Heterogeneous Treatment Effects of Medicaid and Efficient Policies

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June 4, 2020

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• Setting eligibility criteria = segmentation and targeting the population.

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- Treatment variable: Random assignment of Medicaid.
- **Control variables**: Federal Poverty Level, Age, Household size, Insurance status of last 6-months, Education status, Employment status.
- **Drop** the race, gender and residency for non-discriminatory policy/waiver purpose.

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 - **OHP Standard**: for non-categorically eligible Medicaid population.

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- DHS got approved to lottery/randomize OHP standard from CMS.
- Data comprises 74,922 individuals (representing 66,385 households).
- At baseline, data on lottery (*W*) and various demographics (*x*) were collected, then after a year, data on outcome variable (*Y*) were collected.

Heterogeneous Treatment Effects: Why and How?

Why Heterogeneous Treatment Effects?

- Mixed effects of Medicaid \Longrightarrow gap in literature
- Systematically identify subpopulations and estimate treatment effects, valid inference.
- HTEs are mandatory step to understand mechanisms and identify efficient policy.

Causal ML: Causal Tree (Athey and Imbens, 2016)

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- Goal: Minimize MSE of treatment effects on each leaf i.e.

$$-E_{\mathcal{S}^{tr}}\left[\sum_{i\in\mathcal{S}^{tr}}\left(\tau_{i}-\hat{\tau}(X_{i})\right)^{2}\right]$$

- Cross-validate to prune the tree-depths
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- Sample splitting and Honest estimation.


Causal ML: Causal Forest = Many Causal Trees

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Out-of-Sample Prediction

NOW LETS TRY TO CLASSIFY THIS



Age = 30 HH income = \$12000 HH size = 5 Education = 12

Out-of-Sample Prediction



Out-of-Sample Prediction



$$\begin{split} \hat{\Gamma}_{i} &= \hat{\tau}_{-i}(X_{i}) + \frac{1\{W_{i} = \pi(X_{i})\}}{\hat{e}_{-i}(X_{i}, W_{i})} \cdot (Y_{i} - \hat{\mu}_{-i}(X_{i}, W_{i})) \\ \text{Standard errors are Cluster-robust at household level.} \\ \text{Establish asymptotic normality:} \quad \frac{\hat{\tau}(x) - \tau(x)}{\sigma(x)} \Rightarrow N(0, 1). \end{split}$$

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Results: Part 1 of 3

 Table 2: Health Care Utilization

Outcome variables	ITT (1)	LATE (2)	ATE (3)	Heuristic (4)	MFP (5)	DFP (6)
Panel A: Health care utilization Extensive margins						
Currently taking any prescription medications	(0.021^{**})	0.067^{**}	0.007	-0.018	0.801	-0.494
	(0.009)	(0.03)	(0.009)	(0.018)	(1.015)	(0.734)
Outpatient visits last six months	0.07^{***}	0.224^{***}	0.062^{***}	0.055^{***}	1.028^{***}	1.316^{***}
	(0.009)	(0.027)	(0.009)	(0.017)	(0.145)	(0.312)
ER visits last six months	0.009	0.029	0.005	-0.014	0.696	-3.331
	(0.008)	(0.024)	(0.008)	(0.015)	(1.172)	(1.816)
Inpatient hospital admissions last six months	$0.002 \\ (0.004)$	0.005 (0.014)	0.001 (0.005)	-0.006 (0.009)	0.272 (2.322)	-0.626 (1.4)
Intensive margins	0.104^{*}	0.342*	0.042	-0.119	0.899	-0.383
Number of prescription medications currently taking	(0.055)	(0.177)	(0.055)	(0.109)	(1.219)	(1.005)
Number of Outpatient visits last six months	0.335^{***}	1.087^{***}	0.304^{***}	0.426^{***}	1.037^{***}	1.502^{***}
	(0.052)	(0.166)	(0.055)	(0.11)	(0.188)	(0.373)
Number of ER visits last six months	$0.006 \\ (0.016)$	0.018 (0.053)	-0.003 (0.017)	-0.115^{***} (0.035)	1.97 (14.846)	-10.89 (2.98)
Number Inpatient hospital admissions last six months	0.007	0.024	0.007	0.008	0.713	-2.071
	(0.007)	(0.021)	(0.007)	(0.014)	(0.661)	(1.974)

Health Care Utilization

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 $\mathsf{Outpatient}_{i,h} = \beta_0 + \beta_1 \mathsf{Lottery}_{i,h} + x_{ih}\beta_2 + \varepsilon_{it}$



Control group

Health Care Utilization

Outcome variables	ITT	LATE	ATE
	(1)	(2)	(3)
Outpatient visits last six months	0.07^{***} (0.009)	0.224^{***} (0.027)	$\begin{array}{c} 0.062^{***} \\ (0.009) \end{array}$

$$\begin{aligned} & \textit{Medicaid}_{i,h} = \delta_0 + \delta_1 \textit{Lottery}_{i,h} + x_{ih} \delta_2 + \mu_{it} \\ & \textit{Outpatient}_{i,h} = \phi_0 + \phi_1 \widehat{\textit{Medicaid}}_{i,h} + x_{ih} \phi_2 + \nu_{it} \end{aligned}$$



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A Average Treatment Effect Assessed in a Heterogeneous Population



Tests for Heterogeneity.

Heterogeneity: Graphical Intuition



Heterogeneity: Graphical Intuition



Results: Part 2 of 3

Accessing Treatment Heterogeneity

Outcome variables	ITT	LATE	ATE	Heuristic	MFP	DFP
	(1)	(2)	(3)	(4)	(5)	(6)
Outpatient visits last six months	$\begin{array}{c} 0.07^{***} \\ (0.009) \end{array}$	$\begin{array}{c} 0.224^{***} \\ (0.027) \end{array}$	0.062^{***} (0.009)	0.055^{***} (0.017)	$\begin{array}{c} 1.028^{***} \\ (0.145) \end{array}$	$\begin{array}{c} 1.316^{***} \\ (0.312) \end{array}$

Table 2: Health Care Utilization

Notes: The ***, **, and * represent 1%, 5%, and 10% level of significance, respectively. Enclosed in the parenthesis are household-level clustered heteroscedasticity-consistent standard errors. The regressions in Columns (1) and (2) include household size dummies, survey wave dummies, and survey wave interacted with household size dummies. For the LATE estimates in Column (2), the instrumental variable is lottery assignment, and the endogenous variable is "Ever in Medicaid". The ITT and LATE estimates are base on the double-selection post-LASSO.

- Measure of Treatment Heterogeneity: Heuristic*, MFP = 1, and DFP > 0.
- Once we know there is treatment heterogeneity, we would like to know efficient policy that allocates scares resources.

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• We would like to minimize the regret.

• Estimate regret by Q-learning:

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- The regret converges as:

$$\sqrt{n}\left(\hat{R}_{DML}\left(\pi\right)-R\left(\pi\right)\right)\overset{d}{\rightarrow}N\left(0,\sigma^{2}\left(\pi\right)\right)$$

Double Machine Learning VC Dimension

Results: Part 3 of 3

Estimate of the utility improvement of various policies over a random assignment baseline.

Variable	Baseline (1)	Probability rule (2)	CATE rule (3)	Shallow tree (4)	Deeper tree (5)
Panel A: Health care utilization Outpatient visits last six months	0.604^{***} (0.002)	4.74^{***} (0.182)	5.119^{***} (0.17)	4.228^{***} (0.197)	2.898^{***} (0.177)

• **Baseline**: On average 60% of population have outpatient visit in last 6 month.

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Estimate of the utility improvement of various policies over a random assignment baseline.

Probability rule: If the propensity is below average, assign the treatment.

$$\hat{\Gamma}_i = \hat{e}_i^{(-i)}(X_i, W_i)$$

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CATE rule: If the CATE is non-zero and positive, assign the treatment.

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Estimate of the utility improvement of various policies over a random assignment baseline.

Shallow tree: treatment assignment rule base on a simpler tree (2-Level depths).

Deeper tree: treatment assignment rule base on complex or more in-depth tree (up to 6-Level depths).

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Efficient policy to improve outpatient visits



Matching story and policy. VC Dimension

Discussion and Conclusion
Policy evaluation.

- Positive economics
- "What is" the impact of treatment on outcome?
- Random assignment of treatment.
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- False Negative: Not assigning the Medicaid to those who need it.
- Does this method work in an observational setting or panel or IV?

The End Q&A Email: ss0088@mix.wvu.edu

Rubin's Potential Outcome Framework 🔤

We observe a sequence of triples $\{(W_i, Y_i, X_i)\}_i^N$ for N individuals, where:

- W_i represents if individual *i* is treated with lottery insurance or not.
- Y_i is the outcome variable.
- In Rubin (1974) potential outcomes framework,
 - ► Y_i (1) represents potential outcome of subject i if he had received the treatment and
 - Y_i (0) represents potential outcome of subject *i* if he had not had received the treatment
- X_i is vector of observable characteristics.
- Individual treatment effect for i^{th} is: $Y_{i}(1) Y_{i}(0)$
- The average treatment effect (ATE) is: $\tau = E\left[Y_{i}\left(1\right) Y_{i}\left(0\right)\right]$
- Unfortunately, in our data we of course can only observe one of these two potential outcomes, so actually computing this difference for everyone is impossible. This is **fundamental problem of causal inference**.

Rubin's Potential Outcome Framework Duck

The Naive estimator of average treatment effect (ATE) can be expressed as:

$$au = E[au] = E[Y_i(1) - Y_i(0)]$$

With linearity of expectation:

$$\tau = E[Y_{i}(1)] - E[Y_{i}(0)]$$

With independence assumption

$$\tau = E[Y_{i}(1)|W_{i} = 1] - E[Y_{i}(0)|W_{i} = 1]$$

Which is just the simple difference of means:

$$\hat{ au} = rac{1}{n_1}\sum_{i|W_i=1}^{n_1} y_i - rac{1}{n_0}\sum_{i|W_i=0}^{n_0} y_i$$

item This is feasible only if there is no selection and heterogeneity biases.

Rubin's Potential Outcome Framework 🔤

When, λ portion of population is exposed to treatment, the estimator of average treatment effect (ATE) can be expressed as:

$$\underbrace{E[\tau]}_{e} = \lambda \left\{ \underbrace{E[Y_{i}(1) | W_{i} = 1]}_{a} - \underbrace{E[Y_{i}(0) | W_{i} = 1]}_{c} \right\}$$
$$- (1 - \lambda) \left\{ \underbrace{E[Y_{i}(0) | W_{i} = 0]}_{d} - \underbrace{E[Y_{i}(1) | W_{i} = 0]}_{b} \right\}$$

where,

- *a* and *d* can be observed in from the data.
- c is counter factual of a.
- *b* is counter factual of *d*.
- λ is percentage of treated population.
- Odom = (a d) is Observed difference of mean.
- (a c) is Average treatment effect on treated (ATT).
- (b d) is Average treatment effect on untreated (ATU).

Rubin's Potential Outcome Framework The ATE can be expressed as:

$$e = \lambda \left(a - c
ight) - \left(1 - \lambda
ight) \left(d - b
ight)$$

The solution can be expressed as:

$$\underbrace{(a-d)}_{Odom} = \underbrace{e}_{ATE} + \underbrace{(c-d)}_{selection \ bias} + \underbrace{(1-\lambda)\left\{(a-c) - (b-d)\right\}}_{heterogeneous \ treatment \ effect \ bias}$$

- Under assumption of unconfoundedness¹, selection bias and heterogeneous treatment effect bias nullify and causal ATE is simple difference of mean. Technically: Y_i(1), Y_i(0) ⊥ W_i|X_i.
- The overlap assumption guarantees that no sub-population is entirely located in only one of control or treatment groups. Technically: $\forall x \in \text{supp}(X), \ 0 < P(W = 1 | X = x) < 1.$

¹Unconfoundedness implies treatment is randomly assigned, and knowing observable characteristics of individual i, and then treatment status gives no information on the potential outcomes.

Pre-treatment Mean Comparison **Deck**

Female

English preferred

Signed up on first day

PO Box address

MSA

Race as White

Race as Black

Race as Spanish/Hispanic/Latino

4-year college degree or more

High school diploma or GED

Less than high school

Vocational training or 2-year degree

Don't currently work

Work below 20 hours/week

Work 20-29 hours/week

Work 30+ hrs/week



Double Selection Post-LASSO (back)

- LASSO simultaneously performs model selection and coefficient estimation by minimizing the sum of squared residuals plus a penalty term.
- Consider a following linear model

$$\tilde{y}_i = \Theta_i \beta_1 + \varepsilon_i$$

 where Θ is high-dimensional covariates, the LASSO estimator is defined as the solution to:

$$\min_{\beta_1 \in \mathbb{R}^p} E_n \left[\left(\tilde{y}_i - {}_i \Theta \beta_1 \right)^2 \right] + \frac{\lambda}{n} \|\beta_1\|_1$$

- The penalty level λ is a tuning parameter to regularize/controls the degree of penalization and to guard against over-fitting. The penalty term penalizes the size of the model through the sum of absolute values of coefficients.
- The cross-validation technique chooses the best λ in prediction models and $\|\beta\|_1 = \sum_{j=1}^p |\beta_j|$.
- The kinked nature of penalty function induces $\hat{\beta}$ to have many zeros; thus LASSO solution feasible for model selection.

Double Selection Post-LASSO (back)

• The double-post-LASSO procedure comprises the following steps:

- **First**, run LASSO of dependent variables on a large list of potential covariates to select a set of predictors for the dependent variable.
- Second, run LASSO of treatment variable on a large list of potential covariates to select a set of predictors for treatment².
- ▶ Third, run OLS regression of dependent variable on treatment variable, and the union of the sets of regressors selected in the two LASSO runs to estimate the effect of treatment on the dependent variable then correct the inference with usual heteroscedasticity robust OLS standard error.

 $^{^2\}mbox{If}$ the treatment is truly exogenous, I should expect this second step should not select any variables.

Why Causal-ML to estimate heterogeneous effects? Deck

• Problem of the multiple hypothesis testing³.



³The "multiple hypothesis testing problems" leads to the so-called "ex-post selection problem," which is widely recognized in the program evaluation literature. For example, for fifty single hypotheses tests, the probability that at least one test falsely rejects the null hypotheses at the 5% significance level (assuming independent test statistics as an extreme case) is $1 - 0.95^{50} = 0.92$ or 92%.

Why Causal-ML to Estimate Heterogeneous Effects? **Dec**

- Performing ad-hoc searches or p-hacking⁴ to detect the responsive subgroups may lead to false discoveries or may mistake noise for an actual treatment effect.
- To avoid many of the issues associated with data mining or p-hacking, researchers can commit in advance to study only a subgroup by a preregistered analysis plan⁵. However, this may also prevent discovering unanticipated results and developing new hypotheses.
- Using ML algorithms like CARTs, R.F., and N.N, etc. seems practical, but these algorithms are designed for prediction and not for causal inference.

⁴The *p*-hacking is an exhaustive search for statistically significant relations from combinations of variables or combinations of interactions of variables or subgroups. The *p*-hacking could lead to discovering the statistically significant relationship, when, in fact, there could have no real underlying effect.

⁵A preregistered analysis plan is sets of analyses plans released in the public domain by the researchers in advance prior they collect the data and learn about outcomes.

Athey and Imben Causal Tree Lack

Consider a tree or partitioning \prod to a partition of feature space X, with $\#(\prod)$ the numbers of elements in the partition given as:

$$\prod = \big\{\ell_1, \dots, \ell_{\#(\tau)}\big\}$$

Then given a partition \prod , for each observations (Y_i^{obs}, X_i, W_i) , the population average condition mean function $\mu(x; \prod)$ is given as:

$$\mu\left(w,x;\prod\right) \equiv \mathbb{E}\left[Y_{i}\left(w\right)|X_{i} \in \ell\left(x;\prod\right)\right]$$

And its average causal effect is given as:

$$au\left(x;\prod
ight)\equiv\mathbb{E}\left[Y_{i}\left(1
ight)-Y_{i}\left(0
ight)|X_{i}\in\ell\left(x;\prod
ight)
ight]$$

Then the goal is to construct $\pi(.)$ that maximizes the following honest criterion:

$$Q^{H}(\pi) = -\mathbb{E}_{S^{te}, S^{est}}\left[MSE_{\tau}\left(S^{te}, S^{est}, \pi\left(S^{tr}\right)\right)\right]$$

Athey and Imben Causal Tree Lack

The above equation can be rearranged as:

$$-EMSE_{\tau}\left(S^{tr}, N^{est}, \prod\right) = \alpha \frac{1}{N^{tr}} \sum_{i \in S^{tr}} \left(X_i; S^{tr}, \prod, \alpha\right)$$

$$-(1-\alpha)\left(\frac{1}{N^{tr}}+\frac{1}{N^{est}}\right)\cdot\sum_{\ell\in\prod}\left(\frac{s_{\mathcal{S}_{treat}}^{2}\left(\ell\right)}{p}+\frac{s_{\mathcal{S}_{control}}^{2}\left(\ell\right)}{1-p}\right)$$

Which provides a causal tree, where $s_{S_{treat}}^2(\ell)$ is the within-leaf variance on outcomes Y for $S_{control}^{tr}$ in leaf ℓ ; $s_{S_{control}}^2(\ell)$ is the counterpart for S_{treat}^{tr} ; $p = N_{treat}/N$ is the treatment probability, $\alpha \in (0, 1)$ and is a parameter to adjust the portion of *MSE* and the variance of *EMSE*.

Sample Splitting, Honest Estimation and Cross-fitting **Geodese**

• Sample splitting and Honest estimation.



Sample Splitting, Honest Estimation and Cross-fitting **Game**

• Sample splitting and Honest estimation.



Cross-fitting



Bootstrapping and Aggregating (Boosting) **Gack**

Aggregation of Information

Then probability of x out of n choosing correctly follows a bi-nominal distribution given as: $P_n = \sum_{x=\frac{n+1}{2}}^n \binom{n}{x} p^x (1-p)^{n-x}$

Extremely large *n* makes correct choice

 $\lim_{n\to\infty}P_n\to 1.$

This means that the limit results that group competence approaches one as group size approaches infinity. In other words, extremely large committees almost certainty make the correct choice.

Causal Random Forest 🔤

• Random Forest approach makes prediction from an average of *b* CARTs or trees, as follow:

(1) for each tree $b=1,\ldots,B$, draw a subsample $S_b\subseteq\{1,\ldots,n\}$

(2) grow a tree via recursive partitioning on each such subsample of the data; and

(3) make a prediction by averaging the prediction made by individual tree as:

$$\hat{u}(x) = \frac{1}{B} \sum_{b=1}^{B} \sum_{n=1}^{n} \frac{Y_{i} \mathbf{1} \left(\{ X_{i} \in L_{b}(x), i \in S_{b} \} \right)}{|\{i : X_{i} \in L_{b}(x), i \in S_{b} \}|}$$

- where, L_b(x) denotes the leaf of the bth tree containing the training sample x.
- For out-of-bag prediction, one can estimate the average as µ̂⁽⁻ⁱ⁾(x) by only considering those trees b for which i ∉ S_b. (-i) superscript denote "out-of-bag" or "out-of-fold" prediction.

R-Learner Objective Function **back**

• "*R*-learner" objective function for heterogeneous treatment effect estimation as:

$$\hat{\tau}(\cdot) = \arg\min_{\tau} \left\{ \sum_{i=1}^{n} \left(\left(Y_i - \hat{m}^{(-i)}(X_i) \right) - \tau(X_i) \left(W_i - \hat{e}^{(-i)}(X_i) \right) \right)^2 + \right.$$

- where, $\lambda_n(\tau(\cdot))$ is a "regularizer" that controls the complexity of the learned conditional average treatment effect $\hat{\tau}(\cdot)$ function.
- $e(x) = P[W_i | X_i = x]$ is the propensity score or probability of being treated.
- m(x) = E [Y_i|X_i = x] is expected outcomes marginalizing over treatment; (-i) superscript denote "out-of-bag" or "out-of-fold" prediction.

Causal Random Forest **Dack**

- Random Forest ensembles of many trees and provides prediction as an average prediction made by many individual trees.
- A Random Forest can be equivalent as an adaptive kernel method and re-express the random forest from equation:

$$\hat{\mu}(x) = \sum_{i=1}^{n} a_i(x) Y_i; \quad a_i(x) = \frac{1}{B} \sum_{b=1}^{B} \frac{Y_i \mathbf{1}(\{X_i \in L_b(x), i \in S_b\})}{|\{i : X_i \in L_b(x), i \in S_b\}|}$$

- where, $a_i(x)$ is a data-adaptive kernel or simply weights that measure how often the i^{th} training example appears in the same leaf as the test point x.
- The kernel-based perspective on forests suggests a natural way to use them for treatment estimation by first growing a forest to get weights $a_i(x)$, and then set

$$\hat{\tau} = \frac{\sum_{i=1}^{n} a_i(x_i) \left(Y_i - \hat{m}^{(-i)}(X_i)\right) \left(W_i - \hat{e}^{(-i)}(X_i)\right)}{\sum_{i=1}^{n} a_i(x_i) \left(W_i - \hat{e}^{(-i)}(X_i)\right)}$$

Causal Random Forest **back**

- At the implementation level, the causal forest starts by fitting two separate regression forests to estimate m̂(·) and ê(·) and making out-of-bag predictions using these two first-stage forests.
- Then the model uses these out-of-bag predictions as inputs to the causal forest where cross-validation on the "*R*-learner" objective function, chooses the tuning parameters for the causal forest.

Accessing Treatment Heterogeneity **Gack**

- Heuristic test:
 - Group observation based on CATE (above or below median CATE).
 - Test if the ATE of these two groups is different from each other or not.
 - It provides qualitative insights about the strength of heterogeneity.



Below Median CATEs

Accessing Treatment Heterogeneity **Deck**

- Mean Forest Prediction (MFP) by Chernozhukov et al. (2018)
 - Define: $B_i = Y_i \hat{y}_i^{(-i)}$
 - Define: $C_i = \overline{\tau}(W_i \hat{e}_i^{(-i)})$
 - $\bar{\tau}$ is out-of-sample ATE, and $\hat{e}_i^{(-i)}$ is propensity.
 - $\frac{dB_i}{dC_i} = 1$, for calibrated model.
- Differential Forest Prediction (DFP) by Chernozhukov et al. (2018)

•
$$D_i = (\hat{\tau}^{(-i)}(X_i) - \bar{\tau})(W_i - \hat{e}_i^{(-i)})$$

• $\frac{dB_i}{dD_i} > 0$, for existence of treatment heterogeneity.

Accessing Treatment Heterogeneity **Dec**

Outcome variables	ITT	LATE	ATE	Heuristic	MFP	DFP
	(1)	(2)	(3)	(4)	(5)	(6)
Outpatient visits last six months	$\begin{array}{c} 0.07^{***} \\ (0.009) \end{array}$	$\begin{array}{c} 0.224^{***} \\ (0.027) \end{array}$	0.062^{***} (0.009)	0.055^{***} (0.017)	1.028^{***} (0.145)	$\begin{array}{c} 1.316^{***} \\ (0.312) \end{array}$

Table 2: Health Care Utilization

Notes: The ***, **, and * represent 1%, 5%, and 10% level of significance, respectively. Enclosed in the parenthesis are household-level clustered heteroscedasticity-consistent standard errors. The regressions in Columns (1) and (2) include household size dummies, survey wave dummies, and survey wave interacted with household size dummies. For the LATE estimates in Column (2), the instrumental variable is lottery assignment, and the endogenous variable is "Ever in Medicaid". The ITT and LATE estimates are base on the double-selection post-LASSO.

 Measure of Treatment Heterogeneity: Heuristic*, MFP = 1, and DFP > 0. Results: Graphical Intuition (3D Plot)

3D Visualization.

Frisch–Waugh–Lovell (FWL) theorem and Double Machine Learning (DML) [back]

- y = f(d, x).
- Imagine f is a linear function.
- y = f(x) and get residual $e_1 = y \hat{y}$.
- d = f(x) and get residual $e_2 = d \hat{d}$.
- $e_1 = f(e_2)$ the coefficient yields how y changes w.r.t. the variable d.
- This is FWL theorem.
- Now, imagine f as some ML algorithm, hence this is DML.
Training and Testing Error and VC Dimension **Deck**



Training and Testing Error and VC Dimension **Deck**



TestingError \leq TrainingError $+ \sqrt{VC(\Pi)/n}$

Training and Testing Error and VC Dimension **Deck**



TestingError \leq TrainingError $+ \sqrt{VC(\Pi)/n}$

Efficient Policy **back**

Variable	Random assignment	Probability rule	CATE rule	Shallow tree	Deeper tree
	(1)	(2)	(3)	(4)	(5)
Panel A: Health care utilization					
Outpatient visits	0.604^{***}	4.74***	5.119^{***}	4.228***	2.898^{***}
last six months	(0.002)	(0.182)	(0.17)	(0.197)	(0.177)
Panel B: Preventive care utilization					
Blood cholesterol checked (ever)	0.659^{***}	0.575^{***}	3.023^{***}	1.934^{***}	1.59^{***}
	(0.005)	(0.176)	(0.146)	(0.166)	(0.154)
Blood tested for high blood	0.625***	1.066***	3.059***	2.665***	2.068***
sugar/diabetes (ever)	(0.003)	(0.157)	(0.124)	(0.137)	(0.178)
Mammogram within last	0.331***	7.008***	10.228***	9.26***	5.75***
12 months (women + 40)	(0.002)	(0.482)	(0.398)	(0.552)	(0.42)
Pap test within last	0.411***	3.489***	5.682***	4.955***	4.058***
12 months (women)	(0.003)	(0.286)	(0.24)	(0.315)	(0.316)
Panel C: Self-reported health					
Self-reported health good/very	0.579^{***}	1.952^{***}	4.186^{***}	4.225***	2.588^{***}
good/excellent (not fair or poor)	(0.003)	(0.174)	(0.145)	(0.201)	(0.195)
Panel D: Potential mechanism					
Have usual place	0.558^{***}	5.462^{***}	7.44***	7.305***	4.718***
of clinic-based care	(0.002)	(0.227)	(0.203)	(0.237)	(0.202)
Have personal doctor	0.544^{***}	6.114***	6.432***	6.144***	4.576***
• • • • • • • • •	(0.003)	(0.192)	(0.207)	(0.244)	(0.181)
Happiness, very happy or pretty	0.629***	2.137***	4.883***	5.042***	3.306***
happy (vs. not too happy)	(0.002)	(0.196)	(0.174)	(0.218)	(0.166)

Table 6: Estimate of the utility improvement of various policies over a random assignment baseline.

Efficient Policy (back)



Efficient Policy (back)

