap assumption is satisfied when there is a chance of seeing observations in both the control and the treatment groups at each combination of covariate values; see \([TE]\) \texttt{teffects intro} or \([TE]\) \texttt{teffects intro advanced}.

The overlap assumption is required by the estimators implemented in \texttt{teffects}. Intuitively, when the overlap assumption is violated, we cannot predict, or otherwise account for, the unobserved outcomes for some individuals.
There is evidence that the overlap assumption is violated when an estimated density has too much mass around 0 or 1; see Busso, DiNardo, and McCrary (2014). An implication of this point is that when the overlap assumption is violated, the estimated densities will have relatively little mass in the regions in which they overlap.

Example 1: Assumption not violated

Continuing with example 1 of [TE] `teffects ipw`, we estimate the average treatment effect of smoking on birthweight and then draw the overlap plot:

```
. use http://www.stata-press.com/data/r15/cattaneo2
. teffects ipw (bweight) (mbsmoke mmarried c.mage##c.mage fbaby medu, probit)
Iteration 0: EE criterion = 4.621e-21
Iteration 1: EE criterion = 7.358e-26
Treatment-effects estimation Number of obs = 4,642
Estimator : inverse-probability weights
Outcome model : weighted mean
Treatment model: probit

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
<th></th>
<th></th>
<th>[95% Conf. Interval]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coef. Std. Err. z P&gt;</td>
<td>z</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ATE</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mbsmoke (smoker vs nonsmoker)</td>
<td>-230.6886 25.81524 -8.94 0.000 -281.2856 -180.0917</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>POmean</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mbsmoke nonsmoker</td>
<td>3403.463 9.571369 355.59 0.000 3384.703 3422.222</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
```
The graph displays the estimated density of the predicted probabilities that a nonsmoking mother is a nonsmoker and the estimated density of the predicted probabilities that a smoking mother is a nonsmoker.

Neither plot indicates too much probability mass near 0 or 1, and the two estimated densities have most of their respective masses in regions in which they overlap each other. Thus there is no evidence that the overlap assumption is violated.

Example 2: Assumption violated

This example produces an overlap plot that indicates a failure of the overlap assumption. We will use simulated data, so we know that the assumption is not true.

In our simulated dataset, some of the 1,000 adult males were given drug XY1 for high blood pressure and others were not. A scatterplot of systolic blood pressure (systolic) and weight (weight) reveals that heavier men were given the treatment. (The scatterplots corresponding to the treatment group are colored red, while the scatterplots corresponding to the control group are colored blue.)
There are no observations in the treated group for small weights, and there are no observations in the control group for large weights. There is clear evidence that the overlap assumption is violated.

Drawing an overlaid scatterplot is a straightforward way to check the overlap assumption in this example because there is only one covariate. This method is not available when there is more than one covariate. The predicted probability is a one-dimensional measure that captures the relevant multivariate information.

Below we estimate the parameters needed to calculate the predicted probabilities. The `pstolerance(1e-8)` option is specified to ensure that estimation is performed as long as the predicted probabilities are at least as large as 1e–8.

```
. teffects ipw (systolic) (xy1 weight), pstolerance(1e-8)
```

<table>
<thead>
<tr>
<th>Treatment-effects estimation</th>
<th>Number of obs</th>
<th>= 1,000</th>
</tr>
</thead>
<tbody>
<tr>
<td>Estimator</td>
<td>inverse-probability weights</td>
<td></td>
</tr>
<tr>
<td>Outcome model</td>
<td>weighted mean</td>
<td></td>
</tr>
<tr>
<td>Treatment model: logit</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coef.</td>
<td>Std. Err.</td>
<td>z</td>
<td>P&gt;</td>
<td>z</td>
</tr>
<tr>
<td>ATE</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>xy1 (Treated vs Untreated)</td>
<td>-16.23679</td>
<td>2.191703</td>
<td>-7.41</td>
<td>0.000</td>
<td>-20.53245, -11.94114</td>
</tr>
<tr>
<td>POmean</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>xy1 Untreated</td>
<td>127.9094</td>
<td>.7004533</td>
<td>182.61</td>
<td>0.000</td>
<td>126.5365, 129.2822</td>
</tr>
</tbody>
</table>

.T treated ... (mcolor(red))
> (scatter systolic weight if xy1==0, mcolor(blue)),
> legend(label(1 "Treated") label(2 "Untreated"))

```
```
Now we can obtain the overlap plot.

```
tffects overlap
```

![Overlap plot](image)

The estimated density of the predicted probabilities that a treated individual is not assigned to XY1 treatment has most of its mass near 0. The estimated density of the predicted probabilities that an untreated individual is not assigned to XY1 treatment has most of its mass near 1. Note that the two have very little mass in the region in which they overlap. There is clear evidence that the overlap assumption is violated.

### Stored results

`tffects overlap` stores the following in `r()`:

- **Scalars**
  - `r(bwidthj)` kernel bandwidth for treatment level `j`
  - `r(nj)` number of points at which the estimate was evaluated for treatment level `j`
  - `r(scalej)` density bin width for treatment level `j`

- **Macros**
  - `r(kernel)` name of kernel

### References


Also see

[TE] stteffects — Treatment-effects estimation for observational survival-time data
[TE] stteffects ipw — Survival-time inverse-probability weighting
[TE] stteffects ipwra — Survival-time inverse-probability-weighted regression adjustment
[TE] teffects — Treatment-effects estimation for observational data
[TE] teffects aipw — Augmented inverse-probability weighting
[TE] teffects ipw — Inverse-probability weighting
[TE] teffects ipwra — Inverse-probability-weighted regression adjustment
[TE] teffects nnmatch — Nearest-neighbor matching
[TE] teffects psmatch — Propensity-score matching
[TE] teffects ra — Regression adjustment
Postestimation commands

The following postestimation command is of special interest after `teffects`:

<table>
<thead>
<tr>
<th>Command</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>teffects overlap</code></td>
<td>overlap plots</td>
</tr>
<tr>
<td><code>tebalance</code></td>
<td>check balance of covariates</td>
</tr>
</tbody>
</table>

The following standard postestimation commands are also available:

<table>
<thead>
<tr>
<th>Command</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>estat summarize</code></td>
<td>summary statistics for the estimation sample</td>
</tr>
<tr>
<td><code>estat vce</code></td>
<td>variance–covariance matrix of the estimators (VCE)</td>
</tr>
<tr>
<td><code>estimates</code></td>
<td>cataloging estimation results</td>
</tr>
<tr>
<td><code>hausman</code></td>
<td>Hausman’s specification test</td>
</tr>
<tr>
<td><code>lincom</code></td>
<td>point estimates, standard errors, testing, and inference for linear combinations of coefficients</td>
</tr>
<tr>
<td><code>nlcom</code></td>
<td>point estimates, standard errors, testing, and inference for nonlinear combinations of coefficients</td>
</tr>
<tr>
<td><code>predict</code></td>
<td>predictions, residuals, influence statistics, and other diagnostic measures</td>
</tr>
<tr>
<td><code>predictnl</code></td>
<td>point estimates, standard errors, testing, and inference for generalized predictions</td>
</tr>
<tr>
<td><code>test</code></td>
<td>Wald tests of simple and composite linear hypotheses</td>
</tr>
<tr>
<td><code>testnl</code></td>
<td>Wald tests of nonlinear hypotheses</td>
</tr>
</tbody>
</table>
predict

Description for predict

predict creates a new variable containing predictions such as treatment effects, potential outcomes, conditional means, propensity scores, linear predictions, nearest-neighbor distances, and log square root of latent variances.

Menu for predict

Statistics > Postestimation

Syntaxes for predict

Syntaxes are presented under the following headings:

- Syntax for predict after aipw and ipwra
- Syntax for predict after ipw
- Syntax for predict after nnmatch and psmatch
- Syntax for predict after ra

Syntax for predict after aipw and ipwra

```
predict [type] {stub* | newvar | newvarlist} [if] [in]
[, statistic tlevel(treat_level)]
predict [type] {stub* | newvarlist} [if] [in], scores
```

<table>
<thead>
<tr>
<th>statistic</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>te</td>
<td>treatment effect; the default</td>
</tr>
<tr>
<td>cmean</td>
<td>conditional mean at treatment level</td>
</tr>
<tr>
<td>ps</td>
<td>propensity score</td>
</tr>
<tr>
<td>xb</td>
<td>linear prediction</td>
</tr>
<tr>
<td>psxb</td>
<td>linear prediction for propensity score</td>
</tr>
<tr>
<td>lnsigma</td>
<td>log square root of conditional latent variance (for outcome model &quot;hetprobit()&quot;) at treatment level</td>
</tr>
<tr>
<td>pslnsigma</td>
<td>log square root of latent variance (for treatment model &quot;hetprobit()&quot;) for propensity score</td>
</tr>
</tbody>
</table>

If you do not specify tlevel() and only specify one new variable, te and psxb assume tlevel() specifies the first noncontrol treatment level.

If you do not specify tlevel() and only specify one new variable, cmean, ps, xb, and lnsigma assume tlevel() specifies the first treatment level.

You specify one or \( t \) new variables with cmean, ps, xb, and lnsigma, where \( t \) is the number of treatment levels.

You specify one or \( t-1 \) new variables with te, psxb, and pslnsigma.
Syntax for predict after ipw

```
predict [type] { stub* | newvar | newvarlist } [if] [in]
    [, statistic tlevel(treat_level)]
```

```
predict [type] { stub* | newvarlist } [if] [in], scores
```

<table>
<thead>
<tr>
<th>statistic</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>main</td>
<td></td>
</tr>
<tr>
<td>ps</td>
<td>propensity score; the default</td>
</tr>
<tr>
<td>xb</td>
<td>linear prediction for the propensity score</td>
</tr>
<tr>
<td>lnsigma</td>
<td>log square root of latent variance (for treatment model hetprobit())</td>
</tr>
</tbody>
</table>

If you do not specify tlevel() and only specify one new variable, ps assumes tlevel() specifies the first treatment level.
If you do not specify tlevel() and only specify one new variable, xb assumes tlevel() specifies the first noncontrol treatment level.
You specify one or t new variables with ps, where t is the number of treatment levels.
You specify one or t-1 new variables with xb and lnsigma.

Syntax for predict after nnmatch and psmatch

```
predict [type] { stub* | newvarlist } [, statistic tlevel(treat_level)]
```

<table>
<thead>
<tr>
<th>statistic</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>main</td>
<td></td>
</tr>
<tr>
<td>te</td>
<td>treatment effect; the default</td>
</tr>
<tr>
<td>po</td>
<td>potential outcome</td>
</tr>
<tr>
<td>distance</td>
<td>nearest-neighbor distance</td>
</tr>
<tr>
<td>ps</td>
<td>propensity score (psmatch only)</td>
</tr>
<tr>
<td>lnsigma</td>
<td>log square root of latent variance (for treatment model hetprobit())</td>
</tr>
</tbody>
</table>

These statistics are available for the estimation sample only and require the estimation option generate(stub). This is because of the nonparametric nature of the matching estimator.
If you do not specify tlevel() and only specify one new variable, po and ps assume tlevel() specifies the first treatment level.
You specify one new variable with te and lnsigma.
You specify one or two new variables with po and ps.
Syntax for predict after ra

```
predict [type] { stub* | newvar | newvarlist } [if] [in]
    [, statistic tlevel(treat_level) ]
```

```
predict [type] { stub* | newvarlist } [if] [in], scores
```

<table>
<thead>
<tr>
<th>statistic</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Main</td>
<td></td>
</tr>
<tr>
<td>te</td>
<td>treatment effect; the default</td>
</tr>
<tr>
<td>cmean</td>
<td>conditional mean at treatment level</td>
</tr>
<tr>
<td>xb</td>
<td>linear prediction</td>
</tr>
<tr>
<td>lnsigma</td>
<td>log square root of conditional latent variance (for outcome model hetprobit()) at treatment level</td>
</tr>
</tbody>
</table>

If you do not specify tlevel() and only specify one new variable, te assumes tlevel() specifies the first noncontrol treatment level.

If you do not specify tlevel() and only specify one new variable, cmean, xb, and lnsigma assume tlevel() specifies the first treatment level.

You specify one or t new variables with cmean, xb, and lnsigma, where t is the number of treatment levels.

You specify one or t−1 new variables with te.

Options for predict

Options are presented under the following headings:

- Options for predict after aipw and ipwra
- Options for predict after ipw
- Options for predict after nnmatch and psmatch
- Options for predict after ra

Options for predict after aipw and ipwra

```
Main
```

te, the default, calculates the treatment effect for each noncontrol treatment level or the treatment level specified in tlevel(). If you specify the tlevel() option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level (except the control level).

```
cmean
```
cmean calculates the conditional mean for each treatment level or the treatment level specified in tlevel(). If you specify the tlevel() option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

```
ps
```
ps calculates the propensity score of each treatment level or the treatment level specified in tlevel(). If you specify the tlevel() option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

```
xb
```
xb calculates the linear prediction at each treatment level or the treatment level specified in tlevel(). If you specify the tlevel() option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.
psxb calculates the linear prediction for the propensity score at each noncontrol level of the treatment or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level (except the control level).

lnsigma calculates the log square root of the conditional latent variance for each treatment level or the treatment level specified in `tlevel()`. This option is valid when outcome model `hetprobit()` was used. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

pslnsigma calculates the log square root of the latent variance for the propensity score. This option is only valid when treatment model `hetprobit()` was used. Specify only one new variable.

tlevel(`treat_level`) specifies the treatment level for prediction.

scores calculates the score variables. Parameter-level scores are computed for the treatment mean and average treatment-effect equations. Equation-level scores are computed for the propensity-score equations.

The $j$th new variable will contain the scores for the $j$th parameter in the coefficient table if $j \leq t$, where $t$ is the number of treatment levels. Otherwise, it will contain the scores for fitted equation $j - t$ following the first $t$ parameters in the coefficient table.

**Options for predict after ipw**

- **Main**
  - ps, the default, calculates the propensity score of each treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.
  - xb calculates the linear prediction for the propensity score at each noncontrol level of the treatment or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level (except the control level).
  - lnsigma calculates the log square root of the latent variance. This option is only valid when treatment model `hetprobit()` was used. Specify only one new variable.
  - tlevel(`treat_level`) specifies the treatment level for prediction.
  - scores calculates the score variables. Parameter-level scores are computed for the treatment mean and average treatment-effect equations. Equation-level scores are computed for the propensity-score equations.

The $j$th new variable will contain the scores for the $j$th parameter in the coefficient table if $j \leq t$, where $t$ is the number of treatment levels. Otherwise, it will contain the scores for fitted equation $j - t$ following the first $t$ parameters in the coefficient table.

**Options for predict after nnmatch and psmatch**

- **Main**
  - te, the default, calculates the treatment effect.
  - po calculates the predicted potential outcomes for each observation and treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify new variables for the control and treated groups.
distance calculates the distances of the nearest neighbors for each observation. The number of variables generated is equal to the maximum number of nearest-neighbor matches. This is equal to the number of index variables generated by the estimation option `generate(stub)`. You may use the `stub*` syntax to set the distance variable prefix: `stub1, stub2, ...`.

ps calculates the propensity score of each treatment level or the propensity score of the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify new variables for the control and treated groups.

lnsigma calculates the log square root of the latent variance. This option is only valid when treatment model `hetprobit()` was used. Specify only one new variable.

tlevel(treat_level) restricts potential-outcome estimation to either the treated group or the control group. This option may only be specified with options po and ps.

Options for predict after ra

- **te**, the default, calculates the treatment effect for each noncontrol treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level (except the control level).

- cmean calculates the conditional mean for each treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

- xb calculates the linear prediction at each treatment level or the treatment level specified in `tlevel()`. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

- lnsigma calculates the log square root of the conditional latent variance for each treatment level or the treatment level specified in `tlevel()`. This option is valid when outcome model `hetprobit()` was used. If you specify the `tlevel()` option, you need to specify only one new variable; otherwise, you must specify a new variable for each treatment level.

- tlevel(treat_level) specifies the treatment level for prediction.

- scores calculates the score variables. Parameter-level scores are computed for the treatment mean and average treatment-effect equations. Equation-level scores are computed for the regression equations.

  The $j$th new variable will contain the scores for the $j$th parameter in the coefficient table if $j \leq t$, where $t$ is the number of treatment levels. Otherwise, it will contain the scores for fitted equation $j - t$ following the first $t$ parameters in the coefficient table.

Remarks and examples

Checking model specification is the most frequent reason for postestimation computation after `teffects`. `teffects overlap` provides a graphical method for checking the overlap assumption; see [TE] `teffects overlap`. Summarizing the estimated probabilities provides another check. Recall that the reciprocals of these estimated probabilities are used as weights by some of the estimators. If the estimated probabilities are too small, the weights blow up.
We estimate the ATE of maternal smoking on infant birthweight by inverse-probability weighting; see example 1 of [TE] tefects ipw for background.

```
. use http://www.stata-press.com/data/r15/cattaneo2
. tefects ipw (bweight) (mbsmoke mmarried c.mage##c.mage fbaby medu, probit)
Iteration 0:  EE criterion =  4.621e-21
Iteration 1:  EE criterion =  7.358e-26
Treatment-effects estimation Number of obs      =   4,642
Estimator      : inverse-probability weights
Outcome model  : weighted mean
Treatment model: probit

                      Robust
       | Coef.  Std. Err.     z  P>|z|     [95% Conf. Interval]
----------------------------------------------------------------------------
ATE                      
mbsmoke (smoker vs nonsmoker)  
                          -230.6886   25.81524  -8.94  0.000  -281.2856   -180.0917
POmean                   
mbsmoke nonsmoker         
                          3403.463    9.571369  355.59  0.000   3384.703   3422.222
```

Below we compute and summarize the estimated treatment probabilities.

```
. predict pr1
(option ps assumed; propensity score)
. summarize pr1 if mbsmoke==1, detail
                      propensity score, mbsmoke=nonsmoker
                  Percentiles Smallest
1%       .2991634      .2196947
5%       .5441551      .2258079
10%      .5973879      .2258079
25%      .6377700      .2409025
50%      .7601717      Mean       .7456264
75%      .8453946      .9533503
90%      .8943686      .9596144
95%      .9096801      .9610220
99%      .9367017      .9665684
        Largest       Std. Dev.   .1276102
```

The smallest values do not imply very large weights.
Below we compute and summarize the estimated probabilities of not getting the treatment.

```
generate pr0 = 1 -pr1
summarize pr0 if mbsmoke==0, detail
```

<table>
<thead>
<tr>
<th>Percentiles</th>
<th>Smallest</th>
</tr>
</thead>
<tbody>
<tr>
<td>1%</td>
<td>0.0351884</td>
</tr>
<tr>
<td>5%</td>
<td>0.0578012</td>
</tr>
<tr>
<td>10%</td>
<td>0.0674359</td>
</tr>
<tr>
<td>25%</td>
<td>0.0950869</td>
</tr>
<tr>
<td>50%</td>
<td>0.1372589</td>
</tr>
<tr>
<td>75%</td>
<td>0.2211142</td>
</tr>
<tr>
<td>90%</td>
<td>0.3242757</td>
</tr>
<tr>
<td>95%</td>
<td>0.3883457</td>
</tr>
<tr>
<td>99%</td>
<td>0.501537</td>
</tr>
</tbody>
</table>

```
Percentiles Smallest
1% .0351884 .0074551 5% .0578012 .0079309 10% .0674359 .0106305 Obs 3,778
25% .0950869 .0106305 Sum of Wgt. 3,778
50% .1372589 Mean .1698913
75% .2211142 Largest Std. Dev. .1059434
90% .3242757 Variance .011224
95% .3883457 Skewness 1.514456
99% .501537 Kurtosis 6.151114
```

Although there are two small probabilities, overall the small values do not imply large weights.

**Also see**

[TE] teffects overlap — Overlap plots  
[TE] teffects aipw — Augmented inverse-probability weighting  
[TE] teffects ipw — Inverse-probability weighting  
[TE] teffects ipwra — Inverse-probability-weighted regression adjustment  
[TE] teffects nnmatch — Nearest-neighbor matching  
[TE] teffects psmatch — Propensity-score matching  
[TE] teffects ra — Regression adjustment  
[U] 20 Estimation and postestimation commands
**teffects psmatch — Propensity-score matching**

### Description

teffects psmatch estimates the average treatment effect (ATE) and average treatment effect on the treated (ATET) from observational data by propensity-score matching (PSM). PSM estimators impute the missing potential outcome for each subject by using an average of the outcomes of similar subjects that receive the other treatment level. Similarity between subjects is based on estimated treatment probabilities, known as propensity scores. The treatment effect is computed by taking the average of the difference between the observed and potential outcomes for each subject. teffects psmatch accepts a continuous, binary, count, fractional, or nonnegative outcome.

See [TE] teffects intro or [TE] teffects intro advanced for more information about estimating treatment effects from observational data.

### Quick start

ATE of treat on y estimated by PSM using a logistic model for treat on x and indicators for levels of categorical variable a

```
teffects psmatch (y) (treat x i.a)
```

As above, but estimate the ATET

```
teffects psmatch (y) (treat x i.a), atet
```

ATE of treat using a heteroskedastic probit model for treatment

```
teffects psmatch (y) (treat x i.a, hetprobit(x i.a))
```

With 4 matches per observation

```
teffects psmatch (y) (treat x i.a), nneighbor(4)
```
Syntax

teffects psmatch (ovar) (tvar tmvarlist [, tmodel]) [if] [in] [weight]

[ , stat options ]

ovar is a binary, count, continuous, fractional, or nonnegative outcome of interest.
tvar must contain integer values representing the treatment levels.
tmvarlist specifies the variables that predict treatment assignment in the treatment model. Only two
treatment levels are allowed.

<table>
<thead>
<tr>
<th>tmodel</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>logit</td>
<td>logistic treatment model; the default</td>
</tr>
<tr>
<td>probit</td>
<td>probit treatment model</td>
</tr>
<tr>
<td>hetprobit(varlist)</td>
<td>heteroskedastic probit treatment model</td>
</tr>
</tbody>
</table>

_tmodel specifes the model for the treatment variable.

<table>
<thead>
<tr>
<th>stat</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>ate</td>
<td>estimate average treatment effect in population; the default</td>
</tr>
<tr>
<td>atet</td>
<td>estimate average treatment effect on the treated</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Model</th>
<th>Description</th>
<th>Stat</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>logit</td>
<td>logistic treatment model; the default</td>
<td>ate</td>
<td>estimate average treatment effect in population; the default</td>
</tr>
<tr>
<td>probit</td>
<td>probit treatment model</td>
<td>atet</td>
<td>estimate average treatment effect on the treated</td>
</tr>
<tr>
<td>hetprobit(varlist)</td>
<td>heteroskedastic probit treatment model</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
## teffects psmatch — Propensity-score matching

### options

<table>
<thead>
<tr>
<th>Model</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>nneighbor(#)</code></td>
<td>specify number of matches per observation; default is <code>nneighbor(1)</code></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>SE/Robust</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>vce(vcetype)</code></td>
<td><code>vcetype</code> may be</td>
</tr>
<tr>
<td></td>
<td><code>vce(robust [, nn(#)])</code>; use robust Abadie–Imbens standard errors with # matches</td>
</tr>
<tr>
<td></td>
<td><code>vce(iid)</code>; use independent and identically distributed Abadie–Imbens standard errors</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Reporting</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>level(#)</code></td>
<td>set confidence level; default is <code>level(95)</code></td>
</tr>
<tr>
<td><code>display_options</code></td>
<td>control columns and column formats, row spacing, line width, display of omitted variables and base and empty cells, and factor-variable labeling</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Advanced</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>caliper(#)</code></td>
<td>specify the maximum distance for which two observations are potential neighbors</td>
</tr>
<tr>
<td><code>pstolerance(#)</code></td>
<td>set tolerance for in overlap assumption</td>
</tr>
<tr>
<td><code>osample(newvar)</code></td>
<td><code>newvar</code> identifies observations that violate the overlap assumption</td>
</tr>
<tr>
<td>`control(#</td>
<td>label)`</td>
</tr>
<tr>
<td>`tlevel(#</td>
<td>label)`</td>
</tr>
<tr>
<td><code>generate(stub)</code></td>
<td>generate variables containing the observation numbers of the nearest neighbors</td>
</tr>
<tr>
<td><code>coeflegend</code></td>
<td>display legend instead of statistics</td>
</tr>
</tbody>
</table>

`tmvarlist` may contain factor variables; see [U] 11.4.3 Factor variables.

by and `statsby` are allowed; see [U] 11.1.10 Prefix commands.

`fweight`s are allowed; see [U] 11.1.6 weight.

`coeflegend` does not appear in the dialog box.

See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.

### Options

<table>
<thead>
<tr>
<th>Model</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>nneighbor(#)</code></td>
<td>specifies the number of matches per observation. The default is <code>nneighbor(1)</code>. Each individual is matched with at least the specified number of individuals from the other treatment level. <code>nneighbor()</code> must specify an integer greater than or equal to 1 but no larger than the number of observations in the smallest group.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Stat</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>stat</code></td>
<td>is one of two statistics: <code>ate</code> or <code>atet</code>. <code>ate</code> is the default. <code>ate</code> specifies that the average treatment effect be estimated. <code>atet</code> specifies that the average treatment effect on the treated be estimated.</td>
</tr>
</tbody>
</table>
**vce(vcetype)** specifies the standard errors that are reported. By default, `t effects psmatch` uses two matches in estimating the robust standard errors.

**vce(robust [ , nn(#)])** specifies that robust standard errors be reported and that the requested number of matches be used optionally.

**vce(iid)** specifies that standard errors for independent and identically distributed data be reported.

The standard derivative-based standard-error estimators cannot be used by `t effects psmatch`, because these matching estimators are not differentiable. The implemented method were derived by Abadie and Imbens (2006, 2011, 2012); see *Methods and formulas*.

As discussed in Abadie and Imbens (2008), bootstrap estimators do not provide reliable standard errors for the estimator implemented by `t effects psmatch`.

---

**caliper(#)** specifies the maximum distance at which two observations are a potential match. By default, all observations are potential matches regardless of how dissimilar they are.

In `t effects psmatch`, the distance is measured by the estimated propensity score. If an observation has no matches, `t effects psmatch` exits with an error.

**pstolerance(#)** specifies the tolerance used to check the overlap assumption. The default value is `pstolerance(1e-5)`. `t effects` will exit with an error if an observation has an estimated propensity score smaller than that specified by `pstolerance()`.

**osample(newvar)** specifies that indicator variable `newvar` be created to identify observations that violate the overlap assumption. Two checks are made to verify the assumption. The first ensures that the propensity scores are greater than `pstolerance(#)` and less than $1 - \text{pstolerance}()$. The second ensures that each observation has at least `nneighbor(#)` matches in the opposite treatment group within the distance specified by `caliper(#)`.

The **vce(robust, nn(#))** option also requires at least # matches in the same treatment group within the distance specified by `caliper(#)`.

The average treatment effect on the treated, option **atet**, using **vce(iid)** requires only `nneighbor(#)` control group matches for the treated group.

**control(# | label)** specifies the level of `tvar` that is the control. The default is the first treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. **control()** and **tlevel()** may not specify the same treatment level.

**tlevel(# | label)** specifies the level of `tvar` that is the treatment for the statistic **atet**. The default is the second treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. **tlevel()** may only be specified with statistic **atet**. **tlevel()** and **control()** may not specify the same treatment level.

**generate(stub)** specifies that the observation numbers of the nearest neighbors be stored in the new variables **stub1**, **stub2**, …. This option is required if you wish to perform postestimation based
on the matching results. The number of variables generated may be more than `nneighbors(#)` because of tied distances. These variables may not already exist.

The following option is available with `teffects psmatch` but is not shown in the dialog box: `coeflegend`; see [R] estimation options.

Remarks and examples

Propensity-score matching uses an average of the outcomes of similar subjects who get the other treatment level to impute the missing potential outcome for each subject. The ATE is computed by taking the average of the difference between the observed and potential outcomes for each subject. `teffects psmatch` determines how near subjects are to each other by using estimated treatment probabilities, known as propensity scores. This type of matching is known as propensity-score matching (PSM).

PSM does not need bias correction, because PSM matches on a single continuous covariate. In contrast, the nearest-neighbor matching estimator implemented in `teffects nnmatch` uses a bias-correction term when matching on more than one continuous covariate. In effect, the PSM estimator parameterizes the bias-correction term in the treatment probability model. See [TE] `teffects intro` or [TE] `teffects intro advanced` for more information about this estimator.

We will illustrate the use of `teffects psmatch` by using data from a study of the effect of a mother’s smoking status during pregnancy (`mbsmoke`) on infant birthweight (`bweight`) as reported by Cattaneo (2010). This dataset also contains information about each mother’s age (`mage`), education level (`medu`), marital status (`mmarried`), whether the first prenatal exam occurred in the first trimester (`prenatal1`), whether this baby was the mother’s first birth (`fbaby`), and the father’s age (`fage`).

Example 1: Estimating the ATE

We begin by using `teffects psmatch` to estimate the ATE of `mbsmoke` on `bweight`. We use a logistic model (the default) to predict each subject’s propensity score, using covariates `mage`, `medu`, `mmarried`, and `fbaby`. Because the performance of PSM hinges upon how well we can predict the propensity scores, we will use factor-variable notation to include both linear and quadratic terms for `mage`, the only continuous variable in our model:

```
. use http://www.stata-press.com/data/r15/cattaneo2
. teffects psmatch (bweight) (mbsmoke mmarried c.mage##c.mage fbaby medu)
```

```
Treatment-effects estimation Number of obs = 4,642
Estimator : propensity-score matching Matches: requested = 1
Outcome model : matching min = 1
Treatment model: logit max = 74

<table>
<thead>
<tr>
<th></th>
<th>AI Robust</th>
<th></th>
<th></th>
<th></th>
<th>[95% Conf. Interval]</th>
</tr>
</thead>
<tbody>
<tr>
<td>bweight</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ATE</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mbsmoke</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>smoker</td>
<td>-210.9683</td>
<td>32.021</td>
<td>-6.59</td>
<td>0.000</td>
<td>-273.7284        -148.2083</td>
</tr>
<tr>
<td>vs</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>nonsmoker</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
```

The average birthweight if all mothers were to smoke would be 211 grams less than the average that would occur if none of the mothers had smoked.
By default, `teffects psmatch` estimates the ATE by matching each subject to a single subject with the opposite treatment whose propensity score is closest. Sometimes, however, we may want to ensure that matching occurs only when the propensity scores of a subject and a match differ by less than a specified amount. To do that, we use the `caliper()` option. If a match within the distance specified in the `caliper()` option cannot be found, `teffects psmatch` exits.

### Example 2: Specifying the caliper

Here we reconsider the previous example, first specifying that we only want to consider a pair of observations a match if the absolute difference in the propensity scores is less than 0.03:

```stata
. teffects psmatch (bweight) (mbsmoke mmarried c.mage##c.mage fbaby medu), > caliper(0.03)
```

The error arose because there is not a smoking mother whose propensity score is within 0.03 of the propensity score of the nonsmoking mother in observation 4504. If we instead raise the caliper to 0.10, we have matches for all subjects and therefore obtain the same results as in example 1:

```stata
. teffects psmatch (bweight) (mbsmoke mmarried c.mage##c.mage fbaby medu), > caliper(0.1)
```

### Technical note

Example 2 highlights that estimating the ATE requires finding matches for both the treated and control subjects. In contrast, estimating the ATET only requires finding matches for the treated subjects. Because subject 4504 is a control subject, we can estimate the ATET using `caliper(0.03)`. We must also specify `vce(iid)` because the default robust standard errors for the estimated ATET require viable matches for both treated subjects and control subjects. (This requirement comes from the nonparametric method derived by Abadie and Imbens [2012].)
. teffects psmatch (bweight) (mbsmoke mmarried c.mage##c.mage fbaby medu), > atet vce(iid) caliper(0.03)

| Treatment-effects estimation | Number of obs |   = 4,642 |
|-------------------------------|----------------|
| Estimator: propensity-score matching | Matches: requested |   = 1 |
| Outcome model: matching | min |   = 1 |
| Treatment model: logit | max |   = 74 |

| bweight | Coef. | Std. Err. | z   | P>|z| | [95% Conf. Interval] |
|---------|-------|-----------|-----|-----|----------------------|
| ATET mbsmoke (smoker vs nonsmoker) | -236.7848 | 26.11698 | -9.07 | 0.000 | -287.9731 -185.5964 |

In the previous examples, each subject was matched to at least one other subject, which is the default behavior for `teffects psmatch`. However, we can request that `teffects psmatch` match each subject to multiple subjects with the opposite treatment level by specifying the `nneighbor()` option. Matching on more distant neighbors can reduce the variance of the estimator at a cost of an increase in bias.

**Example 3**

Now we request that `teffects psmatch` match a mother to four mothers in the opposite treatment group:

. teffects psmatch (bweight) (mbsmoke mmarried c.mage##c.mage fbaby medu), > nneighbor(4)

| Treatment-effects estimation | Number of obs |   = 4,642 |
|-------------------------------|----------------|
| Estimator: propensity-score matching | Matches: requested |   = 4 |
| Outcome model: matching | min |   = 4 |
| Treatment model: logit | max |   = 74 |

| bweight | Coef. | Std. Err. | z   | P>|z| | [95% Conf. Interval] |
|---------|-------|-----------|-----|-----|----------------------|
| ATE mbsmoke (smoker vs nonsmoker) | -224.006 | 29.88627 | -7.50 | 0.000 | -282.582 -165.43 |

These results are similar to those reported in example 1.

**Video example**

Treatment effects in Stata: Propensity-score matching
Stored results

teffects psmatch stores the following in e():

Scalars
- `e(N)`: number of observations
- `e(n_j)`: number of observations for treatment level \( j \)
- `e(k_levels)`: number of levels in treatment variable
- `e(caliper)`: maximum distance between matches
- `e(treated)`: level of treatment variable defined as treated
- `e(control)`: level of treatment variable defined as control
- `e(k_nneighbor)`: requested number of matches
- `e(k_nnmin)`: minimum number of matches
- `e(k_nnmax)`: maximum number of matches
- `e(k_robust)`: matches for robust VCE

Macros
- `e(cmd)`: teffects
- `e(cmdline)`: command as typed
- `e(depvar)`: name of outcome variable
- `e(tvar)`: name of treatment variable
- `e(subcmd)`: psmatch
- `e(tmodel)`: logit, probit, or hetprobit
- `e(stat)`: statistic estimated, ate or atet
- `e(wtype)`: weight type
- `e(wexp)`: weight expression
- `e(title)`: title in estimation output
- `e(tlevels)`: levels of treatment variable
- `e(psvarlist)`: variables in propensity-score model
- `e(hvarlist)`: variables for variance, only if hetprobit
- `e(vce)`: vcetype specified in vce()
- `e(vctype)`: title used to label Std. Err.
- `e(datasignature)`: the checksum
- `e(datasignaturevars)`: variables used in calculation of checksum
- `e(properties)`: b V
- `e(estat_cmd)`: program used to implement estat
- `e(predict)`: program used to implement predict
- `e(marginsnotok)`: predictions disallowed by margins

Matrices
- `e(b)`: coefficient vector
- `e(V)`: variance–covariance matrix of the estimators
- `e(bps)`: coefficient vector from propensity-score model
- `e(Vps)`: variance–covariance matrix of the estimators from propensity-score model

Functions
- `e(sample)`: marks estimation sample

Methods and formulas

The methods and formulas used by teffects psmatch are documented in the Methods and formulas of [TE] teffects nnmatch.

References


Also see

[TE] *teffects postestimation* — Postestimation tools for teffects
[TE] *teffects* — Treatment-effects estimation for observational data
[TE] *teffects nnmatch* — Nearest-neighbor matching
[U] 20 Estimation and postestimation commands
**teffects ra** estimates the average treatment effect (ATE), the average treatment effect on the treated (ATET), and the potential-outcome means (POMs) from observational data by regression adjustment (RA). RA estimators use contrasts of averages of treatment-specific predicted outcomes to estimate treatment effects. **teffects ra** accepts a continuous, binary, count, fractional, or nonnegative outcome and allows a multivalued treatment.

See [TE] **teffects intro** or [TE] **teffects intro advanced** for more information about estimating treatment effects from observational data.

### Quick start

ATE from a linear model of $y_1$ on $x_1$ and $x_2$ with binary treatment $treat2$

```
teffects ra (y1 x1 x2) (treat2)
```

As above, but estimate the ATET

```
teffects ra (y1 x1 x2) (treat2), atet
```

As above, but estimate the potential-outcome means

```
teffects ra (y1 x1 x2) (treat2), pomeans
```

ATE of $treat2$ using a heteroskedastic probit model for binary outcome $y_2$

```
teffects ra (y2 x1 x2, hetprobit(x1 x2)) (treat2)
```

ATE of $treat2$ using a Poisson model for count outcome $y_3$

```
teffects ra (y3 x1 x2, poisson) (treat2)
```

ATE for each level of three-valued treatment $treat3$

```
teffects ra (y1 x1 x2) (treat3)
```

As above, and specify that $treat3 = 3$ is the control level

```
teffects ra (y1 x1 x2) (treat3), control(3)
```

Same as above, specified using the label “MyControl” corresponding to $treat3 = 3$

```
teffects ra (y1 x1 x2) (treat3), control("MyControl")
```
teffects ra (ovar omvarlist [, omodel noconstant]) (tvar) [if] [in] [weight]
[ , stat options]

`ovar` is a binary, count, continuous, fractional, or nonnegative outcome of interest.
`omvarlist` specifies the covariates in the outcome model.
`tvar` must contain integer values representing the treatment levels.

<table>
<thead>
<tr>
<th><code>omodel</code></th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>linear</code></td>
<td>linear outcome model; the default</td>
</tr>
<tr>
<td><code>logit</code></td>
<td>logistic outcome model</td>
</tr>
<tr>
<td><code>probit</code></td>
<td>probit outcome model</td>
</tr>
<tr>
<td><code>hetprobit(varlist)</code></td>
<td>heteroskedastic probit outcome model</td>
</tr>
<tr>
<td><code>poisson</code></td>
<td>exponential outcome model</td>
</tr>
<tr>
<td><code>flogit</code></td>
<td>fractional logistic outcome model</td>
</tr>
<tr>
<td><code>fprobit</code></td>
<td>fractional probit outcome model</td>
</tr>
<tr>
<td><code>fhetprobit(varlist)</code></td>
<td>fractional heteroskedastic probit outcome model</td>
</tr>
</tbody>
</table>

`omodel` specifies the model for the outcome variable.

<table>
<thead>
<tr>
<th><code>stat</code></th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><code>ate</code></td>
<td>estimate average treatment effect in population; the default</td>
</tr>
<tr>
<td><code>atet</code></td>
<td>estimate average treatment effect on the treated</td>
</tr>
<tr>
<td><code>pomeans</code></td>
<td>estimate potential-outcome means</td>
</tr>
</tbody>
</table>
**t effects ra — Regression adjustment**

### Options

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>SE/Robust</strong></td>
</tr>
<tr>
<td><strong>vce(vcetype)</strong></td>
</tr>
<tr>
<td><strong>Reporting</strong></td>
</tr>
<tr>
<td><strong>level(#)</strong></td>
</tr>
<tr>
<td><strong>aequations</strong></td>
</tr>
<tr>
<td><strong>display_options</strong></td>
</tr>
<tr>
<td><strong>Maximization</strong></td>
</tr>
<tr>
<td><strong>maximize_options</strong></td>
</tr>
<tr>
<td><strong>Advanced</strong></td>
</tr>
<tr>
<td>**control(#</td>
</tr>
<tr>
<td>**tlevel(#</td>
</tr>
<tr>
<td><strong>coeflegend</strong></td>
</tr>
</tbody>
</table>

---

**omvarlist** may contain factor variables; see [U] 11.4.3 Factor variables.

*bootstrap*, by, *jackknife*, and *statsby* are allowed; see [U] 11.1.10 Prefix commands.

Weights are not allowed with the *bootstrap* prefix; see [R] bootstrap.

*fweights*, *iweights*, and *pweights* are allowed; see [U] 11.1.6 weight.

*coeflegend* does not appear in the dialog box.

See [U] 20 Estimation and postestimation commands for more capabilities of estimation commands.

### Options

- **Model**
  - `noconstant`; see [R] estimation options.

- **Stat**
  - `stat` is one of three statistics: *ate*, *atet*, or *pomeans*. *ate* is the default.
    - `ate` specifies that the average treatment effect be estimated.
    - `atet` specifies that the average treatment effect on the treated be estimated.
    - `pomeans` specifies that the potential-outcome means for each treatment level be estimated.

- **SE/Robust**
  - `vce(vcetype)` specifies the type of standard error reported, which includes types that are robust to some kinds of misspecification (*robust*), that allow for intragroup correlation (*cluster clustvar*), and that use bootstrap or jackknife methods (*bootstrap*, *jackknife*); see [R] `vce_option`.

- **Reporting**
  - `level(#)`; see [R] estimation options.
  - `aequations` specifies that the results for the outcome-model or the treatment-model parameters be displayed. By default, the results for these auxiliary parameters are not displayed.
display_options: noci, nopvalues, noomitted, vsquish, noemptycells, baselevels, allbaselevels, nofvlabel, fvwrap(#), fvwrapon(style), cformat(%fmt), pformat(%fmt), sformat(%fmt), and nolstretch; see [R] estimation options.

Maximization

maximize_options: iterate(#), [no]log, and from(init_specs); see [R] maximize. These options are seldom used.

init_specs is one of

matname [ , skip copy ]
# [ , # ...], copy

Advanced

control(# | label) specifies the level of tvar that is the control. The default is the first treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. control() may not be specified with statistic pmeans. control() and tlevel() may not specify the same treatment level.

tlevel(# | label) specifies the level of tvar that is the treatment for the statistic atet. The default is the second treatment level. You may specify the numeric level # (a nonnegative integer) or the label associated with the numeric level. tlevel() may only be specified with statistic atet. tlevel() and control() may not specify the same treatment level.

The following option is available with teffects ra but is not shown in the dialog box:

coefflegend; see [R] estimation options.

Remarks and examples

Remarks are presented under the following headings:

Overview
Video example

Overview

Regression adjustment (RA) estimators use the contrasts of the averages of treatment-specific predicted outcomes to estimate treatment effects. RA estimators use a two-step approach to estimating treatment effects:

1. They fit separate regression models of the outcome on a set of covariates for each treatment level.
2. They compute the averages of the predicted outcomes for each subject and treatment level. These averages reflect the POMs. The contrasts of these averages provide estimates of the ATEs. By restricting the computations of the means to the subset of treated subjects, we obtain the ATETs.

RA estimators are consistent as long as the treatment is independent of the potential outcomes after conditioning on the covariates. In fact, teffects ra uses an estimation technique that implements both steps at once so that we do not need to correct the standard errors in the second step to reflect the uncertainty surrounding the predicted outcomes.
We will illustrate the use of teffects ra by using data from a study of the effect of a mother’s smoking status during pregnancy (mbsmoke) on infant birthweight (bweight) as reported by Cattaneo (2010). This dataset also contains information about each mother’s age (mage), education level (medu), marital status (mmarried), whether the first prenatal exam occurred in the first trimester (prenatal1), and whether this baby was the mother’s first birth (fbaby).

Example 1: Estimating the ATE

We begin by using teffects ra to estimate the average treatment effect of smoking, controlling for first-trimester exam status, marital status, mother’s age, and first-birth status. In Stata, we type

```
use http://www.stata-press.com/data/r15/cattaneo2
.teffects ra (bweight prenatal1 mmarried mage fbaby) (mbsmoke)
```

<table>
<thead>
<tr>
<th>bweight</th>
<th>Robust</th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coef.</td>
<td>Std. Err.</td>
<td>z</td>
<td>P&gt;</td>
<td>z</td>
</tr>
<tr>
<td>ATE</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mbsmoke</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(smoker vs nonsmoker)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>-239.6392</td>
<td>23.82402</td>
<td>-10.06</td>
<td>0.000</td>
<td>-286.3334</td>
<td>-192.945</td>
</tr>
<tr>
<td>POmean</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>mbsmoke</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(nonsmoker)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3403.242</td>
<td>9.525207</td>
<td>357.29</td>
<td>0.000</td>
<td>3384.573</td>
<td>3421.911</td>
</tr>
</tbody>
</table>

The average birthweight if all mothers were to smoke would be 240 grams less than the average of 3,403 grams that would occur if none of the mothers had smoked.

The previous results showed us the average amount by which infants’ weights are affected by their mothers’ decision to smoke. We may instead be interested in knowing the average amount by which the weight of babies born to smoking mothers was decreased as a result of smoking. The ATET provides us with the answer.
To obtain the ATET rather than the ATE, we use the \texttt{atet} option:

```plaintext
. teffects ra (bweight prenatall mmarried mage fbaby) (mbsmoke), atet
```

Iteration 0:  
EE criterion = 7.629e-24
Iteration 1:  
EE criterion = 2.697e-26

Treatment-effects estimation  
Estimator : regression adjustment  
Outcome model : linear  
Treatment model: none

| bweight | Robust Coef. | Std. Err. | z | P>|z| | [95% Conf. Interval] |
|---------|--------------|-----------|---|-------|----------------------|
| ATET    |              |           |   |       |                      |
| mbsmoke | smoker vs   |           |   |       |                      |
|         | nonsmoker   |           |   |       |                      |
| -223.3017 | 22.7422    | -9.82   | 0.000 | -267.8755 | -178.7278 |
| POmean  |              |           |   |       |                      |
| mbsmoke | nonsmoker   |           |   |       |                      |
| 3360.961 | 12.75749    | 263.45  | 0.000 | 3335.957  | 3385.966 |

The average birthweight is 223 grams less when all the mothers who smoke do so than the average of 3,361 grams that would have occurred if none of these mothers had smoked.

The ATET differs from the ATE because the distribution of the covariates among mothers who smoke differs from the distribution for nonsmoking mothers. For example, in \texttt{TE} \texttt{teffects intro}, we remarked that in our sample, mothers who smoked tended to be older than those who did not. The differing distributions of covariates also affect the estimated POMs.

By default, \texttt{teffects ra} reports the ATE, which is the difference between the two POMs in the case of a binary treatment variable. Sometimes, we want to know the estimated POMs themselves. We might also want to see the actual regression equations used to estimate the POMs. Obtaining this information is easy, as the next example illustrates.
Example 3: Estimating the POMs

Here we use the `pomeans` option to display the POMs and the `aequations` option to display the estimated regression coefficients for the treated and untreated subjects.

```
. teffects ra (bweight prenatal1 mmarried mage fbaby) (mbsmoke),
    > pomeans aequations
Iteration 0:  EE criterion = 7.734e-24
Iteration 1:  EE criterion = 2.850e-26
```

Treatment-effects estimation

Number of obs = 4,642
Estimator : regression adjustment
Outcome model : linear
Treatment model: none

|                | Coef.  | Std. Err. | z     | P>|z|  | [95% Conf. Interval] |
|----------------|--------|-----------|-------|------|---------------------|
| **POMs**       |        |           |       |      |                     |
| mbsmoke        |        |           |       |      |                     |
| nonsmoker      | 3403.242 | 9.525207  | 357.29| 0.000| 3384.573 - 3421.911 |
| smoker         | 3163.603 | 21.86351  | 144.70| 0.000| 3120.751 - 3206.455 |
| **OME0**       |        |           |       |      |                     |
| prenatal1      | 64.40859 | 27.52699  | 2.34  | 0.019| 10.45669 - 118.3605 |
| mmarried       | 160.9513 | 26.6162   | 6.05  | 0.000| 108.7845 - 213.1181 |
| mage           | 2.546828 | 2.084324  | 1.22  | 0.222| -1.538373 - 6.632028 |
| fbaby          | -71.3286 | 19.64701  | -3.63 | 0.000| -109.836 -32.82117  |
| _cons          | 3202.746 | 54.01082  | 59.30 | 0.000| 3096.886 - 3308.605 |
| **OME1**       |        |           |       |      |                     |
| prenatal1      | 25.11133 | 40.37541  | 0.62  | 0.534| -54.02302 - 104.2457 |
| mmarried       | 133.6617 | 40.86443  | 3.27  | 0.001| 53.5689 - 213.7545  |
| mage           | -7.370881| 4.21817   | -1.75 | 0.081| -15.63834 - .8965804 |
| fbaby          | 41.43991 | 39.70712  | 1.04  | 0.297| -36.38461 - 119.2644 |
| _cons          | 3227.169 | 104.4059  | 30.91 | 0.000| 3022.537 - 3431.801 |

The nonsmoker POM for infant birthweight is 3,403 grams; that means that if none of the women in our sample smoked during pregnancy, the expected average birthweight would be 3,403 grams. The POM if all mothers did smoke during pregnancy is 3,164 grams, a difference of 240 grams, as we established in example 1. The coefficients for the equation labeled OME0 represent the linear equation used to estimate the nontreated POM, and the coefficients for the equation labeled OME1 represent the linear equation used to estimate the treated POM. The coefficients are identical to those we would obtain using `regress`, but the standard errors differ slightly because `teffects ra` does not make the small-sample adjustment that `regress` does.

We often express statistics as percentages to alleviate scaling issues and aid interpretation. In the present context, we may wish to express an ATE as a percentage of the untreated POM to gain a more intuitive measure of efficacy.
Example 4: Reporting the ATE as a percentage

Sometimes, we are interested in reporting the estimated treatment effect as a percentage of the untreated POM. We continue to use the same model as in the previous examples, but we specify the coeflegend option so that teffects ra reports the names of the parameters. Knowing the correct names to use, we can then use nlcom to obtain the percentage change along with its delta-method-based standard error. We type

```
use http://www.stata-press.com/data/r15/cattaneo2
teffects ra (bweight prenatal1 mmarried mage fbaby) (mbsmoke), coeflegend
```

```
Iteration 0: EE criterion = 7.734e-24
Iteration 1: EE criterion = 1.196e-25
```

```
Treatment-effects estimation
Number of obs = 4,642
Estimator : regression adjustment
Outcome model : linear
Treatment model: none

<table>
<thead>
<tr>
<th>bweight</th>
<th>Coef. Legend</th>
</tr>
</thead>
<tbody>
<tr>
<td>ATE</td>
<td></td>
</tr>
<tr>
<td>mbsmoke (smoker vs nonsmoker)</td>
<td>-239.6392 _b[ATE:r1vs0.mbsmoke]</td>
</tr>
<tr>
<td>POmean mbsmoke nonsmoker</td>
<td>3403.242 _b[POmean:0.mbsmoke]</td>
</tr>
</tbody>
</table>

```
.nlcom _b[ATE:r1vs0.mbsmoke] / _b[POmean:0.mbsmoke]
_nl_1: _b[ATE:r1vs0.mbsmoke] / _b[POmean:0.mbsmoke]
```

```
| bweight | Coef. Std. Err. z P>|z| [95% Conf. Interval] |
|---------|------------------|--------|----------|----------------|
| _nl_1   | -.070415 .0069245 -10.17 0.000 -.0839867 -.0568433 |
```

The average birthweight falls by an estimated 7.0% when every mother smokes relative to the case when no mothers smoke. We also obtain a 95% confidence interval of a 5.7% to 8.4% reduction.

Birthweights cannot be negative, though it is possible for a linear regression model to make negative predictions. A common way to enforce nonnegative predictions is to use an exponential conditional-mean model, which is commonly fitted using the Poisson quasimaximum-likelihood estimator, as discussed in Cameron and Trivedi (2005, sec. 5.7), Wooldridge (2010, sec. 18.2), and Pawitan (2001, chap. 14). teffects ra provides an option to use this model rather than linear regression for the outcomes.
Example 5: Modeling nonnegative outcomes

Now we refit our model of smoking behavior on birthweight, but we specify the poisson option in the outcome-model equation so that teffects ra uses the Poisson exponential model rather than linear regression:

```
. teffects ra (bweight prenatal1 mmarrried mage fbaby, poisson) (mbsmoke)
```

| Coef. | Std. Err. | z     | P>|z|  | [95% Conf. Interval] |
|-------|-----------|-------|------|----------------------|
| ATE mbsmoke (smoker vs nonsmoker) | -239.6669 | 23.83757 | -10.05 | 0.000 | -286.3877 -192.9462 |

In this case, using a model that forces outcomes to be nonnegative did not make any substantive difference. In this dataset, nearly 90% of babies weigh at least 2,700 grams, and even the smallest baby weighs 340 grams. When the dependent variable is so large, the predictions from Poisson and linear regression models are remarkably similar.

We now consider models for fractional outcomes. Fractional responses concern outcomes between 0 and 1. For instance, averaged 0/1 outcomes such as participation rates, but can also include variables that are naturally on a 0 to 1 scale such as pollution levels, patient oxygen saturation, and Gini coefficients (income inequality measures).

Example 6: Modeling fractional outcomes

We will illustrate the use of teffects ra with the outcome-model option fprobit by using simulated data. The observations are 543 cities at least 200 miles apart. The data contain information about each city’s level of industrialization (industrial), average annual rainfall in millimeters (rainfall), whether or not the city has a metro or train (train), and traffic congestion measured by an index (traffic).

Our outcome is the level of pollution (pollution) measured on a 0 to 1 scale. Values of pollution between 0 and 0.3 have no public health implications, but values greater than 0.7 imply that people with breathing or health problems should remain indoors. We study the effect of a tax on gas-guzzler cars on air pollution (guzzler). A tax that is effective in reducing pollution improves public health.

We estimate the ATE of a gas-guzzler tax on pollution, controlling for average yearly rainfall, traffic congestion, the level of industrialization, and whether the city has a train or a metro by using a fractional probit model.
. use http://www.stata-press.com/data/r15/pollution
(Simulated Urban Pollution Data)
. teffects ra (pollution rainfall i.traffic industrial i.train, fprobit) (guzzler)
Iteration 0:   EE criterion =  3.023e-16
Iteration 1:   EE criterion =  1.043e-31
Treatment-effects estimation           Number of obs   =   534
Estimator      : regression adjustment
Outcome model  : fractional probit
Treatment model: none

<table>
<thead>
<tr>
<th></th>
<th>Robust</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coef.  Std. Err.</td>
</tr>
<tr>
<td>pollution</td>
<td></td>
</tr>
<tr>
<td>ATE guzzler</td>
<td>-.0960214 .0113896</td>
</tr>
<tr>
<td>(tax vs no tax)</td>
<td></td>
</tr>
<tr>
<td>POmean guzzler</td>
<td>.3879346 .0101733</td>
</tr>
<tr>
<td>(no tax)</td>
<td></td>
</tr>
</tbody>
</table>

The POM if no city were to implement a gas-guzzler tax is an air pollution index of 0.39. If all cities implement a gas-guzzler tax, the air pollution index would decrease by 0.096 relative to a scenario where no city implements the tax.

Video example

Treatment effects: Regression adjustment
Stored results

t/effects ra stores the following in e():

Scalar
    e(N) number of observations
    e(nj) number of observations for treatment level j
    e(N_clust) number of clusters
    e(k_eq) number of equations in e(b)
    e(k_levels) number of levels in treatment variable
    e(treated) level of treatment variable defined as treated
    e(control) level of treatment variable defined as control
    e(converged) 1 if converged, 0 otherwise

Macro
    e(cmd) teffects
    e(cmdline) command as typed
    e(depvar) name of outcome variable
    e(tvar) name of treatment variable
    e(subcmd) ra
    e(omodel) linear, logit, probit, hetprobit, poisson, flogit, fprobit, or fhetprobit
    e(stat) statistic estimated, ate, atet, or pomeans
    e(ute) weight type
    e(wexp) weight expression
    e(title) title in estimation output
    e(clustvar) name of cluster variable
    e(tlevels) levels of treatment variable
    e(vcetype) vcetype specified in vce()
    e(properties) b V
    e(estat_cmd) program used to implement estat
    e(predict) program used to implement predict
    e(marginsnotok) predictions disallowed by margins
    e(asbalanced) factor variables fvset as asbalanced
    e(asobserved) factor variables fvset as asobserved

Matrix
    e(b) coefficient vector
    e(V) variance–covariance matrix of the estimators

Function
    e(sample) marks estimation sample

Methods and formulas

t/effects ra implements a smooth treatment-effects estimator. All smooth treatment-effects estimators are documented in [TE] teffects aipw.

References


Also see

[TE] **teffects postestimation** — Postestimation tools for teffects  
[TE] **teffects** — Treatment-effects estimation for observational data  
[ERM] **eregress** — Extended linear regression  
[U] **20 Estimation and postestimation commands**
Glossary

**AIPW estimator.** See **augmented inverse-probability-weighted estimator.**

**analysis time.** Analysis time is like time, except that 0 has a special meaning: \( t = 0 \) is the time of onset of risk, the time when failure first became possible.

Analysis time is usually not what is recorded in a dataset. A dataset of patients might record calendar time. Calendar time must then be mapped to analysis time.

The letter \( t \) is reserved for time in analysis-time units. The term time is used for time measured in other units.

The origin is the time corresponding to \( t = 0 \), which can vary subject to subject. Thus \( t = \text{time} - \text{origin} \).

**ATE.** See **average treatment effect.**

**ATET.** See **average treatment effect on the treated.**

**augmented inverse-probability-weighted estimator.** An augmented inverse-probability-weighted (AIPW) estimator is an inverse-probability-weighted estimator that includes an augmentation term that corrects the estimator when the treatment model is misspecified. When the treatment is correctly specified, the augmentation term vanishes as the sample size becomes large. An AIPW estimator uses both an outcome model and a treatment model and is a doubly robust estimator.

**average treatment effect.** The average treatment effect is the average effect of the treatment among all individuals in a population.

**average treatment effect on the treated.** The average treatment effect on the treated is the average effect of the treatment among those individuals who actually get the treatment.

**censored, left-censored, and right-censored.** An observation is left-censored when the exact time of failure is not known; it is merely known that the failure occurred before \( t_l \). Suppose that the event of interest is becoming employed. If a subject is already employed when first interviewed, his outcome is left-censored.

An observation is right-censored when the time of failure is not known; it is merely known that the failure occurred after \( t_r \). If a patient survives until the end of a study, the patient’s time of death is right-censored.

In common usage, censored without a modifier means right-censored.

Also see **truncation, left-truncation, and right-truncation.**

**CI assumption.** See **conditional-independence assumption.**

**conditional mean.** The conditional mean expresses the average of one variable as a function of some other variables. More formally, the mean of \( y \) conditional on \( x \) is the mean of \( y \) for given values of \( x \); in other words, it is \( E(y|x) \).

A conditional mean is also known as a regression or as a conditional expectation.

**conditional-independence assumption.** The conditional-independence assumption requires that the common variables that affect treatment assignment and treatment-specific outcomes be observable. The dependence between treatment assignment and treatment-specific outcomes can be removed by conditioning on these observable variables.
This assumption is also known as a selection-on-observables assumption because its central tenet is the observability of the common variables that generate the dependence.

**counterfactual.** A counterfactual is an outcome a subject would have obtained had that subject received a different level of treatment. In the binary-treatment case, the counterfactual outcome for a person who received treatment is the outcome that person would have obtained had the person instead not received treatment; similarly, the counterfactual outcome for a person who did not receive treatment is the outcome that person would have obtained had the person received treatment.

Also see *potential outcome*.

**doubly robust estimator.** A doubly robust estimator only needs one of two auxiliary models to be correctly specified to estimate a parameter of interest.

Doubly robust estimators for treatment effects are consistent when either the outcome model or the treatment model is correctly specified.

**EE estimator.** See *estimating-equation estimator*.

**estimating-equation estimator.** An estimating-equation (EE) estimator calculates parameters estimates by solving a system of equations. Each equation in this system is the sample average of a function that has mean zero.

These estimators are also known as $M$ estimators or $Z$ estimators in the statistics literature and as generalized method of moments (GMM) estimators in the econometrics literature.

**failure event.** Survival analysis is really time-to-failure analysis, and the failure event is the event under analysis. The failure event can be death, heart attack, myopia, or finding employment. Many authors—including Stata—write as if the failure event can occur only once per subject, but when we do, we are being sloppy. Survival analysis encompasses repeated failures, and all of Stata’s survival analysis features can be used with repeated-failure data.

**hazard, cumulative hazard, and hazard ratio.** The hazard or hazard rate at time $t$, $h(t)$, is the instantaneous rate of failure at time $t$ conditional on survival until time $t$. Hazard rates can exceed 1. Say that the hazard rate were 3. If an individual faced a constant hazard of 3 over a unit interval and if the failure event could be repeated, the individual would be expected to experience three failures during the time span.

The cumulative hazard, $H(t)$, is the integral of the hazard function $h(t)$, from 0 (the onset of risk) to $t$. It is the total number of failures that would be expected to occur up until time $t$, if the failure event could be repeated. The relationship between the cumulative hazard function, $H(t)$, and the survivor function, $S(t)$, is

$$S(t) = \exp\{-H(t)\}$$

$$H(t) = -\ln\{S(t)\}$$

The hazard ratio is the ratio of the hazard function evaluated at two different values of the covariates: $h(t \mid x)/h(t \mid x_0)$. The hazard ratio is often called the relative hazard, especially when $h(t \mid x_0)$ is the baseline hazard function.

**i.i.d. sampling assumption.** See *independent and identically distributed sampling assumption*.

**independent and identically distributed sampling assumption.** The independent and identically distributed (i.i.d.) sampling assumption specifies that each observation is unrelated to (independent of) all the other observations and that each observation is a draw from the same (identical) distribution.
individual-level treatment effect. An individual-level treatment effect is the difference in an individual’s outcome that would occur because this individual is given one treatment instead of another. In other words, an individual-level treatment effect is the difference between two potential outcomes for an individual.

For example, the blood pressure an individual would obtain after taking a pill minus the blood pressure an individual would obtain had that person not taken the pill is the individual-level treatment effect of the pill on blood pressure.

inverse-probability-weighted estimators. Inverse-probability-weighted (IPW) estimators use weighted averages of the observed outcome variable to estimate the potential-outcome means. The weights are the reciprocals of the treatment probabilities estimated by a treatment model.

inverse-probability-weighted regression-adjustment estimators. Inverse-probability-weighted regression-adjustment (IPWRA) estimators use the reciprocals of the estimated treatment probability as weights to estimate missing-data-corrected regression coefficients that are subsequently used to compute the potential-outcome means.

IPW estimators. See inverse-probability-weighted estimators.

IPWRA estimators. See inverse-probability-weighted regression-adjustment estimators.

left-censored. See censored, left-censored, and right-censored.

left-truncation. See truncation, left-truncation, and right-truncation.

matching estimator. An estimator that compares differences between the outcomes of similar—that is, matched—individuals. Each individual that receives a treatment is matched to a similar individual that does not get the treatment, and the difference in their outcomes is used to estimate the individual-level treatment effect. Likewise, each individual that does not receive a treatment is matched to a similar individual that does get the treatment, and the difference in their outcomes is used to estimate the individual-level treatment effect.

multiple-record st data. See st data.

multivalued treatment effect. A multivalued treatment refers to a treatment that has more than two values. For example, a person could have taken a 20 mg dose of a drug, a 40 mg dose of the drug, or not taken the drug at all.

nearest-neighbor matching. Nearest-neighbor matching uses the distance between observed variables to find similar individuals.

observational data. In observational data, treatment assignment is not controlled by those who collected the data; thus some common variables affect treatment assignment and treatment-specific outcomes.

outcome model. An outcome model is a model used to predict the outcome as a function of covariates and parameters.

overlap assumption. The overlap assumption requires that each individual have a positive probability of each possible treatment level.

POMs. See potential-outcome means.

potential outcome. The potential outcome is the outcome an individual would obtain if given a specific treatment.

For example, an individual has one potential blood pressure after taking a pill and another potential blood pressure had that person not taken the pill.

potential-outcome means. The potential-outcome means refers to the means of the potential outcomes for a specific treatment level.
The mean blood pressure if everyone takes a pill and the mean blood pressure if no one takes a pill are two examples.

The average treatment effect is the difference between potential-outcome mean for the treated and the potential-outcome mean for the not treated.

**propensity score.** The propensity score is the probability that an individual receives a treatment.

**propensity-score matching.** Propensity-score matching uses the distance between estimated propensity scores to find similar individuals.

**regression-adjustment estimators.** Regression-adjustment estimators use means of predicted outcomes for each treatment level to estimate each potential-outcome mean.

**right-censored.** See *censored, left-censored, and right-censored.*

**right-truncation.** See *truncation, left-truncation, and right-truncation.*

**selection-on-observables.** See *conditional-independence assumption.*

**shape parameter.** A shape parameter governs the shape of a probability distribution. One example is the parameter $p$ of the Weibull model.

**single-record st data.** See *st data.*

**smooth treatment-effects estimator.** A smooth treatment-effects estimator is a smooth function of the data so that standard methods approximate the distribution of the estimator. The RA, IPW, AIPW, and IPWRA estimators are all smooth treatment-effects estimators while the nearest-neighbor matching estimator and the propensity-score matching estimator are not.

**st data.** st stands for survival time. In survival-time data, each observation represents a span of survival, recorded in variables $t_0$ and $t$. For instance, if in an observation $t_0$ were 3 and $t$ were 5, the span would be $(t_0, t]$, meaning from just after $t_0$ up to and including $t$.

Sometimes variable $t_0$ is not recorded; $t_0$ is then assumed to be 0. In such a dataset, an observation that had $t = 5$ would record the span $(0, 5]$.

Each observation also includes a variable $d$, called the failure variable, which contains 0 or nonzero (typically, 1). The failure variable records what happened at the end of the span: 0, the subject was still alive (had not yet failed) or 1, the subject died (failed).

Sometimes variable $d$ is not recorded; $d$ is then assumed to be 1. In such a dataset, all time-span observations would be assumed to end in failure.

Finally, each observation in an st dataset can record the entire history of a subject or each can record a part of the history. In the latter case, groups of observations record the full history. One observation might record the period $(0, 5]$ and the next, $(5, 8]$. In such cases, there is a variable ID that records the subject for which the observation records a time span. Such data are called multiple-record st data. When each observation records the entire history of a subject, the data are called single-record st data. In the single-record case, the ID variable is optional.

See [ST] *stset.*

**survival-time data.** See *st data.*

**survivor function.** Also known as the survivorship function and the survival function, the survivor function, $S(t)$, is 1) the probability of surviving beyond time $t$, or equivalently, 2) the probability that there is no failure event prior to $t$, 3) the proportion of the population surviving to time $t$, or equivalently, 4) the reverse cumulative distribution function of $T$, the time to the failure event: $S(t) = \Pr(T > t)$. Also see *hazard, cumulative hazard, and hazard ratio.*
treatment model. A treatment model is a model used to predict treatment-assignment probabilities as a function of covariates and parameters.

truncation, left-truncation, and right-truncation. In survival analysis, truncation occurs when subjects are observed only if their failure times fall within a certain observational period of a study. Censoring, on the other hand, occurs when subjects are observed for the whole duration of a study, but the exact times of their failures are not known; it is known only that their failures occurred within a certain time span.

Left-truncation occurs when subjects come under observation only if their failure times exceed some time $t_l$. It is only because they did not fail before $t_l$ that we even knew about their existence. Left-truncation differs from left-censoring in that, in the censored case, we know that the subject failed before time $t_l$, but we just do not know exactly when.

Imagine a study of patient survival after surgery, where patients cannot enter the sample until they have had a post-surgical test. The patients’ survival times will be left-truncated. This is a “delayed entry” problem, one common type of left-truncation.

Right-truncation occurs when subjects come under observation only if their failure times do not exceed some time $t_r$. Right-truncated data typically occur in registries. For example, a cancer registry includes only subjects who developed a cancer by a certain time, and thus survival data from this registry will be right-truncated.

unconfoundedness. See conditional-independence assumption.

weighted-regression-adjustment estimator. Weighted-regression-adjustment estimators use means of predicted outcomes for each treatment level to estimate each potential-outcome mean. The weights are used to estimate censoring-adjusted regression coefficients.
Subject and author index

See the combined subject index and the combined author index in the Glossary and Index.