Debiased Causal Tree: Heterogeneous Treatment Effects Estimation with Unmeasured Confounding

Abstract

Unmeasured confounding poses a significant threat to the validity of causal infer-
ence. Despite that various ad hoc methods are developed to remove confounding
effects, they are subject to certain fairly strong assumptions. In this work, we
consider the estimation of conditional causal effects using observational data and
historical controls, in the presence of unmeasured confounding. Under an inter-
pretable transportability condition, we prove the partial identifiability of conditional
average treatment effect on the treated group (CATT). For tree-based models, a new
notion, *confounding entropy*, is proposed to measure the discrepancy introduced
by unobserved confounders between the conditional outcome distribution of the
treated and control groups. The confounding entropy generalizes conventional
confounding bias, and can be estimated effectively using historical controls. We
develop a new method, debiased causal tree, whose splitting rule is to minimize
the empirical risk regularized by the confounding entropy. Notably, our method
integrates current observational data (for empirical risk) and their historical controls
(for confounding entropy) harmoniously. We highlight that, debiased causal tree
can not only estimate CATT well in the presence of unmeasured confounding, but
also is a robust estimator of conditional average treatment effect (CATE) against
the imbalance of the treated and control populations when all confounders are
observed. An extension of combining multiple debiased causal trees to further
reduce biases by gradient boosting is considered. The computational feasibility
and statistical power of our method are evidenced by simulations and a study of a
credit card balance dataset.

1 Introduction

Conditional average treatment effect (CATE), also known as heterogeneous treatment effect, can
characterize the causal effect of certain treatments on the outcome over the whole population as a
function of measured covariates. Another closely related concept, conditional average treatment
effect on the treated group (CATT), is more relevant when we focus on the effect of treatments for
the sub-population which is actually exposed to certain specific treatments; for instance, we are more
interested in the effect on children of taking anti-depressants during pregnancy for the people who
actually took the drug [7]. The interpretable and informative nature has made it a great importance to
estimate CATE and CATT from observational or experimental data in a wide spectrum of applications,
including medical decision-making and personalized medicine [19, 34], evaluating advertising and
socioeconomic policies [10, 2, 21] and large-scale randomized experiments for recommendations
[29, 18], etc.

In the absence of unmeasured confounding, there exist lots of methods developed to estimate CATE
(CATT), including tree-based methods [3, 47, 4], meta-learners [27], R-learners [33], doubly robust

learners [25], and so on. Basically, such an unconfoundedness condition hinges on an investigator's ability to accurately measure all related covariates, which is questionable in practice, especially when there exist covariate-dependent measurement errors [46, 42], biased treatment allocations [23, 43] and batch effects [17, 20]. In the presence of unmeasured confounding, however, the identifiability of CATE or CATT cannot be ensured in general, leading to failures of the aforementioned estimation procedures.

To control the effect introduced by confounders, various methods have been developed, e.g., sensitivity analyses [36], instrumental variable (IV) approaches [1, 31], and proximal causal learning [28, 32, 39]. Nonetheless, several limitations of these methods hinder their direct application in CATE/CATT estimation. For example, we cannot derive point estimation from sensitivity analyses, and even valid IVs can only partially identify CATE in principle [41]. Although ATE and ATT can be identified in proximal causal learning [15], the assumption that the measured covariates are correctly classified into three types of proxies requires prior knowledge.

Recently, a line of work proposed to combine randomized trials [14] and observational studies to adjust confounders, which typically requires transportability [35] of a randomized controlled trial to external observational studies [37]. For example, [50] proposed to use the confounding function which summarizes the impact of confounders to estimate heterogeneous treatment effects, where the confounding function is assumed to be transportable. For a comprehensive review, see [13]. Problematically, many observational studies do not have related randomized trials due to practical and methodological issues such as cost and restrictive inclusion/exclusion criteria.

In this work, we move further in the direction of data fusion, without requiring experimental data. Instead, we consider the case where covariates and the outcome are collected at multiple timestamps, before and after the treatment. We take two timestamps $t \in \{t_1, t_2\}(t_1 < t_2)$, as an example, where the treatment is imposed at the time t_2. Following the potential outcome framework [38], denote by $Y_t = DY_t^{(1)} + (1-D)Y_t^{(0)}$ the observed outcomes at the time t, where $Y_t^{(d)}$ is the potential outcome for the treatment $D = d \in \{0, 1\}$. We remark that $Y_{t_1} = Y_{t_1}^{(0)}$ since the treatment does not take place at t_1. Let X and U be time-invariant covariates and unmeasured confounders, respectively. Our work is motivated by the following identity

$$
E(Y_t^{(1)} \mid D = 1, X = x) - E(Y_t^{(0)} \mid D = 0, X = x) = \underbrace{E(Y_t^{(1)} - Y_t^{(0)} \mid D = 1, X = x)}_{\text{CATT}}
$$
$$
+ \underbrace{E(Y_t^{(0)} \mid D = 1, X = x) - E(Y_t^{(0)} \mid D = 0, X = x)}_{\text{Confounding Bias, } \mathbf{b}_t}, \qquad \forall t
\tag{1}
$$

which holds regardless of U. In general, the confounding bias [6, 50] $\mathbf{b}_t$ does not vanish due to U. However, the smaller $\mathbf{b}_t$ is, the closer CATT could be to the difference in conditional expectations. We then propose a tree-based method, which, at high level, aims to (i) find an optimal partition of the feature space such that the confounding bias is smallest within each small region, and (ii) estimate the left hand of (1). Despite that the confounding bias is unavailable in general, we can represent it with tree-based models using data collected at $t = t_1$ since $Y_{t_1} = Y_{t_1}^{(0)}$ and it is known which individual would receive the treatment. If boundaries of the optimal partition that leads to the smallest historical confounding bias change "mildly" over time, i.e., it is transportable, the partition derived at time t_1 is also nearly optimal at t_2. The two steps can proceed simultaneously, and we calibrate and formalize the basic idea in Section 2.2.

Our contribution is summarized as follows: (i) we introduce a new notion, confounding entropy, to characterize the discrepancy between the conditional distributions of the treated and control groups that generalizes the confounding bias, and propose a tractable and scalable method, debiased causal tree, to estimate CATT from observational data before and after the treatment by developing a new criterion for node splitting, which minimizes both empirical risk and confounding entropy at the same time; (ii) to further reduce the bias, we integrate multiple causal trees by gradient boosting, which we refer to as Gradient Boosting Causal Tree (GBCT); (iii) under the unconfoundedness condition, we show that GBCT is robust to imbalance of the treated and control populations and outperforms other compared methods, which is obvious in severely imbalanced cases.

 ## 2 Methodology

 We first introduce the problem setup, related notation, and basic ideas behind our proposed method.

 ### 2.1 Problem Setup

 To begin with, we consider the case of binary treatment for simplicity. Our method can be easily
 extended to multiple treatments. Let $Y_t^{(d)} \in \mathbb{R}$ be the potential outcome at the time $t, t \in \{t_1, \ldots, t_m\}$
 for the treatment $D = d \in \{0, 1\}$. We impose the treatment at the time $t = t_m$ and let $X \in [0, 1]^p$
 be a time-invariant p-dimensional covariate. Before the treatment occurs, the historical outcomes
 are denoted by $Y_{t_k} = Y_{t_k}^{(0)}, 1 \leq k \leq m - 1, t_1 < \cdots < t_{m-1} < t_m$. Given n triplets of i.i.d.
 samples $\{(X_i, \{Y_{i,t_k}\}_{k=1}^m, D_i)\}_{i=1}^n$ with $Y_{i,t_m} = D_i Y_{i,t_m}^{(1)} + (1 - D_i) Y_{i,t_m}^{(0)}$, our goal is to estimate
 $\eta^*(x) \equiv E(Y_{t_m}^{(1)} - Y_{t_m}^{(0)} \mid D = 1, X = x)$. In general, η^* is not identifiable due to unmeasured
 confounding; however we can prove that it is partially identifiable under certain transportability
 condition. In the following, we denote the indicator function of event A by $\mathrm{I}(A)$.

 Equation (1) implies that, $\eta^*(x) \approx \delta_{t_m}(x) \equiv E(Y_{t_m}^{(1)} \mid D = 1, X = x) - E(Y_t^{(0)} \mid D = 0, X = x)$
 if the confounding bias $\boldsymbol{b}_{t_m}(x) \equiv E(Y_{t_m}^{(0)} \mid D = 1, X = x) - E(Y_{t_m}^{(0)} \mid D = 0, X = x) \approx 0$. In
 general, $\left| E_X[\boldsymbol{b}_{t_m}(X)] \right| \geq c_0 > 0$ due to unmeasured confounding. Proper stratification [26], notably,
 can help mitigate confounding effects by introducing multiple subgroups within which confounders
 remain almost constant. Therefore, we propose to find the optimal partition of the feature space, e.g.,
 $\{Q_j\}_{j=1}^q$ with $\cup_{j=1}^q Q_j = [0, 1]^p$, such that

$$\boldsymbol{b}_{t_m}(Q_j) \equiv E(Y_{t_m}^{(0)} \mid D = 1, X \in Q_j) - E(Y_{t_m}^{(0)} \mid D = 0, X \in Q_j) \approx 0,$$

 for each $j = 1, \ldots, q$. Alternatively, we can obtain such a partition by the following optimization

$$(Q_1^*, \ldots, Q_q^*) = \underset{Q_j : 1 \leq j \leq q}{\operatorname{argmin}} \sum_{j=1}^q \boldsymbol{b}_{t_m}^2(Q_j). \tag{2}$$

 Once $(Q_1^*, \ldots, Q_q^*)$ being derived, by (1), we can choose

$$\eta(x) = \sum_{j=1}^q \delta_{t_m}(Q_j^*) \mathrm{I}\{x \in Q_j^*\} \tag{3}$$

 to approximate $\eta^*(\cdot)$, where $\delta_{t_m}(Q_j^*) = E(Y_{t_m}^{(1)} \mid D = 1, X \in Q_j^*) - E(Y_t^{(0)} \mid D = 0, X \in Q_j^*)$
 is a constant function that approximates $\delta_{t_m}(x)$ for $x \in Q_j^*$. In (3), choosing a constant function
 to approximate $\delta_{t_m}(x)$ within Q_j^* is only for simplicity, and local polynomials of any degree are
 applicable. As expected, $\eta(x)$ is close to $\eta^*(x)$ provided that (2) implies $\max_{1 \leq j \leq q} \boldsymbol{b}_{t_m}^2(Q_j^*) \approx 0$.

 A major difficulty of solving (2) is the unavailability of $\boldsymbol{b}_{t_m}(Q_j)$, $1 \leq j \leq q$, since $Y_{t_m}^{(0)}$ is a
 counterfactual for $D = 1$. Fortunately, since D_i is known for each i and all units stay untreated
 before t_m, $\boldsymbol{b}_t(Q_j)$ for all $t \in \{t_1, \ldots, t_{m-1}\}$ are identifiable. Instead of (2), consider the following
 optimization

$$(Q_1^{**}, \ldots, Q_q^{**}) = \underset{Q_j : 1 \leq j \leq q}{\operatorname{argmin}} \frac{1}{m-1} \sum_{k=1}^{m-1} \sum_{j=1}^q \boldsymbol{b}_{t_k}^2(Q_j). \tag{4}$$

 The partition $\boldsymbol{Q}^{**} = \{Q_j^{**}\}_{j=1}^q$ of the feature space yields minimal historical confounding bias. If
 confounding effects are independent of time, $Q_j^{**} = Q_j^*$ holds for each $j = 1, \ldots, q$. We impose
 the following condition to ensure transportability from the historical optimal partition to its current
 counterpart.

 Assumption 1. *For the partition $\{Q_j^{**}\}_{j=1}^q$ defined in (4), $B(t) \equiv \log \left\{ \sum_{j=1}^q \boldsymbol{b}_t^2(Q_j^{**}) \right\} : t \mapsto$*
 $B(t) \in \mathbb{R}$ is a Lipschitz continuous function with parameter L; that is,

$$\left| B(\tau_1) - B(\tau_2) \right| \leq L|\tau_1 - \tau_2|, \quad \text{for any } \tau_1, \tau_2 \geq 0.$$

Assumption 1 is testable empirically for $t \in [0, t_m)$, provided that historical controls are collected at sufficiently many timestamps. Under Assumption 1, we prove in the Appendix that $Q_j^* \approx Q_j^{**}$ *in effect*; that is, the solution to (4) is transferrable or transportable to the current data source. The partial identifiability of $\eta^*(x)$ is justified by the following theorem.

Theorem 1. *Under Assumption 1, for the partition $\{Q_j^{**}\}_{j=1}^q$ of $[0,1]^p$ defined in (4), it holds that*

$$\sup_{x \in [0,1]^p} \left| \eta(x) - \eta^*(x) \right| \le \left\{ \frac{C_L}{m-1} \sum_{k=1}^{m-1} \sum_{j=1}^q \boldsymbol{b}_{t_k}^2(Q_j^{**}) \right\}^{1/2} + \max_{1 \le j \le q} \sup_{x \in Q_j^{**}} \left| \eta^*(x) - \eta^*(Q_j^{**}) \right|, \quad (5)$$

where $\eta^*(Q_j^{**}) = E\big(Y_{t_m}^{(1)} - Y_{t_m}^{(0)} \mid D = 1, X \in Q_j^{**}\big)$ and C_L depends only on L and $t_m - t_1$.

The first term in (5) denotes the confounding bias, and hinges on the transportability assumed in Assumption 1, while the second term measures the error of using a piecewise constant function to approximate $\eta^*(\cdot)$. Theorem 1 shows that $\eta^*(\cdot)$ is identifiable up to a band, which is similar in spirit to CATE estimation by IV [41]. Interestingly, the width of band in our case can be close enough to zero under stronger assumptions. For example, the confounding bias can be small if the stratification $\{Q_j^{**}\}_{j=1}^q$ can effectively adjust for confounding, and under certain smoothness condition, the approximation error decays to zero rapidly as the maximal volume of Q_j^{**} tends to zero. *Our goal is to develop a computational feasible method to solve* (4) *and estimate $\eta^*(x)$ as in* (3) *substituting Q_j^* with Q_j^{**}, for each j.*

2.2 Debiased Causal Tree

When it comes to stratifying or segmenting the feature space, the perhaps mostly used algorithms are tree-based methods [22], including classification and regression tree (CART) [9], random forest [8], and gradient boosting trees [16]. A regression tree $T(x; \boldsymbol{Q}, \boldsymbol{\mu})$ consists of two components: a set of leafs that partition the feature space, and the corresponding parameter $\boldsymbol{\mu} = \boldsymbol{\mu}(\boldsymbol{Q}) = (\mu(Q_j))$, where Q_j is the region partitioned by the j-th leaf, $1 \le j \le q$, and q is the total number of leafs. Typically, given a partition $\boldsymbol{Q}$ and n pairs of i.i.d samples $\{(X_i, y_i)\}_{i=1}^n$, the optimal data driven estimator is $T(x; \boldsymbol{Q}, \widehat{\boldsymbol{\mu}}) = \sum_{j=1}^q \mathrm{I}\{x \in Q_j\} \widehat{\mu}(Q_j)$, where

$$\widehat{\mu}(Q_j) = \frac{1}{|\{i : X_i \in Q_j\}|} \sum_{i=1}^n y_i \, \mathrm{I}\{X_i \in Q_j\}. \quad (6)$$

We define the auxiliary trees $T(x; \boldsymbol{Q}, \boldsymbol{\theta}_t^{(d)})$ to represent confounding biases, where $\boldsymbol{\theta}_t^{(d)} = (\theta_{t,j}^{(d)})$ such that $\theta_{t,j}^{(d)} = \theta_t^{(d)}(Q_j)$ fits the conditional mean $E(Y_t^{(0)} \mid D = d, X \in Q_j)$ for $d = 0, 1, t < t_m$, and define the main tree $T(x; \boldsymbol{Q}, \boldsymbol{\mu}^{(d)})$ to learn CATT, where $\boldsymbol{\mu}^{(d)} = (\mu_j^{(d)})$ such that $\mu_j^{(d)} = \mu^{(d)}(Q_j)$ fits the conditional mean $E(Y_{t_m}^{(d)} \mid D = d, X \in Q_j)$ for $d = 0, 1$. Auxiliary and main trees share the same splitting rules.

Confounding entropy. Treating the partition $\{Q_j\}_{j=1}^q$ as a parameter to be optimized, by (6), we estimate $\boldsymbol{b}_t(Q_j)$ for each $t \in \{t_1, \ldots, t_{m-1}\}$ by

$$\widehat{\boldsymbol{b}}_t(Q_j) = \widehat{\theta}_t^{(1)}(Q_j) - \widehat{\theta}_t^{(0)}(Q_j), \quad \text{where} \quad \widehat{\theta}_t^{(d)}(Q_j) = \sum_{i=1}^n \frac{\mathrm{I}\{X_i \in Q_j, D_i = d\} Y_{i,t}}{|\{i : X_i \in Q_j, D_i = d\}|}. \quad (7)$$

For simplicity, we write $\widehat{\theta}_t^{(d)}(Q_j)$ as $\widehat{\theta}_{t,j}^{(d)}$. Plugging $\widehat{\boldsymbol{b}}_t(Q_j)$ in (7) into (4), we obtain the estimator of the optimal partitions. Nonetheless, $\widehat{\boldsymbol{b}}_t(Q_j)$ as well as its population counterpart $\boldsymbol{b}_t(Q_j)$ only summarizes information of the first-order moment, and thus can give rise to partitions that yield estimators with large variances. This motivates us to model the discrepancy between $P_{t,1,j} = P(Y_t^{(0)} \mid D = 1, X \in Q_j)$ and $P_{t,0,j} = P(Y_t^{(0)} \mid D = 0, X \in Q_j)$ using symmetric cross entropy [49], defined as $\mathrm{H}(P_{t,1,j}, P_{t,0,j}) = -E_{P_{t,1}}[\log(P_{t,0,j})] - E_{P_{t,0,j}}[\log(P_{t,1,j})]$, which we refer to as the *confounding entropy* for time t and region Q_j. For each $t < t_m$, parametrize the working densities of $P_{t,1,j}$ and $P_{t,0,j}$ by $p(y_t; \theta_{t,j}^{*(1)})$ and $p(y_t; \theta_{t,j}^{*(0)})$, respectively. In our case, the parameter is the conditional mean $\theta_{t,j}^{*(d)} = E(Y_t^{(0)} \mid D = d, X \in Q_j)$ for both $d = 0, 1$. The estimator of symmetric

cross entropy between $P_{t,1,j}$ and $P_{t,0,j}$ can be obtained by cross-fitting the negative log-likelihood; that is,

$$\widehat{\mathrm{H}}_t(Q_j) \equiv - \sum_{i:X_i \in Q_j} \left\{ \frac{D_i \log(p(Y_{i,t}; \widehat{\theta}_{t,j}^{(0)}))}{|\{i : X_i \in Q_j, D_i = 1\}|} + \frac{(1 - D_i) \log(p(Y_{i,t}; \widehat{\theta}_{t,j}^{(1)}))}{|\{i : X_i \in Q_j, D_i = 0\}|} \right\}. \tag{8}$$

In particular, for Gaussian distributions, $\log(p(y_t; \theta_{t,j}^{*(d)})) = -\left(y_t - \theta_{t,j}^{*(d)}\right)^2 + \text{const}$, $d = 0, 1$,

$$\widehat{\mathrm{H}}_t(Q_j) = 2\left(\widehat{\theta}_{t,j}^{(0)} - \widehat{\theta}_{t,j}^{(1)}\right)^2 + \sum_{i:X_i \in Q_j} \left\{ \frac{D_i(Y_{i,t} - \widehat{\theta}_{t,j}^{(1)})^2}{|\{i : X_i \in Q_j, D_i = 1\}|} + \frac{(1 - D_i)(Y_{i,t} - \widehat{\theta}_{t,j}^{(0)})^2}{|\{i : X_i \in Q_j, D_i = 0\}|} \right\}, \tag{9}$$

where, the first term is exactly $\widehat{\boldsymbol{b}}_t^2(Q_j)$ defined in (7), and the second term is the sum of sample variances of $\widehat{\theta}_{t,j}^{(0)}$ and $\widehat{\theta}_{t,j}^{(1)}$. Therefore, both confounding bias and variance can be emphasized. In the end, assuming historical data collected at distinct timestamps are independent, the confounding entropy of all historical controls $E(\Pi_{t<t_m}\Pi_{j=1}^q P_{t,1,j}, \Pi_{t<t_m}\Pi_{j=1}^q P_{t,0,j})$ is estimated by

$$\widehat{\mathrm{H}}(\boldsymbol{Q}) \equiv \frac{1}{m-1} \sum_{k=1}^{m-1} \sum_{j=1}^{q} \widehat{\mathrm{H}}_{t_k}(Q_j), \quad \widehat{\mathrm{H}}_{t_k}(Q_j) \text{ is defined in (9).} \tag{10}$$

We remark that (10) can be interpreted as a marginally independent composite likelihood estimator [44], which can be readily generalized to model potential temporal correlation among $\{Y_{t_k,i}\}_{k,i=1}^{(m-1),n}$. However, for simplicity, we presume no temporal dependencies. We also remark that, under this composite likelihood framework, $\widehat{\boldsymbol{b}}_t(Q_j)$ is exactly the empirical cross entropy with working densities $\log(p(y_t; \theta_{t,j}(d))) = -(y_t - \mu_{t,j}(d)) + \text{const}$ for both $d = 0, 1$ (i.e., exponential distribution). As illustrated, such a specification of density functions, under which high-order moments equal to the first moment up to constants, is prone to high-order variations.

Estimating CATT. On the basis of (6), we then can estimate $\eta^*(x)$ with

$$\widehat{\eta}_h(x) = T(x; \widehat{\boldsymbol{Q}}_h, \widehat{\boldsymbol{\mu}}^{(1)}(\widehat{\boldsymbol{Q}}_h))) - T(x; \widehat{\boldsymbol{Q}}_h, \widehat{\boldsymbol{\mu}}^{(0)}(\widehat{\boldsymbol{Q}}_h)), \quad \widehat{\boldsymbol{\mu}}^{(d)}(\boldsymbol{Q}) = \left(\widehat{\mu}^{(d)}(Q_j)\right), \tag{11}$$

where by (6), $\widehat{\mu}^{(d)}(Q_j) = \sum_{i:X_i \in Q_j}(Y_{i,t_m}^{(d)} \mathrm{I}\{D_i = d\})/|\{i : X_i \in Q_j, D_i = d\}|, d = 0, 1$ for any given $\boldsymbol{Q}$, and $\widehat{\boldsymbol{Q}}_h = \{\widehat{Q}_j^h\}_{j=1}^q$ denotes the minimizer of $\widehat{\mathrm{H}}(\boldsymbol{Q})$ in (10). Here, the superscript "h" denotes the fact that only historical data are used to learn $\boldsymbol{Q}^{**}$. This two-stage estimator $\widehat{\eta}_h(x)$ may suffer from potential heterogeneity between historical and current data sources, since splitting mechanism of $\widehat{\eta}_h(x)$ is independent of current data set $\{(X_i, Y_{i,t_m}, D_i)\}_{i=1}^n$. It relies crucially on the transportability of historical optimal partitions $\{Q_j^{**}\}_{j=1}^q$. To leverage the information of current data, we aim to find a partition in which $E(Y_{t_m}^{(d)} \mid D = d, X = x)$ can be accurately learned by corresponding trees. Let $\widehat{m}(x, D; \boldsymbol{Q}) = DT(x; \boldsymbol{Q}, \widehat{\boldsymbol{\mu}}^{(1)}(\boldsymbol{Q})) + (1 - D)T(x; \boldsymbol{Q}, \widehat{\boldsymbol{\mu}}^{(0)}(\boldsymbol{Q}))$. We aim to find a $\boldsymbol{Q}$ such that $\widehat{m}(x, D; \boldsymbol{Q}) \approx DY_{t_m}^{(1)}(x) + (1 - D)Y_{t_m}^{(0)}(x)$ meanwhile $\widehat{\mathrm{H}}(\boldsymbol{Q})$ being small.

Regularized Regression Tree Estimator. For any problem-specific loss function $\ell(\cdot, \cdot)$, e.g., squared loss for regression and cross-entropy for classification, we propose the following regularized empirical risk minimization:

$$\widehat{\boldsymbol{Q}} = \operatorname*{argmin}_{\boldsymbol{Q}} \left\{ \frac{1}{n} \sum_{i=1}^{n} \ell\left(\widehat{m}(X_i, D_i; \boldsymbol{Q}), Y_{i,t_m}\right) + \lambda\widehat{\mathrm{H}}(\boldsymbol{Q}) \right\}, \tag{12}$$

where λ denotes the tuning parameter, and the CATT estimator is given by $\widehat{\eta}(x) = T(x; \widehat{\boldsymbol{Q}}, \widehat{\boldsymbol{\mu}}^{(1)}(\widehat{\boldsymbol{Q}})) - T(x; \widehat{\boldsymbol{Q}}, \widehat{\boldsymbol{\mu}}^{(0)}(\widehat{\boldsymbol{Q}}))$.

We remark that there exists a trade-off between data fidelity of the current source and transportability of historical sources for the choice of λ. When $\lambda \to \infty$, $\widehat{\eta}(\cdot)$ is reduced to the naive estimator proposed in (11), ignoring the information of current data. In contrast, $\lambda = 0$ means that we only use current data to learn CATT, which, under the unconfoundedness condition, is similar in spirit to causal tree estimators proposed in [47] and [3]. For the optimal choice of λ, our method minimizes the confounding bias introduced by unmeasured confounders that deteriorate the performance of conventional causal trees, and thus the name "debiased causal tree" follows.

2.3 Ensemble of Debiased Causal Trees by Gradient Boosting

Ensemble is a commonly used method to improve the performance of tree-based models. Boosting and bagging are two representative methods, where the former is designed to reduce bias, and the latter focuses on variance reduction. Existing tree-based methods developed for treatment effect estimation combine multiple trees to make a random forest, since unregularized decision trees are prone to overfitting, especially when the tree in employment is deep. In our case, the objective function (12) includes a regularization term. As a consequence, our estimator tends to be underfitting instead of overfitting, particularly when the working densities that define (8) are misspecified and/or the tuning parameter λ is relatively large. This motivates us to choose the boosting framework to improve the performance of single debiased causal tree. We define the boosting of k trees as $F(x; \boldsymbol{\mathcal{Q}}_k, \boldsymbol{M}_k) = \sum_{i=1}^{k} T(x; \boldsymbol{Q}_i, \boldsymbol{\mu}_i)$, where $\boldsymbol{\mathcal{Q}}_k = \{\boldsymbol{Q}_1, \ldots, \boldsymbol{Q}_k\} \equiv \boldsymbol{\mathcal{Q}}_{k-1} \cup \{\boldsymbol{Q}_k\}$ denotes the set of partitions of each tree, and $\boldsymbol{M}_k = \{\boldsymbol{\mu}_1, \ldots, \boldsymbol{\mu}_k\} \equiv \boldsymbol{M}_{k-1} \cup \{\boldsymbol{\mu}_k\}$ denotes the set of corresponding parameters. To highlight the difference, we use $\boldsymbol{\Theta}_k = \{\boldsymbol{\theta}_1, \ldots, \boldsymbol{\theta}_k\}$ to denote parameters of auxiliary tree models that measure confounding entropy.

At the k-th iteration, we have already obtained $k - 1$ trees, $F^{(d)}(x; \widehat{\boldsymbol{\mathcal{Q}}}_{k-1}, \widehat{\boldsymbol{M}}_{k-1}^{(d)})$, to learn $E(Y_{t_m}^{(d)} \mid D = d, X = x), d = 0, 1$, whose difference yields CATT, and $k - 1$ trees, $F_{\text{bias}}^{(d)}(x; \widehat{\boldsymbol{\mathcal{Q}}}_{k-1}, \widehat{\boldsymbol{\Theta}}_{k-1}^{(d)})$, to learn $E(Y_t \mid D = d, X = x), t < t_m$, whose difference yields confounding bias. Notice that the trees for confounding bias share the same splitting regions with trees for CATT. In the following, we illustrate how to obtain the k-th tree.

Parallel to a single debiased causal tree, we first show a refined representation of confounding entropy by integrating multiple trees. Following the gradient boosting algorithm [16], the k-th tree $T(x; \boldsymbol{Q}_k, \boldsymbol{\theta}_k^{(d)})$ manages to fit the residual, $Y_{i,t}(X_i) - F_{\text{bias}}^{(d)}(X_i; \widehat{\boldsymbol{\mathcal{Q}}}_{k-1}, \widehat{\boldsymbol{\Theta}}_{k-1}^{(d)})$ with $D_i = d$ for each $d = 0, 1$. Denote by $\widehat{\boldsymbol{\theta}}_k^{(d)}(\boldsymbol{Q}_k) = (\widehat{\theta}_k^{(d)}(Q_{k,j}))$ the local mean for any partition $\boldsymbol{Q}_k$, where $\widehat{\theta}_k^{(d)}(Q_{k,j})$ is defined similarly to (7) with $Y_{i,t}$ being replaced by residuals. Similar to (8), we use the cross mean squared error to represent the confounding entropy

$$\widehat{\mathrm{H}}_t(\boldsymbol{Q}_k) \equiv \sum_{i=1}^{n} \left\{ \frac{\left(Y_{i,t}^{(0)} - F_{\text{bias}}^{(1)}(X_i; \widetilde{\boldsymbol{\mathcal{Q}}}_k, \widehat{\boldsymbol{\Theta}}_k^{(1)})\right)^2}{|\{i : D_i = 0\}|} + \frac{\left(Y_{i,t}^{(1)} - F_{\text{bias}}^{(0)}(X_i; \widetilde{\boldsymbol{\mathcal{Q}}}_k, \widehat{\boldsymbol{\Theta}}_k^{(0)})\right)^2}{|\{i : D_i = 1\}|} \right\}, \quad (13)$$

where $\widetilde{\boldsymbol{\mathcal{Q}}}_k = \widehat{\boldsymbol{\mathcal{Q}}}_{k-1} \cup \{\boldsymbol{Q}_k\}$, $\widehat{\boldsymbol{\Theta}}_k^{(d)} = \widehat{\boldsymbol{\Theta}}_{k-1}^{(d)} \cup \{\widehat{\boldsymbol{\theta}}_k^{(d)}(\widehat{\boldsymbol{Q}}_k)\}$, and $Y_{i,t}^{(d)}$ denotes $Y_{i,t}$ with $D_i = d$.

It suffices to consider the integration of trees to learn $\delta_{t_m}(x)$. Again, the k-th tree $T(x; \boldsymbol{Q}_k, \boldsymbol{\mu}_k^{(d)})$ manages to fit the residual, $\widehat{R}_i^{(d)} = Y_{i,t_m}^{(d)} - F^{(d)}(X_i; \widehat{\boldsymbol{\mathcal{Q}}}_{k-1}, \widehat{\boldsymbol{M}}_{k-1}^{(d)})$ with $D_i = d$. Let $\widehat{\boldsymbol{\mu}}_k^{(d)}(\boldsymbol{Q}_k) = (\widehat{\mu}_k^{(d)}(Q_{k,j}))$ be the local mean for any partition $\boldsymbol{Q}_k$, where $\widehat{\mu}_k^{(d)}(Q_{k,j})$ is defined similarly to (11) with $Y_{i,t_m}^{(d)}$ being replaced by $\widehat{R}_i^{(d)}$. The objective function for the k-th tree is displayed as follows:

$$\widehat{\boldsymbol{Q}}_k = \underset{\boldsymbol{Q}_k}{\operatorname{argmin}} \left\{ \frac{1}{n} \sum_{i=1}^{n} \ell(\widehat{R}_i^{(D_i)}, T(X_i; \boldsymbol{Q}_k, \widehat{\boldsymbol{\mu}}_k^{(D_i)}(\boldsymbol{Q}_k))) + \frac{\lambda}{m-1} \sum_{u=1}^{m-1} \widehat{\mathrm{H}}_{t_u}(\boldsymbol{Q}_k) \right\}. \quad (14)$$

Update $\widehat{\boldsymbol{\mathcal{Q}}}_k = \widehat{\boldsymbol{\mathcal{Q}}}_{k-1} \cup \{\widehat{\boldsymbol{Q}}_k\}$, and $\widehat{\boldsymbol{M}}_k^{(d)} = \widehat{\boldsymbol{M}}_{k-1}^{(d)} \cup \{\widehat{\boldsymbol{\mu}}_k^{(d)}(\widehat{\boldsymbol{Q}}_k)\}, d = 0, 1$. The CATT estimator at the k-th iteration is $\widehat{\eta}(x) = F^{(1)}(x; \widehat{\boldsymbol{\mathcal{Q}}}_k, \widehat{\boldsymbol{M}}_k^{(1)}) - F^{(0)}(x; \widehat{\boldsymbol{\mathcal{Q}}}_k, \widehat{\boldsymbol{M}}_k^{(0)})$.

We refer to our proposed method as Gradient Boosting Causal Tree (GBCT). We remark that the objective function (14) is greedily minimized by gradually adding the tree. Also, to speed up the computation and stabilize the algorithm, we follow XGBoost to approximate the objective function with a second-order Taylor expansion and exploit a squared ℓ_2 regularization. See [11] for details.

3 Applicability of Debiased Causal Trees Without Confounders

In this section, we discuss the applicability of debiased causal trees without confounders. Despite that the confounding bias $\boldsymbol{b}_t(x) \equiv 0$ for any t and x without unmeasured confounding, numerically, the term $\widehat{\boldsymbol{b}}_t(Q_j)$ in (7) may not vanish, especially when the minimal sample size is small. This issue is substantive when treated and untreated units are imbalanced. Notably, the imbalance can significantly

degrade the performance of T-learners and S-learners [e.g., see 27]. Given the historical training set $\{(X_i, Y_{i,t}, D_i)\}_{i=1}^n$, the smaller empirical confounding entropy is for the partition $\boldsymbol{Q}$, the more balanced samples can be within each small region. Here, for simplicity, we consider $m = 2$ such that only one historical data source is used. Let $n_d(Q_j) = |\{i : D_i = d, X_i \in Q_j\}|$ be the number of samples in Q_j with $D_i = d$. We formalize the above intuition with the empirical confounding bias $\widehat{\mathrm{H}}(\boldsymbol{Q})$ defined in (10) with working distribution being Gaussian.

Proposition 1. *Suppose there are no unobserved confounders, and if $Y_{i,t} \sim Y_t$ with mean μ_t and variance σ_t^2 independently for a given $t < t_m$, then, under the Gaussian specification of working densities, it holds that $E\left[\widehat{\mathrm{H}}(\boldsymbol{Q})\right] = 2\sigma_t^2 + \sigma_t^2 \sum_{j=1}^q (n_0(Q_j) + n_1(Q_j))/(n_0(Q_j)n_1(Q_j))$.*

By Proposition 1 and the harmonic mean inequality, to reduce $\widehat{\mathrm{H}}(\boldsymbol{Q})$, it suffices to find a Q_j such that $n_0(Q_j) \approx n_1(Q_j)$, especially for Q_j such that $\max\{n_0(Q_j), n_1(Q_j)\}$ is small. Numerical analysis can be found in Figure 1 in the Appendix, which indicates that, our proposed debiased causal tree is robust to imbalance of treatments.

4 Experiments

We conduct experiments with both simulated and real data for a better understanding of our proposed strategy. We investigate two scenarios in simulation studies: one with and one without unobserved confounding. When implementing GBCT, we subtract the empirical pre-treatment confounding bias from effect estimates as a trick to improve performance, i.e., $\widehat{\eta}(x) = F^{(1)}(x; \widehat{\boldsymbol{\mathcal{Q}}}_k, \widehat{\boldsymbol{M}}_k^{(1)}) - F^{(0)}(x; \widehat{\boldsymbol{\mathcal{Q}}}_k, \widehat{\boldsymbol{M}}_k^{(0)}) - \frac{1}{m-1}\sum_{u=1}^{m-1}\sum_{Q \in \widehat{\boldsymbol{\mathcal{Q}}}_k} \widehat{\boldsymbol{b}}_{t_u}(Q)$. We evaluate GBCT against state-of-the-art causal inference algorithms: (i) meta learners [27], which adopt machine learning models to fit the response surfaces on the treated and control populations separately, including TLearner, SLearner and Xlearner; (ii) causal forests, which are forest-based methods to model the treatment effects, including generalized causal forest [GRF, 4], doubly robust forest [DR-RF, 5] and double machine learning forest [DML-RF, 12]. These benchmarks are implemented by the third-party library ECOML [24].

To make a fair experimental comparison, we set all hyperparameters using the same strategy. The number of trees in ensemble models (including boosting and bagging) is 200, the sub-sample ratios of instance and feature are 0.8 and 0.6 respectively, and the learning rate is 0.3. The maximum depth of each tree in forest-based (GRF, DML-RF and DR-RF) and boosting-based (meta learners and GBCT) methods is 10 and 3, respectively, where it should be noted that due to the respective characteristics of the bagging and boosting frameworks, the trees in random forests are generally deeper. Meta learners employ LightGBM as base models and causal forests utilize sub-sampled honest random forest [3]. Both covariates X and pre-treatment outcomes are used as input features for benchmark methods. However, we only use covariates X to build GBCT, and the pre-treatment outcomes are treated as labels to de-bias in training phrases.

4.1 Simulation Studies

Data Generating Process. We start by generating a p-dimensional confounder vector X from $\sum_{s=1}^S \mathrm{I}(G = s)\mathcal{N}(\mu_s, \Sigma_s)$, where $G \in \{1, 2, \cdots, S\}$ is a hidden variable indicating which latent cluster each subject belongs to. The treatment-independent feature vector W is developed similarly. We then generate the potential outcomes by the following factor augmented auto regressive process of order one, for each $1 \le k \le m$,

$$Y_{t_k}^{(0)} = \rho Y_{t_{k-1}}^{(0)} + (1 - \rho)\{\alpha(X, W) + f(X, W, \lambda_t) + \varepsilon\}, \quad Y_{t_m}^{(1)} = Y_{t_m}^{(0)} + g(X) + \varepsilon_1,$$

where $\rho \in [0, 1]$ is the individual-specific auto-correlation coefficient following a truncated normal distribution $\bar{\mathcal{N}}(\mu_\rho, \sigma_\rho^2; [0, 1])$, λ_t is the time-varying factor, the initial value $Y_{t_0}^{(0)} = \beta(X, W)$, $\{\alpha(\cdot), \beta(\cdot), f(\cdot), g(\cdot)\}$ are specified functions, and $\varepsilon, \varepsilon_1$ are mutually independent noise items subject to $\mathcal{N}(0, 1)$. To ensure the samples are drawn from the stationary distribution of the data generating process, we use outcomes with time larger than a threshold t_T as observations.

We assign treatment for each unit according to its latent cluster, such that $P(D = 1 \mid G = s) - P(D = 0 \mid G = s) = (-1)^s\phi$ for $s = 1, 2, \cdots, S$, where $\phi \in [0, 1]$ is a specified value. The parameter ϕ measures the degree of balance between the treated and control populations; as ϕ rises from zero

to one, the imbalance gradually increases. In particular, $\phi = 0$ means the treated group has the same population as the control group (i.e., RCT), whereas $\phi = 1$ means all treated group's units are from different latent clusters than the control group's. In following experiments, we set $S = 2$, $\mu_\rho = 0.7, \sigma_\rho = 0.2$, and the dimension of feature space $2p = 20$. A total of 20000 samples are generated and randomly split into training and validation sets 10 times.

No Unmeasured Confounding Scenario. The confounding vector X blocks the backdoor path from treatment to potential outcomes. Thus, given X, potential outcomes $Y_{t_m}^{(d)}$ is independent of D for $d \in \{0, 1\}$. Suppose that we observe all confounders X, and apply `GBCT` as well as benchmarks to estimate the conditional treatment effects. We compare our proposed method with benchmark algorithms on the mean absolute error (MAE) of conditional average treatment effects, i.e., $\text{MAE}_{\text{CATE}} = \frac{1}{n} \sum_{i=1}^n |\hat{\eta}(X_i) - \eta_i|$, where η_i is the true value of the ith individual's treatment effect. Table 1 shows the results. As we can see, `GBCT` outperforms the benchmark algorithms in most cases, especially when ϕ is larger, suggesting that it is beneficial to reduce the empirical confounding bias in case of no unmeasured confounding. As ϕ increases towards one, except for `GBCT` and `GRF`, other methods generally lead to severe bias and significant performance degradation due to low overlap. This confirms the robustness of `GBCT` against imbalance between the treated and control populations. Moreover, we perform ablation experiments to compare `GBCT` with its variants without using de-bias loss. The results presented in the Appendix demonstrate the important role of the strategies to reduce confounding bias in making the estimates less sensitive to the degree of imbalance.

Table 1: MAE (mean±s.d.) of each algorithm on the simulated data. Scenario I represents the absence of unmeasured confounding, whereas II represents the presence of unmeasured confounding. In addition, •/∘ indicates whether `GBCT` is statistically superior/inferior to the benchmark algorithms (pairwise t-test at 0.05 significance level).

| | ϕ | meta learners | | | | causal forests | | |
		Slearner	Tlearner	Xlearner	DR-RF	DML-RF	GRF	GBCT
	0.2	1.26 ± 0.09•	1.16 ± 0.07•	1.06 ± 0.05∘	1.23 ± 0.01•	1.28 ± 0.01•	1.26 ± 0.01•	1.10 ± 0.01
	0.5	1.46 ± 0.13•	1.50 ± 0.17•	1.19 ± 0.10	1.29 ± 0.02•	1.34 ± 0.03•	1.31 ± 0.02•	1.25 ± 0.03
I	0.8	1.52 ± 0.11•	2.02 ± 0.46•	1.75 ± 0.53•	1.51 ± 0.23•	1.48 ± 0.08•	1.39 ± 0.04•	1.31 ± 0.02
	0.9	1.48 ± 0.03•	1.84 ± 0.22•	1.43 ± 0.21•	1.80 ± 0.44•	1.44 ± 0.08•	1.41 ± 0.12•	1.28 ± 0.03
	1	4.56 ± 0.13•	3.28 ± 1.59•	2.46 ± 0.99•	4.73 ± 0.08•	16.02 ± 9.39•	1.96 ± 0.20•	1.71 ± 0.06
	0.2	1.48 ± 0.07•	1.97 ± 0.53•	1.54 ± 0.28•	1.60 ± 0.02•	1.66 ± 0.02•	1.28 ± 0.01•	1.07 ± 0.01
	0.5	1.59 ± 0.08•	3.54 ± 1.90•	2.81 ± 1.75•	1.70 ± 0.09•	1.72 ± 0.05•	1.31 ± 0.02•	1.25 ± 0.02
II	0.8	1.72 ± 0.09•	4.10 ± 1.72•	4.07 ± 2.07•	3.94 ± 3.31•	2.14 ± 0.43•	1.40 ± 0.06•	1.29 ± 0.04
	0.9	2.30 ± 0.07•	2.63 ± 0.29•	2.35 ± 0.23•	2.13 ± 0.03•	3.10 ± 0.12•	1.72 ± 0.03•	1.65 ± 0.03
	1	3.84 ± 0.24•	6.32 ± 3.49•	4.64 ± 2.94•	5.20 ± 0.06•	2.84 ± 0.48•	2.38 ± 0.17•	1.81 ± 0.10

Unmeasured Confounding Scenario. We consider the scenario where the confounding vector X are not fully observed. For estimation and evaluation, we retain only the first dimension of the confounding vector and discard the rest. When $\phi = 0$, the treatment is assigned randomly, resulting in no confounding bias. However, as ϕ approaches one, the impact of confounding on treatment assignment grows stronger. Therefore, a large ϕ suggests a substantial confounding bias due to unmeasured confounding. In this case, most traditional methods may fail to adjust for bias without further assumptions, while in contrast, `GBCT` can approximate CATT well. To support the superiority of our approach, we use $\text{MAE}_{\text{CATT}} = \left(\sum_{i=1}^n D_i\right)^{-1} \sum_{i=1}^n D_i |\hat{\eta}(X_i) - \eta_i|$ as the criterion to evaluate the performance of algorithms, and compare `GBCT` with the benchmarks in numerical studies. We report the results in Table 1. As expected, `GBCT` yields smaller mean absolute errors than other methods, and the gap becomes more pronounced as ϕ increases, indicating that `GBCT` can offer a more robust and compelling solution for estimating conditional treatment effects in the presence of unmeasured confounding. We also compare the performance of `GBCT` with `GRF` at the single-tree scale on removing confounding bias, showing that our method can be more effective, see the Appendix.

4.2 Real Data: Credit Card Balance Dataset

The dataset comes from a randomized controlled trial (RCT) by a commercial finance company aimed at assessing users' heterogeneous responses to increasing credit line of credit card. The trial employs a stratified random assignment design with strata based on risk, dividing users into two groups: low-risk and medium-risk. Within each group, users are randomly assigned to one of four treatment groups:

increasing credit line by 0, 2000, 3000 or 6000 converted to some currency. In this study, the outcome of interest is the balance of the credit card account in the following month after treatment, i.e., the total amount of debt owed to the credit card. In addition, we observe 87-dimensional covariates at the individual level, as well as outcome values over the last eight months.

In real-world financial scenarios, however, increasing credit line always favors low-risk users; such preference can lead to confounding bias in estimation. In order to assess the performance of different methods in the real-world scenario, we artificially construct a biased observational dataset by increasing credit line for the low-risk group but not for the medium-risk group. See Table 3 in the Appendix for details of the trial dataset and the biased observational dataset.

We are interested in the causal effects of increasing credit line on balance within each risk group. Unlike simulated data, we do not know all potential outcomes for each user, so we cannot use $\mathrm{MAE}_{\mathrm{CATE}}$ for evaluation. Fortunately, due to the randomness of treatment allocation, the ground-truth of the average treatment effect is approximately the mean difference between any two treatment groups in the trial dataset, denoted as η_{ATE}. Note that in RCT, the average treatment effect is equal to the average treatment effect on the treated group. Therefore, we compare our proposed method GBCT with the benchmarks on the mean absolute error of the average treatment effect, $\mathrm{MAE}_{\mathrm{ATE}} = \left| n^{-1} \sum_{i=1}^n \hat{\eta}(X_i) - \eta_{\mathrm{ATE}} \right|$.

Table 2: The mean absolute error $\mathrm{MAE}_{\mathrm{ATE}}$ (mean$\pm$ s.d.) of each algorithm on the credit card balance dataset. The left and right sides are trained on the biased observational and the trial data, respectively. We treat the users with no increase in credit line as the control group.

	risk	Biased			RCT		
		2k	3k	6k	2k	3k	6k
TLearner	medium-risk	49.1 ± 39.8	106.5 ± 119.7	178.6 ± 116.4	92.6 ± 75.3	75.2 ± 55.6	76.9 ± 39.1
	low-risk	57.0 ± 39.0	104.8 ± 122.2	197.7 ± 87.4	87.8 ± 76.9	51.8 ± 55.4	53.9 ± 34.7
XLearner	medium-risk	88.7 ± 129.0	85.0 ± 139.3	99.6 ± 132.9	98.8 ± 106.4	93.0 ± 81.2	93.0 ± 48.7
	low-risk	62.4 ± 129.0	99.4 ± 139.3	127.1 ± 132.9	97.0 ± 106.4	81.5 ± 81.2	98.1 ± 48.7
SLearner	medium-risk	207.3 ± 26.3	253.4 ± 37.8	261.9 ± 59.4	12.9 ± 18.8	23.4 ± 9.7	24.7 ± 10.9
	low-risk	95.1 ± 28.4	100.4 ± 33.7	45.1 ± 39.6	41.3 ± 8.4	59.0 ± 29.2	100.9 ± 34.9
DR-RF	medium-risk	249.6 ± 9.3	317.7 ± 12.2	408.6 ± 9.4	8.6 ± 7.5	14.2 ± 7.73	48.4 ± 8.8
	low-risk	149.0 ± 6.5	162.8 ± 8.9	151.2 ± 11.1	35.0 ± 5.0	31.7 ± 6.9	55.5 ± 9.4
DML-RF	medium-risk	> 999.9	> 999.9	> 999.9	$\mathbf{8.5 \pm 6.7}$	15.7 ± 8.6	56.4 ± 8.6
	low-risk	> 999.9	> 999.9	> 999.9	34.5 ± 4.9	32.8 ± 6.4	62.4 ± 9.3
GBCT	medium-risk	$\mathbf{10.6 \pm 6.1}$	$\mathbf{27.5 \pm 13.6}$	$\mathbf{61.5 \pm 29.5}$	13.2 ± 7.6	$\mathbf{8.0 \pm 6.0}$	$\mathbf{14.2 \pm 9.2}$
	low-risk	$\mathbf{8.4 \pm 5.8}$	$\mathbf{11.5 \pm 8.6}$	$\mathbf{19.3 \pm 22.5}$	$\mathbf{21.7 \pm 5.2}$	$\mathbf{11.6 \pm 7.4}$	$\mathbf{22.8 \pm 9.9}$

The results are reported in Table.2. It is impressive to observe that on both trial and biased data, GBCT significantly outperforms the benchmarks in terms of mean error and standard deviation. Although the benchmarks DR-RF and DML-RF do not perform poorly on the trial data, they suffer severely from bias regarding the observational data; a possible reason is the low overlap between the low-risk and medium-risk groups. As a comparison, GBCT suffers substantially less from bias than others. These findings are in line with those of the prior simulation studies. Therefore, we recommend using GBCT in practice to gain a more stable and robust estimate of causal effects.

5   Discussion

In this work, we propose a simple yet powerful method, debiased causal tree, to remove confounding bias caused by unmeasured confounding in the estimation of causal effects (e.g., CATT). Our method only hinges on the availability and transportability of historical controls, i.e., outcomes of units before the treatment, to ensure CATT to be partially identifiable. This allows our method to be readily applicable in a wide variety of scenarios, especially in clinical trials [45]. Moreover, our method can be generalized to integrate observational data and external control [30], with a much weaker transportability condition than existing work [40]. Theoretically, our identifiability result provides the possibility of statistical justification for the proposed estimators with finite samples. For example, provided that the optimal partition Q^{**} is estimated accurately, the convergence rate of $\left| \hat{\eta}(\cdot) - \eta^*(\cdot) \right|$ can be derived using the adaptive concentration results developed in [48], which will be left for future work.

References

[1] Joshua D. Angrist, Guido W. Imbens, and Donald B. Rubin. Identification of causal effects using instrumental variables. *Journal of the American Statistical Association*, 91(434):444–455, 1996.

[2] Susan Athey. Beyond prediction: Using big data for policy problems. *Science*, 355(6324):483–485, 2017.

[3] Susan Athey and Guido Imbens. Recursive partitioning for heterogeneous causal effects. *Proceedings of the National Academy of Sciences*, 113(27):7353–7360, 2016.

[4] Susan Athey, Julie Tibshirani, and Stefan Wager. Generalized random forests. *The Annals of Statistics*, 47(2):1148–1178, 2019.

[5] Heejung Bang and James M Robins. Doubly robust estimation in missing data and causal inference models. *Biometrics*, 61(4):962–973, 2005.

[6] Elias Bareinboim and Judea Pearl. Controlling selection bias in causal inference. In Neil D. Lawrence and Mark Girolami, editors, *Proceedings of the Fifteenth International Conference on Artificial Intelligence and Statistics*, volume 22 of *Proceedings of Machine Learning Research*, pages 100–108, La Palma, Canary Islands, 21–23 Apr 2012. PMLR.

[7] Takoua Boukhris, Odile Sheehy, Laurent Mottron, and Anick Bérard. Antidepressant use during pregnancy and the risk of autism spectrum disorder in children. *JAMA pediatrics*, 170(2):117–124, 2016.

[8] Leo Breiman. Bagging predictors. *Machine learning*, 24(2):123–140, 1996.

[9] Leo Breiman, Jerome H Friedman, Richard A Olshen, and Charles J Stone. *Classification and regression trees*. Routledge, 2017.

[10] Lester Breslow and Michael Johnson. California's proposition 99 on tobacco, and its impact. *Annual Review of Public Health*, 14(1):585–604, 1993.

[11] Tianqi Chen and Carlos Guestrin. Xgboost: A scalable tree boosting system. In *Proceedings of the 22nd acm sigkdd international conference on knowledge discovery and data mining*, pages 785–794, 2016.

[12] Victor Chernozhukov, Denis Chetverikov, Mert Demirer, Esther Duflo, Christian Hansen, Whitney Newey, and James Robins. Double/debiased machine learning for treatment and structural parameters. *The Econometrics Journal*, 21(1):C1–C68, 01 2018.

[13] Bénédicte Colnet, Imke Mayer, Guanhua Chen, Awa Dieng, Ruohong Li, Gaël Varoquaux, Jean-Philippe Vert, Julie Josse, and Shu Yang. Causal inference methods for combining randomized trials and observational studies: a review. *arXiv preprint arXiv:2011.08047*, 2020.

[14] John Concato, Nirav Shah, and Ralph I Horwitz. Randomized, controlled trials, observational studies, and the hierarchy of research designs. *New England journal of medicine*, 342(25):1887–1892, 2000.

[15] Yifan Cui, Hongming Pu, Xu Shi, Wang Miao, and Eric Tchetgen Tchetgen. Semiparametric proximal causal inference. *arXiv preprint arXiv:2011.08411*, 2020.

[16] Jerome H Friedman. Greedy function approximation: a gradient boosting machine. *Annals of statistics*, pages 1189–1232, 2001.

[17] Johann A Gagnon-Bartsch and Terence P Speed. Using control genes to correct for unwanted variation in microarray data. *Biostatistics*, 13(3):539–552, 2012.

[18] Alexandre Gilotte, Clément Calauzènes, Thomas Nedelec, Alexandre Abraham, and Simon Dollé. Offline A/B testing for recommender systems. In *Proceedings of the Eleventh ACM International Conference on Web Search and Data Mining*, pages 198–206, 2018.

[19] Thomas A. Glass, Steven N. Goodman, Miguel A. Hernán, and Jonathan M Samet. Causal inference in public health. *Annual review of public health*, 34:61–75, 2013.

[20] Wilson Wen Bin Goh, Wei Wang, and Limsoon Wong. Why batch effects matter in omics data, and how to avoid them. *Trends in Biotechnology*, 35(6):498–507, 2017. Special Issue: Computation and Modeling.

[21] Brett R Gordon, Florian Zettelmeyer, Neha Bhargava, and Dan Chapsky. A comparison of approaches to advertising measurement: Evidence from big field experiments at facebook. *Marketing Science*, 38(2):193–225, 2019.

[22] Gareth James, Daniela Witten, Trevor Hastie, and Robert Tibshirani. Tree-based methods. In *An introduction to statistical learning*, pages 327–365. Springer, 2021.

[23] Leslie A. Kalish and Colin B. Begg. Treatment allocation methods in clinical trials: A review. *Statistics in Medicine*, 4(2):129–144, 1985.

[24] Battocchi Keith, Dillon Eleanor, Hei Maggie, Lewis Greg, Oka Paul, Oprescu Miruna, and Syrgkanis Vasilis. EconML: A Python Package for ML-Based Heterogeneous Treatment Effects Estimation. https://github.com/microsoft/EconML, 2019. Version 0.x.

[25] Edward H Kennedy. Optimal doubly robust estimation of heterogeneous causal effects. *arXiv preprint arXiv:2004.14497*, 2020.

[26] W Kernan. Stratified randomization for clinical trials. *Journal of Clinical Epidemiology*, 52(1):19–26, January 1999.

[27] Sören R Künzel, Jasjeet S Sekhon, Peter J Bickel, and Bin Yu. Metalearners for estimating heterogeneous treatment effects using machine learning. *Proceedings of the national academy of sciences*, 116(10):4156–4165, 2019.

[28] Manabu Kuroki and Judea Pearl. Measurement bias and effect restoration in causal inference. *Biometrika*, 101(2):423–437, 03 2014.

[29] Lihong Li, Wei Chu, John Langford, and Robert E Schapire. A contextual-bandit approach to personalized news article recommendation. In *Proceedings of the 19th international conference on World wide web*, pages 661–670, 2010.

[30] Xinyu Li, Wang Miao, Fang Lu, and Xiao-Hua Zhou. Improving efficiency of inference in clinical trials with external control data. *Biometrics*, November 2021.

[31] Wei Lin, Rui Feng, and Hongzhe Li. Regularization methods for high-dimensional instrumental variables regression with an application to genetical genomics. *Journal of the American Statistical Association*, 110(509):270–288, 2015. PMID: 26392642.

[32] Wang Miao, Zhi Geng, and Eric J Tchetgen Tchetgen. Identifying causal effects with proxy variables of an unmeasured confounder. *Biometrika*, 105(4):987–993, 08 2018.

[33] Xinkun Nie and Stefan Wager. Quasi-oracle estimation of heterogeneous treatment effects. *Biometrika*, 108(2):299–319, 2021.

[34] Ziad Obermeyer and Ezekiel J Emanuel. Predicting the future—big data, machine learning, and clinical medicine. *The New England journal of medicine*, 375(13):1216, 2016.

[35] Judea Pearl and Elias Bareinboim. Transportability of causal and statistical relations: A formal approach. In *Twenty-fifth AAAI conference on artificial intelligence*, 2011.

[36] P. R. Rosenbaum and D. B. Rubin. Assessing sensitivity to an unobserved binary covariate in an observational study with binary outcome. *Journal of the Royal Statistical Society: Series B (Methodological)*, 45(2):212–218, 1983.

[37] Peter M Rothwell. External validity of randomised controlled trials:"to whom do the results of this trial apply?". *The Lancet*, 365(9453):82–93, 2005.

[38] Donald B Rubin. Bayesian inference for causal effects: The role of randomization. *The Annals of statistics*, pages 34–58, 1978.

[39] Xu Shi, Wang Miao, Jennifer C. Nelson, and Eric J. Tchetgen Tchetgen. Multiply robust causal inference with double-negative control adjustment for categorical unmeasured confounding. *Journal of the Royal Statistical Society: Series B (Statistical Methodology)*, 82(2):521–540, 2020.

[40] Xu Shi, Ziyang Pan, and Wang Miao. Data integration in causal inference. *WIREs Computational Statistics*, April 2022.

[41] Sonja A. Swanson, Miguel A. Hernán, Matthew Miller, James M. Robins, and Thomas S. Richardson. Partial identification of the average treatment effect using instrumental variables: Review of methods for binary instruments, treatments, and outcomes. *Journal of the American Statistical Association*, 113(522):933–947, 2018. PMID: 31537952.

[42] Carmen D. Tekwe, Roger S. Zoh, Miao Yang, Raymond J. Carroll, Gilson Honvoh, David B. Allison, Mark Benden, and Lan Xue. Instrumental variable approach to estimating the scalar-on-function regression model with measurement error with application to energy expenditure assessment in childhood obesity. *Statistics in Medicine*, 38(20):3764–3781, 2019.

[43] Christopher Tran and Elena Zheleva. Learning triggers for heterogeneous treatment effects. In *Proceedings of the AAAI Conference on Artificial Intelligence*, volume 33, pages 5183–5190, 2019.

[44] Cristiano Varin, Nancy Reid, and David Firth. An overview of composite likelihood methods. *Statistica Sinica*, pages 5–42, 2011.

[45] Kert Viele, Scott Berry, Beat Neuenschwander, Billy Amzal, Fang Chen, Nathan Enas, Brian Hobbs, Joseph G. Ibrahim, Nelson Kinnersley, Stacy Lindborg, Sandrine Micallef, Satrajit Roychoudhury, and Laura Thompson. Use of historical control data for assessing treatment effects in clinical trials. *Pharmaceutical Statistics*, 13(1):41–54, August 2013.

[46] Sholom Wacholder. When measurement errors correlate with truth: Surprising effects of nondifferential misclassification. *Epidemiology*, 6(2):157–161, 1995.

[47] Stefan Wager and Susan Athey. Estimation and inference of heterogeneous treatment effects using random forests. *Journal of the American Statistical Association*, 113(523):1228–1242, 2018.

[48] Stefan Wager and Guenther Walther. Adaptive concentration of regression trees, with application to random forests. *arXiv preprint arXiv:1503.06388*, 2015.

[49] Yisen Wang, Xingjun Ma, Zaiyi Chen, Yuan Luo, Jinfeng Yi, and James Bailey. Symmetric cross entropy for robust learning with noisy labels. In *Proceedings of the IEEE/CVF International Conference on Computer Vision*, pages 322–330, 2019.

[50] Shu Yang, Donglin Zeng, and Xiaofei Wang. Improved inference for heterogeneous treatment effects using real-world data subject to hidden confounding. *arXiv preprint arXiv:2007.12922*, 2020.

A Appendix

In this section, we include related proofs, additional simulation plots that demonstrate the necessity of our proposed debiased causal tree, and a brief introductive table of the real-data.

A.1 Related Proofs

Proof of Theorem 1. By the triangular inequality, we obtain

$$|\eta(x) - \eta^*(x)| \leq \sum_{j=1}^{q} |\delta_{t_m}(Q_j^{**}) - \eta^*(Q_j^{**})| \mathrm{I}\{x \in Q_j^{**}\} + \sum_{j=1}^{q} |\eta^*(Q_j^{**}) - \eta^*(x)\mathrm{I}\{x \in Q_j^{**}\}|$$

$$\leq \sum_{j=1}^{q} |\boldsymbol{b}_{t_m}(Q_j^{**})| \mathrm{I}\{x \in Q_j^{**}\} + \max_{1 \leq j \leq q} \sup_{x \in Q_j^{**}} |\eta^*(Q_j^{**}) - \eta^*(x)|. \qquad (15)$$

By the Cauchy–Schwarz inequality, we further obtain

$$\sum_{j=1}^{q} |\boldsymbol{b}_{t_m}(Q_j^{**})| \mathrm{I}\{x \in Q_j^{**}\} \leq \left\{ \sum_{j=1}^{q} |\boldsymbol{b}_{t_m}(Q_j^{**})|^2 \right\}^{1/2}. \qquad (16)$$

Next, we bound $\sum_{j=1}^{q} |\boldsymbol{b}_{t_m}(Q_j^{**})|^2 = \exp\{B(t_m)\}$. By Assumption 1, it holds that

$$B(t_m) \leq B(t_k) + L|t_m - t_k|, \quad \text{i.e.,} \quad \sum_{j=1}^{q} |\boldsymbol{b}_{t_m}(Q_j^{**})|^2 \leq e^{L|t_m - t_k|} \sum_{j=1}^{q} |\boldsymbol{b}_{t_k}(Q_j^{**})|^2,$$

which implies that

$$\sum_{j=1}^{q} |\boldsymbol{b}_{t_m}(Q_j^{**})|^2 \leq e^{L(t_m - t_1)} \frac{1}{m-1} \sum_{k=1}^{m-1} \sum_{j=1}^{q} |\boldsymbol{b}_{t_k}(Q_j^{**})|^2. \qquad (17)$$

Choosing $C_L \equiv e^{L(t_m - t_1)}$, and combining (15)–(17) give

$$\sup_{x \in [0,1]^p} |\eta(x) - \eta^*(x)| \leq \left\{ \frac{C_L}{m-1} \sum_{k=1}^{m-1} \sum_{j=1}^{q} \boldsymbol{b}_{t_k}^2(Q_j^{**}) \right\}^{1/2} + \max_{1 \leq j \leq q} \sup_{x \in Q_j^{**}} |\eta^*(x) - \eta^*(Q_j^{**})|.$$

$$\square$$

Proof of Proposition 1. By (10), under the Gaussian specification of working densities,

$$\widehat{\mathrm{H}}(\boldsymbol{Q}) = \sum_{j=1}^{q} \left\{ (\widehat{\theta}_{t,j}^{(0)} - \widehat{\theta}_{t,j}^{(1)})^2 + \sum_{i: X_i \in Q_j} \left\{ \frac{D_i(Y_{i,t} - \widehat{\theta}_{t,j}^{(1)})^2}{|\{i : X_i \in Q_j, D_i = 1\}|} + \frac{(1-D_i)(Y_{i,t} - \widehat{\theta}_{t,j}^{(0)})^2}{|\{i : X_i \in Q_j, D_i = 0\}|} \right\} \right\}.$$

Under the unconfoundedness condition, if $Y_{i,t} \sim Y_t$ with mean μ_t and variance σ_t^2 independently, for each given Q_j,

$$\widehat{\theta}_{t,j}^{(d)} = \sum_{i=1}^{n} \frac{Y_{i,t}\mathrm{I}\{D_i = d, X_i \in Q_j\}}{n_d(Q_j)}, \quad \text{where} \quad n_d(Q_j) = |\{i : X_i \in Q_j, D_i = d\}|,$$

is the mean of i.i.d random variables with finite second moment and $\widehat{\theta}_{t,j}^{(0)}$ is independent of $\widehat{\theta}_{t,j}^{(1)}$. Thus,

$$E\left[(\widehat{\theta}_{t,j}^{(0)} - \widehat{\theta}_{t,j}^{(1)})^2\right] = \sigma_t^2 \left(\frac{1}{n_1(Q_j)} + \frac{1}{n_0(Q_j)} \right),$$

and

$$E\left[\sum_{i: X_i \in Q_j} \left\{ \frac{D_i(Y_{i,t} - \widehat{\theta}_{t,j}^{(1)})^2}{|\{i : X_i \in Q_j, D_i = 1\}|} + \frac{(1-D_i)(Y_{i,t} - \widehat{\theta}_{t,j}^{(0)})^2}{|\{i : X_i \in Q_j, D_i = 0\}|} \right\} \right] = \sigma_t^2 \left(\frac{n_1(Q_j) - 1}{n_1(Q_j)} + \frac{n_0(Q_j) - 1}{n_0(Q_j)} \right).$$

We then conclude that

$$E\left[\widehat{\mathrm{H}}(\boldsymbol{Q})\right] = 2\sigma_t^2 + \sigma_t^2 \sum_{j=1}^{q} \frac{n_0(Q_j) + n_1(Q_j)}{n_0(Q_j)n_1(Q_j)}.$$

$$\square$$

 A.2 Additional Simulation Plots

(a) ϕ=0.9   (b) ϕ=1

Figure 1: The confounding bias on the validation data under different depths of a single causal tree in the unmeasured confounding scenario. The y-axis represents the empirical confounding bias and x-axis represents the depth of the causal tree. The results show that GBCT performs better than GRF at the single-tree scale in reducing confounding bias.

Figure 2: Ablation experiment: $\mathrm{MAE}_{\mathrm{CATE}}$ changes as ϕ increases from zero to one. "NR" indicates that the pre-treatment confounding bias is not subtracted, and "ND" indicates that $\widehat{\mathrm{H}}(\boldsymbol{Q})$ is not used when splitting tree nodes with $\lambda = 0$. It can be seen that the confounding entropy plays an important role in improving performance.

 A.3   Additional Real Data Description

Table 3: Description of the credit card balance dataset. The biased observational dataset only consists of the underlined instances. The features consist of the time-invariant covariates and the balance of credit card account over the last eight months.

Risk	Number of Instances				Number of Features
	0	2k	3k	6k	
medium-risk	389477	390928	391211	391215	87+8
low-risk	456773	459762	459443	460352	