

tesensitivity: A Stata Package for Assessing Sensitivity to the Unconfoundedness Assumption

Linqi Zhang* Paul Diegert† Matthew A. Masten‡ Alexandre Poirier§

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Introduction

A standard question in causal inference is to identify and estimate the effect of some treatment variable X on an outcome variable Y . A common assumption used to identify such effects is *unconfoundedness*, also known as selection on observables, conditional independence, ignorability, or exogenous selection. This assumption is not refutable, meaning that the data alone cannot tell us whether it is true. Nonetheless, researchers may wonder: How important is this assumption in their analyses? Put differently: How sensitive are results obtained under the unconfoundedness assumption?

Masten and Poirier (2018) provide a set of theoretical results used to answer this question. They define conditional *partial* independence as any assumption weaker than full conditional independence. They specifically consider a class of assumptions called *conditional c-dependence*. This class measures relaxations of conditional independence by a single parameter $c \in [0, 1]$. When $c = 0$, this corresponds to the conditional independence assumption. For any $c > 0$, conditional independence only partially holds, and hence we cannot exactly learn the value of our treatment effect parameters, such as the average treatment effect (ATE). Instead, we only get bounds. Masten and Poirier (2018) characterize these bounds as a function of c . Small values of c give narrow bounds while larger values of c give wider bounds. Just how wide these bounds are, and hence how sensitive one's results are, depends on the data.

Specifically, define the following random variables:

- Y_x : the potential outcome for a given treatment $x \in \{0, 1\}$
- X : the treatment
- W : a vector of covariates
- Y : the observed outcome

The observed outcome satisfies

$$Y = (1 - X)Y_0 + XY_1.$$

Rather than observing the full data generating process, (Y_0, Y_1, X, Y) , we only observe (Y, X, W) .

We say that X is *conditionally c-dependent* with Y_x given W if:

$$\sup_{y_x \in \text{supp}(Y_x|W=w)} |\mathbb{P}(X = 1|Y_x = y_x, W = w) - \mathbb{P}(X = 1|W = w)| \leq c$$

*Department of Economics, Boston College, linqi.zhang@bc.edu

†Department of Economics, Duke University, paul.diegert@duke.edu

‡Department of Economics, Duke University, matt.masten@duke.edu

§Department of Economics, Georgetown University, alexandre.poirier@georgetown.edu

for all $w \in \text{supp}(W)$.

Under this assumption, the identified set for a treatment effect statistic will be a closed interval, which depends on c and the distribution of (Y, X, W) . The main purpose of the **tesensitivity** package is to calculate these intervals and show how the identified set for treatment effect statistics varies with the sensitivity parameter c .

In addition to estimating these bounds for a range of values of c , this package also calculates a *breakdown point* relative to a conclusion about a treatment effect statistic. As discussed in Masten and Poirier (2020), the *breakdown point* is the maximum value of c under which the conclusion still holds. For example, if we consider the conclusion that the average treatment effect is positive, then the breakdown point is the minimum value of c such that 0 is included in the identified set.

The package also includes tools to visualize the analysis, interpret the scale of c -dependence, and compare results of multiple sensitivity analyses.

This vignette provides a tutorial for the package based on the empirical illustration in Masten, Poirier, and Zhang (2020).

Getting Started

We will illustrate usage of the package using data from LaLonde (1986) as reconstructed by Dehejia and Wahba (1999) and used in the empirical illustration in Masten, Poirier, and Zhang (2020).

The dataset is included with the package, and can be loaded using the `sysuse` command:

```
. sysuse lalonde1986, clear
```

These will be the variables used in the analysis:

```
. local outcome "re78"
. local treatment "treat"
. local controls "married age black hispanic education re74 re75 re74pos re75pos"
```

The dataset includes all of the samples referenced in Masten, Poirier, and Zhang (2020). `sample1` flags the experimental sample using the treatment and control groups from the National Supported Work (NSW) demonstration project. `sample3` flags the sample using the NSW treatment group and a control group constructed from the PSID and trimmed to exclude workers with earnings above \$5,000 3 or 4 years before the program began. (`sample2` is the same as `sample3` but without trimming.)

For reference, we start by estimating the average treatment effect of these experimental and non-experimental samples using the **teffects** package.

For the experimental sample:

```
. teffects ipw (`outcome`) (`treatment` `controls`) if sample1
Iteration 0:  EE criterion = 1.061e-15
Iteration 1:  EE criterion = 5.245e-23

Treatment-effects estimation      Number of obs      =      445
Estimator      : inverse-probability weights
Outcome model  : weighted mean
Treatment model: logit
```

| | re78 | Coef. | Robust Std. Err. | z | P> z | [95% Conf. Interval] |
|-------------------|------|---------|---------------------|------|-------|----------------------|
| ATE | | | | | | |
| treat (1 vs 0) | | 1632.93 | 649.886 | 2.51 | 0.012 | 359.1766 2906.683 |

| | | | | | | |
|--------|----------|----------|-------|-------|----------|---------|
| P0mean | | | | | | |
| treat | | | | | | |
| 0 | 4578.176 | 343.7891 | 13.32 | 0.000 | 3904.362 | 5251.99 |

For the trimmed observational sample:

```
. teffects ipw (`outcome`) (`treatment` `controls`) if sample3
Iteration 0: EE criterion = 3.624e-16
Iteration 1: EE criterion = 2.284e-22
Treatment-effects estimation      Number of obs      =      390
Estimator      : inverse-probability weights
Outcome model   : weighted mean
Treatment model : logit
```

| | | Coef. | Robust Std. Err. | z | P> z | [95% Conf. Interval] |
|----------|--|----------|---------------------|------|-------|----------------------|
| re78 | | | | | | |
| ATE | | | | | | |
| treat | | | | | | |
| (1 vs 0) | | 3336.676 | 769.1799 | 4.34 | 0.000 | 1829.112 4844.241 |
| P0mean | | | | | | |
| treat | | | | | | |
| 0 | | 2868.201 | 331.8332 | 8.64 | 0.000 | 2217.82 3518.582 |

Under the standard unconfoundedness assumption, the treatment effect is positive and statistically significant in both samples. Now we will use the **tesensitivity** package to analyze how sensitive these results are to this assumption.

The main subcommand of the **tesensitivity** package is **cp**i, i.e., conditional partial independence. This command estimates bounds on treatment effect statistics given a set of *c*-dependence values and calculates the breakdown point for the conclusion that the treatment effect statistic is above a given threshold.

First, we calculate bounds on the ATE. By default, the command calculates bounds for a uniform grid of 40 values of *c*, and the breakdown point for the conclusion that $ATE > \{ \}$ 0.

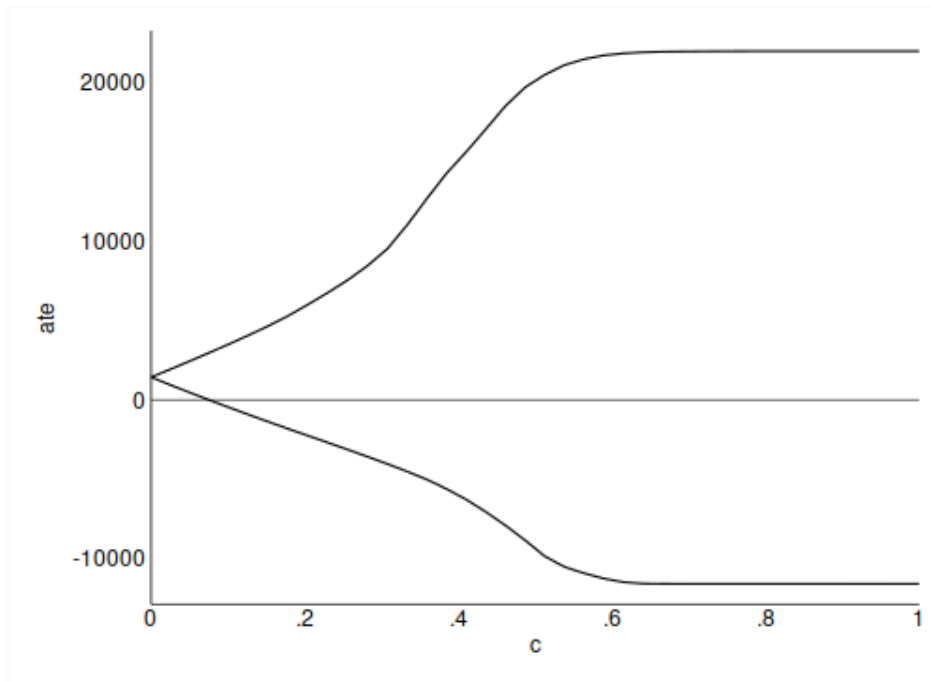
```
. tesensitivity cpi (`outcome` `controls`) (`treatment` `controls`) if sample1, ate
Treatment effects sensitivity
Analysis      : cond. partial independence  Number of obs =      445
Outcome model : linear quantile             Breakdown   =      0.075
Treatment model : logistic                  Conclusion   =      ate > 0
Outcome variable : re78
```

| c | ate |
|-------|-------------------|
| 0.000 | [1,419, 1,419] |
| 0.026 | [920, 1,930] |
| 0.051 | [433, 2,454] |
| 0.077 | [-43, 2,986] |
| 0.103 | [-508, 3,529] |
| 0.128 | [-963, 4,091] |
| 0.154 | [-1,411, 4,674] |
| 0.179 | [-1,851, 5,313] |
| 0.205 | [-2,282, 6,023] |
| 0.231 | [-2,714, 6,765] |
| 0.256 | [-3,150, 7,554] |
| 0.282 | [-3,594, 8,449] |
| 0.308 | [-4,047, 9,513] |
| 0.333 | [-4,520, 11,004] |

| | |
|-------|--------------------|
| 0.359 | [-5,026, 12,662] |
| 0.385 | [-5,616, 14,257] |
| 0.410 | [-6,291, 15,611] |
| 0.436 | [-7,067, 17,048] |
| 0.462 | [-7,926, 18,514] |
| 0.487 | [-8,862, 19,680] |
| 0.513 | [-9,871, 20,477] |
| 0.538 | [-10,532, 21,081] |
| 0.564 | [-10,934, 21,454] |
| 0.590 | [-11,265, 21,687] |
| 0.615 | [-11,489, 21,811] |
| 0.641 | [-11,582, 21,881] |
| 0.667 | [-11,591, 21,915] |
| 0.692 | [-11,594, 21,928] |
| 0.718 | [-11,594, 21,937] |
| 0.744 | [-11,594, 21,945] |
| 0.769 | [-11,594, 21,950] |
| 0.795 | [-11,594, 21,951] |
| 0.821 | [-11,594, 21,951] |
| 0.846 | [-11,594, 21,952] |
| 0.872 | [-11,594, 21,952] |
| 0.897 | [-11,594, 21,952] |
| 0.923 | [-11,594, 21,952] |
| 0.949 | [-11,594, 21,952] |
| 0.974 | [-11,594, 21,952] |
| 1.000 | [-11,594, 21,952] |

We can see this graphically using the `cpiplot` subcommand, which can be called after `tesensitivity cpi`.

```
. tesensitivity cpiplot
```



By default, this command plots a horizontal line for the lower bound in the conclusion used for the breakdown point computation.

Interpreting the c -dependence parameter

In our ATE example, the bounds get very wide for moderately large values of c , but it isn't immediately clear what would be a reasonable assumption to maintain about the value of c . The `cscale` subcommand provides tools to interpret the scale of c -dependence.

The main tool provided in this subcommand is the leave-one-out analysis discussed in section 2 of Masten and Poirier (2018). Also see section 2 of Masten, Poirier, and Zhang (2020). This compares the propensity scores obtained when leaving out each covariate in the analysis compared to the propensity scores including all covariates. This provides a reference point for thinking about the level of c -dependence that might be caused by other omitted variables.

Since the deviation in propensity scores varies across the values of the covariate left out, this induces a distribution of deviations. The `cscale` subcommand provides options to calculate the supremum over this distribution and quantiles of the distribution for each covariate.

By default, `tesensitivity cscale` calculates the maximum and the 50th, 75th, and 90th percentiles of the distribution for each covariate.

```
. tesensitivity cscale
```

| | | | | |
|--|-----------------------------------|---------------|---|-----|
| Treatment-effects sensitivity analysis | | | | |
| Analysis | : leave one out prop. score diff. | Number of obs | = | 445 |
| Treatment model | : logistic | | | |

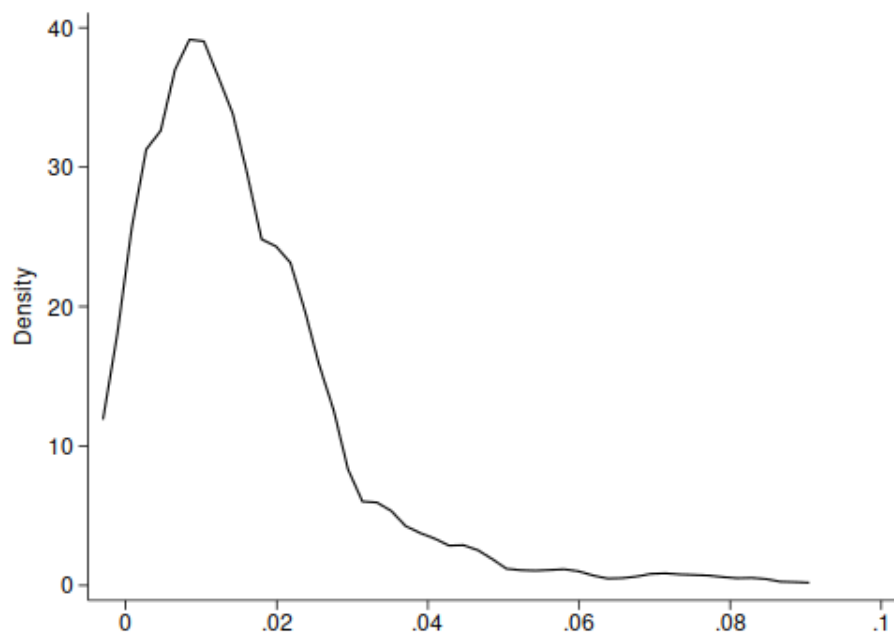
| quantile | 0.500 | 0.750 | 0.900 | max |
|-----------|-------|-------|-------|-------|
| married | 0.006 | 0.012 | 0.032 | 0.042 |
| age | 0.015 | 0.024 | 0.034 | 0.099 |
| black | 0.007 | 0.009 | 0.014 | 0.082 |
| hispanic | 0.007 | 0.017 | 0.099 | 0.124 |
| education | 0.012 | 0.022 | 0.031 | 0.087 |
| re74 | 0.002 | 0.011 | 0.035 | 0.209 |
| re75 | 0.001 | 0.004 | 0.008 | 0.053 |
| re74pos | 0.002 | 0.010 | 0.018 | 0.034 |
| re75pos | 0.013 | 0.017 | 0.062 | 0.082 |

Other quantiles can be calculated with the `quantiles` option.

Note that this command uses the specification of the last call to `tesensitivity cpi`. So it uses the covariates and the sample selection criteria given in the previous call.

We also can plot the density of propensity score deviations in the leave-one-out analysis. For example, for education, we have:

```
. tesensitivity cscale education, density
```



To see the bounds on the ATE at the maximum deviations for each covariate, use the `reference` option.

```
. tesensitivity cpi (`outcome` `controls`) (`treatment` `controls`) if sample1, ate cref
```

Treatment effects sensitivity

```
Analysis      : cond. partial independence  Number of obs =      445
Outcome model  : linear quantile             Breakdown    =      0.075
Treatment model : logistic                   Conclusion   =      ate > 0
Outcome variable : re78
```

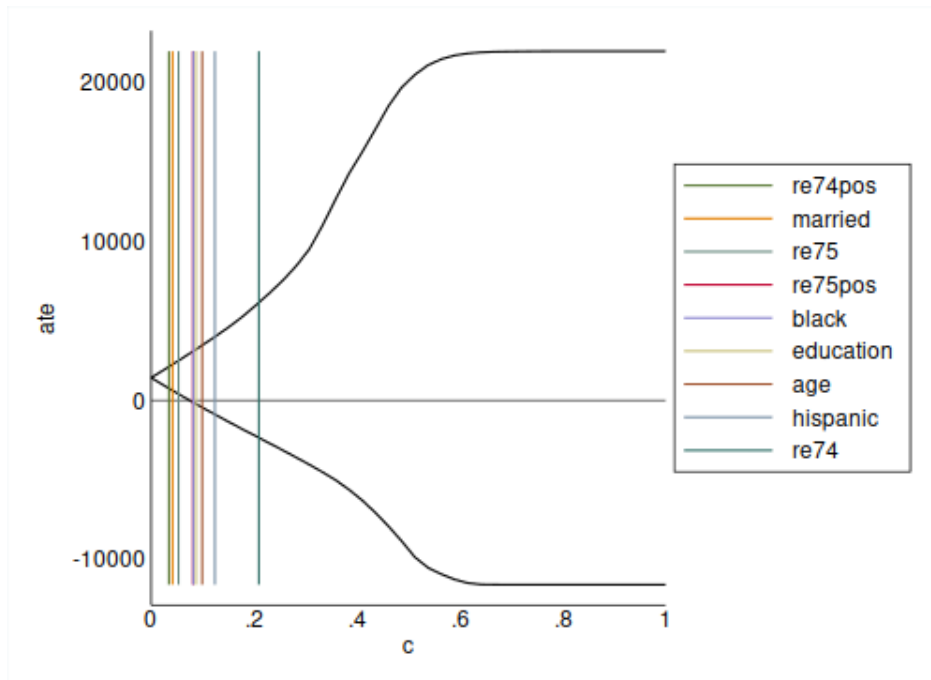
| | c | ate |
|-----------|-------|-------------------|
| | 0.000 | [1,419, 1,419] |
| | 0.026 | [920, 1,930] |
| re74pos | 0.034 | [751, 2,109] |
| married | 0.042 | [616, 2,254] |
| | 0.051 | [433, 2,454] |
| re75 | 0.053 | [398, 2,492] |
| | 0.077 | [-43, 2,986] |
| re75pos | 0.082 | [-128, 3,083] |
| black | 0.082 | [-134, 3,090] |
| education | 0.087 | [-234, 3,206] |
| age | 0.099 | [-448, 3,458] |
| | 0.103 | [-508, 3,529] |
| hispanic | 0.124 | [-888, 3,996] |
| | 0.128 | [-963, 4,091] |
| | 0.154 | [-1,411, 4,674] |
| | 0.179 | [-1,851, 5,313] |
| | 0.205 | [-2,282, 6,023] |
| re74 | 0.209 | [-2,354, 6,146] |
| | 0.231 | [-2,714, 6,765] |
| | 0.256 | [-3,150, 7,554] |
| | 0.282 | [-3,594, 8,449] |
| | 0.308 | [-4,047, 9,513] |
| | 0.333 | [-4,520, 11,004] |
| | 0.359 | [-5,026, 12,662] |
| | 0.385 | [-5,616, 14,257] |

| | |
|-------|--------------------|
| 0.410 | [-6,291, 15,611] |
| 0.436 | [-7,067, 17,048] |
| 0.462 | [-7,926, 18,514] |
| 0.487 | [-8,862, 19,680] |
| 0.513 | [-9,871, 20,477] |
| 0.538 | [-10,532, 21,081] |
| 0.564 | [-10,934, 21,454] |
| 0.590 | [-11,265, 21,687] |
| 0.615 | [-11,489, 21,811] |
| 0.641 | [-11,582, 21,881] |
| 0.667 | [-11,591, 21,915] |
| 0.692 | [-11,594, 21,928] |
| 0.718 | [-11,594, 21,937] |
| 0.744 | [-11,594, 21,945] |
| 0.769 | [-11,594, 21,950] |
| 0.795 | [-11,594, 21,951] |
| 0.821 | [-11,594, 21,951] |
| 0.846 | [-11,594, 21,952] |
| 0.872 | [-11,594, 21,952] |
| 0.897 | [-11,594, 21,952] |
| 0.923 | [-11,594, 21,952] |
| 0.949 | [-11,594, 21,952] |
| 0.974 | [-11,594, 21,952] |
| 1.000 | [-11,594, 21,952] |

This table includes the values of c -dependence equal to the maximum propensity scores deviation for each of the covariates, which are displayed on the left of the table.

We can also plot these with a line for the maximum deviation of propensity scores for each covariate.

```
. tesensitivity cpiplot, creflines
```



Note that covariates are ordered in the legend by their maximum propensity score deviation.

Storing and comparing estimates

Results from calls to `tesensitivity cpi` are stored in `e()`, and are compatible with several built-in postestimation commands to store and combine results.

First, to see descriptive statistics of the variables used in a call to `tesensitivity cpi`, we can use the `estat summarize` command:

```
. estat summarize
```

| | | | | |
|---------------------------------|--|-----------------|--|-----|
| Estimation sample tesensitivity | | Number of obs = | | 445 |
|---------------------------------|--|-----------------|--|-----|

| Variable | Mean | Std. Dev. | Min | Max |
|-----------|----------|-----------|-----|----------|
| re78 | 5300.764 | 6631.492 | 0 | 60307.93 |
| treat | .4157303 | .4934022 | 0 | 1 |
| married | .1685393 | .3747658 | 0 | 1 |
| age | 25.37079 | 7.100282 | 17 | 55 |
| black | .8337079 | .3727617 | 0 | 1 |
| hispanic | .0876404 | .2830895 | 0 | 1 |
| education | 10.19551 | 1.792119 | 3 | 16 |
| re74 | 2102.265 | 5363.582 | 0 | 39570.68 |
| re75 | 1377.138 | 3150.961 | 0 | 25142.24 |
| re74pos | .2674157 | .4431092 | 0 | 1 |
| re75pos | .3505618 | .4776829 | 0 | 1 |

Note that a matrix of the covariates is also saved in `e(covsupp)`.

We can also store results using `estimates store` and see the output from any stored results with `replay`.

```
. qui tesensitivity cpi (`outcome' `controls') (`treatment' `controls') if sample1, ate
. estimates store atel
. estimates replay atel
```

Model atel

Treatment effects sensitivity

| | | | | |
|------------------|------------------------------|---------------|---|---------|
| Analysis | : cond. partial independence | Number of obs | = | 445 |
| Outcome model | : linear quantile | Breakdown | = | 0.075 |
| Treatment model | : logistic | Conclusion | = | ate > 0 |
| Outcome variable | : re78 | | | |

| c | ate |
|-------|-------------------|
| 0.000 | [1,419, 1,419] |
| 0.026 | [920, 1,930] |
| 0.051 | [433, 2,454] |
| 0.077 | [-43, 2,986] |
| 0.103 | [-508, 3,529] |
| 0.128 | [-963, 4,091] |
| 0.154 | [-1,411, 4,674] |
| 0.179 | [-1,851, 5,313] |
| 0.205 | [-2,282, 6,023] |
| 0.231 | [-2,714, 6,765] |
| 0.256 | [-3,150, 7,554] |
| 0.282 | [-3,594, 8,449] |
| 0.308 | [-4,047, 9,513] |
| 0.333 | [-4,520, 11,004] |
| 0.359 | [-5,026, 12,662] |
| 0.385 | [-5,616, 14,257] |
| 0.410 | [-6,291, 15,611] |
| 0.436 | [-7,067, 17,048] |

| | |
|-------|-------------------|
| 0.462 | [-7,926, 18,514] |
| 0.487 | [-8,862, 19,680] |
| 0.513 | [-9,871, 20,477] |
| 0.538 | [-10,532, 21,081] |
| 0.564 | [-10,934, 21,454] |
| 0.590 | [-11,265, 21,687] |
| 0.615 | [-11,489, 21,811] |
| 0.641 | [-11,582, 21,881] |
| 0.667 | [-11,591, 21,915] |
| 0.692 | [-11,594, 21,928] |
| 0.718 | [-11,594, 21,937] |
| 0.744 | [-11,594, 21,945] |
| 0.769 | [-11,594, 21,950] |
| 0.795 | [-11,594, 21,951] |
| 0.821 | [-11,594, 21,951] |
| 0.846 | [-11,594, 21,952] |
| 0.872 | [-11,594, 21,952] |
| 0.897 | [-11,594, 21,952] |
| 0.923 | [-11,594, 21,952] |
| 0.949 | [-11,594, 21,952] |
| 0.974 | [-11,594, 21,952] |
| 1.000 | [-11,594, 21,952] |

.

We can also compare multiple sets of estimates. To illustrate, we'll compare the results from the experimental sample stored in `ate1` with results from the observational, trimmed sample.

```
. tesensitivity cpi (`outcome` `controls`) (`treatment` `controls`) if sample3, ate
```

Treatment effects sensitivity

```
Analysis      : cond. partial independence  Number of obs =      390
Outcome model  : linear quantile             Breakdown    =      0.020
Treatment model : logistic                   Conclusion    =      ate > 0
Outcome variable : re78
```

| c | ate |
|-------|-------------------|
| 0.000 | [2,865, 2,865] |
| 0.026 | [-663, 11,806] |
| 0.051 | [-2,879, 15,385] |
| 0.077 | [-4,389, 17,794] |
| 0.103 | [-5,356, 19,300] |
| 0.128 | [-6,121, 20,318] |
| 0.154 | [-6,764, 21,055] |
| 0.179 | [-7,296, 21,687] |
| 0.205 | [-7,746, 22,256] |
| 0.231 | [-8,232, 22,747] |
| 0.256 | [-8,678, 23,234] |
| 0.282 | [-9,092, 23,642] |
| 0.308 | [-9,485, 23,984] |
| 0.333 | [-9,798, 24,278] |
| 0.359 | [-10,127, 24,599] |
| 0.385 | [-10,438, 24,905] |
| 0.410 | [-10,752, 25,231] |
| 0.436 | [-10,993, 25,478] |
| 0.462 | [-11,247, 25,734] |
| 0.487 | [-11,463, 25,929] |
| 0.513 | [-11,692, 26,126] |
| 0.538 | [-11,935, 26,314] |
| 0.564 | [-12,120, 26,469] |
| 0.590 | [-12,327, 26,655] |
| 0.615 | [-12,465, 26,788] |
| 0.641 | [-12,602, 26,900] |
| 0.667 | [-12,711, 27,007] |

| | | |
|-------|-----------|---------|
| 0.692 | [-12,834, | 27,071] |
| 0.718 | [-12,936, | 27,130] |
| 0.744 | [-13,044, | 27,189] |
| 0.769 | [-13,114, | 27,241] |
| 0.795 | [-13,175, | 27,275] |
| 0.821 | [-13,242, | 27,302] |
| 0.846 | [-13,316, | 27,326] |
| 0.872 | [-13,345, | 27,348] |
| 0.897 | [-13,358, | 27,368] |
| 0.923 | [-13,367, | 27,383] |
| 0.949 | [-13,374, | 27,392] |
| 0.974 | [-13,376, | 27,397] |
| 1.000 | [-13,376, | 27,399] |

```
. estimates store ate3
```

The `tesensitivity cpitable` command provides a simple way to compare results:

```
. tesensitivity cpitable ate1 ate3
```

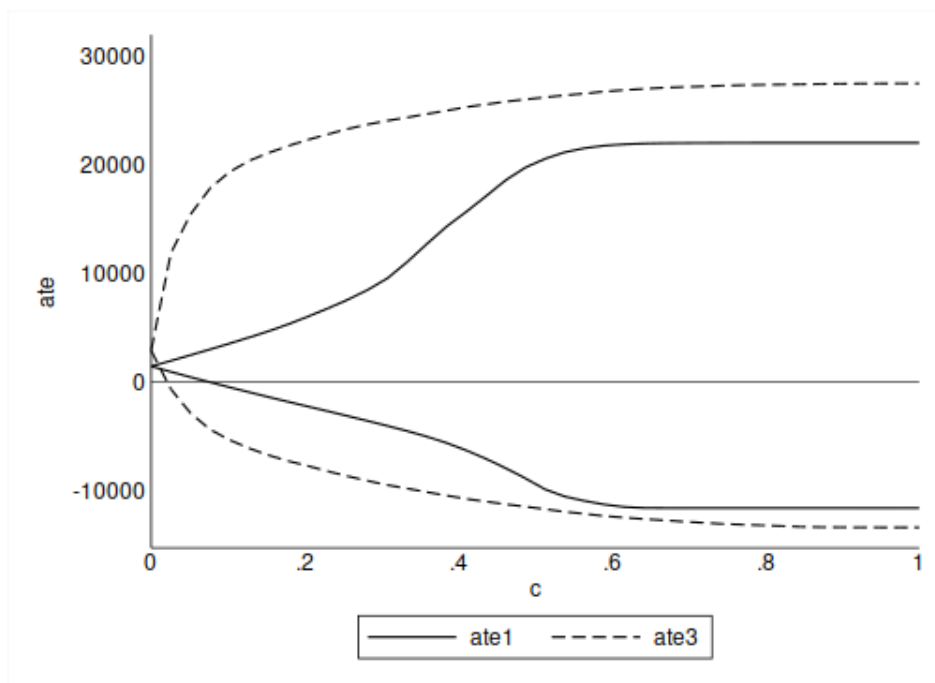
Treatment effects sensitivity
Analysis : conditional partial independence, multiple results

| c | ate1 | ate3 |
|-------|--------------------|--------------------|
| 0.000 | [1,419, 1,419] | [2,865, 2,865] |
| 0.026 | [920, 1,930] | [-663, 11,806] |
| 0.051 | [433, 2,454] | [-2,879, 15,385] |
| 0.077 | [-43, 2,986] | [-4,389, 17,794] |
| 0.103 | [-508, 3,529] | [-5,356, 19,300] |
| 0.128 | [-963, 4,091] | [-6,121, 20,318] |
| 0.154 | [-1,411, 4,674] | [-6,764, 21,055] |
| 0.179 | [-1,851, 5,313] | [-7,296, 21,687] |
| 0.205 | [-2,282, 6,023] | [-7,746, 22,256] |
| 0.231 | [-2,714, 6,765] | [-8,232, 22,747] |
| 0.256 | [-3,150, 7,554] | [-8,678, 23,234] |
| 0.282 | [-3,594, 8,449] | [-9,092, 23,642] |
| 0.308 | [-4,047, 9,513] | [-9,485, 23,984] |
| 0.333 | [-4,520, 11,004] | [-9,798, 24,278] |
| 0.359 | [-5,026, 12,662] | [-10,127, 24,599] |
| 0.385 | [-5,616, 14,257] | [-10,438, 24,905] |
| 0.410 | [-6,291, 15,611] | [-10,752, 25,231] |
| 0.436 | [-7,067, 17,048] | [-10,993, 25,478] |
| 0.462 | [-7,926, 18,514] | [-11,247, 25,734] |
| 0.487 | [-8,862, 19,680] | [-11,463, 25,929] |
| 0.513 | [-9,871, 20,477] | [-11,692, 26,126] |
| 0.538 | [-10,532, 21,081] | [-11,935, 26,314] |
| 0.564 | [-10,934, 21,454] | [-12,120, 26,469] |
| 0.590 | [-11,265, 21,687] | [-12,327, 26,655] |
| 0.615 | [-11,489, 21,811] | [-12,465, 26,788] |
| 0.641 | [-11,582, 21,881] | [-12,602, 26,900] |
| 0.667 | [-11,591, 21,915] | [-12,711, 27,007] |
| 0.692 | [-11,594, 21,928] | [-12,834, 27,071] |
| 0.718 | [-11,594, 21,937] | [-12,936, 27,130] |
| 0.744 | [-11,594, 21,945] | [-13,044, 27,189] |
| 0.769 | [-11,594, 21,950] | [-13,114, 27,241] |
| 0.795 | [-11,594, 21,951] | [-13,175, 27,275] |
| 0.821 | [-11,594, 21,951] | [-13,242, 27,302] |
| 0.846 | [-11,594, 21,952] | [-13,316, 27,326] |
| 0.872 | [-11,594, 21,952] | [-13,345, 27,348] |
| 0.897 | [-11,594, 21,952] | [-13,358, 27,368] |
| 0.923 | [-11,594, 21,952] | [-13,367, 27,383] |
| 0.949 | [-11,594, 21,952] | [-13,374, 27,392] |
| 0.974 | [-11,594, 21,952] | [-13,376, 27,397] |
| 1.000 | [-11,594, 21,952] | [-13,376, 27,399] |

| Analysis : breakdown point, multiple | | |
|--------------------------------------|-------|-------|
| conclusion | ate1 | ate3 |
| stat > 0 | 0.075 | 0.020 |

We can also see this graphically by passing multiple results to `tesensitivity cpiplot`

```
. tesensitivity cpiplot ate1 ate3
```



Other treatment effect statistics

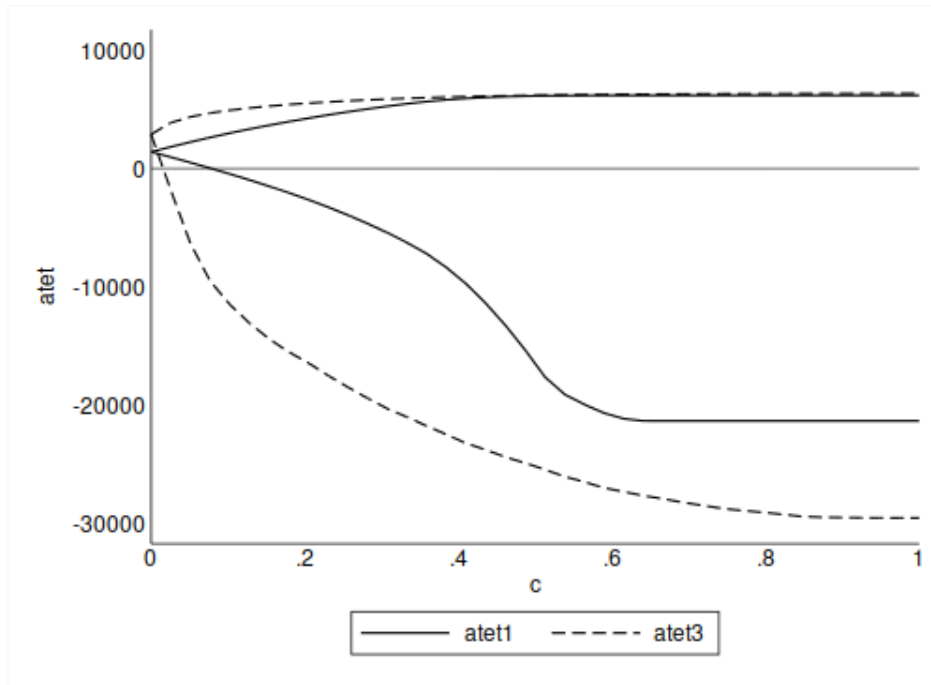
The `tesensitivity` package includes support for five treatment effect statistics:

1. Average treatment effect (ATE)
2. Average treatment effect on the treated (ATET)
3. Quantile treatment effect (QTE)
4. Conditional average treatment effect (CATE)
5. Conditional quantile treatment effect (CQTE)

Like the ATE, bounds on the ATET can be calculated without further options. To illustrate, let's compare the sensitivity of ATET in the two samples graphically.

```
. qui tesensitivity cpi (`outcome` `controls`) (`treatment` `controls`) if sample1, atet
. qui estimates store atet1
. qui tesensitivity cpi (`outcome` `controls`) (`treatment` `controls`) if sample3, atet
. qui estimates store atet3
. tesensitivity cpiplot atet1 atet3
```

```
. qui graph export ATET_compare.png, width(500) replace
```



For QTE, we also have to specify the quantile using the `quantiles` option. The median is used by default. Let's compare the QTE at the median to the ATE for the experimental sample:

```
. qui tesensitivity cpi (`outcome` `controls`) (`treatment` `controls`) if sample1, qte
. qui estimates store qte1_50
. tesensitivity cpitable atel qte1_50
```

Treatment effects sensitivity
Analysis : conditional partial independence, multiple results

| c | atel | qte1_50 |
|-------|--------------------|--------------------|
| 0.000 | [1,419, 1,419] | [988, 988] |
| 0.026 | [920, 1,930] | [321, 2,009] |
| 0.051 | [433, 2,454] | [-541, 2,857] |
| 0.077 | [-43, 2,986] | [-1,051, 3,835] |
| 0.103 | [-508, 3,529] | [-1,989, 4,724] |
| 0.128 | [-963, 4,091] | [-3,036, 6,029] |
| 0.154 | [-1,411, 4,674] | [-3,497, 6,755] |
| 0.179 | [-1,851, 5,313] | [-4,082, 7,877] |
| 0.205 | [-2,282, 6,023] | [-4,824, 8,882] |
| 0.231 | [-2,714, 6,765] | [-5,209, 9,643] |
| 0.256 | [-3,150, 7,554] | [-6,073, 9,973] |
| 0.282 | [-3,594, 8,449] | [-6,910, 10,784] |
| 0.308 | [-4,047, 9,513] | [-7,171, 11,177] |
| 0.333 | [-4,520, 11,004] | [-7,345, 12,558] |
| 0.359 | [-5,026, 12,662] | [-7,643, 60,308] |
| 0.385 | [-5,616, 14,257] | [-8,271, 60,308] |
| 0.410 | [-6,291, 15,611] | [-8,795, 60,308] |
| 0.436 | [-7,067, 17,048] | [-9,379, 60,308] |
| 0.462 | [-7,926, 18,514] | [-9,737, 60,308] |
| 0.487 | [-8,862, 19,680] | [-10,211, 60,308] |
| 0.513 | [-9,871, 20,477] | [-10,313, 60,308] |
| 0.538 | [-10,532, 21,081] | [-60,308, 60,308] |

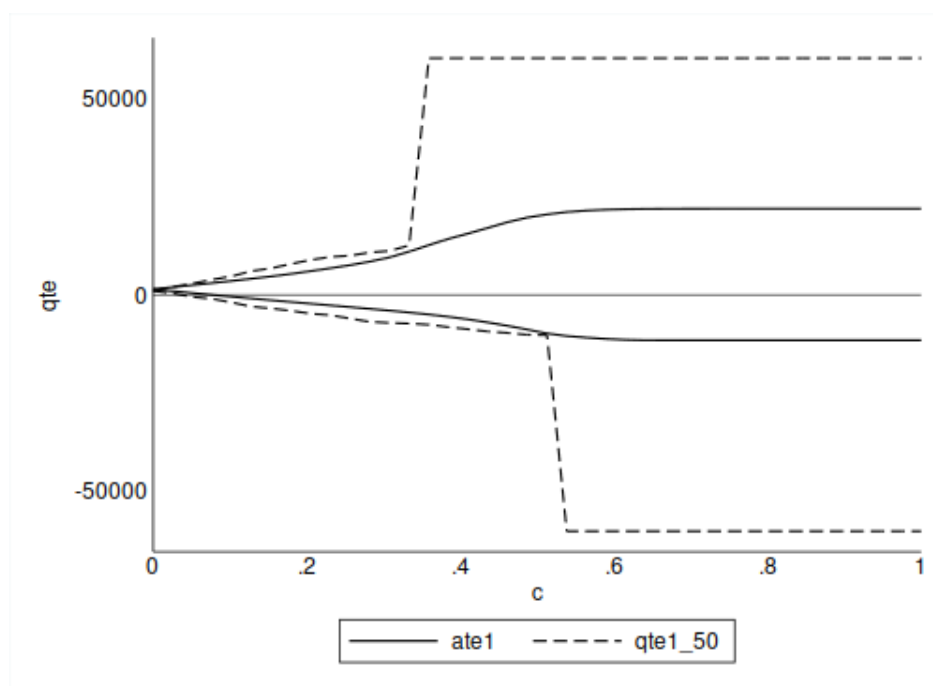
| | | |
|-------|-------------------|-------------------|
| 0.564 | [-10,934, 21,454] | [-60,308, 60,308] |
| 0.590 | [-11,265, 21,687] | [-60,308, 60,308] |
| 0.615 | [-11,489, 21,811] | [-60,308, 60,308] |
| 0.641 | [-11,582, 21,881] | [-60,308, 60,308] |
| 0.667 | [-11,591, 21,915] | [-60,308, 60,308] |
| 0.692 | [-11,594, 21,928] | [-60,308, 60,308] |
| 0.718 | [-11,594, 21,937] | [-60,308, 60,308] |
| 0.744 | [-11,594, 21,945] | [-60,308, 60,308] |
| 0.769 | [-11,594, 21,950] | [-60,308, 60,308] |
| 0.795 | [-11,594, 21,951] | [-60,308, 60,308] |
| 0.821 | [-11,594, 21,951] | [-60,308, 60,308] |
| 0.846 | [-11,594, 21,952] | [-60,308, 60,308] |
| 0.872 | [-11,594, 21,952] | [-60,308, 60,308] |
| 0.897 | [-11,594, 21,952] | [-60,308, 60,308] |
| 0.923 | [-11,594, 21,952] | [-60,308, 60,308] |
| 0.949 | [-11,594, 21,952] | [-60,308, 60,308] |
| 0.974 | [-11,594, 21,952] | [-60,308, 60,308] |
| 1.000 | [-11,594, 21,952] | [-60,308, 60,308] |

Analysis : breakdown point, multiple

| conclusion | ate1 | qte1_50 |
|------------|-------|---------|
| stat > 0 | 0.075 | 0.038 |

. tesensitivity cpiplot ate1 qte1_50

And graphically:



The two conditional statistics, CATE and CQTE, are evaluated at a single point in the support of the covariates. By default the mean of each covariate is used. The `median` option can be selected to use median values instead.

There are two ways to specify a custom covariate value: it can be specified directly using the `covariates` option, or quantiles of the covariates can be specified using the `qcovariates` option.

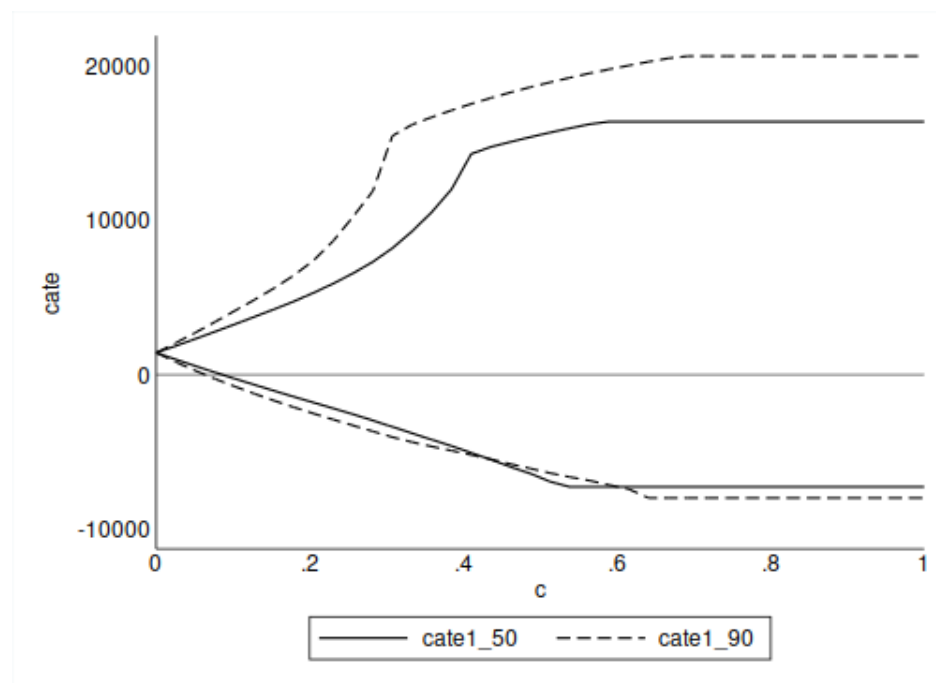
Both options can accept input in three forms:

1. a list of values separated with spaces.
2. a $1 \times K$ unlabeled matrix.
3. a $1 \times S$ ($S \leq K$) matrix with column names corresponding to variable names.

If formats 1 or 2 are used, values must be provided for all covariates and they must be in the same order as they were in the call to `tesensitivity cpi`. If format 3 is used, it is not necessary to specify the support for all covariates. Covariates not included will be evaluated at the mean by default or at the median if the `median` option is selected. If format 3 is used, it is also possible to use both the `covariates` and `qcovariates` options for different covariates.

In this example, we'll compare the CATE for married people at the median income to the CATE for unmarried people at the 90th percentile of earnings in 1974. For both analyses, all other covariates are set at their median values.

```
. matrix cov = (1)
. matrix colnames cov = married
. qui tesensitivity cpi (`outcome' `controls') (`treatment' `controls') ///
>   if sample1, cate median cov(cov)
. qui estimates store cate1_50
.
. matrix cov[1,1] = 0
. matrix qcov = (.9)
. matrix colnames qcov = re74
. qui tesensitivity cpi (`outcome' `controls') (`treatment' `controls') ///
>   if sample1, cate median qcov(qcov) cov(cov)
. qui estimates store cate1_90
.
. tesensitivity cpiplot cate1_50 cate1_90
```



We can see what the values of the covariates in this specification where by using the stored result `e(covsupp)`

```
. estimates restore cate1_50
(results cate1_50 are active now)

. matrix list e(covsupp)
e(covsupp)[1,10]
      married      age      black      hispanic      education      re74      re75      re74pos
r1          1         24          1           0           10          0          0          0
      re75pos      _const
r1          0          1

.
. estimates restore cate1_90
(results cate1_90 are active now)

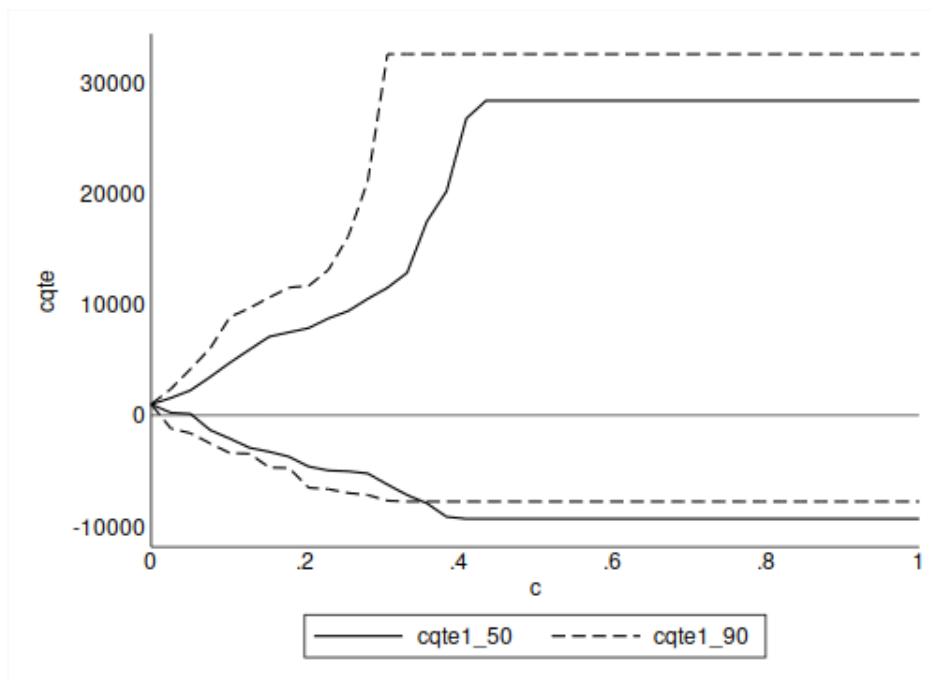
. matrix list e(covsupp)
e(covsupp)[1,10]
      married      age      black      hispanic      education      re74      re75      re74pos
r1          0         24          1           0           10 7914.1309          0          0
      re75pos      _const
r1          0          1
```

And we can consider consider the same analysis for the conditional quantile treatment effect for the 50th percentile.

```
. matrix cov = (1)
. matrix colnames cov = married
. qui tesensitivity cpi (`outcome` `controls`) (`treatment` `controls`) ///
>     if sample1, cqte median cov(cov)
. qui estimates store cqte1_50

.
. matrix cov[1,1] = 0
. matrix qcov = (.9)
. matrix colnames qcov = re74
. qui tesensitivity cpi (`outcome` `controls`) (`treatment` `controls`) ///
>     if sample1, cqte median qcov(qcov) cov(cov)
. qui estimates store cqte1_90

.
. tesensitivity cpiplot cqte1_50 cqte1_90
```



Binary outcome variables

In all of the previous analysis, we have assumed that the outcome variable has continuous support. For binary outcomes, only the ATE is currently supported. To illustrate, we'll look at the outcome of having positive earnings (i.e., $re78 > 0$).

```
. gen re78pos = re78 > 0
. tesensitivity cpi (re78pos `controls`) (`treatment` `controls`) if sample1, ate
```

Treatment effects sensitivity

| | | | | |
|------------------|------------------------------|---------------|---|---------|
| Analysis | : cond. partial independence | Number of obs | = | 445 |
| Outcome model | : logistic | Breakdown | = | 0.059 |
| Treatment model | : logistic | Conclusion | = | ate > 0 |
| Outcome variable | : re78pos | | | |

| c | ate |
|-------|-------------------|
| 0.000 | [0.1121, 0.1121] |
| 0.026 | [0.0654, 0.1431] |
| 0.051 | [0.0162, 0.1741] |
| 0.077 | [-0.0346, 0.2057] |
| 0.103 | [-0.0909, 0.2378] |
| 0.128 | [-0.1490, 0.2666] |
| 0.154 | [-0.1886, 0.2928] |
| 0.179 | [-0.2249, 0.3174] |
| 0.205 | [-0.2562, 0.3405] |
| 0.231 | [-0.2824, 0.3628] |
| 0.256 | [-0.3067, 0.3845] |
| 0.282 | [-0.3283, 0.4052] |
| 0.308 | [-0.3483, 0.4243] |
| 0.333 | [-0.3670, 0.4420] |
| 0.359 | [-0.3833, 0.4579] |
| 0.385 | [-0.3983, 0.4717] |
| 0.410 | [-0.4120, 0.4838] |
| 0.436 | [-0.4255, 0.4945] |

| | |
|-------|-------------------|
| 0.462 | [-0.4370, 0.5025] |
| 0.487 | [-0.4464, 0.5080] |
| 0.513 | [-0.4545, 0.5121] |
| 0.538 | [-0.4613, 0.5152] |
| 0.564 | [-0.4668, 0.5175] |
| 0.590 | [-0.4711, 0.5192] |
| 0.615 | [-0.4744, 0.5203] |
| 0.641 | [-0.4763, 0.5208] |
| 0.667 | [-0.4773, 0.5211] |
| 0.692 | [-0.4778, 0.5211] |
| 0.718 | [-0.4782, 0.5212] |
| 0.744 | [-0.4785, 0.5212] |
| 0.769 | [-0.4787, 0.5212] |
| 0.795 | [-0.4787, 0.5212] |
| 0.821 | [-0.4787, 0.5213] |
| 0.846 | [-0.4787, 0.5213] |
| 0.872 | [-0.4787, 0.5213] |
| 0.897 | [-0.4787, 0.5213] |
| 0.923 | [-0.4787, 0.5213] |
| 0.949 | [-0.4787, 0.5213] |
| 0.974 | [-0.4787, 0.5213] |
| 1.000 | [-0.4787, 0.5213] |

Displaying progress updates

The computation of some statistics, particularly `ate`, `atet`, and `qte` can take some time. To see progress updates, use the `verbose` option. This produces output as follows:

```
. tesensitivity cpi (re78pos `controls`) (`treatment` `controls`) if sample1, ate verbose
calculating ate (41)
-----|----- 1 -----|----- 2 -----|----- 3 -----|----- 4 -----|----- 5
.....
calculating breakdown point...
```

```
Treatment effects sensitivity
Analysis      : cond. partial independence  Number of obs =          445
Outcome model : logistic                    Breakdown      =          0.059
Treatment model : logistic                  Conclusion     =          ate > 0
Outcome variable : re78pos
```

| c | ate |
|-------|-------------------|
| 0.000 | [0.1121, 0.1121] |
| 0.026 | [0.0654, 0.1431] |
| 0.051 | [0.0162, 0.1741] |
| 0.077 | [-0.0346, 0.2057] |
| 0.103 | [-0.0909, 0.2378] |
| 0.128 | [-0.1490, 0.2666] |
| 0.154 | [-0.1886, 0.2928] |
| 0.179 | [-0.2249, 0.3174] |
| 0.205 | [-0.2562, 0.3405] |
| 0.231 | [-0.2824, 0.3628] |
| 0.256 | [-0.3067, 0.3845] |
| 0.282 | [-0.3283, 0.4052] |
| 0.308 | [-0.3483, 0.4243] |
| 0.333 | [-0.3670, 0.4420] |
| 0.359 | [-0.3833, 0.4579] |
| 0.385 | [-0.3983, 0.4717] |
| 0.410 | [-0.4120, 0.4838] |
| 0.436 | [-0.4255, 0.4945] |
| 0.462 | [-0.4370, 0.5025] |
| 0.487 | [-0.4464, 0.5080] |

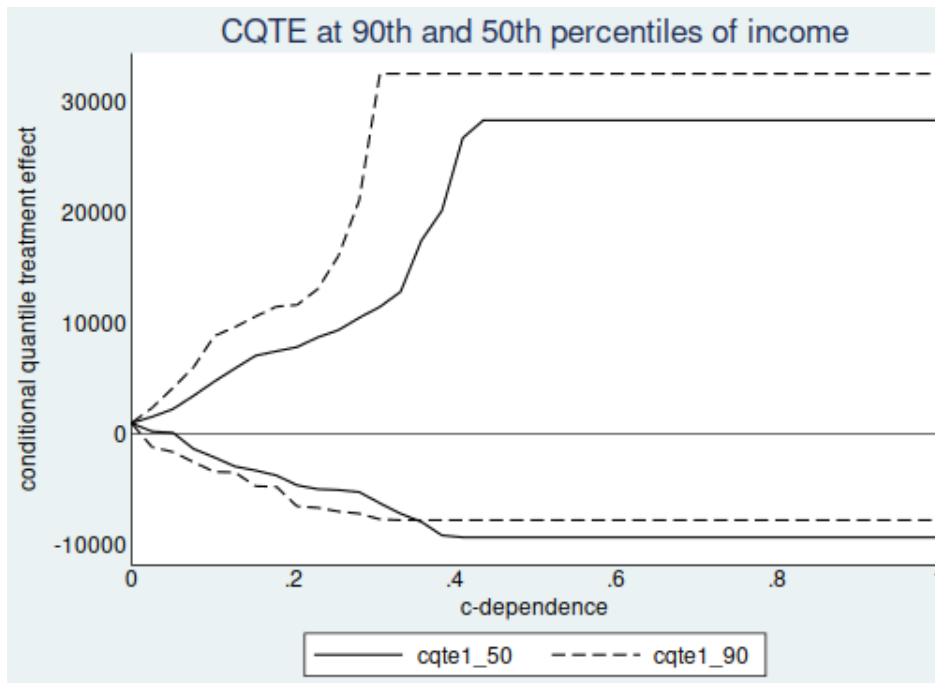
| | | |
|-------|-----------|---------|
| 0.513 | [-0.4545, | 0.5121] |
| 0.538 | [-0.4613, | 0.5152] |
| 0.564 | [-0.4668, | 0.5175] |
| 0.590 | [-0.4711, | 0.5192] |
| 0.615 | [-0.4744, | 0.5203] |
| 0.641 | [-0.4763, | 0.5208] |
| 0.667 | [-0.4773, | 0.5211] |
| 0.692 | [-0.4778, | 0.5211] |
| 0.718 | [-0.4782, | 0.5212] |
| 0.744 | [-0.4785, | 0.5212] |
| 0.769 | [-0.4787, | 0.5212] |
| 0.795 | [-0.4787, | 0.5212] |
| 0.821 | [-0.4787, | 0.5213] |
| 0.846 | [-0.4787, | 0.5213] |
| 0.872 | [-0.4787, | 0.5213] |
| 0.897 | [-0.4787, | 0.5213] |
| 0.923 | [-0.4787, | 0.5213] |
| 0.949 | [-0.4787, | 0.5213] |
| 0.974 | [-0.4787, | 0.5213] |
| 1.000 | [-0.4787, | 0.5213] |

Controlling the formatting of plots

The formatting of graphs produced by `tesensitivity cpiplot` can be specified by the user.

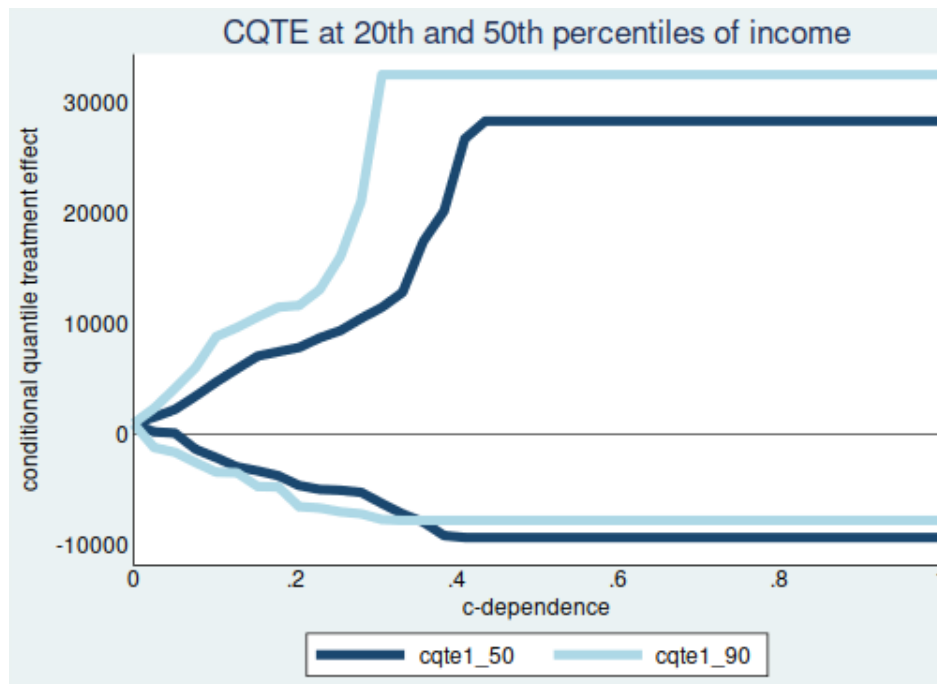
To change the overall formatting of the figure, any of the *twoway_options* to Stata's `graph twoway` command can be passed to `tesensitivity cpiplot`. Let's change some of the titles, background colors, and margins of the CQTE comparison in the previous section.

```
. qui tesensitivity cpiplot cqte1_50 cqte1_90, ///
>         xtitle(c-dependence) ytitle(conditional quantile treatment effect) ///
>         graphregion(color(%8) margin(vsmall)) ///
>         title(CQTE at 90th and 50th percentiles of income)
```



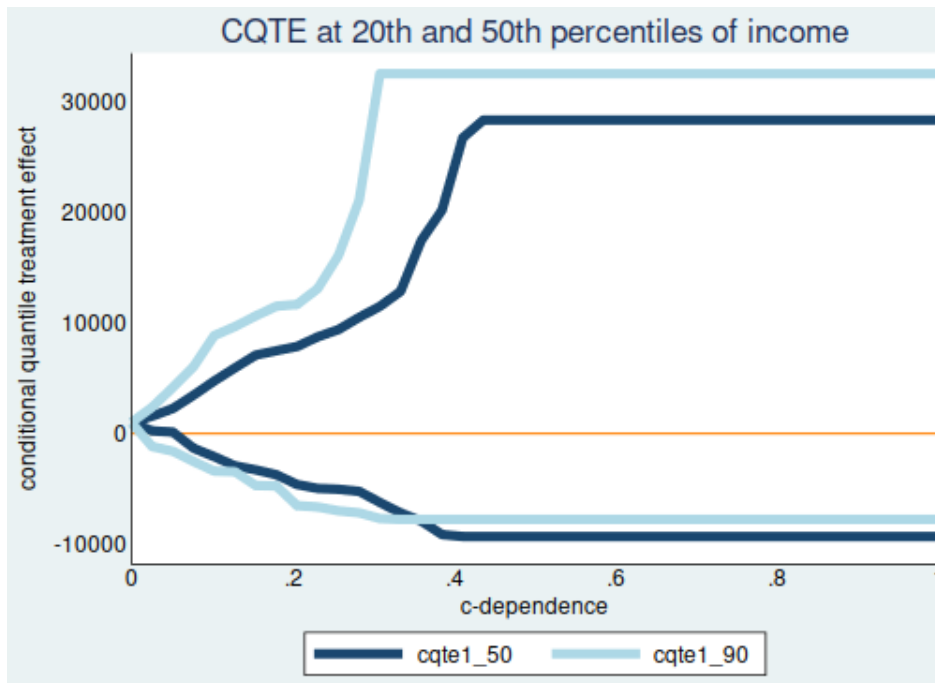
By default, the bound lines are black and line patterns are used to mark the different estimates included in the plot. This can be customized using the `boundpatterns` and `boundcolors` options. If only one color or pattern is specified, then this is applied to bound lines for all estimates. Any other options to Stata's `graph twoway line` command can be passed to `tesensitivity cpiplot` in the `boundoptions` options. These are applied to the bounds lines for all estimates. In the following example, we use colors rather than patterns to indicate the different estimates and change the width of the lines to be thicker.

```
. qui tesensitivity cpiplot cqte1_50 cqte1_90, ///
> xtitle(c-dependence) ytitle(conditional quantile treatment effect) ///
> graphregion(color(%8) margin(vsmall)) ///
> title(CQTE at 20th and 50th percentiles of income) ///
> boundcolors(navy ltblue) boundpatterns(solid) boundoptions(lwidth(vthick))
```



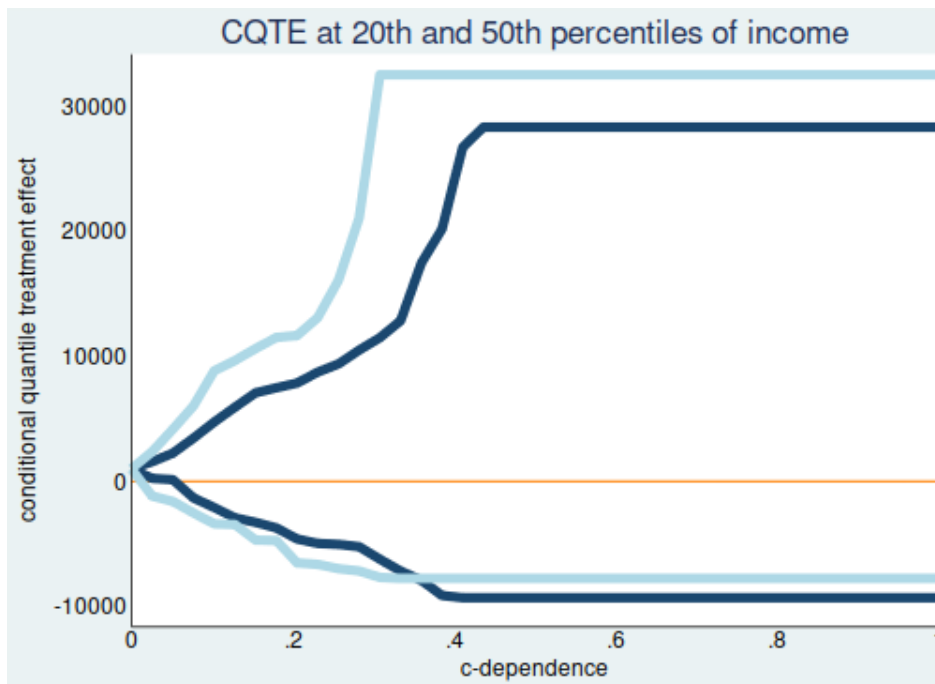
Finally, by default the plot includes a horizontal line at the conclusion for the breakdown point. This can be removed with the `nobreakdown` option and the formatting can be controlled with the `breakdownoptions` option. Any suboptions of the `added_line_options` options to Stata's `yline` option can be included here. In the following example, we change the color of the breakdown line.

```
. qui tesensitivity cpiplot cqte1_50 cqte1_90, ///
> xtitle(c-dependence) ytitle(conditional quantile treatment effect) ///
> graphregion(color(%8) margin(vsmall)) ///
> title(CQTE at 20th and 50th percentiles of income) ///
> boundcolors(navy ltblue) boundpatterns(solid) boundoptions(lwidth(vthick)) ///
> breakdownoptions(lcolor(orange))
```



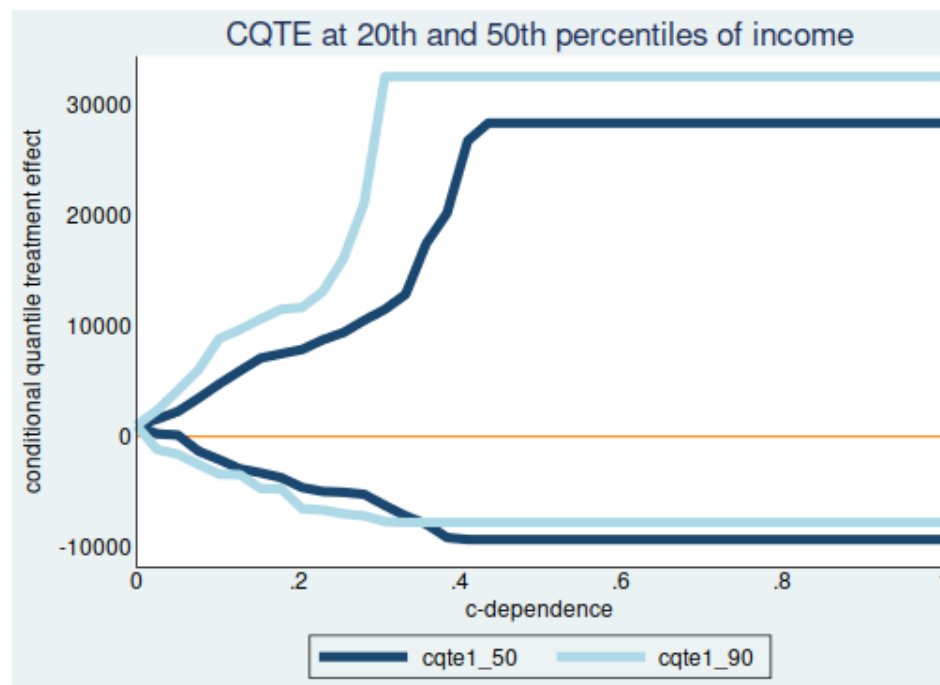
To suppress the legend, simply use the `nolegend` option

```
. qui tesensitivity cpiplot cqte1_50 cqte1_90, ///
> xtitle(c-dependence) ytitle(conditional quantile treatment effect) ///
> graphregion(color(%8) margin(vsmall)) ///
> title(CQTE at 20th and 50th percentiles of income) ///
> boundcolors(navy ltblue) boundpatterns(solid) boundoptions(lwidth(vthick)) ///
> breakdownoptions(lcolor(orange)) nolegend
```



Alternatively, any *legend_option* to Stata's `legend` option can be specified in `legoptions` except for `order` and `label` which are handled automatically:

```
. qui tesensitivity cpiplot cqte1_50 cqte1_90, ///
> xtitle(c-dependence) ytitle(conditional quantile treatment effect) ///
> graphregion(color(%8) margin(vsmall)) ///
> title(CQTE at 20th and 50th percentiles of income) ///
> boundcolors(navy ltblue) boundpatterns(solid) boundoptions(lwidth(vthick)) ///
> breakdownoptions(lcolor(orange)) legoptions(region(fcolor(%8)))
```



References

- Dehejia, Rajeev, and Sadek Wahba. 1999. “Causal Effects in Nonexperimental Studies: Reevaluating the Evaluation of Training Programs.” *Journal of the American Statistical Association* 94 (448): 1053–62.
- LaLonde, Robert. 1986. “Evaluating the Econometric Evaluations of Training Programs with Experimental Data.” *The American Economic Review* 76 (4). American Economic Association: 604–20.
- Masten, Matthew, and Alexandre Poirier. 2018. “Identification of Treatment Effects Under Conditional Partial Independence.” *Econometrica* 86 (1): 317–51.
- . 2020. “Inference on Breakdown Frontiers.” *Quantitative Economics* 11 (1): 41–111.
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