

Causal Impact of Campaign Strategies Analysis

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1 Average Causal Effect (ATE)

Assumptions:

- 1) Consistency: $D_i = d \rightarrow Y_i(d) = Y_i$
- 2) Unfoundness: given X , if $\{Y_i(1), Y_i(0)\} \perp D_i | X_i$, D_i is conditionally ignorable.

In our question, $D_i = 1$: have the specific VSA; $D_i = 0$: don't have the specific VSA

$$ATE = E[E[Y_i | D_i = 1, X_i]] - E[E[Y_i | D_i = 0, X_i]]$$

$$A\hat{T}E = \frac{1}{n} \sum_{i=1}^n [\hat{E}[Y_i | D_i = 1, X_i] - \hat{E}[Y_i | D_i = 0, X_i]]$$

Consider $E[Y_i | D_i, X_i] = \tau D_i + X_i^\top \beta$, then $ATE = \tau \rightarrow \tau$ has a causal interpretation.

Now we need to do linear regression to fit $E[Y_i | D_i = 1, X_i]$

In this setting, Assumption 2 is relatively difficult to fulfill, because Regression is more dependent on having a comprehensive enough set of covariates to satisfy "conditional ignorability".

Randomized Controlled Trials (RCT)

RCT is a well-known method to calculate causal effect. In our settings, we cannot randomize D_i , and it's also hard to determine which variable should we include in regression so that conditional ignorability holds; what will happen when X_i is assumed to be linear in model but the CEF is not linear on X_i ?

However, because our data collected before we change the campaign strategy in different age groups (16-23; other) are almost parallel, we consider to use difference in differences to analysis the causal effect.

2 Difference-in Differences

Assumptions:

- 1) SUTVA: Stable Unit Treatment Value Assumption \rightarrow the potential outcomes of unit i are not relevant to the treatment of other units \rightarrow we should choose the units of analysis to minimize interference across units.
- 2) No-Anticipation Assumption: $Y_{i,t}(g) = Y_{i,t}(\infty)$ for all $t < g \rightarrow$ Before the start of treatment, potential outcomes for units were consistent with those that did not receive treatment
(Selection Bias might be introduced here)
- 3) Parallel Trends Assumption: $E[Y_{i,t=2}(\infty) | G_i = 2] - E[Y_{i,t=1}(\infty) | G_i = 2] = E[Y_{i,t=2}(\infty) | G_i = \infty] - E[Y_{i,t=1}(\infty) | G_i = \infty] \rightarrow$ In the absence of treatment, the evolution of the outcome among treated units is the same as the evolution among the untreated units on average.

Under 1) 2) 3) assumptions,

$$ATT = (E[Y_{i,t=2} | G_i = 2] - E[Y_{i,t=1} | G_i = 2]) - (E[Y_{i,t=2} | G_i = \infty] - E[Y_{i,t=1} | G_i = \infty])$$

ATT represents the average treatment effect on the treated group at time $t=2$.

We have two methods to estimate ATT in population,

- 1) plug-in estimator: $\hat{\theta}^{DID} = (\bar{Y}_{g=2,t=2} - \bar{Y}_{g=2,t=1}) - (\bar{Y}_{g=\infty,t=2} - \bar{Y}_{g=\infty,t=1})$
- 2) two-way fixed-effects (TWFE) regression: $Y_{i,t} = \alpha_0 + \gamma_0 \mathbf{1}\{G_i = 2\} + \lambda_0 \mathbf{1}\{T_i = 2\} + \beta_0^{twfe} (\mathbf{1}\{G_i = 2\} \mathbf{1}\{T_i = 2\}) + \epsilon_{i,t}$, where β_0^{twfe} is the estimate of $\hat{\theta}^{DID}$

In our setting, we use Repeated Cross-section data $\{Y_i, G_i, T_i\}_{i=1}^n$ i.i.d. draw from the mixture distribution:

$$P(Y \leq y, G = g, T = t) = \mathbf{1}\{t = 2\} \lambda P(Y_{t=2} \leq y, G = g | T = 2) + \mathbf{1}\{t = 1\} (1 - \lambda) P(Y_{t=1} \leq y, G = g | T = 1)$$

which means select the pre-intervention time period ($T = 1$) with probability $(1 - \lambda)$ Select the post-intervention time period ($T = 2$) with probability λ . Collect data for $G = 1, G = 2$, respectively.