Alternative Specifications for Instrumental Variable Analysis in Structural Equation Modeling: First Steps Toward Latent Analysis of Symmetrically Predicted Endogenous Subgroups

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Background

- Dissertation was on a method called Analysis of Symmetrically Predicted Endogenous Subgroups (ASPES).
- ASPES came from the evaluation literature (Peck, 2003; Peck, 2013) and has been called a "cousin" of principal stratification.
- It leverages random assignment of an intervention in order to identify average causal effects for each level of a post-random assignment (endogenous) variable of interest that was <u>only observed in one group</u> (e.g., compliance, quality of implementation).

Setting

- Let's call this post-treatment variable *M* since it is an intermediate variable. If it had observations in both the treatment and control groups, we could treat it as a causal mediator.
- However, when M is only observable in one group, it's impossible to treat it as a causal mediator because we have no information on one of the potential mediators (M_0 or M_1).
- Let's imagine *M* is a measure of implementation quality in the treatment group.

If *M* is discrete (e.g., low vs. high quality)...

- ASPES very much resembles well known methods of estimating principal stratum causal effects such as the complier average causal effect (CACE).
- Peck (2003) nonparametrically identified analogues to the CACE and NACE that can be estimated by ASPES.
- ASPES accomplishes this identification via a mean independence assumption, which I argue is best thought of as an exclusion restriction (ER) assumption, because it requires an instrumental variable.

If *M* is discrete (e.g., low vs. high quality)...

- ASPES essentially uses an instrument, *V*, for the interaction effect between treatment assignment (*T*) and *M* on the outcome, *Y*.
- The assumption used in ASPES ends up being mathematically equivalent to an assumption that there is no interaction effect between the chosen instrument and treatment assignment on the outcome.
- I call this the "interaction exclusion restriction" (IER) assumption because it is equivalent to using *T*V* as an instrument for *T*M*, requiring the usual ER assumption on the effect of *TV* on *Y*.

If *M* is discrete (e.g., low vs. high quality)...

• In this setting, ASPES is a nonparametric version of the approach to principal stratification via mixture modeling proposed by Jo (2002).

ASPES (Peck, 2003)

Stage 1:

$$M = f(V, X)$$

(estimate this in the treatment group using cross-validation method to prevent overfitting, then compute \widehat{M} for each unit)

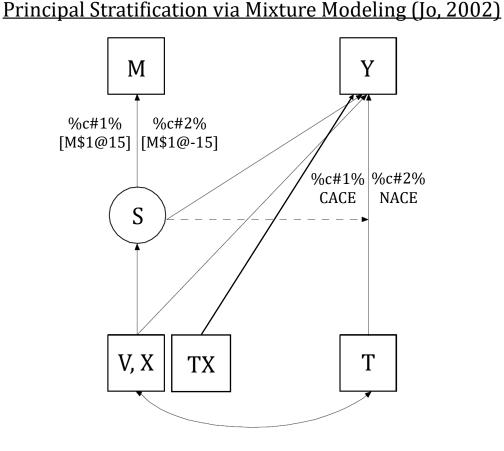
Stage 2:

$$CACE^{P} = E[Y|T = 1, \hat{M} = 1] - E[Y|T = 0, \hat{M} = 1]$$

$$NACE^{P} = E[Y|T = 1, \hat{M} = 0] - E[Y|T = 0, \hat{M} = 0]$$

$$CACE = \frac{E[1 - \hat{M}|M = 0]CACE^{P} - (1 - E[\hat{M}|M = 1])NACE^{P}}{E[1 - \hat{M}|M = 0] + E[\hat{M}|M = 1] - 1}$$

$$NACE = \frac{E[\hat{M}|M = 1]NACE^{P} - (1 - E[1 - \hat{M}|M = 0])CACE^{P}}{E[1 - \hat{M}|M = 0] + E[\hat{M}|M = 1] - 1}$$



- There isn't a lot of guidance on how to appropriately incorporate *M* in our analyses when it's continuous.
- The principal stratification framework struggles to accommodate an *M* with many levels (even many discrete levels).
- Often, researchers will artificially discretize *M* in this case so that they can apply the well-established methods for the discrete *M* case.

• ASPES can actually be used with a continuous *M* similarly to how two stage least squares (TSLS) methods for IV often handle continuous causal variables of interest.

Stage 1:

 $M = f(V,X) + \delta$ $\delta \sim N(0, \sigma_M^2)$ (estimate this in the treatment group using cross-validation method to prevent overfitting,

then compute \widehat{M} for each unit)

Stage 2:

$$Y = g(T, \widehat{M}, X) + \epsilon$$

$$\epsilon \sim N(0, \sigma_Y^2)$$

This can be done with TSLS, using linear models in both stages if those functional form assumptions are reasonable.

Stage 1:

$$M = \alpha_0 + \alpha_1 V + \alpha_2 X + \delta$$
$$\delta \sim N(0, \sigma_M^2)$$

(estimate this in the treatment group using cross-validation method to prevent overfitting, then compute \widehat{M} for each unit)

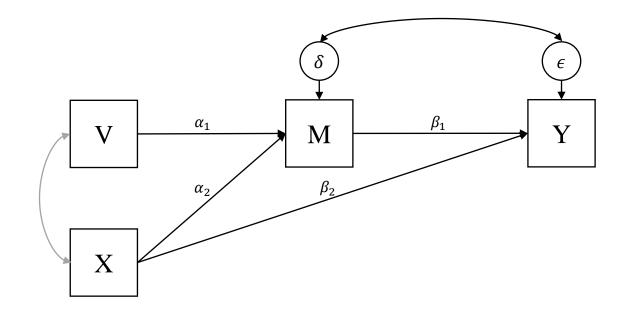
Stage 2:

$$Y = \beta_0 + \beta_1 T + \beta_2 \widehat{M} + \beta_3 T \widehat{M} + \beta_4 X + \beta_5 T X + \epsilon$$

$$\epsilon \sim N(0, \sigma_Y^2)$$

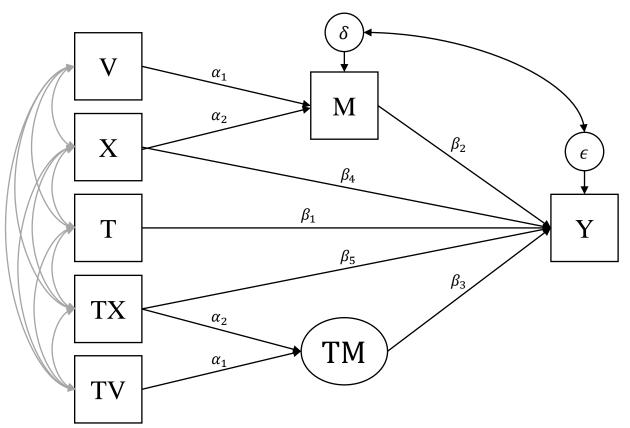
Translating to FIML-estimated Model

 $M = \alpha_0 + \alpha_1 V + \alpha_2 X + \delta$ $Y = \beta_0 + \beta_1 \widehat{M} + \beta_2 X + \epsilon$



Translating to FIML-estimated Model

$$\begin{split} M &= \alpha_0 + \alpha_1 V + \alpha_2 X + \delta \\ Y &= \beta_0 + \beta_1 T + \beta_2 \widehat{M} + \beta_3 T \widehat{M} + \beta_4 X + \beta_5 T X + \epsilon \end{split}$$



Translating to FIML-estimated Model

- Full information maximum likelihood (FIML) handles the missing data in *M* under the assumption of missingness at random (MAR), which is satisfied by design due to the missingness in *M* being perfectly explained by *T*.
- Obviously, this limits the ASPES approach by making parametric assumptions, but these could be relaxed, and estimating the model with FIML circumvents the cumbersome stage-1 cross-validation and opens ASPES up to all the benefits/possibilities of latent variable modeling.

Preliminary Simulation Results

- Preliminary simulations have thus far been confirming that this FIML approach reproduces the estimates found with the original ASPES two-stage implementation using linear models, and that both approaches accurately recover the *TM* interaction effect when the IER assumption holds.
- Additionally, as long as the IER assumption was satisfied, the *TM* effect is accurately recovered even when *M* and *Y* share a confounder that mediates and/or moderates the treatment effect.

Causal Moderation?

- Although this clearly cannot be considered causal mediation, the question still stands: is this *TM* interaction effect an average causal *moderation* effect?
- Or is it simply an unbiased estimate of the heterogeneity in the average causal effect that happens to relate to *M*?
- In potential outcomes terms, is this the unbiased average causal effect of M on $Y_1 Y_0$? Or is this just a measure of the relationship between M and $Y_1 Y_0$ that may solely be due to an omitted confounder?

Causal Moderation?

• Let's consider VanderWeele's (2014) decomposition of causal mediation/interaction:

$$ITT = Y_1 - Y_0 = CDE(0) + INT_{ref} + INT_{med} + PIE$$

Controlled Direct Effect $(CDE(0)) = Y_{10} - Y_{00}$ Reference Interaction $(INT_{ref}) = ((Y_{11} - Y_{10}) - (Y_{01} - Y_{00}))(M_0)$ Mediated Interaction $(INT_{med}) = ((Y_{11} - Y_{10}) - (Y_{01} - Y_{00}))(M_1 - M_0)$ Pure Indirect Effect (PIE) = $(Y_{01} - Y_{00})(M_1 - M_0)$

Causal Moderation?

• If we define *M* as the value that would be realized when assigned to the treatment group, then there is no difference between M_0 and M_1 ; $M_0 = M_1 = M$

$$ITT = Y_1 - Y_0 = CDE(0) + INT_{ref} + INT_{med} + PIE$$

Controlled Direct Effect: $(CDE(0)) = Y_{10} - Y_{00}$ Reference Interaction: $(INT_{ref}) = ((Y_{11} - Y_{10}) - (Y_{01} - Y_{00}))(M)$ Mediated Interaction: $(INT_{med}) = ((Y_{11} - Y_{10}) - (Y_{01} - Y_{00}))(M - M)$ Pure Indirect Effect: $(PIE) = (Y_{01} - Y_{00})(M - M)$

Causal Moderation, kind of?

- What we end up with is a kind of one-sided moderation effect.
- We haven't actually identified the causal effect of *M* on *Y* given *T*, but we have identified the causal effect of *T* on *Y* given *M* (obviously through random assignment) and the causal effect of *M* on *Y* for the treatment group.
- It's arguably causal moderation in the sense that M causing changes in Y_1 while causing no changes in Y_0 naturally means that M causes changes in $Y_1 Y_0$.

Conclusions

- Still working on these simulations
- Next steps include exploring ASPES with a latent variables using the FIML approach, as well as other extensions such as multilevel versions of the approach.
- Thanks for listening!

References

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