Interpretation and robustness

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$$Dem_{it} = \alpha + \beta_1 GDP_{it} + \beta_2 Oil_{it} + \beta_3 GDP_{it} \times Oil_{it} + \delta X_{it} + \mu_i + \gamma_t + \varepsilon_{it}.$$

• β_1 is the effect of the treatment conditional on Oil = 0:

$$E(Dem_{it}|GDP_{it} = 1, Oil_{it} = 0, X) - E(Dem_{it}|GDP_{it} = 0, Oil_{it} = 0, X)$$

• $\beta_1 + \beta_3 Oil$ is the effect of the treatment conditional on Oil = C:

$$E(Dem_{it}|GDP_{it} = 1, Oil_{it} = C, X) - E(Dem_{it}|GDP_{it} = 0, Oil_{it} = C, X)$$

• β_3 is the additional effect of the treatment for *Oil* when goes from 0 to *C*.



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β₃ is the additional effect of the treatment for Oil when goes from 0 to C.



- When we use observational data we may not observe all confounders.
 - We may be able to control for observable/measurable ones.
 - We can also use the tools we've learned: diff-in-diff, instrumental variables, RDD, and so on.
- Is this sufficient? Oftentimes it's not!
 - Robustness to small and sensible manipulations?
 - Robustness to potential confounders?
 - Robustness to alternative explanations?
- What's the purpose of addressing the previous issues?
 - To at least provide robust conclusions about the direction of the causal effect.
 - To show the estability of the magnitude/significance of the effect.

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Robustness tests are ways to

- Robustness means two things in general:
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 - Introduce sets of confounders at a time to check for stability.
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- Robustness tests can be also clever exercises! (Placebo tests)
 - Theoretical prior indicates where should find an effect and where we shouldn't.

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 - Classify confounders in sets, e.g.: fixed effects, individual-level, municipality-level.
 - \bullet Time \times area fixed effects control s for time-variant confounders at higher level than the level of the treatment assignment.
- Drop observations/areas/sets of observations with replacement.
 - Similar to bootstrapping, but not bootstrapping.
 - One at a time, recompute, plot coefficient with SE.
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 - Generate a variable that has different levels of correlation with treatment and outcome.
 - Plot it against all observable confounders it should look extraneous/implausible.
- Manski bounds/partial identification (won't be covered).
 - Assume worst case scenario and generate the bounds.
 - Trim bounds if warranted.
- Additional: For multiple treatments/outcomes if measurement is similar:
 - Equivalence testing. Evaluate all four patterns:
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Theoretical robustness tests: measurement and placebos

- No empirical paper is good without a theory (or model) driving data analysis.
- Solid theory gives guidelines to define measurement of dependent variable, treatment and confounders.
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 - It can also guide the selection of an instrumental variable.
- Solid theory provides use with guidelines for computing interaction effects.
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