# Heterogeneity in voter preferences for female candidates

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10-06-2024

1. Run an OLS regression of picked\_cand\_a on cand\_a\_female. What is the average preference for or against female candidates?

```
load("politician gender prefs.rdata")
data <- df
data$picked_cand_a <- as.numeric(data$picked_cand_a)</pre>
data$cand_a_female <- as.numeric(data$cand_a_female)</pre>
summary(lm(picked_cand_a ~ cand_a_female, data = data))
##
## Call:
## lm(formula = picked_cand_a ~ cand_a_female, data = data)
## Residuals:
##
      Min
               1Q Median
                               3Q
## -0.5716 -0.4723 0.4284 0.4284 0.5277
##
## Coefficients:
                Estimate Std. Error t value Pr(>|t|)
## (Intercept) 0.47230 0.01899 24.871 < 2e-16 ***
## cand a female 0.09933
                            0.02674 3.715 0.000212 ***
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 0.4974 on 1382 degrees of freedom
## Multiple R-squared: 0.009886,
                                   Adjusted R-squared: 0.009169
## F-statistic: 13.8 on 1 and 1382 DF, p-value: 0.0002116
```

Run regressions of picked\_cand\_a on cand\_a\_female and one of each of the other included regressions, main effects and interactions.

```
regressors <- setdiff(names(data), c("picked_cand_a", "cand_a_female"))
regression_results <- list()
for (regressor in regressors) {
  formula <- as.formula(paste("picked_cand_a ~ cand_a_female *", regressor))
  model <- lm(formula, data = data)
  regression_results[[regressor]] <- broom::tidy(model)
}</pre>
```

### What is the most significant heterogeneity?

```
most significant <- NULL
lowest_p_value <- Inf</pre>
for (regressor in regressors) {
 interaction_row <- regression_results[[regressor]] |>
   dplyr::filter(term == paste("cand_a_female:", regressor, sep = ""))
 if (nrow(interaction_row) > 0) {
   p_value <- interaction_row$p.value</pre>
   if (p_value < lowest_p_value) {</pre>
     lowest_p_value <- p_value</pre>
     most_significant <- regressor
   }
 }
}
most_significant_model <- lm(as.formula(paste("picked_cand_a ~ cand_a_female *", most_significant)), da
summary(most_significant_model)
##
## Call:
## lm(formula = as.formula(paste("picked_cand_a ~ cand_a_female *",
      most_significant)), data = data)
##
## Residuals:
##
      Min
               10 Median
                               3Q
                                      Max
## -0.6830 -0.5238 0.3806 0.4443 0.7494
##
## Coefficients:
##
                                Estimate Std. Error t value Pr(>|t|)
## (Intercept)
                                 0.04459 -1.121
## cand a female
                                -0.04997
                                                              0.263
## considerwoman n
                                -0.06464
                                            0.01655 -3.906 9.85e-05 ***
                                                    4.141 3.67e-05 ***
## cand_a_female:considerwoman_n 0.09648
                                         0.02330
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 0.4943 on 1380 degrees of freedom
## Multiple R-squared: 0.02335,
                                   Adjusted R-squared: 0.02123
## F-statistic:
                  11 on 3 and 1380 DF, p-value: 3.881e-07
```

#### Why can't we use the standard errors from these estimates for analyzing hetergeneity?

We need to correct for the fact that we are testing for multiple hypotheses, or we will end up with many false positives. We can use Bonferroni or Romano-Wolf correction to adjust the p-values for multiple testing.

#### 2. Estimate a causal forest model

```
Y = as.numeric(df$picked_cand_a)
W = as.numeric(df$cand_a_female)
X <- mltools::one_hot(
data.table::data.table(</pre>
```

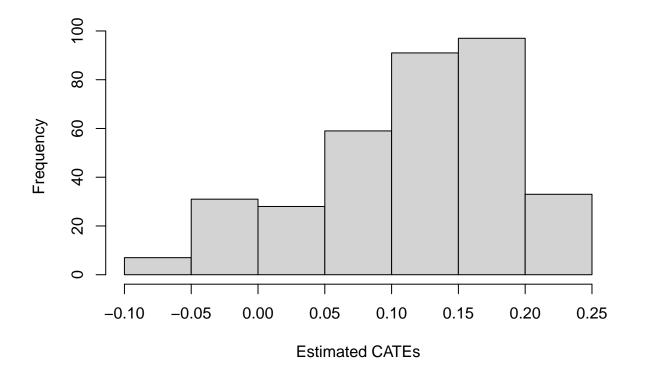
```
df |>
dplyr::select(-cand_a_female, -picked_cand_a) ) )

# Split data into a train and test sample.
set.seed(42)
train = sample(nrow(X), 0.75 * nrow(X))
test = -train

# Fit a CATE function on training data.
cate.forest = causal_forest(X[train, ], Y[train], W[train], num.trees = 5000, seed = 42)
```

## 3. Predict the heterogenous treatment effect

```
X.test = X[test, ]
tau.hat.test = predict(cate.forest, X.test, estimate.variance=TRUE)$predictions
# A histogram of CATE estimates.
hist(tau.hat.test, xlab = "Estimated CATEs", main = "")
```

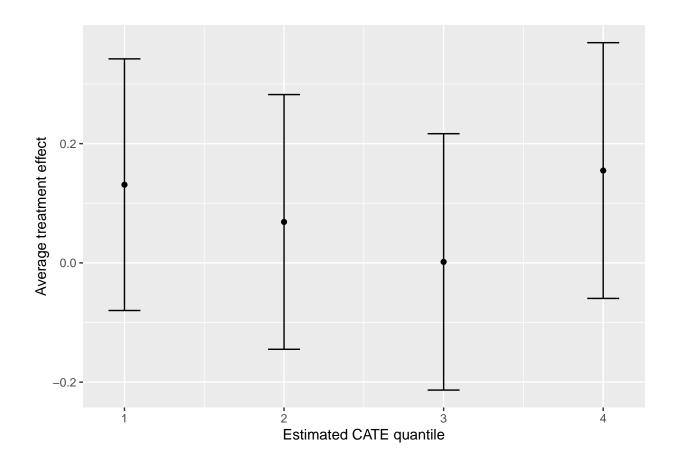


Does the predicted treatment effects predict treatment heterogeneity in the test data?

```
summary(lm(Y[test] ~ W[test] + tau.hat.test + W[test] * tau.hat.test))
```

```
## Call:
## lm(formula = Y[test] ~ W[test] + tau.hat.test + W[test] * tau.hat.test)
## Residuals:
                1Q Median
##
      Min
                                3Q
                                       Max
## -0.5749 -0.4531 -0.4287 0.5124 0.5750
## Coefficients:
                        Estimate Std. Error t value Pr(>|t|)
##
## (Intercept)
                         0.43206
                                    0.07726
                                             5.592 4.59e-08 ***
                                    0.10095
## W[test]
                         0.05413
                                              0.536
                                                       0.592
## tau.hat.test
                         0.10399
                                    0.56377
                                              0.184
                                                       0.854
## W[test]:tau.hat.test 0.31334
                                    0.74535
                                              0.420
                                                       0.674
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 0.5003 on 342 degrees of freedom
## Multiple R-squared: 0.009789,
                                   Adjusted R-squared: 0.001103
## F-statistic: 1.127 on 3 and 342 DF, p-value: 0.3381
No, the interaction is not significant.
# Vizualize
num.groups = 4
quartile = cut(tau.hat.test,
               quantile(tau.hat.test, seq(0, 1, by = 1 / num.groups)),
              labels = 1:num.groups,
              include.lowest = TRUE)
# Create a list of test set samples by CATE quartile.
samples.by.quartile = split(seq_along(quartile), quartile)
# Look at ATEs in each of these quartiles. To calculate these we fit a separate evaluation forest.
eval.forest = causal_forest(X.test, Y[test], W[test])
# Calculate ATEs for each group.
ate.by.quartile = lapply(samples.by.quartile, function(samples) {
  average_treatment_effect(eval.forest, subset = samples)
})
# Plot group ATEs along with 95% confidence bars.
df.plot.ate = data.frame(
 matrix(unlist(ate.by.quartile), num.groups, byrow = TRUE, dimnames = list(NULL, c("estimate", "std.err
 group = 1:num.groups
ggplot(df.plot.ate, aes(x = group, y = estimate)) +
  geom_errorbar(aes(ymin = estimate - 1.96 * std.err, ymax = estimate + 1.96 * std.err, width = 0.2)) +
 xlab("Estimated CATE quantile") +
 ylab("Average treatment effect")
```

##



### 4. What are the ten most important variables for hetergeneity?

```
varimp.cate = variable_importance(cate.forest)
ranked.variables = order(varimp.cate, decreasing = TRUE)
print(colnames(X)[ranked.variables[1:10]])

## [1] "considerwoman_n" "considerwoman" "age" "sdo"
## [5] "progressive" "candwarm" "believe_Believe" "horserace_Warren"
## [9] "favmetoo" "suppclean"
```

### Plot the predicted treatment effect as a function of two most important variables.

```
top.varnames = colnames(X)[ranked.variables[1:2]]

# Select the test set samples predicted to have low/high CATEs.
low = samples.by.quartile[[1]]
high = samples.by.quartile[[num.groups]]

# Make some long format data frames for ggplot.
df.lo = data.frame(
    covariate.value = unlist(as.vector(X.test[low, ..top.varnames])),
    covariate.name = rep(top.varnames, each = length(low)),
    cate.estimates = "Low"
)
```

```
df.hi = data.frame(
    covariate.value = unlist(as.vector(X.test[high, ..top.varnames])),
    covariate.name = rep(top.varnames, each = length(high)),
    cate.estimates = "High"
)
df.plot.hist = rbind(df.lo, df.hi)

# Plot overlaid histograms of the selected covariates by low/high classification.
ggplot(df.plot.hist, aes(x = covariate.value, fill = cate.estimates)) +
    geom_histogram(alpha = 0.7, position = "identity") +
    facet_wrap(~ covariate.name, scales = "free", ncol = 2)
```

## `stat\_bin()` using `bins = 30`. Pick better value with `binwidth`.

