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August 2008

# Quantile Treatment Effects in the Regression Discontinuity Design

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## **ABSTRACT**

### **Quantile Treatment Effects in the Regression Discontinuity Design**

This paper shows nonparametric identification of quantile treatment effects (QTE) in the regression discontinuity design (RDD) and proposes simple estimators. Quantile treatment effects are a very helpful tool to characterize the effects of certain interventions on the outcome distribution. The distributional impacts of social programs such as welfare, education, training programs and unemployment insurance are of large interest to economists.

JEL Classification: C13, C14, C21

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# 1 Introduction

In recent years, the regression discontinuity design (RDD) has received tremendous attention in applied economic research.<sup>1</sup> All these applications focus on the estimation of average treatment effect. In many research areas, one is not only interested in mean impacts, but also in the *distributional* consequences of treatment interventions. In the field of education (e.g. Angrist and Lavy (1999), Puhani and Weber (2007)), educational inequality e.g. in cognitive achievement is of large public interest. When examining the effects of training (e.g. Black, Galdo, and Smith (2005)), policy makers are often more interested in the effects at the lower quantiles than at the upper quantiles. When analyzing the effects of unemployment insurance on unemployment durations (e.g. Lalive (2008)), the distribution of the unemployment durations is of interest, e.g. the risk of becoming long-term unemployed.<sup>2</sup>

Quantile treatment effects (QTE) are a convenient tool to characterize the potentially heterogeneous impacts of variables on different points of an outcome distribution. In this paper we show how QTE can be identified *nonparametrically* in the regression discontinuity design and propose nonparametric estimators.<sup>3</sup> We also discuss the identification of the potential outcome distributions.

## 2 Identification of QTE in the RDD

Following the setup of Hahn, Todd, and van der Klaauw (2001), let  $D_i \in \{0, 1\}$  be a binary treatment variable, let  $Y_i^0, Y_i^1$  be the individual potential outcomes. The potential outcomes as well as the treatment effect are permitted to vary freely across individuals, i.e. no constant treatment effect is assumed. In the examples mentioned,  $D$  may represent school quality, class

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<sup>1</sup>For an incomplete list see e.g. Angrist and Lavy (1999), Battistin and Rettore (2002), Battistin and Rettore (2008), Black (1999), Black, Galdo, and Smith (2005), Black, Jang, and Kim (2006), Buddelmeyer and Skoufias (2003), Brügger, Lalive, and Zweimüller (2008), Chay and Greenstone (2005), Chay, McEwan, and Urquiola (2005), DiNardo and Lee (2004), Fredriksson and Öckert (2006), Forslund and NordströmSkans (2006), Imbens and Lemieux (2008), Jacob and Lefgren (2004a), Jacob and Lefgren (2004b), Gormley and Phillips (2005), Guryan (2001), Lalive (2008), Lalive, Wüllrich, and Zweimüller (2008), Leuven, Lindahl, Oosterbeek, and Webbink (2007), Matsudaira (2008), NordströmSkans and Lindqvist (2005), Öckert (2008), Puhani and Weber (2007), van der Klaauw (2002), van der Klaauw (2008) and the special issue of the Journal of Econometrics 2008.

<sup>2</sup>Note that this distribution also identifies the hazard rates.

<sup>3</sup>In future work we are going to derive the asymptotic properties of these estimators.

size, participation in training etc. Let  $Z_i$  be a variable that influences the treatment variable in a discontinuous way, e.g. total school enrollment, profiling risk score etc.

In the literature, often two different designs are examined: the *sharp* design where  $D_i$  changes for everyone at a known threshold  $z_0$ , e.g.

$$D_i = 1(Z_i \geq z_0). \quad (1)$$

In this sharp design, *all* individuals change programme participation status exactly at  $z_0$ . The *fuzzy* design, on the other hand, permits  $D$  to also depend on other (unobserved) factors but assumes that the treatment probability changes discontinuously at  $z_0$ :

$$\lim_{\varepsilon \rightarrow 0} E[D|Z = z_0 + \varepsilon] - \lim_{\varepsilon \rightarrow 0} E[D|Z = z_0 - \varepsilon] \neq 0. \quad (2)$$

This fuzzy design includes the sharp design as a special case when the left hand side of (2) is equal to one. Therefore the following discussion focusses on the more general fuzzy design.<sup>4</sup>

Identification of treatment effects requires two assumptions. First, the conditional distribution of  $Y^0$  has to be continuous at  $z_0$ . Furthermore, an assumption on the treatment effect is required. Hahn, Todd, and van der Klaauw (2001) consider two different versions: In the first, they assume that the treatment effect is independent of  $D$  conditional on  $Z$  being near  $z_0$ . This is some kind of selection on observables assumption. As an alternative, they consider an instrumental variable type assumption, which assumes for the local *compliers* that the potential outcomes and the potential treatment status are independent of  $Z$ , near  $z_0$ . In the sharp design, both assumptions are equivalent. In the fuzzy design, the IV type assumption is much more frequently used in applications. We therefore focus on the IV type approach.

For stating the identification results, it is helpful to introduce more precise notation first. Let  $\mathcal{N}_\varepsilon$  be an  $\varepsilon$  neighbourhood about  $z_0$  and partition  $\mathcal{N}_\varepsilon$  into  $\mathcal{N}_\varepsilon^+ = \{z : z \geq z_0, z \in \mathcal{N}_\varepsilon\}$  and  $\mathcal{N}_\varepsilon^- = \{z : z < z_0, z \in \mathcal{N}_\varepsilon\}$ . According to their reaction to the instrument  $z$  over  $\mathcal{N}_\varepsilon$  we can

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<sup>4</sup>The fuzzy design may apply when the treatment decision contains some element of discretion. Case workers may have some discretion about whom they offer a programme, or they may base their decision also on criteria that are unobserved to the econometrician.

partition the population into five subpopulations:

$$\begin{aligned}
\mathcal{T}_{i,\varepsilon} &= a & \text{if } D_i(z) = 1 \quad \forall z \in \mathcal{N}_\varepsilon^- & \text{ and } D_i(z) = 1 \quad \forall z \in \mathcal{N}_\varepsilon^+ \\
\mathcal{T}_{i,\varepsilon} &= n & \text{if } D_i(z) = 0 \quad \forall z \in \mathcal{N}_\varepsilon^- & \text{ and } D_i(z) = 0 \quad \forall z \in \mathcal{N}_\varepsilon^+ \\
\mathcal{T}_{i,\varepsilon} &= c & \text{if } D_i(z) = 0 \quad \forall z \in \mathcal{N}_\varepsilon^- & \text{ and } D_i(z) = 1 \quad \forall z \in \mathcal{N}_\varepsilon^+ \\
\mathcal{T}_{i,\varepsilon} &= d & \text{if } D_i(z) = 1 \quad \forall z \in \mathcal{N}_\varepsilon^- & \text{ and } D_i(z) = 0 \quad \forall z \in \mathcal{N}_\varepsilon^+ \\
\mathcal{T}_{i,\varepsilon} &= i & \text{if } D_i(z) \text{ is nonconstant over } \mathcal{N}_\varepsilon^- & \text{ or over } \mathcal{N}_\varepsilon^+.
\end{aligned}$$

These subpopulations are a straightforward extension of the LATE concept of Imbens and Angrist (1994). The first group contains those units that will *always* be treated, the second contains those that will *never* be treated, and the third and fourth group contains the units that are treated only on one side of  $z_0$ . The fifth group (labelled indefinite) contains all units that switch at other values than  $z_0$  and that react non-monotonously, e.g. they may first switch from  $D = 0$  to 1 and then back for increasing values of  $z$ . We will assume that in the limit only the first three groups exist, and that the fraction of compliers is positive. Note that in the sharp design, everyone is a complier for  $\varepsilon \rightarrow 0$ .

**Assumption 1:**

- i) Existence of compliers  $\lim_{\varepsilon \rightarrow 0} \Pr(\mathcal{T}_\varepsilon = c | Z = z_0) > 0$
- ii) Monotonicity  $\lim_{\varepsilon \rightarrow 0} \Pr(\mathcal{T}_\varepsilon = t | Z \in \mathcal{N}_\varepsilon) = 0 \quad \text{for } t \in \{d, i\}$
- iii) Independent IV  $\lim_{\varepsilon \rightarrow 0} \Pr(\mathcal{T}_\varepsilon = t | Z \in \mathcal{N}_\varepsilon^+) - \Pr(\mathcal{T}_\varepsilon = t | Z \in \mathcal{N}_\varepsilon^-) = 0 \quad \text{for } t \in \{a, n, c\}$
- iv) IV Exclusion  $\lim_{\varepsilon \rightarrow 0} F_{Y^1|Z \in \mathcal{N}_\varepsilon^+, \mathcal{T}_\varepsilon=t}(u) - F_{Y^1|Z \in \mathcal{N}_\varepsilon^-, \mathcal{T}_\varepsilon=t}(u) = 0 \quad \text{for } t \in \{a, c\}$   
 $\lim_{\varepsilon \rightarrow 0} F_{Y^0|Z \in \mathcal{N}_\varepsilon^+, \mathcal{T}_\varepsilon=t}(u) - F_{Y^0|Z \in \mathcal{N}_\varepsilon^-, \mathcal{T}_\varepsilon=t}(u) = 0 \quad \text{for } t \in \{n, c\}$
- v) Density at threshold  $F_Z(z)$  is differentiable at  $z_0$  and  $f_Z(z_0) > 0$

In words, Assumption 1 requires the existence of some compliers and the absence of defiers near  $z_0$ . In addition, the potential outcomes and the type are jointly independent of  $Z$  near  $z_0$ , that is  $(Y^0, Y^1, \mathcal{T}) \perp\!\!\!\perp Z$  in a neighbourhood of  $z_0$ .

Define the  $\tau$ -th quantile of  $Y$  as  $Q_Y^\tau = \inf \{y : F_Y(y) \geq \tau\}$ . Define  $Q_{Y^d|c}^\tau = \lim_{\varepsilon \rightarrow 0} Q_{Y^d|Z \in \mathcal{N}_\varepsilon, \mathcal{T}_\varepsilon=c}^\tau$  as the limit for the local compliers. The quantile treatment effect (QTE) for the compliers is then defined as

$$\Delta_{QTE}^\tau = Q_{Y^1|c}^\tau - Q_{Y^0|c}^\tau.$$

The following theorem shows that  $Q_{Y^1|c}^\tau$  and  $Q_{Y^0|c}^\tau$  are identified under Assumption 1 by a very simple weighted quantile regression with weights 1 and  $-1$ . Define  $I^+ = 1(Z \geq z_0)$  and  $I^- = 1 - I^+$ .

**Theorem 1 (Quantiles of potential outcomes)** *Under Assumption 1, the quantiles of the potential outcomes for the local compliers are identified as the solution of the following optimization problem*

$$\begin{aligned} Q_{Y^1|c}^\tau &= \lim_{\varepsilon \rightarrow 0} \arg \min_q E [\rho_\tau(Y - q) (2I^+ - 1) | Z \in \mathcal{N}_\varepsilon, D = 1] \\ Q_{Y^0|c}^\tau &= \lim_{\varepsilon \rightarrow 0} \arg \min_q E [\rho_\tau(Y - q) (2I^- - 1) | Z \in \mathcal{N}_\varepsilon, D = 0] \end{aligned}$$

where  $\rho_\tau(u) = u \cdot \{\tau - 1(u < 0)\}$ . (All proofs are given in the appendix.)

Based on this representation, a straightforward estimator of the quantiles is obtained as

$$\begin{aligned} \hat{Q}_{Y^1|c}^\tau &= \arg \min_q \sum_{i:D_i=1} \rho_\tau(Y_i - q) (2 \cdot 1(Z_i \geq z_0) - 1) K\left(\frac{Z_i - z_0}{h}\right) \\ \hat{Q}_{Y^0|c}^\tau &= \arg \min_q \sum_{i:D_i=0} \rho_\tau(Y_i - q) (2 \cdot 1(Z_i \geq z_0) - 1) K\left(\frac{Z_i - z_0}{h}\right), \end{aligned}$$

where  $K$  is a kernel function. These are simple univariate quantile regression with weights 1 and  $-1$ , multiplied with kernel weights. Note that  $\hat{Q}_{Y^1|c}^\tau$  and  $\hat{Q}_{Y^0|c}^\tau$  are estimated from independent observations, since the former uses only the  $D_i = 1$  observations and the latter only the  $D_i = 0$  observations.

Despite its simplicity one should note that the objective function of the weighted quantile regression estimator is not convex since some of the weights are negative. This complicates the optimization problem a little because local optima could exist and conventional linear programming algorithms cannot be used. The problem is, however, not very serious because we have two one-dimensional estimation problems in the treated and non-treated populations. In addition, the objective function can change only at the values of  $Y_i$  observed in the sample such that only  $n$  weighted means need to be computed to find the global minimum.

We explore now an alternative approach that may be fruitful when we want to estimate many (or all) QTE. To this end we estimate the cumulative distribution function. Define

$$F_{Y^d|c}(u) = \lim_{\varepsilon \rightarrow 0} F_{Y^d|Z \in \mathcal{N}_\varepsilon, \mathcal{T}_\varepsilon=c}(u).$$

Theorem 2 shows that the distribution functions of the potential outcomes for compliers are identified by the ratio of two weighted means.

**Theorem 2 (Distribution of potential outcomes)** *Under Assumption 1, the distribution of the potential outcomes for the local compliers are identified as*

$$\begin{aligned} F_{Y^1|c}(u) &= \lim_{\varepsilon \rightarrow 0} \frac{E[1(Y \leq u) \cdot (2I^+ - 1) | Z \in \mathcal{N}_\varepsilon, D = 1]}{E[2I^+ - 1 | Z \in \mathcal{N}_\varepsilon, D = 1]} \\ F_{Y^0|c}(u) &= \lim_{\varepsilon \rightarrow 0} \frac{E[1(Y \leq u) \cdot (2I^- - 1) | Z \in \mathcal{N}_\varepsilon, D = 0]}{E[2I^- - 1 | Z \in \mathcal{N}_\varepsilon, D = 0]}. \end{aligned}$$

Note that in the *sharp* design, everyone is a complier at  $z_0$ , such that the cdf of the potential outcomes in the population is identified in this case as

$$\begin{aligned} \lim_{\varepsilon \rightarrow 0} F_{Y^1|Z \in \mathcal{N}_\varepsilon}(u) &= \lim_{\varepsilon \rightarrow 0} E[1(Y \leq u) | Z \in \mathcal{N}_\varepsilon, D = 1] \\ \lim_{\varepsilon \rightarrow 0} F_{Y^0|Z \in \mathcal{N}_\varepsilon}(u) &= \lim_{\varepsilon \rightarrow 0} E[1(Y \leq u) | Z \in \mathcal{N}_\varepsilon, D = 0]. \end{aligned}$$

Based on Theorem 2, straightforward estimators of the distribution functions are obtained as

$$\begin{aligned} \hat{F}_{Y^1|c}(u) &= \frac{\sum_{i:D_i=1} 1(Y_i \leq u) (2 \cdot 1(Z_i \geq z_0) - 1) K\left(\frac{Z_i - z_0}{h}\right)}{\sum_{i:D_i=1} (2 \cdot 1(Z_i \geq z_0) - 1) K\left(\frac{Z_i - z_0}{h}\right)}, \\ \hat{F}_{Y^0|c}(u) &= \frac{\sum_{i:D_i=0} 1(Y_i \leq u) (2 \cdot 1(Z_i \geq z_0) - 1) K\left(\frac{Z_i - z_0}{h}\right)}{\sum_{i:D_i=0} (2 \cdot 1(Z_i \geq z_0) - 1) K\left(\frac{Z_i - z_0}{h}\right)}. \end{aligned}$$

The estimated distribution function is well-behaved for all types of outcome variables while the quantiles identified in Theorem 1 will be well-behaved only when  $Y$  is continuously distributed. Therefore, Theorem 2 is interesting for discrete or mixed outcome variables. Furthermore, since we have a closed-form solution for the distribution function, its estimation may be a first step towards the estimation of the QTE. The negativity of some of the weights, however, implies that the estimated distribution function will not be monotonously increasing in finite samples. This problem can be solved by monotonizing the estimated distribution function using the method of Chernozhukov, Fernandez-Val, and Galichon (2007) and finally inverting it to obtain the quantiles.

### 3 QTE with estimated threshold probability

In the preceding section we have used the fact that  $\lim_{\varepsilon \rightarrow 0} \Pr(Z \geq z_0 | Z \in \mathcal{N}_\varepsilon) = \frac{1}{2}$ . This result follows by differentiability of  $F_Z$  at  $z_0$  and led to the very simple formulae of Theorems 1 and 2. In small samples, however, we may not have very many data points available at  $z_0$  and therefore have to rely on a larger smoothing window. In this case, the number of data points could be asymmetric around  $z_0$ , and we could obtain more precise estimates by estimating the probability of being above the threshold (within the smoothing area). Define  $p_\varepsilon = \Pr(Z \geq z_0 | Z \in \mathcal{N}_\varepsilon)$  for a given  $\varepsilon$ .

#### Theorem 3 (Quantiles of potential outcomes with estimated threshold probability)

*Under Assumption 1, the quantiles of the potential outcomes for the local compliers are identified as*

$$\begin{aligned} Q_{Y^1|c}^\tau &= \lim_{\varepsilon \rightarrow 0} \arg \min_q E \left[ \rho_\tau(Y - q) \frac{I^+ - p_\varepsilon}{p_\varepsilon(1 - p_\varepsilon)} | Z \in \mathcal{N}_\varepsilon, D = 1 \right] \\ Q_{Y^0|c}^\tau &= \lim_{\varepsilon \rightarrow 0} \arg \min_q E \left[ \rho_\tau(Y - q) \frac{I^+ - p_\varepsilon}{p_\varepsilon(1 - p_\varepsilon)} | Z \in \mathcal{N}_\varepsilon, D = 0 \right]. \end{aligned}$$

This result simplifies to Theorem 1 by using that  $\lim_{\varepsilon \rightarrow 0} p_\varepsilon = \frac{1}{2}$ . The representation of Theorem 3, nevertheless, suggests a different estimation strategy where one plugs in an estimate of  $\Pr(Z \geq z_0 | Z \in \mathcal{N}_\varepsilon)$  instead of the value  $\frac{1}{2}$ . Using the estimated  $p_\varepsilon$  often performed better in Monte Carlo simulations in small samples than when using  $\lim_{\varepsilon \rightarrow 0} p_\varepsilon = \frac{1}{2}$ . In some sense this result appears to be related to the well-known result in the propensity score matching literature that estimators which use the estimated propensity score are more efficient than estimators that use the true propensity score. This result might not be directly transferable here, though, since we are in a nonparametric context.

For completeness, we give the identification results for the distribution function.

#### Theorem 4 (Distribution of potential outcomes with estimated threshold probability)

*Under Assumption 1, the distribution of the potential outcomes for the local compliers are*

identified as

$$\begin{aligned}
 F_{Y^1|c}(u) &= \lim_{\varepsilon \rightarrow 0} \frac{E \left[ 1(Y \leq u) \frac{I^+ - p_\varepsilon}{p_\varepsilon(1-p_\varepsilon)} | Z \in \mathcal{N}_\varepsilon, D = 1 \right]}{E \left[ \frac{I^+ - p_\varepsilon}{p_\varepsilon(1-p_\varepsilon)} | Z \in \mathcal{N}_\varepsilon, D = 1 \right]} \\
 F_{Y^0|c}(u) &= \lim_{\varepsilon \rightarrow 0} \frac{E \left[ 1(Y \leq u) \frac{I^+ - p_\varepsilon}{p_\varepsilon(1-p_\varepsilon)} | Z \in \mathcal{N}_\varepsilon, D = 0 \right]}{E \left[ \frac{I^+ - p_\varepsilon}{p_\varepsilon(1-p_\varepsilon)} | Z \in \mathcal{N}_\varepsilon, D = 0 \right]}.
 \end{aligned}$$

## 4 QTE in RDD with covariates

In this section, we extend the regression discontinuity design to incorporate additional covariates  $X$  in a fully nonparametric way, and suppose that Assumption 1 holds conditionally on  $X$ .

There are several reasons why one might want to control for  $X$ . To mention a few: Covariates can help to eliminate small sample biases, especially if the number of observations close to the threshold  $z_0$  is rather small such that one also has to include observations in the estimation process that are further apart. This point is emphasized particularly in Black, Galdo, and Smith (2005). We also permit that the density  $f_{X|Z}$  is discontinuous at  $z_0$ . This can occur when the variable  $Z$  itself is confounded, e.g. in a situation of dynamic treatment assignment as in van der Klaauw (2008). It can also occur when different data collection schemes have been used for individuals above the threshold  $z_0$  versus those below  $z_0$ , e.g. if those above  $z_0$  have been hospitalized while those below  $z_0$  received outpatient care with restricted follow-up data collection. Another reason for incorporating covariates applies when the threshold crossing at  $z_0$  itself affects various  $X$  variables that one would like to control for. For example,  $Z$  may represent proximity to a state or regional border and crossing the border is associated with certain changes in laws or regulations that one is interested in. At the same time, a few other covariates may change in distribution as well, which one would like to control for. For a recent example see Brügger, Lalive, and Zweimüller (2008). As a final example for a discontinuity in  $f_{X|Z}$  we consider the decomposition between direct and indirect effects of the treatment effect.  $X$  is here a post-treatment variable, and a change in treatment status  $D$  may have an effect on  $Y$  via  $X$  as well as a direct effect on  $Y$ . While RDD estimation without covariates estimates the total effect, in various situations one is interested in disentangling the direct from the indirect effect, which under certain conditions

can be done by controlling for  $X$ .<sup>5</sup>

We assume in the following that Assumption 1 holds conditionally on  $X$ . Theorems 1 to 4 now apply immediately to the treatment effect conditionally on  $X$ . In many situations we are however more interested in the unconditional effect, i.e. the effect on all local compliers irrespective of their value of  $X$ . There are at least three reasons why unconditional effects are interesting. First, for the purpose of evidence-based policy making a small number of summary measures can be more easily conveyed to the policy makers and the public than a large number of estimated effects for each and every value of  $X$ . Second, unconditional effects can be estimated more precisely than conditional effects. Third, the definition of the unconditional effects does not depend on the variables included in  $X$ .<sup>6</sup> One can therefore consider different sets of control variables  $X$  and still estimate the same object, which is useful for examining robustness of the results to the set of control variables.

The following results identify the unconditional effects, which are obtained by first conditioning on  $X$  and thereafter integrating with respect to  $X$ . For identification we need a common support restriction with respect to  $X$  and we also assume the existence of a density. (At the expense of more complex notation we could also easily permit discrete  $X$ .)

**Assumption 2:** Assume Assumption 1(i), (ii), (v) and Assumption 1(iii) and (iv) conditionally on  $X$ . Further assume:

- Common support  $\lim_{\varepsilon \rightarrow 0} \text{Supp}(X|Z \in \mathcal{N}_\varepsilon^+) = \lim_{\varepsilon \rightarrow 0} \text{Supp}(X|Z \in \mathcal{N}_\varepsilon^-)$
- Density at threshold  $\lim_{\varepsilon \rightarrow 0} F_{X|Z \in \mathcal{N}_\varepsilon^+}(x)$  and  $\lim_{\varepsilon \rightarrow 0} F_{X|Z \in \mathcal{N}_\varepsilon^-}(x)$  exist and are differentiable in  $x$  at  $z_0$  with pdf  $f^+(x|z_0)$  and  $f^-(x|z_0)$ , respectively.

(Regarding notation:  $f^+(x, z_0) = f^+(x|z_0)f(z_0)$  refers to the joint distribution of  $X$  and  $Z$  whereas  $f^+(x|z_0)$  refers to the conditional distribution of  $X$ . Analogously for the limit from below.)

Note that we permit  $f(x, z)$  to be continuous at  $z_0$ , i.e.  $f^+(x|z_0) = f^-(x|z_0)$ , or to be discontinuous, i.e.  $f^+(x|z_0) \neq f^-(x|z_0)$ .

With these additional assumptions, we can identify the quantile and cumulative distribution functions of the potential outcomes. The formulae, however, are not so neat as in Section 2.

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<sup>5</sup>Even if  $f_{X|Z}$  is not discontinuous at  $z_0$ , we conjecture that there may be efficiency gains by incorporating  $X$ . We will analyze this issue in more detail in future work.

<sup>6</sup>This, of course, is only true if  $X$  contains only pre-treatment variables.

Define  $p_\varepsilon(x) = \Pr(Z \geq z_0 | X = x, Z \in \mathcal{N}_\varepsilon)$ .

**Theorem 5 (Quantiles of the potential outcomes)** *Under Assumption 2,  $Q_{Y^1|c}^\tau$  and  $Q_{Y^0|c}^\tau$  are the solutions of the following optimization problem*

$$\begin{aligned} Q_{Y^1|c}^\tau &= \lim_{\varepsilon \rightarrow 0} \arg \min_q E \left[ \rho_\tau(Y - q) \frac{I^+ - p_\varepsilon(X)}{p_\varepsilon(X)(1 - p_\varepsilon(X))} | Z \in \mathcal{N}_\varepsilon, D = 1 \right] \\ Q_{Y^0|c}^\tau &= \lim_{\varepsilon \rightarrow 0} \arg \min_q E \left[ \rho_\tau(Y - q) \frac{I^+ - p_\varepsilon(X)}{p_\varepsilon(X)(1 - p_\varepsilon(X))} | Z \in \mathcal{N}_\varepsilon, D = 0 \right]. \end{aligned}$$

An analogous result is obtained for the distribution function.

**Theorem 6 (Distribution of potential outcomes)** *Under Assumption 2, the distribution of the potential outcomes for the local compliers are identified as*

$$\begin{aligned} F_{Y^1|c}(u) &= \lim_{\varepsilon \rightarrow 0} \frac{E \left[ \mathbf{1}(Y \leq u) \frac{I^+ - p_\varepsilon(X)}{p_\varepsilon(X)(1 - p_\varepsilon(X))} | Z \in \mathcal{N}_\varepsilon, D = 1 \right]}{E \left[ \frac{I^+ - p_\varepsilon(X)}{p_\varepsilon(X)(1 - p_\varepsilon(X))} | Z \in \mathcal{N}_\varepsilon, D = 1 \right]} \\ F_{Y^0|c}(u) &= \lim_{\varepsilon \rightarrow 0} \frac{E \left[ \mathbf{1}(Y \leq u) \frac{I^+ - p_\varepsilon(X)}{p_\varepsilon(X)(1 - p_\varepsilon(X))} | Z \in \mathcal{N}_\varepsilon, D = 0 \right]}{E \left[ \frac{I^+ - p_\varepsilon(X)}{p_\varepsilon(X)(1 - p_\varepsilon(X))} | Z \in \mathcal{N}_\varepsilon, D = 0 \right]}. \end{aligned}$$

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