The Insidious Problem of Selection Biases: How to Recognize Them, and What to

Do about Them

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A loose outline

- Introduction, basic concepts
- The selection bias problem
- The endogenous switching regression solution
- An application: the impact of teleworking on vehicle-miles driven
- Summary and conclusions



Introduction: the scenario

• There is a *discrete event* of interest

- "Treatment", "experiment", "intervention"
- Extreme event (earthquake, bridge collapse, etc.)
- "New" technologies (e.g. medical, agricultural, telecommunications)
- Policies (e.g. promoting transportation safety, or sustainable transportation)
- Individuals are "untreated" or "treated" (there can be multiple categorical/ordered treatments)
- Individuals may actively choose, or passively experience, the treatment (or not)
- We want to estimate the effect of that event on an (individually-experienced, then aggregated) **outcome** of interest, i.e., the *"treatment effect"*
 - Example outcomes: Blood pressure, crop yield, vehicle-miles driven
 - Assume we observe the outcome whether a person is treated or untreated

Some transportation examples

Treatment

- Wearing a helmet while (motor)cycling
- Buying a fuel-efficient car
- Living in a transit-friendly, mixeduse neighborhood
- Adopting teleworking (TWing)

Outcome

Injury severity

- CO₂ emissions
- Vehicle-miles driven (VMD)



What "treatment effects" (TES) might we be interested in?

• The treatment effect on the treated (TT)

 Difference between outcome if treated versus not, for those who have been treated

The treatment effect on the untreated (TUT)

• Difference between outcome if treated versus not, *for those who have NOT been treated* (i.e. *how would the outcome have been different if they had been treated?*)

• The average treatment effect (ATE)

- The appropriately-weighted average of TT and TUT
- The average impact of treatment for a person *randomly-selected from the population*, which contains both treated and untreated individuals
- It's not always obvious which TE is most relevant, nor even which the authors intend!

Hm – how hard can it be to estimate a TE?

- The problem: with cross-sectional data, we only observe people as treated or untreated (their factual state at the time of measurement) – we don't observe them in the opposite condition (their counterfactual state)
- Well, fine, but can't we just model an outcome Y as, say,
 - $Y = X\beta + \delta T + \eta$, where (throughout, individual-level subscripts are omitted for simplicity)
 - X = explanatory variables; T = treatment dummy; { β , δ } = coefficients; $\eta \sim N(0, \sigma^2)$?
 - $\hat{\delta}$ would be the estimated TE, wouldn't it?
 - But *which* TE? In essence, this approach would force TT = TUT = ATE
- OK, then how about separate equations for the treated and untreated?
 - $Y_T = X \beta_T + \eta_T$ and $Y_U = X \beta_U + \eta_U$
 - ATE = avg over the entire (*representative*) sample of $(\hat{Y}_T \hat{Y}_U) = X(\hat{\beta}_T \hat{\beta}_U)$
 - TT = avg over the treated of $(\hat{Y}_T \hat{Y}_U)$ (where \hat{Y}_U applies $\hat{\beta}_U$ to the treated people's Xs)
 - TUT = avg over the untreated of $(\hat{Y}_T \hat{Y}_U)$ (where \hat{Y}_T applies $\hat{\beta}_T$ to the untreated people's Xs)

This would be fine, *IF*...

- The treatment were *randomly assigned* across the population
 - Then the treated and untreated cases in a (large enough, and representative) sample would each be random samples from the population of outcomes
- But what if, instead, the treated group differs from the untreated group in ways that are relevant to the outcome? I.e. we have a **selection bias**
 - Those receiving the new medical treatment may be sicker
 - Those trying the new fertilizer may have also adopted other innovations
 - Those wearing a helmet (before it became mandatory) may be safer (motor)cyclists (or, they may compensate with *more* risky behaviors)
 - Those buying a fuel-efficient car may want to travel more
- This is likely to happen when the individual voluntarily adopts the treatment!

What's the problem with that (in English)?

- Before treatment, if the to-be-treated people already differ from the to-remain-untreated people, then how can we separate the effect of the treatment from the effect of those pre-existing differences?
 - We can account for the effect of differences in *observed variables* (the Xs) those are controlled for in the model
 - But the naïve model shown previously **does not account for the effect of differences in unobserved variables** (the η s)
- We should not expect the treatment to have the same effect if imposed on, or adopted by, a currently untreated person
- Even the estimated *treatment effect on the treated (TT)* will be biased!
- That's because our estimated TT confounds the true effect of the treatment with the effects due to pre-existing (unobserved) inclinations/traits that may cause a portion of the observed effect to occur even without the treatment
- In other words, our estimates of β_T and β_U will be biased, by virtue of being based on non-random samples

What's the problem (in math)?

- We have two components to the model system (the "endogenous switching regression model", or ESRM, AKA Roy's model, mover-stayer model, Tobit type 5 model, etc.)
- The selection model (binary probit) governs selection into treatment:
 - $Z^* = W\gamma + \varepsilon$; treated if $Z^* > 0$, untreated if $Z^* \le 0$
 - W = explanatory variables, γ = coefficients, ε = error term
- Two outcome models (linear regressions):
 - If treated: $Y_T = X\beta_T + \eta_T$

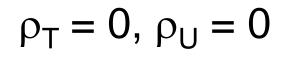
 $\rho_i = Corr(\varepsilon, \eta_i)$

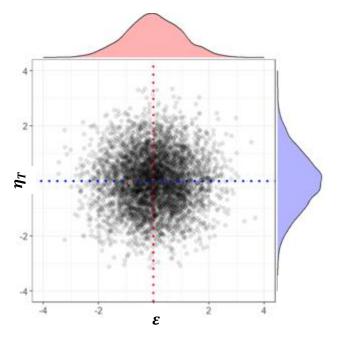
- If untreated: $Y_U = X \beta_U + \eta_U$
- X = explanatory variables, β_T , β_U = coefficients, η_T , η_U = error terms
- Trivariate normal assumption for the error term distribution:

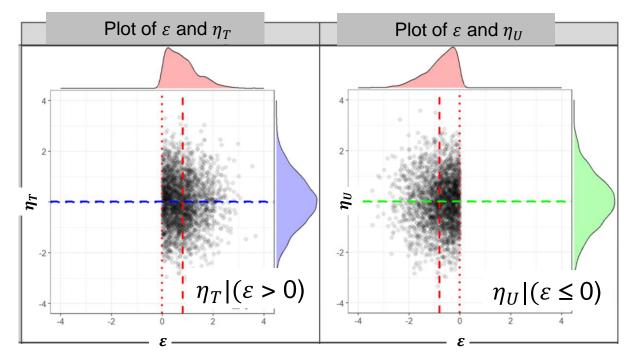
$$\begin{bmatrix} \varepsilon \\ \eta_T \\ \eta_U \end{bmatrix} \sim N \left(\begin{bmatrix} 0 \\ 0 \\ 0 \end{bmatrix}, \begin{bmatrix} 1 & \rho_T \sigma_T & \rho_U \sigma_U \\ \rho_T \sigma_T & \sigma_T^2 & 0 \\ \rho_U \sigma_U & 0 & \sigma_U^2 \end{bmatrix} \right)$$

The distribution of ϵ is truncated for each subsample (T) and (U) (but if ϵ and η are *uncorrelated*, it doesn't matter)

Corr (ε , η_T) = ρ_T = 0







Unconditional expected value (before truncation; by color)

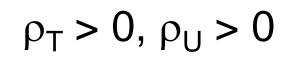
– – Conditional expected value (after truncation; by color)

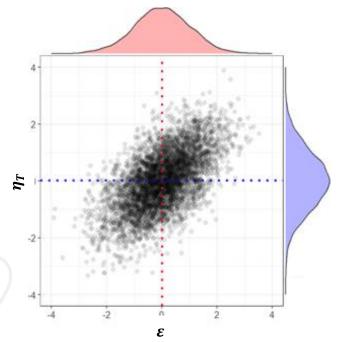
Source: Kim & Mokhtarian, 2023b

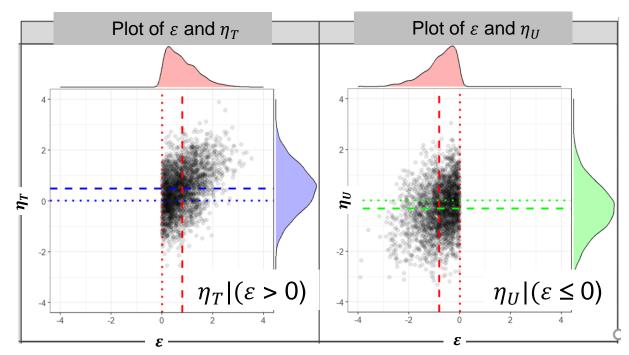
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If ε and η are correlated, truncation for ε propagates to the outcome model error terms, η_T and η_U

Corr (ε , η_T) = ρ_T > 0







Unconditional expected value (before truncation; by color)

--- Conditional expected value (after truncation; by color)

Source: Kim & Mokhtarian, 2023b

Georgia

What's the problem (in math-y English)?

- We only observe people in one state (treated or untreated)
- Therefore, we are sampling from *truncated distributions* of the error terms (unobserved influences, ε) of the *model determining which state people are in* (i.e. the *selection model*)
- Because those ϵ error terms are *correlated* with the η error terms of the outcome equations, that truncates the distributions of the error terms of the *outcome equations* as well
- Thus, although across the population we can assume the outcome errors have mean 0, in our biased sample they have a non-zero mean
- If we do a regular regression we are assuming a 0 mean of the outcome errors, so that non-zero mean (the selectivity bias) interferes with (i.e. biases) the estimates of the true parameters of the model

What's the solution (in math and English)?

- The ESRM "knows" the amount of the selectivity bias, and corrects for it, to produce unbiased estimates of the model parameters, and thus the treatment effect:
 - We want the **unconditional** (i.e. the population-wide) averages of Y:
 - $E(Y_T) = \mathbf{X}\boldsymbol{\beta}_T + E[\varepsilon_T] = \mathbf{X}\boldsymbol{\beta}_T$ (because the pop.-wide $E[\varepsilon_T] = E[\varepsilon_U] = 0$)
 - $E(Y_U) = X \beta_U + E[\varepsilon_U] = X \beta_U$ $\checkmark Cov(\varepsilon, \eta_T)$
 - Instead, we have the (factual) conditional expectations (i.e. separately for T and U):
 - $E(Y_T \text{ for } a T) = X\beta_T + E[\varepsilon_T | Z^* > 0] = X\beta_T + \rho_T \sigma_T \frac{\phi(W\gamma)}{\Phi(W\gamma)}$ Pr[Z* > 0], i.e. Pr[treated]
 - $E(Y_{U \text{ for } a \ U}) = X \beta_U + E[\varepsilon_U | Z^* \le \mathbf{0}] = X \beta_U + \rho_U \sigma_U \left[\frac{-\phi(W\gamma)}{1 \Phi(W\gamma)} \right]$
 - And we can also have the *counterfactual* **conditional** expectations:
 - $E(Y_{T \text{ for } a \text{ } \textit{U}}) = X\beta_T + E[\varepsilon_T | \textbf{Z}^* \leq \textbf{0}] = X\beta_T + \rho_T \sigma_T \left[\frac{-\phi(W\gamma)}{1 \Phi(W\gamma)} \right]$
 - $E(Y_{U \text{ for } a T}) = X \beta_{U} + E[\varepsilon_{U} | Z^{*} > 0] = X \beta_{U} + \rho_{U} \sigma_{U} \frac{\phi(W\gamma)}{\Phi(W\gamma)}$

(what E[Y] would be for an *un*treated person with traits **X**, **W**, *if she were to be treated*)

To restate the problem (with the additional math we've now seen)

- If we estimate an equation like
 - $Y_{T \text{ for } a T} = X \beta_T + [\varepsilon_T | Z^* > 0],$

• Assuming that $E[\varepsilon_T | Z^* > 0] = 0$ when it is really $\rho_T \sigma_T \frac{\phi(W\gamma)}{\Phi(W\gamma)}$, the nonzero bias term gets absorbed into the $X\beta_T$ term, which biases the estimates of β_T (and similarly for β_U)

What's the solution (in math and English)? (cont'd)

- Once we properly account for the selection bias (giving us unbiased estimates of β_T and β_U), then we can compute the treatment effects as:
 - Average treatment effect (ATE):
 - $E(Y_T) E(Y_U) = \boldsymbol{X}(\boldsymbol{\beta}_T \boldsymbol{\beta}_U)$
 - Treatment effect on the treated (TT):
 - $E(Y_{T \text{ for } a T}) E(Y_{U \text{ for } a T}) = X\beta_T + E[\varepsilon_T | Z^* > 0] (X\beta_U + E[\varepsilon_U | Z^* > 0])$
 - Treatment effect on the untreated (TUT):
 - $E(Y_{T \text{ for } a \text{ } U}) E(Y_{U \text{ for } a \text{ } U}) = X\beta_T + E[\varepsilon_T]Z^* \le 0] (X\beta_U + E[\varepsilon_U|Z^* \le 0])$
 - Whereas the observed difference is:
 - $E(Y_{T \text{ for } a T}) E(Y_{U \text{ for } a U}) = X\beta_T + E[\varepsilon_T | Z^* > 0] (X\beta_U + E[\varepsilon_U | Z^* \le 0])$
 - (Where, in estimation, the selection bias terms are invisibly absorbed into the β estimates, thereby biasing them)

counterfactual

• Which is different from all those other quantities

Example application: impact of working from home (WFH) on vehiclemiles driven (VMD)

Acronyms:

WFH = work(ing) from home TW = telework, TWer = teleworker, TWing = teleworking For this talk, TW ≈ WFH NTWer: non-TWer, does not WFH NUTWer: non-usual TWer, WFH < 3 days / wk UTWer: usual TWer, WFH 3+ days / wk VMD = (weekly) vehicle-miles driven (key travel indicator)

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Some potential impacts of WFH on vehicle travel

- There will be local variations
- Multiple factors will counteract each other
- What will the overall net impact be?

Expected (net) impact on \rightarrow due to changes in \downarrow	vehicle-miles	vehicle-trips
Commute travel		
Nonwork travel	?	?
Residential relocation		
Vehicle ownership	?	?
Mode choice (e.g. transit to solo driving)		

A tale of two types of travel diary studies of TWing

TW program evaluations

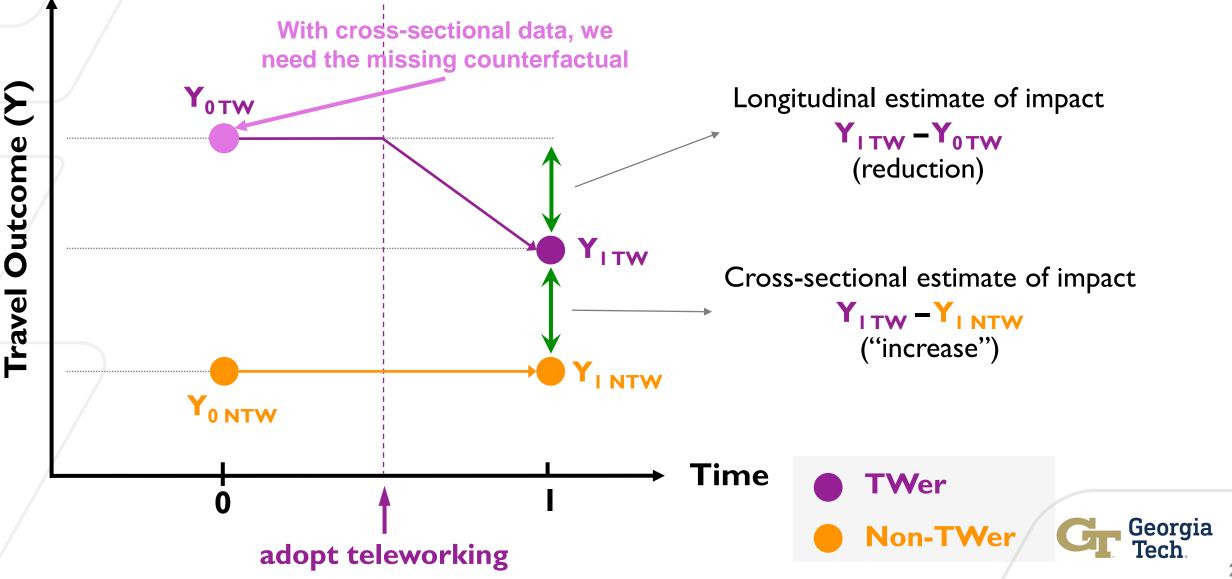
- Early (1980s 1990s)
- Small, unrepresentative samples
- Focused on TWing
- Panel data (before-after)
- Found travel reductions (TW decreased travel)

General travel surveys

- Later (2000s 2010s)
- Large, representative samples
- No emphasis on TWing
- Cross-sectional data
- Finding complementarity (TW increases travel)



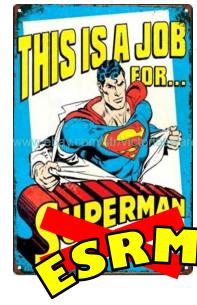
Selection bias: longitudinal vs. cross-sectional inference



This is a job for ESRM!

- Some of Xinyi's contributions:
 - Two TWing treatments (NUTW & UTW), plus untreated (NTW)
 - Estimated ordinal as well as multinomial selection models
 - With log-transformed VMD (then back-transformed treatment effects)
 - Innovative visualizations of aggregate and disaggregate treatment effect components
 - **R scripts** to improve the flexibility and statistical efficiency of previously-existing estimation software
 - Careful exposition, including clearing up some critical ambiguities in Dubin & McFadden (1984)





Data overview (N = 1,584)

Funded by Cintra (Ferrovial)

Impact of COVID-influenced TW on toll revenues

Survey focus

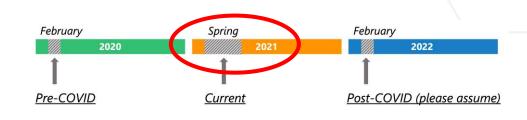
Telework and work patterns before during, and after COVID-19

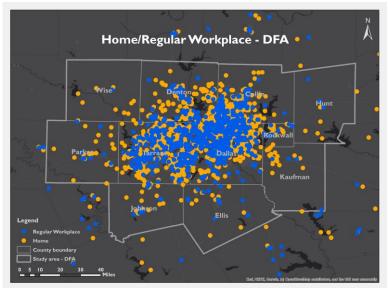
Study areas

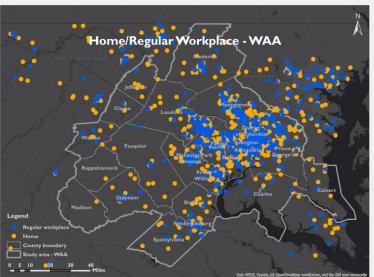
- Dallas-Fort Worth-Arlington (DFA)
- Washington-Arlington-Alexandria (WAA)

Respondent sources

- Cintra database (DB): current and potential customers who consented to be surveyed
- Online panel (OP): three vendor companies
- Data collection Feb. 24 April 30, 2021







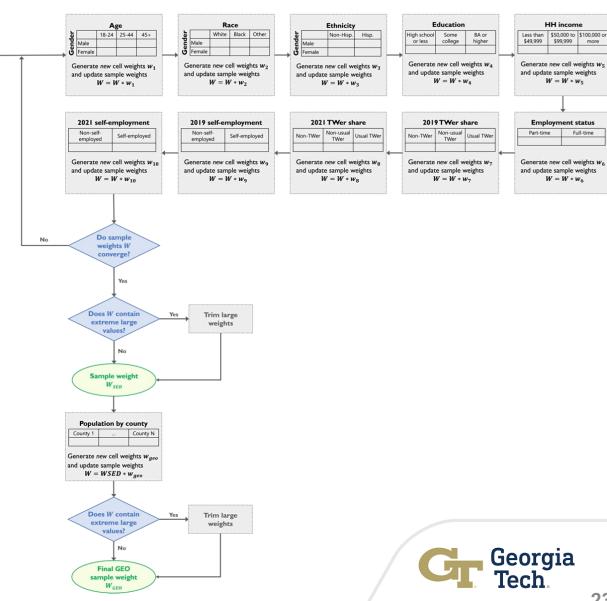
Sample weights

Sample was weighted (by region) to reflect pop. distributions on:

Initial sample

weights (W = 1)

- Gender
- Age
- Race
- Ethnicity
- Education
- Household income
- Employment status
- 2019 (pre-COVID) and 2021 (during-COVID) shares of
 - Non-TWers
 - Non-usual TWers (< 3 days/wk)
 - Usual TWers (3+ days/wk)
- 2019 (pre-COVID) and 2021 (during COVID) self-employment shares
- Employed population by county



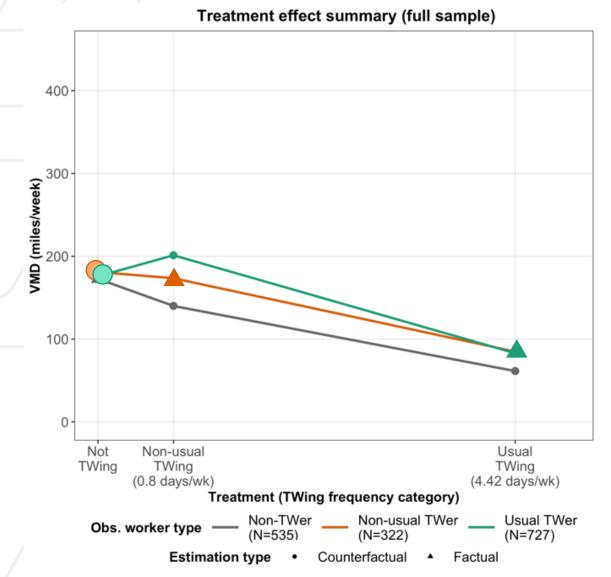
Components of treatment effects (TEs)

 $\mathbb{E}[VMD_{j'}|Z = j]$: Expected weekly VMD if {in, moved to} state j' given observed to belong to group j

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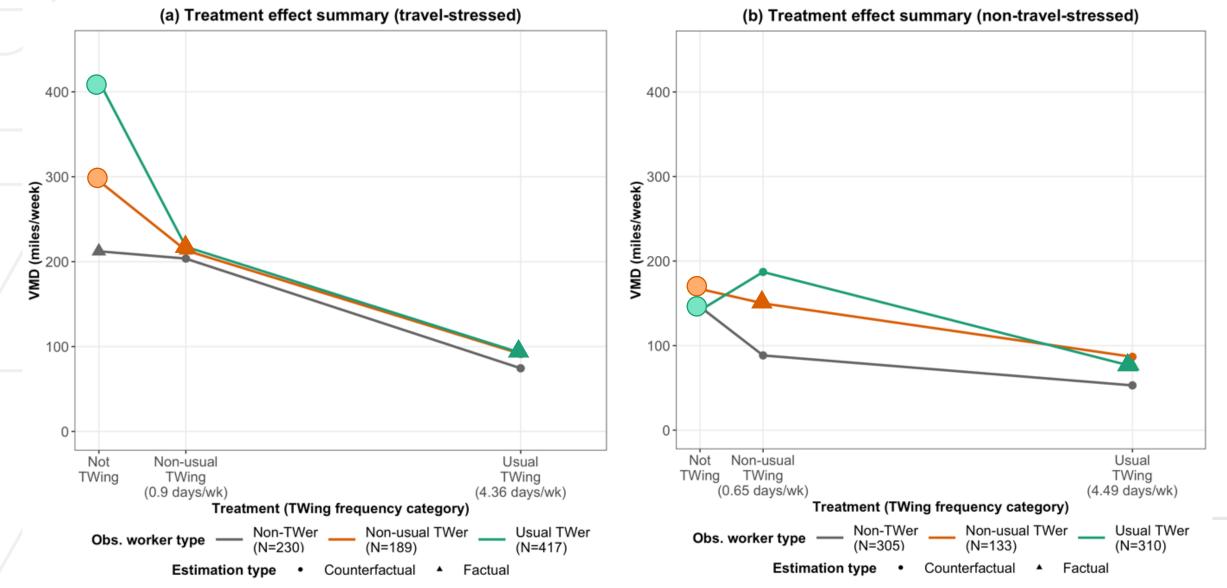
		Potential status		
Observe status	A: If untreated (NTW)	B: If NUTW-treated	C: If UTW-treated	
NTWe	$\mathbb{E}[VMD_{NTW} Z = NTW]$ Expected VMD of a NTWer	$\mathbb{E}[VMD_{NUTW} Z = NTW]$ Expected VMD of a NTWer if TWing less than 3 days/week	$\mathbb{E}[VMD_{UTW} Z = NTW]$ Expected VMD of a NTWer if TWing 3 or more days/week	
	NUTWing treatme	ent effect on the untreated UTWing treatment e	ffect on the untreated	
NUTW	$\mathbb{E}[VMD_{NTW} Z = NUTW]$ Expected VMD of a NUTWer if not TWing	$\mathbb{E}[VMD_{NUTW} Z = NUTW]$ Expected VMD of a NUTWer	$\mathbb{E}[VMD_{UTW} Z = NUTW]$ Expected VMD of a NUTWer if TWing 3 or more days/week	
	Treatment effect	on the NUTW-treated		
UTWer $ \begin{array}{l} \mathbb{E}[VMD_{NTW} Z = UTW] \\ \text{Expected VMD of a UTWer if not} \\ \text{TWing} \end{array} $		$\mathbb{E}[VMD_{NUTW} Z = UTW]$ Expected VMD of a UTWer if TWing less than 3 days/week	$\mathbb{E}[VMD_{UTW} Z = UTW]$ Expected VMD of a UTWer	
		Treatment effect on the UTW-treated	f Georgi Tech	

Full sample model (2 treatments: NUTWing & UTWing)

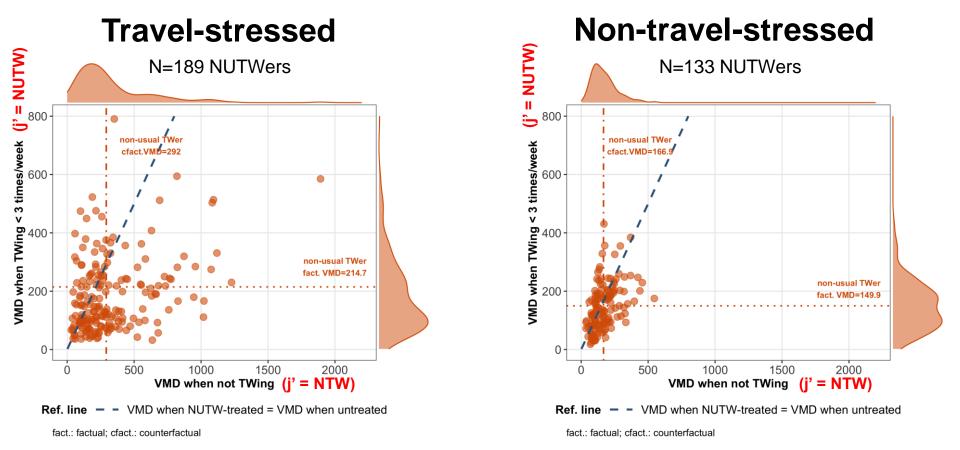


- Focusing on the TEs (compared to not TWing) for the two observed TWer groups:
 - A = factual
 - • , = (NTW) counterfactual
- Ave. VMD of non-usual TWers (15% of the sample) barely declines (not statistically significant)
 - -7.5 mi/wk, or, -9.4 mi/TWing occasion
- Ave. VMD of usual TWers (28% of the sample) declines substantially (statistically significant)
 - -92.9 mi/wk, or, -21.0 mi/TWing occ.

Comparison of travel-stressed and non-travel-stressed



E[*VMD*_{j'}|*Z* = *NUTW*] **Disaggregate TE components for NUTW treatment**



- Individuals below the reference line generate more VMD when they do not telework
- Individuals above the reference line generate more VMD when they adopt (non-usual) teleworking
- 29.6% of travel-stressed NUTWers increase VMD after adopting non-usual TWing
- 38.3% of non-travel-stressed NUTWers increase VMD after adopting non-usual TWing/

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Summary: empirical

- We quantified and compared the impact of teleworking on vehiclemiles driven (VMD) for different types of teleworkers
 - By teleworking **frequency categories**: non-TWer, non-usual TWer, usual TWer
 - By teleworking-related **motive**: travel-stressed or not
- In all models, TWing reduced VMD on average, for its adopters
- TWing reduced VMD most, for travel-stressed TWers
- Substantially more non-travel-stressed TWers increased VMD after beginning to TW than travel-stressed TWers did



Summary: methodological

- The endogenous switching regression model (ESRM) is designed to deal with selection biases in a cross-sectional setting, where we only observe people in one state (the factual state)
- ESRMs can generate estimates of the outcome of interest as if those people were in **other states (counterfactual states)**
- Since selection biases are controlled, ESRMs generate unbiased estimates of treatment effects in a cross-sectional setting
- In doing this, the cross-sectional results *can* be consistent with the longitudinal ones



Conclusion (1)

- What are the generic ingredients of a selection bias like the ones we have been discussing?
 - Discrete intervention / experiment / treatment
 - Need to evaluate the impact of the treatment (for the treated, the untreated, and/or the population at large) on an outcome of interest
 - Cross-sectional data, containing untreated (non-adopters, those who didn't experience the intervention) and one or more types of treated groups
 - Treated and untreated cases differ from each other in unobserved ways that influence the outcome
- Generic question: could (part of) the (apparent) effect of the treatment on the outcome be due to *unobserved pre-treatment differences* between the treated and the untreated??? (E.g., differences in the *willingness to be treated*???)
 - If so, then (without correcting for the bias) you cannot assume that future treatment/ adoption (especially if becoming *mandatory*) will provide the same results

Conclusion (2)

Other transportation applications

	Treatment	Outcome	Implication of a selection bias
•	Built environment (residential self-selection)	Travel behavior (various)	People who choose a neighborhood consistent with their travel predispositions will probably travel differently from those who are nudged into it for other reasons (policy, constraints, other objectives)
	Mode choice	Satisfaction with trip	The fact that mode " x " users are the most satisfied doesn't mean that non-users would have the same satisfaction if they were to use that mode
	Voluntary safety measure (seat belt, helmet)	Accident propensity/ rate	People who opt in may be more risk averse in other ways, or may instead compensate with riskier behavior; either way, their outcomes may not match those of people forced (by law) to adopt
	Voluntary behavior change program	Travel behavior (various)	Results for people who opt in will probably not match those for people who are pushed (via policy, law, cost) to change

Conclusion (3)

- There are many other applications beyond transportation
- We very often only have cross-sectional data, so...



- We hope that our collection of papers explicating, demonstrating, and visually illustrating variants of this approach will lead to greater interest in it and adoption of it
 - Kim SH & PL Mokhtarian (2023a) Comparisons of observed and unobserved parameter heterogeneity in modeling vehicle-miles driven. *Transportation Research Part A* **172**, 103614, 2023.
 - Kim SH & PL Mokhtarian (2023b) A note on the sample selection (switching regression) model and treatment effects for a log-transformed outcome variable, in the context of residential self-selection. *Transportation*. <u>https://link.springer.com/article/10.1007/s11116-023-10384-2</u>.
 - Wang X & PL Mokhtarian (2024) Examining the treatment effect of teleworking on vehicle-miles driven: Applying an ordered probit selection model and incorporating the role of travel stress. *Transportation Research A*, in press. Available from the authors.
 - Wang X & PL Mokhtarian (2023) Reintroducing multinomial logit switching regression models: Examining the treatment effect of two teleworking frequency categories on vehicle-miles driven. Paper # TRBAM-S-23-02767 presented at the Annual Meeting of the Transportation Research Board, January 2024. Available from the authors.



Thank you! Questions?

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Happy to share the slides



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