

Bayesian regression tree models for causal inference: Regularization, confounding, and heterogeneous effects

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Joint work with Jared Murray, P. Richard Hahn, David Yeager, et al.

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Policy Analytics (moving past the 1900's)

The Effect of Corporate Governance on Investment 363

Table 5 ■ Firm value as a function of governance.

Dependent Variable: <i>Firm q</i>	(1)	(2)	(3)	(4)	(5)
<i>Property Type q</i>	0.497 (14.33)***	0.418 (7.64)***	0.403 (7.65)***	0.412 (7.16)***	0.382 (10.44)***
<i>EBITDA</i>	0.403 (5.31)***	0.456 (5.17)***	0.444 (5.11)***	0.444 (5.00)***	0.163 (7.21)***
<i>UPREIT</i>	-0.001 (0.02)	-0.006 (0.09)	-0.023 (0.36)	-0.018 (0.28)	
<i>Interest Coverage</i>	0.057 (0.74)	0.060 (0.83)	0.043 (0.63)	0.038 (0.57)	-0.004 (0.15)
<i>Mkt Cap</i>	0.127 (2.73)***	0.078 (1.85)*	0.087 (1.96)*	0.096 (2.11)**	0.014 (0.39)
<i>Excess Comp</i>		-0.002 (0.03)	0.000 (0.01)	-0.002 (0.05)	-0.020 (0.85)
<i>Instl Ownership</i>		0.053 (1.00)	0.078 (1.48)	0.085 (1.50)	0.101 (2.55)**
<i>Block Ownership</i>			-0.046 (1.38)	-0.041 (1.23)	0.013 (0.59)
<i>D&O Ownership</i>			0.106 (1.57)	0.105 (1.55)	0.072 (2.08)**
<i>Ln(Board Size)</i>				-0.044 (0.77)	-0.097 (2.86)***
<i>Outside Board</i>				0.029 (0.75)	0.021 (0.93)
<i>Maryland</i>				-0.026 (0.53)	
Fixed Effects?	No	No	No	No	Yes
Observations	882	882	882	882	882
R ²	0.53	0.55	0.56	0.56	0.60
<i>p</i> value from <i>F</i> test of null that all governance coefficients are zero		0.61	0.21	0.50	0.00***

Moving from this...

$$y = \alpha Z + \mathbf{X}\beta + \epsilon$$

Regularization and Confounding in Linear Regression for Treatment Effect Estimation

P. Richard Hahn^{*}, Carlos M. Carvalho[†], David Puelz[†], and Jingyu He^{*}

Abstract. This paper investigates the use of regularization priors in the context of treatment effect estimation using observational data where the number of control variables is large relative to the number of observations. First, the phenomenon of “regularization-induced confounding” is introduced, which refers to the tendency of regularization priors to adversely bias treatment effect estimates by over-shrinking control variable regression coefficients. Then, a simultaneous regression model is presented which permits regularization priors to be specified in a way that avoids this unintentional “re-confounding”. The new model is illustrated on synthetic and empirical data.

Keywords: causal inference, observational data, shrinkage estimation.

... to that!

$$y = f(Z, \mathbf{X}) + \epsilon$$

$$y = f(Z, \mathbf{X}) + \epsilon$$

Today:

- a general, “default” framework
- a rich output that allows you to ask lots of different questions

Our setting

We'll assume:

- **Observational and experimental data**
- **Conditional unconfoundedness/ignorability** (we've measured all the factors causally influencing treatment and response),
- **Covariate-dependent treatment effects** (individuals can have different responses to treatment according to their covariates)
- **Binary treatments**

Our assumptions, more formally

Strong ignorability:

$$Y_i(0), Y_i(1) \perp\!\!\!\perp Z_i \mid \mathbf{X}_i = \mathbf{x}_i,$$

Positivity:

$$0 < \Pr(Z_i = 1 \mid \mathbf{X}_i = \mathbf{x}_i) < 1$$

for all i . Therefore

$$\mathbb{E}(Y_i(z) \mid \mathbf{x}_i) = \mathbb{E}(Y_i \mid \mathbf{x}_i, Z_i = z),$$

so the conditional average treatment effect (CATE) is

$$\begin{aligned} \tau(\mathbf{x}_i) &:= \mathbb{E}(Y_i(1) - Y_i(0) \mid \mathbf{x}_i) \\ &= \mathbb{E}(Y_i \mid \mathbf{x}_i, Z_i = 1) - \mathbb{E}(Y_i \mid \mathbf{x}_i, Z_i = 0). \end{aligned}$$

Modeling assumptions

We write

$$E(Y_i | \mathbf{x}_i, z_i) = f(\mathbf{x}_i, z_i),$$

so that

$$\tau(\mathbf{x}_i) := f(\mathbf{x}_i, 1) - f(\mathbf{x}_i, 0).$$

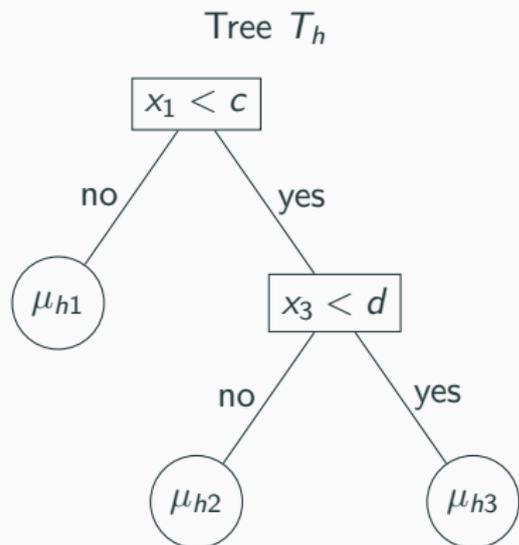
We assume iid Gaussian errors:

$$Y_i = f(\mathbf{x}_i, z_i) + \epsilon_i, \quad \epsilon_i \sim N(0, \sigma^2)$$

nb: Strong ignorability means $\epsilon_i \perp\!\!\!\perp Z_i | \mathbf{x}_i$.

How do we regularize estimates of f ? (What prior on f ?)

Regression Trees

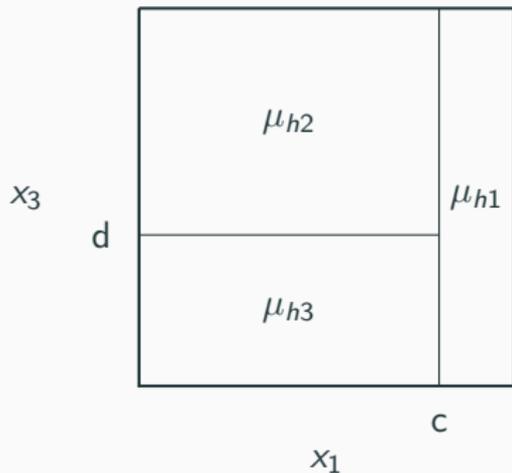


Leaf/End node parameters

$$M_h = (\mu_{h1}, \mu_{h2}, \mu_{h3})$$

$$g(\mathbf{x}, T_h, M_h) = \mu_{ht} \text{ if } \mathbf{x} \in \mathcal{A}_{ht} \text{ (for } 1 \leq t \leq b_h).$$

$g(\mathbf{x}, T_h, M_h)$



Partition $\mathcal{A}_h = \{\mathcal{A}_{h1}, \mathcal{A}_{h2}, \mathcal{A}_{h3}\}$

Bayesian Additive Regression Trees (BART)

Bayesian additive regression trees (Chipman, George, & McCulloch, 2008):

$$y_i = f(\mathbf{x}_i, z_i) + \epsilon_i, \quad \epsilon_i \sim N(0, \sigma^2)$$

$$f(\mathbf{x}, z) = \sum_{h=1}^m g(\mathbf{x}, z, T_h, M_h)$$

Hill (2011) proposes adopting Bayesian additive regression trees (BART) for causal inference.

Making BART great for causal inference

BART is great as a prior over regression functions! It also works great for causal inference, but in some settings it can be problematic:

1. With strong confounding, estimates of individual/average treatment effects from BART can exhibit severe bias.
2. With homogenous effects/moderate heterogeneity, BART's treatment effect estimates are highly variable.

We can fix both of these!

Problem 1: Strong confounding can lead to high bias

Suppose that:

- Y : measure of heart distress,
- Z : took heart medication,
- x_1 and x_2 are blood pressure measurements.

Let's make this easy: $p = 2$, $n = 1,000$, with homogeneous treatment effects ($\tau = 1$).

Problem 1: Strong confounding can lead to high bias

Assume the true model:

$$Y_i = \mu(\mathbf{x}_i) - Z_i + \epsilon_i,$$

$$\mu(\mathbf{x}_i) = 1 \text{ if } x_{i1} < x_{i2}, \quad -1 \text{ otherwise.}$$

$$\Pr(Z_i = 1 \mid x_{i1}, x_{i2}) = \Phi(\mu(\mathbf{x}_i)),$$

$$\epsilon_i \stackrel{iid}{\sim} N(0, 0.7^2), \quad x_{i1}, x_{i2} \stackrel{iid}{\sim} N(0, 1).$$

This example demonstrates targeted selection into treatment:

Patients with $x_{i1} < x_{i2}$ are 5 times as likely to receive the new drug precisely because they are more likely to have higher levels of heart distress.

Despite low noise, low dimension, and homogeneous effects, BART has problems...

BART is badly biased here

Across 250 simulated datasets with $n = 1000$, BART is badly biased:

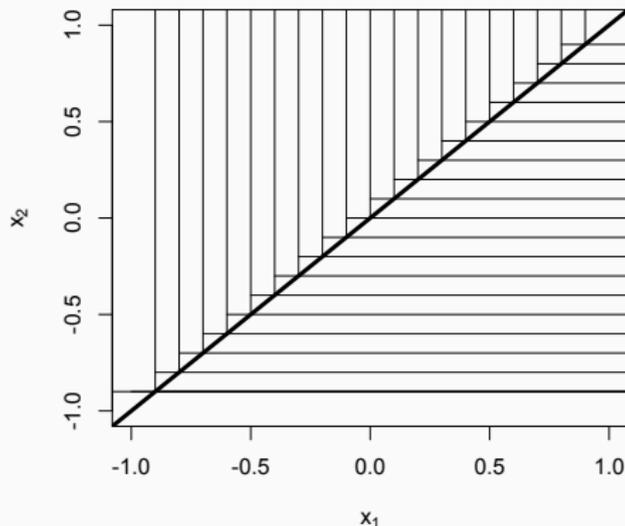
Prior	Bias	Coverage	RMSE
BART	0.14	31%	0.15

This is due to a phenomenon called **regularization induced confounding** (see Hahn, Carvalho, Puelz and He, 2018).

Targeted selection induces regularization induced confounding

Why is BART biased in this example?

- $\pi(\mathbf{x}) = \Pr(Z = 1 \mid \mathbf{x})$ is a noisy function of $\mu(\mathbf{x})$, so $\mu(\mathbf{x})$ “looks like” $\pi(\mathbf{x})$, and $\mu(\mathbf{x})$ is hard to approximate with trees



Targeted selection induces regularization induced confounding

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- Strong confounding means $Z \approx \pi(\mathbf{x})$, and targeted selection means $\mu(\mathbf{x})$ is a function of $\pi(\mathbf{x})$, so $\mu(\mathbf{x})$ can be approximated by a tree that splits on Z
- The BART prior over f penalizes the total number of splits, so to fit $\mu(\mathbf{x})$ BART would rather split on Z once than x_1 and x_2 many times – confusing confounding for treatment effects: **regularization induced confounding** (Hahn et al (2016))

A fix: Propensity Score BART

We can fix this by estimating $\pi(\mathbf{x}) = \Pr(Z = 1 \mid \mathbf{x})$ (here using BART) and including $\hat{\pi}(\mathbf{x})$ as an extra predictor variable

Prior	Bias	Coverage	RMSE
BART	0.14	31%	0.15
Oracle BART	0.00	98%	0.05
ps-BART	0.06	85%	0.08

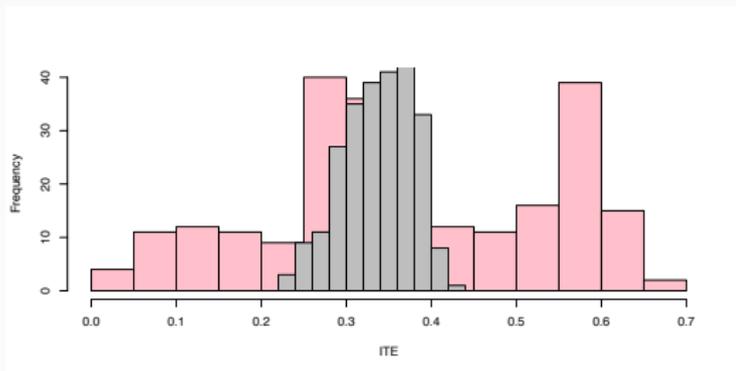
(With an ensemble estimate of $\hat{\pi}$, ps-BART \approx Oracle BART)

Problem 2: Naive priors give high variance estimates

In the model

$$y_i = f(\mathbf{x}_i, \hat{\pi}(\mathbf{x}_i), z_i) + \epsilon_i$$

a BART prior on f provides no direct mechanism to regularize the treatment effect function $\tau(\mathbf{x})$



ps-BART is in pink; our fix is in grey

The fix: Bayesian causal forests

Reparameterize!

$$f(\mathbf{x}_i, z_i) = \mu(\mathbf{x}_i, \hat{\pi}(\mathbf{x}_i)) + \tau(\mathbf{x}_i)z_i,$$

where m and τ have independent BART priors.

Now the treatment effect is

$$\tau(\mathbf{x}_i)$$

and we can “shrink towards homogeneity” with stronger regularization on τ , independent of regularization on m

Tweaking priors in BCF

Several adjustments to the BART prior on τ :

- Higher probability on smaller τ trees (than BART defaults)
- Higher probability on “stumps” (all stumps = homogeneous effects)
- $N^+(0, \nu)$ Hyperprior on the scale of leaf parameters in τ

RIC in the wild: 2017 ACIC Data Analysis Challenge

Treatment-response pairs were simulated according to 32 distinct data generating processes (DGPs), given fixed covariates ($n = 4,302$, $p = 58$) from an empirical study.

We varied three parameters among two levels

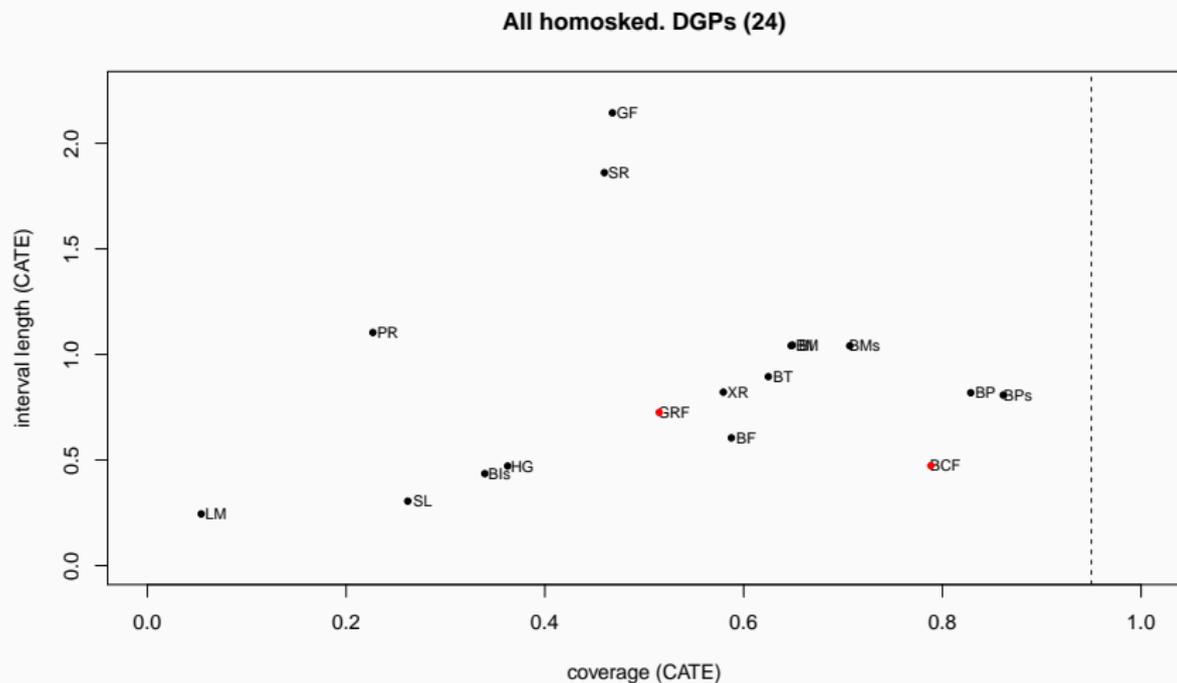
- **High** or **Low** *noise level*,
- **Strong** or **Weak** *confounding*,
- **Small** or **Large** *effect size*.

The error distributions were one of four types

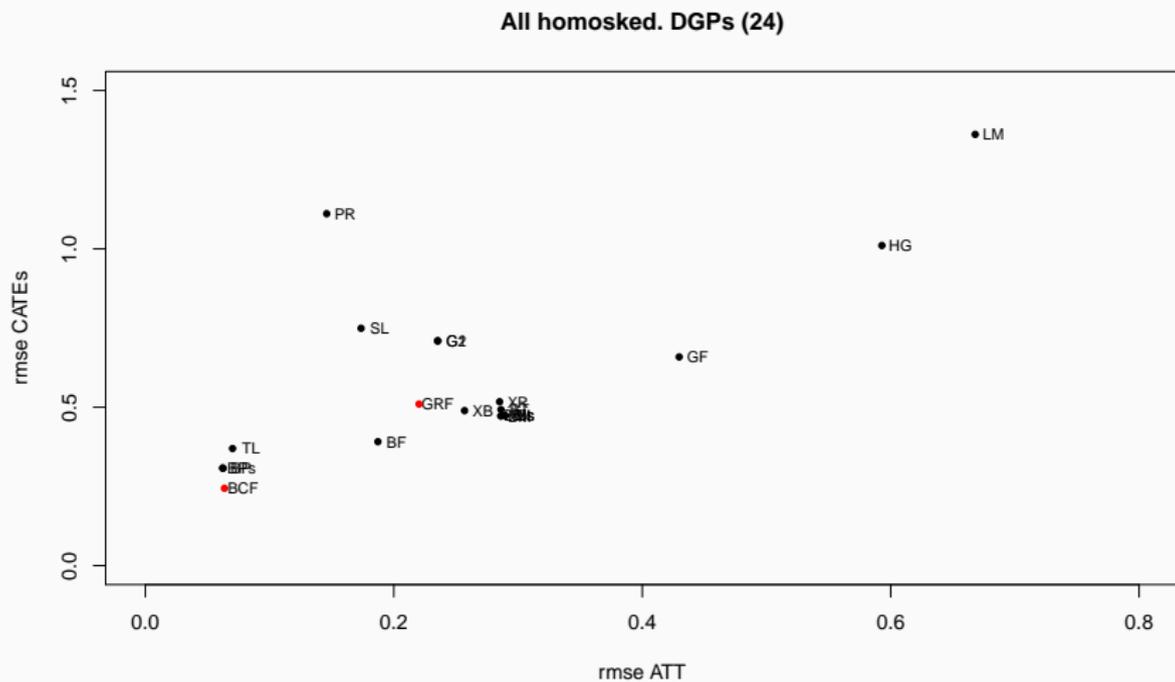
- Additive, homoskedastic, independent,
- Nonadditive, homoskedastic, independent,
- Additive, heteroskedastic, independent.

To assess coverage, 250 replicate data sets were generated for each DGP.

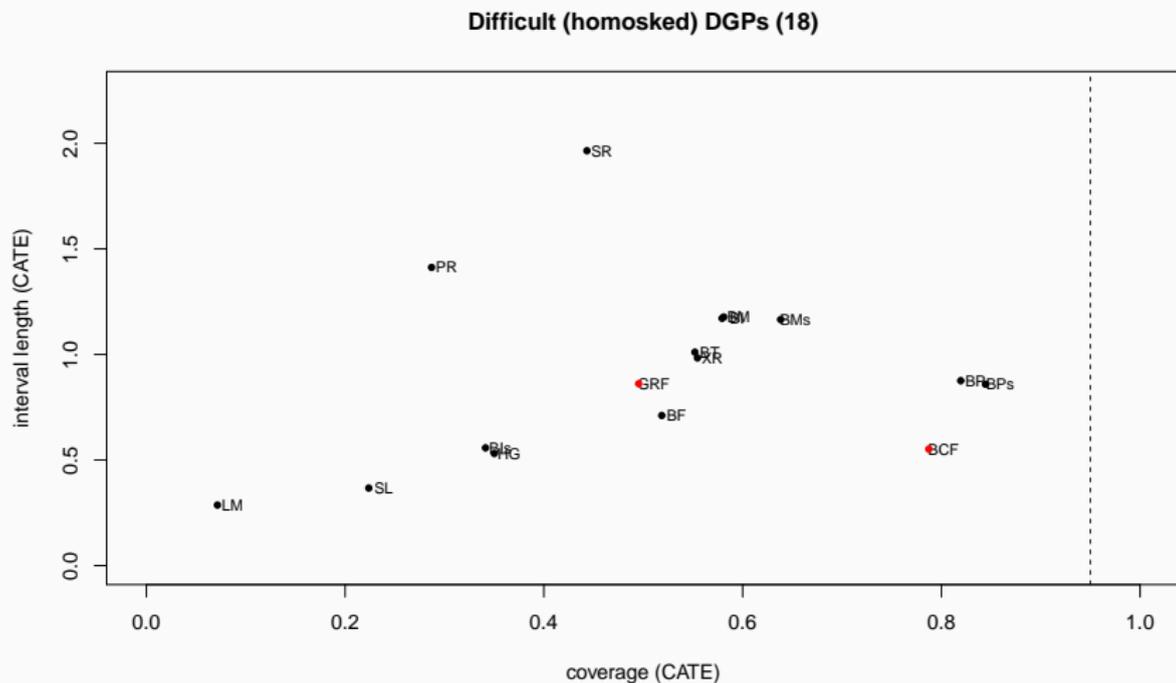
Results: Inference for CATE on homoskedastic DGPs



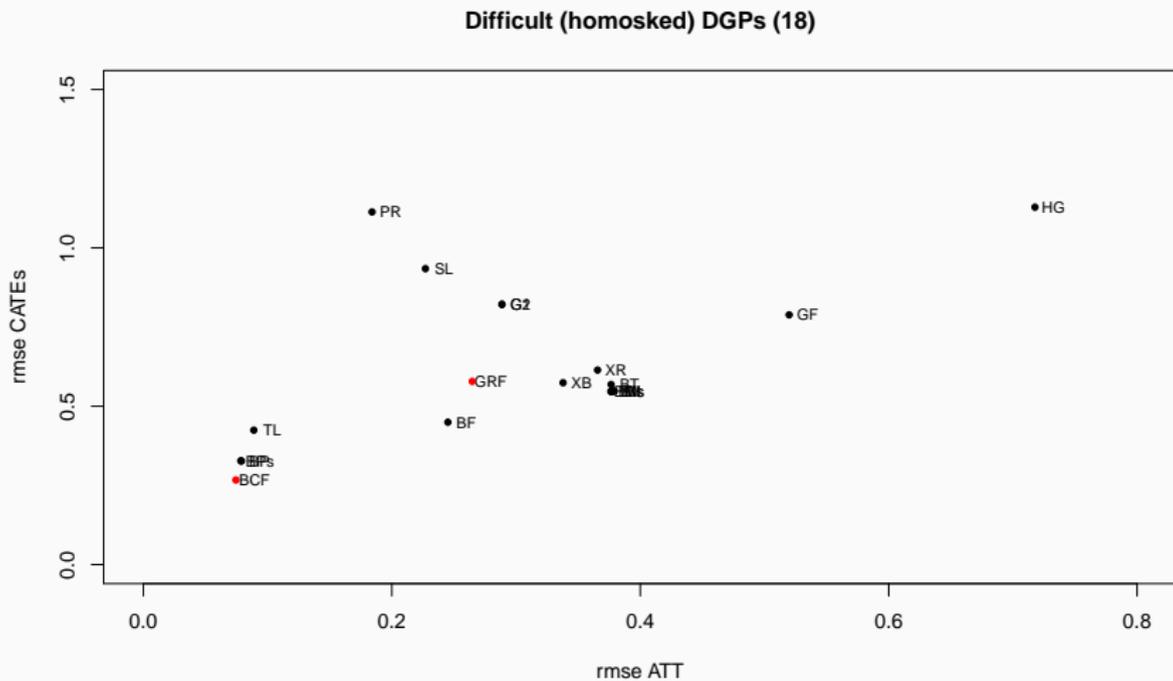
Results: Estimation for homoskedastic DGPs



Results: Inference for CATE on difficult homoskedastic DGPs

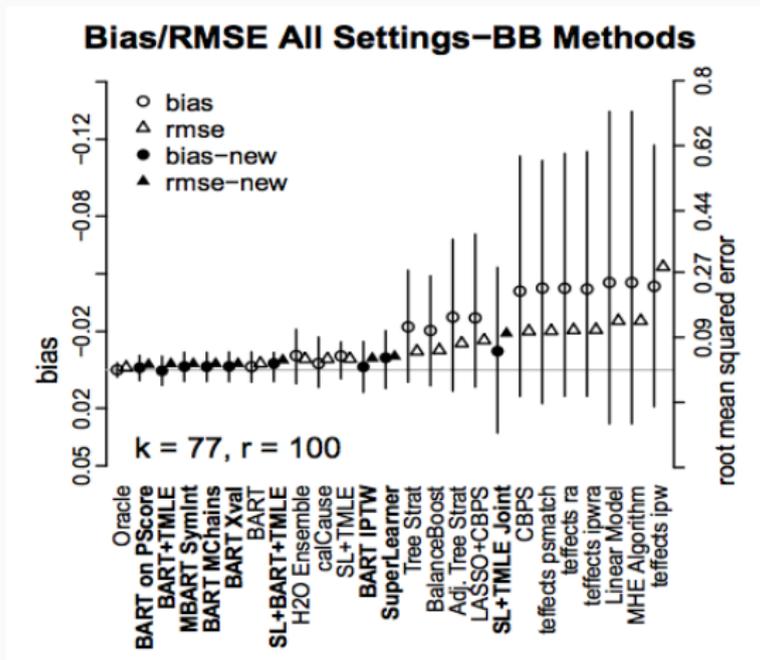


Results: Estimation on difficult homoskedastic DGPs



ACIC 2016 Redux

After our a preprint of our paper the ACIC 2016 challenge organizers ran ps-BART...



Large, highly variable treatment effects and no explicit targeted selection!

Adding BCF and causal RF:

	Cov	IL	Bias	(SD)	Bias	(SD)	PEHE	(SD)
BCF	0.82	0.026	-0.0009	(0.01)	0.008	0.010	0.33	0.18
ps-BART	0.88	0.038	-0.0011	(0.01)	0.010	0.011	0.34	0.16
BART	0.81	0.040	-0.0016	(0.02)	0.012	0.013	0.36	0.19
Causal RF	0.58	0.055	-0.0155	(0.04)	0.029	0.027	0.45	0.21

Average differences relative to BCF, pairwise permutation test p-value:

	Diff Bias	p	Diff Bias	p	Diff PEHE	p
ps-BART	-0.00020	0.146	0.0011	$< 1e^{-6}$	0.010	$< 1e^{-6}$
BART	-0.00070	$< 1e^{-6}$	0.0031	$< 1e^{-6}$	0.037	$< 1e^{-6}$
Causal RF	-0.01453	$< 1e^{-6}$	0.0204	$< 1e^{-6}$	0.125	$< 1e^{-6}$

Takeaways

In observational data regularization-induced confounding can adversely bias treatment effect estimates **from any method that uses regularization**. Explicitly modeling selection is necessary for robust inference.

BART is an impressive response surface method for causal inference; our new BCF models improve on “vanilla” BART in key respects:

- Propensity score estimates as covariates mitigate RIC
- Reparameterization allows regularization to be imposed robustly and directly on the estimand of interest.
- **It also facilitates extensions to multilevel models!**

Multilevel BCF: The National Study of Learning Mindsets

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National Study of Learning Mindsets

- National Study of Learning Mindsets (Yeager et. al., 2017):
Randomized controlled trial of a low-cost mindset intervention
- Probability sample of 76 schools (\approx 14,000 students)
- Specifically designed to assess treatment effect heterogeneity
- Preregistration plan included specific subgroups of interest, **and a blinded exploratory analysis of heterogeneity**

Desiderata for our analysis:

- Avoid model selection/specification search
- School-level effect heterogeneity explained and unexplained by covariates
- Interpretable model summaries for communicating results

Multilevel Linear Models for Heterogeneous Treatment Effects

$$y_{ij} = \alpha_j + \sum_{h=1}^p \beta_h x_{ijh} + \left[\sum_{\ell=1}^k \tau_{\ell} w_{ij\ell} + \gamma_j \right] z_{ij} + \epsilon_{ij}$$

School-specific intercepts/fixed/random effects

Controls at the student and/or school level

Moderators at the student and/or school level

School-specific "unexplained" heterogeneity

Coloring outside the lines: Multilevel Bayesian Causal Forests

We replace linear terms with Bayesian additive regression trees (BART)



$$y_{ij} = \alpha_j + \beta(\mathbf{x}_{ij}) + [\tau(\mathbf{w}_{ij}) + \gamma_j] z_{ij} + \epsilon_{ij}$$

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BART in causal inference: Hill (2011), Green & Kern (2012), ...

Parameterizing treatment effect heterogeneity with BART is due to Hahn, Murray and Carvalho (2017)

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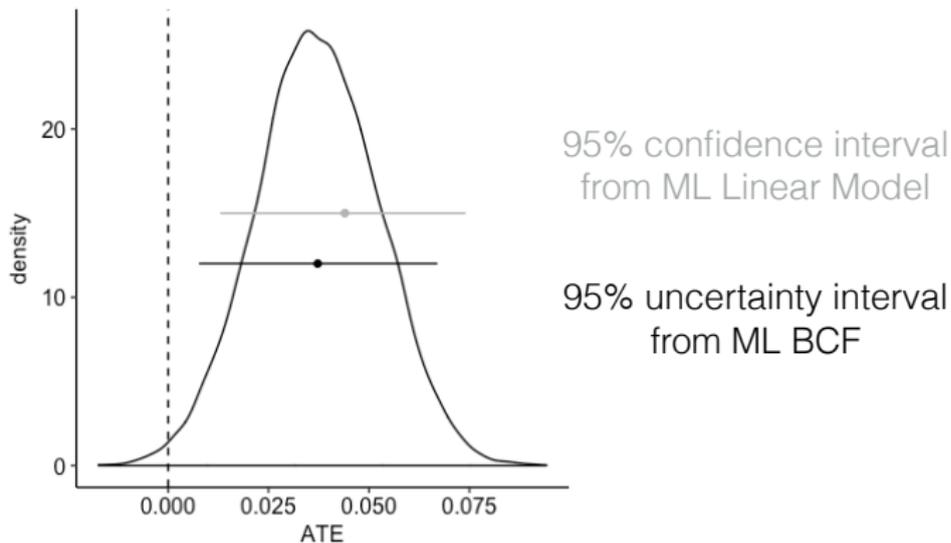
Allows for complicated functional forms (nonlinearity, interactions, etc) without pre-specification...

...while carefully regularizing estimates with prior distributions (shrinkage toward additive structure and discouraging implausibly large treatment effects)

Analyzing data with ML BCF

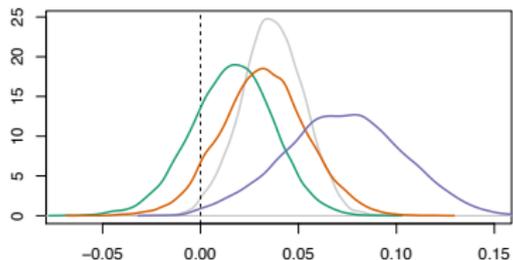
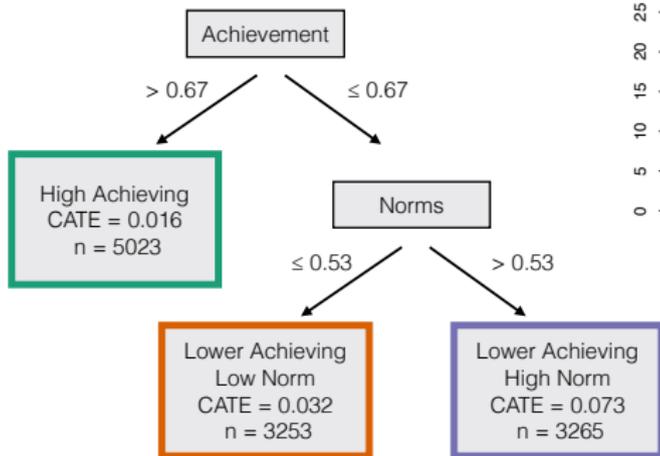
- Obtain posterior samples for all the parameters, compute treatment effect estimates for each unit/school/etc.
- The challenge: How do we summarize these complicated objects?
 - “Roll up” treatment effect estimates to ATE
 - Subgroup search
 - Counterfactual treatment effect predictions/“partial effects of moderators”

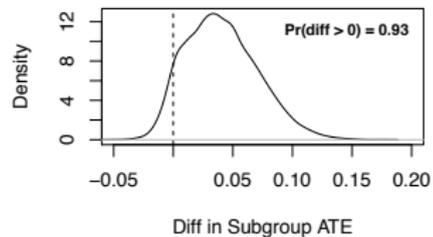
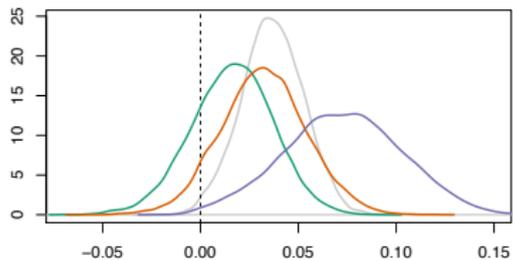
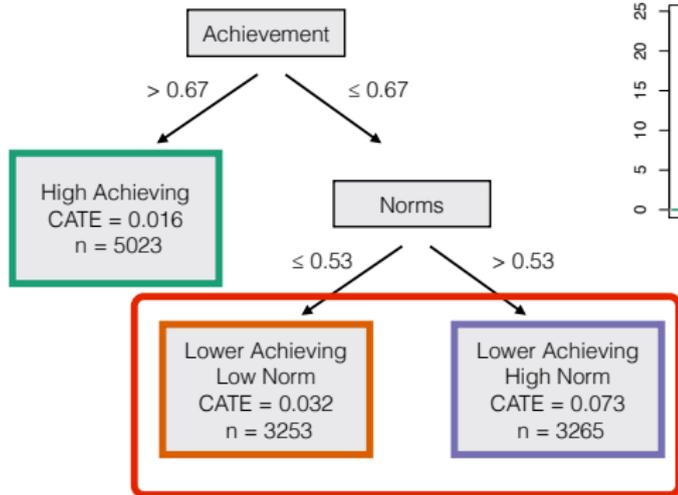
Inference for the Average Treatment Effect

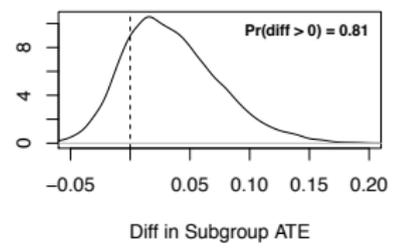
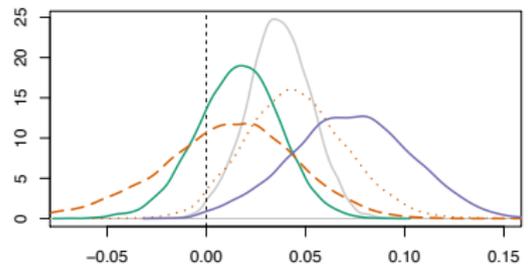
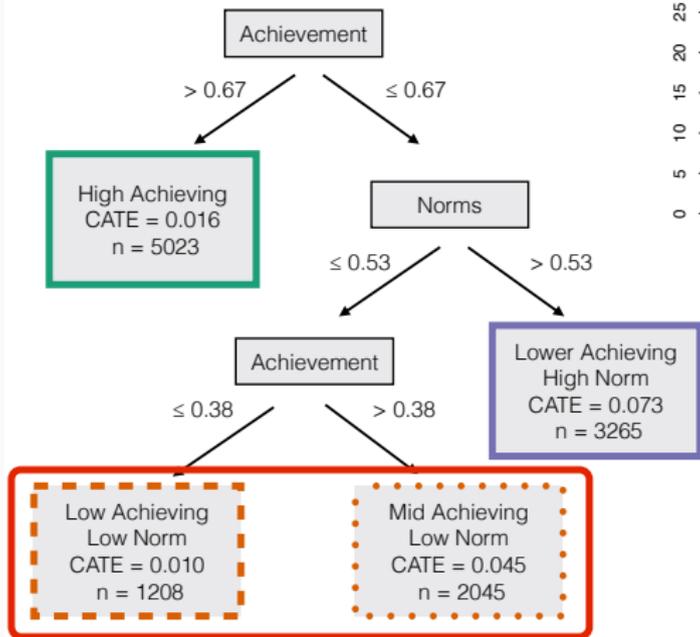


Subgroup search

- Obtain posterior mean of treatment effects
- Use recursive partitioning (CART) **on the posterior mean** to find moderator-determined subgroups with high variation across subgroup ATE
 - Statistically kosher! We use the data once (prior -> posterior)
 - Can be formalized as the Bayes estimate under a particular loss function

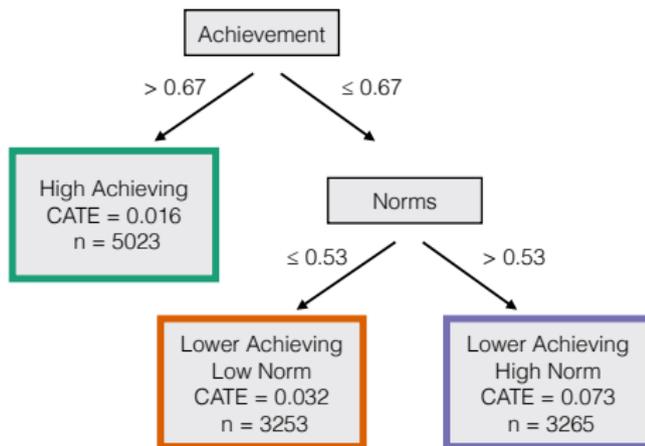






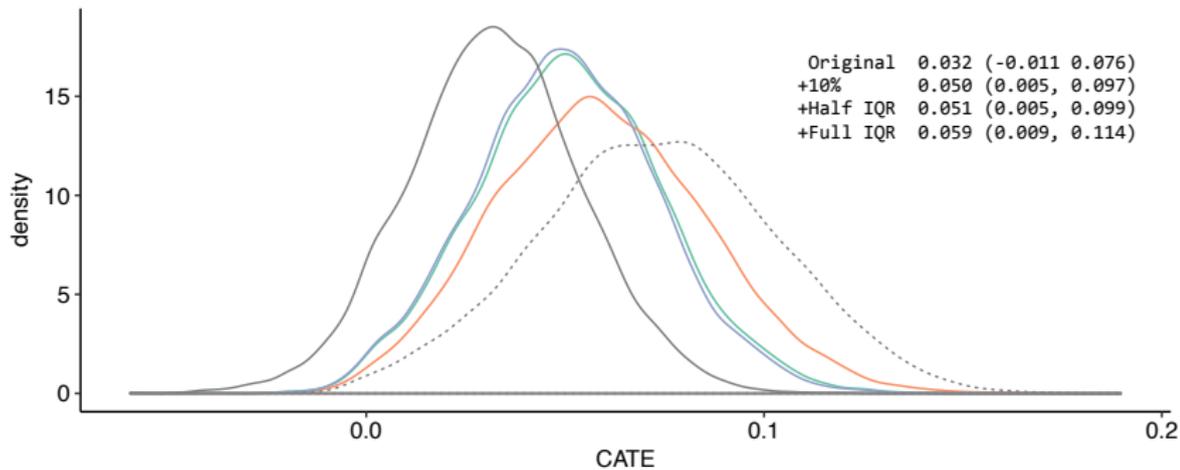
Counterfactual treatment effect predictions

- How do estimated treatment effects change in lower achieving/low norm schools if norms increase, holding constant school minority comp & achievement?
- Not a formal causal mediation analysis (roughly, we would need “no unmeasured moderators correlated with norms”)



1 IQR = 0.6 extra problems
on worksheet task

Increase +0.5 IQR +1 IQR +10% Orig Group Low Norm/Lower Ach High Norm/Lower Ach



Conclusion

- Flexible models + careful regularization + posterior summarization is a winning combination
- Our approach takes the best parts of linear models with lots of researcher degrees of freedom and “black box” machine learning methods that only afford bankshot regularization and summarization
 - Many “degrees of freedom” in the summarization step, but these depend on the data only through the posterior
 - Unlike many ML methods, we can handle multilevel structure and prior knowledge with ease

Thank you!