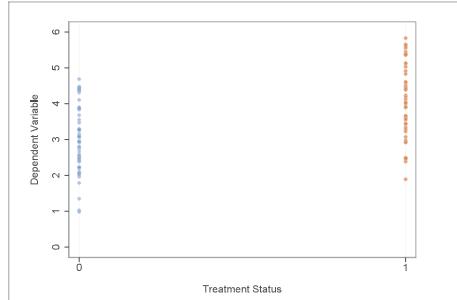




Dummy Variables

## OLS Regression on a Binary Independent Variable

$$Y = \alpha + \beta D$$



control	treatment
$D = 0$	$D = 1$
$\hat{\alpha}$	$\hat{\alpha} + \hat{\beta}$

$$\hat{\alpha} = \bar{Y}_C \text{ (control group mean)}$$

$$\hat{\beta} = \bar{Y}_T - \bar{Y}_C \text{ (difference in means)}$$

## OLS Regression on a Binary Independent Variable

You may or may not remember that in a bivariate regression:

$$\begin{aligned}\hat{\beta}_{OLS} &= \frac{\text{COV}(X, Y)}{\text{VAR}(X)} \\ &= \frac{\sum_i (X_i - \bar{X})(Y_i - \bar{Y})}{\sum_i (X_i - \bar{X})^2}\end{aligned}$$

Notice that the numerator can be re-organized:

$$\sum_i (X_i - \bar{X})(Y_i - \bar{Y}) = \sum_i X_i Y_i - \sum_i \bar{X} Y_i$$

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$$\sum_i (X_i - \bar{X})(Y_i - \bar{Y}) = \sum_i X_i Y_i - \sum_i \bar{X} Y_i$$
$$= \sum_i [Y_i (X_i - \bar{X})]$$

## OLS Regression on a Binary Independent Variable

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When independent variable is binary:  
 $\bar{X} = \frac{n_T}{N}$  ( $n_T$  is # of treated observations)

## OLS Regression on a Binary Independent Variable

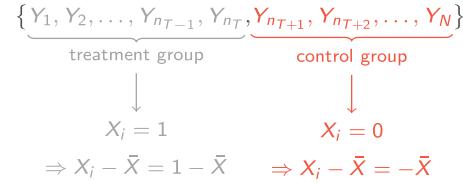
You may or may not remember that in a bivariate regression:

$$\hat{\beta}_{OLS} = \frac{COV(X, Y)}{VAR(X)} = \frac{\sum_i [Y_i(X_i - \bar{X})]}{\sum_i (X_i - \bar{X})^2}$$

When independent variable is binary:

$$\bar{X} = \frac{n_T}{N} \quad (n_T \text{ is } \# \text{ of treated observations})$$

Assume observations are ordered:



## OLS Regression on a Binary Independent Variable

You may or may not remember that in a bivariate regression:

$$\hat{\beta}_{OLS} = \frac{COV(X, Y)}{VAR(X)} = \frac{\sum_i [Y_i(X_i - \bar{X})]}{\sum_i (X_i - \bar{X})^2} = \frac{\sum_i [Y_i(X_i - \bar{X})]}{N\bar{X}(1-\bar{X})}$$

Re-write denominator:

$$\begin{aligned} \sum_i (X_i - \bar{X})^2 &= \sum_{i=1}^{n_T} (1 - \bar{X})^2 + \sum_{i=n_T+1}^N (-\bar{X})^2 \\ &= n_T (1 - \bar{X})^2 + (N - n_T) (-\bar{X})^2 \\ &= \dots = n_T - n_T \bar{X} = N \bar{X} (1 - \bar{X}) \end{aligned}$$

## OLS Regression on a Binary Independent Variable

You may or may not remember that in a bivariate regression:

$$\begin{aligned}\hat{\beta}_{OLS} &= \frac{COV(X, Y)}{VAR(X)} && \sum_i [Y_i (X_i - \bar{X})] \\ &= \frac{\sum_i [Y_i (X_i - \bar{X})]}{\sum_i (X_i - \bar{X})^2} \\ &= \frac{\sum_i [Y_i (X_i - \bar{X})]}{N\bar{X}(1 - \bar{X})} && \text{"linear combination of } Y\text{s"}$$

## OLS Regression on a Binary Independent Variable

You may or may not remember that in a bivariate regression:

$$\begin{aligned}\hat{\beta}_{OLS} &= \frac{COV(X, Y)}{VAR(X)} && \sum_i [Y_i (X_i - \bar{X})] \\ &= \frac{\sum_i [Y_i (X_i - \bar{X})]}{\sum_i (X_i - \bar{X})^2} && = \sum_{i=1}^{n_T} [Y_i (1 - \bar{X})] + \sum_{i=n_T+1}^N [Y_i (-\bar{X})] \\ &= \frac{\sum_i [Y_i (X_i - \bar{X})]}{N\bar{X}(1 - \bar{X})} && = \sum_{i=1}^{n_T} Y_i - \sum_{i=1}^N [Y_i (\bar{X})] \\ &&& = n_T \bar{Y}_T - \bar{X} (N\bar{Y}) \\ &&& = N\bar{X} \bar{Y}_T - N\bar{X} [\bar{X} \bar{Y}_T + (1 - \bar{X}) \bar{Y}_C] \\ &&& = N\bar{X} (1 - \bar{X}) (\bar{Y}_T - \bar{Y}_C)\end{aligned}$$

## OLS Regression on a Binary Independent Variable

You may or may not remember that in a bivariate regression:

$$\begin{aligned}
 \hat{\beta}_{OLS} &= \frac{COV(X, Y)}{VAR(X)} & \sum_i [Y_i (X_i - \bar{X})] \\
 &= \frac{\sum_i [Y_i (X_i - \bar{X})]}{\sum_i (X_i - \bar{X})^2} & = \sum_{i=1}^{n_T} [Y_i (1 - \bar{X})] + \sum_{i=n_T+1}^N [Y_i (-\bar{X})] \\
 &= \frac{\sum_i [Y_i (X_i - \bar{X})]}{N\bar{X}(1 - \bar{X})} & = \sum_{i=1}^{n_T} Y_i - \sum_{i=1}^N [Y_i (\bar{X})] \\
 &= \frac{N\bar{X}(1 - \bar{X})(\bar{Y}_T - \bar{Y}_C)}{N\bar{X}(1 - \bar{X})} & = n_T \bar{Y}_T - \bar{X} (N\bar{Y}) \\
 && \uparrow \\
 && = N\bar{X} \bar{Y}_T - N\bar{X} [\bar{X} \bar{Y}_T + (1 - \bar{X}) \bar{Y}_C] \\
 && \curvearrowleft = N\bar{X} (1 - \bar{X}) (\bar{Y}_T - \bar{Y}_C)
 \end{aligned}$$

## OLS Regression on a Binary Independent Variable

$$\begin{aligned}
 \hat{\beta}_{OLS} &= \frac{COV(X, Y)}{VAR(X)} \\
 &= \frac{\sum_i [Y_i (X_i - \bar{X})]}{\sum_i (X_i - \bar{X})^2} \\
 &= \frac{\sum_i [Y_i (X_i - \bar{X})]}{N\bar{X}(1 - \bar{X})} \\
 &= \frac{N\bar{X}(1 - \bar{X})(\bar{Y}_T - \bar{Y}_C)}{N\bar{X}(1 - \bar{X})} \\
 &= \bar{Y}_T - \bar{Y}_C
 \end{aligned}$$

## OLS Regression on a Binary Independent Variable

When we regress  $Y_i$  on (only) a dummy variable:

$$\hat{\beta}_{OLS} = \bar{Y}_T - \bar{Y}_C$$

- Estimated constant  $\hat{\alpha}_{OLS}$  is control group mean, also  $\hat{Y}_i$
- Predicted  $\hat{Y}_i$  for treated individuals/units is  $\hat{\alpha}_{OLS} + \hat{\beta}_{OLS}$

## OLS Regression on Mutually Exclusive Dummy Variables

$$Y = \alpha + \beta_1 T_1 + \beta_2 T_2 + \beta_3 T_3$$

control	treatment 1	treatment 2	treatment 3
$T_1 = T_2 = T_3 = 0$			

## OLS Regression on Mutually Exclusive Dummy Variables

$$Y = \alpha + \beta_1 T_1 + \beta_2 T_2 + \beta_3 T_3$$

control	treatment 1	treatment 2	treatment 3
$T_1 = T_2 = T_3 = 0$	$T_1 = 1$	$T_2 = 1$	$T_3 = 1$
$\hat{\alpha}$	$\hat{\alpha} + \hat{\beta}_1$	$\hat{\alpha} + \hat{\beta}_2$	$\hat{\alpha} + \hat{\beta}_3$

$\hat{\alpha} = \bar{Y}_C$  (control group mean)

$\hat{\beta}_i = \bar{Y}_{T_i} - \bar{Y}_C$  (difference in means between treatment  $i$  and control)

## Pooling Treatments

- If we pool treatments to estimate an average effect across treatment arms:
  - ▶ Estimated coefficient (and treatment effect)  $\beta_{pooled}$  is average of impacts across treatments
  - ▶ Average depends on  $n_{T_i}$  values: share of treated observations in each treatment arm
  - ▶ Also depends on treatment effect of each arm (pooling arms with no impact will matter)
- Estimates of pooled effect more precise because sample size is larger
  - ▶ When is pooled effect policy relevant?

## Cross-Cutting Designs

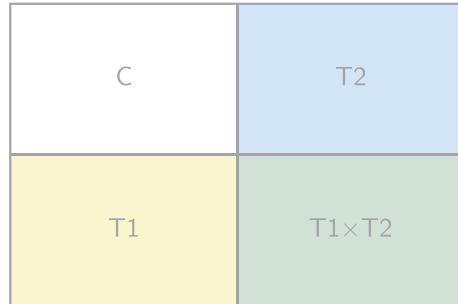
- We often want to estimate the impact of treatments that may work best together
  - ▶ Access to credit and vocational training for unemployed youth
  - ▶ Nutrition supplements and parenting education for at-risk babies and children
  - ▶ Increased enforcement and information campaigns or behavioral nudges for tax compliance
  - ▶ Teacher training and additional materials for under-performing schools
  - ▶ Management consulting and subsidies for exporting firms
- Cross-cutting designs allow us to estimate impacts of each treatment in isolation as well as the pooled impact, to see whether any program effects are additively separable

## Cross-Cutting Designs

		control	treatment 2
		C	T2
control	treatment 1	T1	T1×T2

## Cross-Cutting Designs and Interaction Terms

$$Y = \alpha + \beta_1 T_1 + \beta_2 T_2 + \beta_3 (T_1 \times T_2)$$



## Challenge Problem: Triple Interactions

$$Y = \alpha + \beta_1 T_1 + \beta_2 T_2 + \beta_3 T_3 + \gamma_1 T_1 \times T_2 + \gamma_2 T_2 \times T_3 + \gamma_3 T_1 \times T_3 + \theta T_1 \times T_2 \times T_3$$

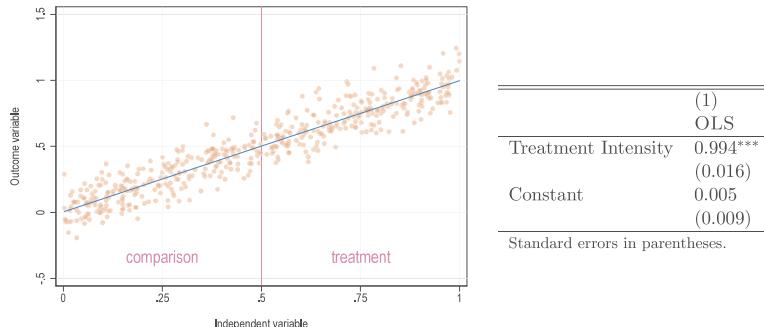
## Continuous Variables

### When Is Treatment a Continuous Variable?

- Sometimes we vary treatment intensity across treatment arms
  - ▶ Subsidies for malaria treatment (Cohen, Dupas, and Schaner 2015)
  - ▶ Varying the size of grants to entrepreneurs/firms, schools, etc.
  - ▶ Proportion treated within clusters (CCTs, job training, etc.)
- Binary treatments might also impact units differently, based on pre-existing conditions
  - ▶ Law banning traditional birth attendants in Malawi (Godlonton and Okeke 2016)
  - ▶ Impact of eliminating primary school fees on completion (cf. Lucas and Mbiti 2012)
- Should we dichotomize treatment variable or exploit continuous variation in intensity?

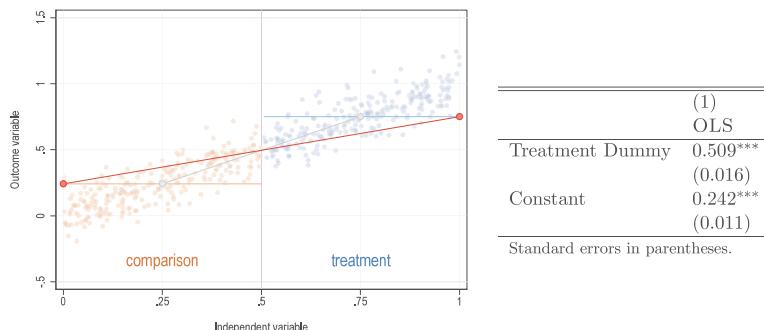
## Continuous Variation in Treatment

$$Y_i = X_i + \varepsilon$$



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$$Y_i = X_i + \varepsilon$$

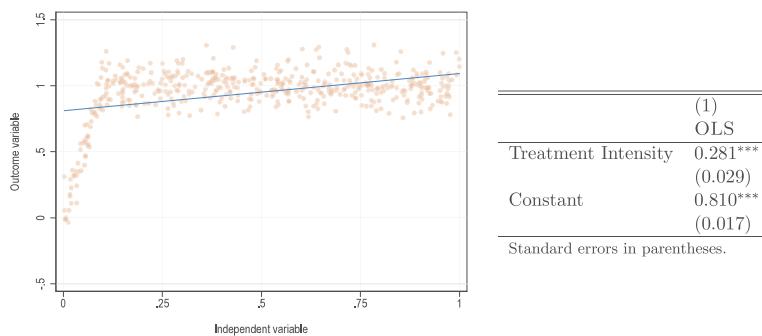


## Dichotomous vs. Continuous Treatment Variables

- When the true dose-response relationship is linear:
  - ▶ OLS w/ a continuous treatment variable is correctly specified
  - ▶ Uses observed variation increase statistical power (odds of finding an effect)
- The estimand is different when we dichotomize treatment
  - ▶ OLS coefficient captures impact of moving from average level of treatment intensity in the control group to average level of treatment intensity in the treatment group
    - ▶ Not the same as impact of moving from treatment intensity 0 to treatment intensity 1
- When true dose-response relationship is linear, OLS with continuous  $X$  is optimal

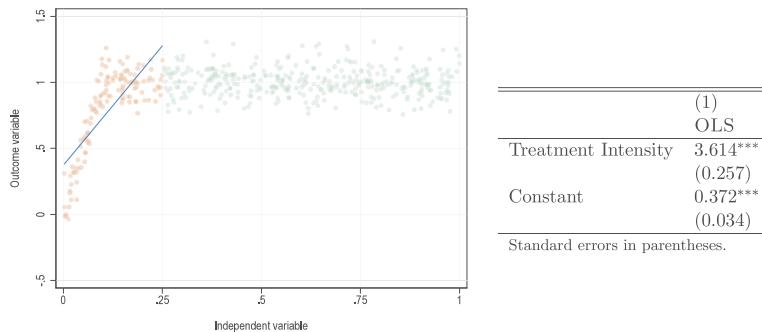
Economics 523 (Professor Jakielo)      Regression, Slide 50

## OLS when the Dose-Response Relationship Is Not Linear



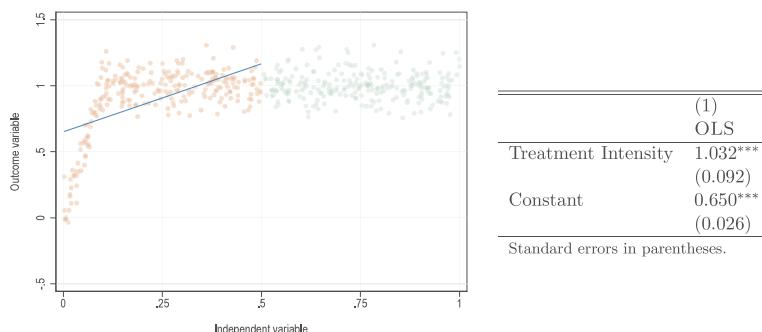
Economics 523 (Professor Jakielo)      Regression, Slide 52

## OLS when the Dose-Response Relationship Is Not Linear



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## OLS when the Dose-Response Relationship Is Not Linear



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## Continuous Variation in Treatment Intensity

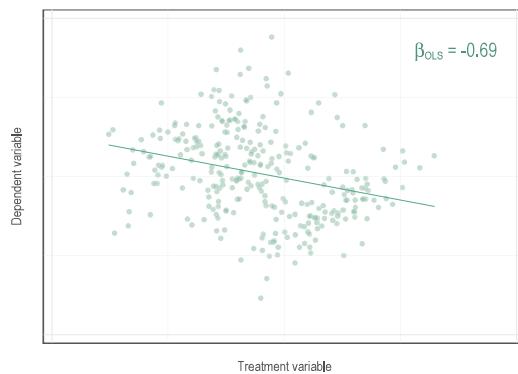
- When the true relationship (i.e. conditional expectation function) is **not** linear:
  - ▶ OLS w/ a continuous treatment variable is incorrectly specified (may or may not matter)
  - ▶ Estimated treatment effect (i.e. coefficient) depends on choice of sample (values of  $X$ )
- Graph your data (though often true relationship obscured by noise)
  - ▶ Choose to dichotomize (and where to dichotomize)
  - ▶ Vary your sample to assess the robustness of your estimates
- Be skeptical of results when treatment assignment process is unclear (observational data!) and you cannot observe the relevant empirical relationships in your data graphically

## Fixed Effects

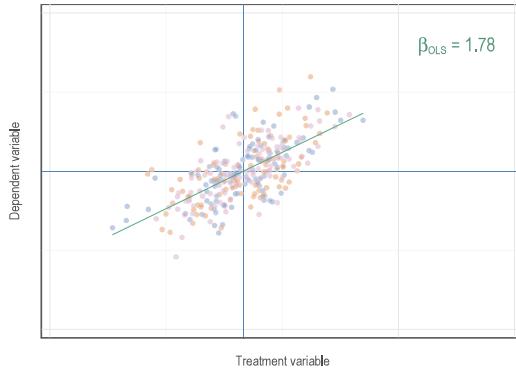
## What Are Fixed Effects?

- Individual dummy variables for mutually exclusive groups in your data
  - ▶ Dummy for male or female
  - ▶ Age (or age group) fixed effects
  - ▶ Continent/country/state/district fixed effects
  - ▶ Year fixed effects
- Why use fixed effects?
  - ▶ Estimation using **within** rather than **between** variation
- We often use multiple sets of fixed effects in empirical work

## Simpson's Paradox



## What Do Fixed Effects Do?



## What Do Fixed Effects Do?

- Fixed effects are equivalent to:
  - ▶ Transforming both independent and dependent variables by subtracting off the mean in each group and adding back the mean in the omitted category (the blue group in the figure)
  - ▶ Equivalently: subtract off the difference in means between group and omitted group
  - ▶ Run OLS in your transformed (i.e. re-centered) data
- Fixed effects mattered because treatment varied across groups
  - ▶ When treatment doesn't vary, FEs can improve precision but won't change slope estimate
  - ▶ When randomized/exogenous treatment probability/intensity varies across groups, you must include fixed effects or control directly for probability of treatment (the propensity score)
- If you regress  $X$  and  $Y$  on FEs, de-meanned variables are the residuals

## The Frisch-Waugh-Lovell Theorem

$$Y = \alpha + \beta X + \gamma Z$$

is equivalent to

$$\tilde{Y} = \alpha + \beta \tilde{X}$$

where

$\tilde{Y}$  = residuals from regressing  $Y$  on  $Z$

$\tilde{X}$  = residuals from regressing  $X$  on  $Z$

## Frisch-Waugh-Lovell: Why It Matters

- When treatment is binary and plausibly exogenous, the difference in outcome means between the treatment and comparison groups provides an unbiased estimate of impact
  - ▶ All treated observations get equal positive weight, all untreated get equal negative weight
- With controls that are correlated with treatment, treatment is (in effect) no longer binary
  - ▶ Untreated observations with covariates that predict a high likelihood of treatment get very low negative weights in linear regression; while treated observations with covariates that predict a low likelihood treatment get very high positive weights in multivariate regression
  - ▶ Everything is still fine if treatment effects are homogeneous: treatment effect is the same for everyone, so it doesn't matter what weights we use to calculate average treatment effect
  - ▶ If effects vary with covariates that predict treatment, mis-specification problems arise
- What to do: show results with and without covariates, residualize and then plot your data

## Empirical Exercise