

*A review of modern methods
of estimating the size of health disparities*

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Emil Coman¹
Helen Wu²

¹ [UConn Health Disparities Institute](#), ² UConn Health

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Health Disparities (HD):
It's just about comparing two groups

Goals

1. Simplify and reposition common analytic methods
2. Compare methods to estimate HDs
3. Suggest cross-pollinations
4. Encourage disparities investigations

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One intuitive model of Health Disparities

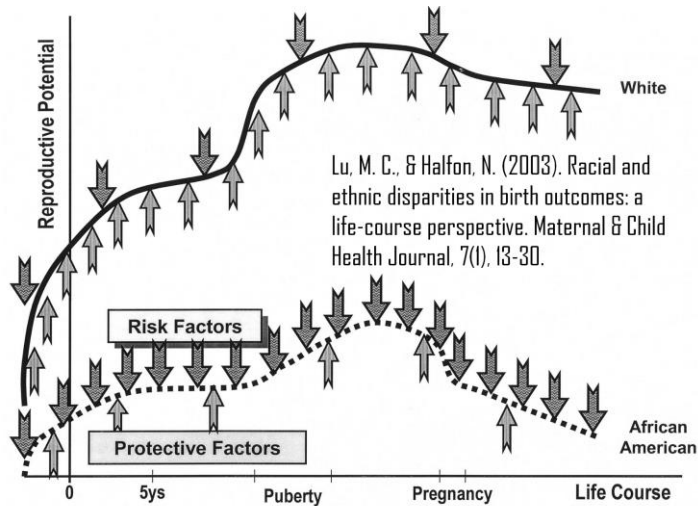


Fig. 1. How differential exposures to risk factors (downward arrows) and protective factors (upward arrows) over the life course affect developmental trajectories and contribute to disparities in birth outcomes. The lower reproductive potential of African American women, relative to White women, results from their cumulative exposure to more risk factors and less protective factors across the life span, particularly during sensitive periods of development.

Lu, M. C., & Halfon, N. (2003). Racial and ethnic disparities in birth outcomes: a life-course perspective. *Maternal & Child Health Journal*, 7(1), 13-30.

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One intuitive model of Health Disparities

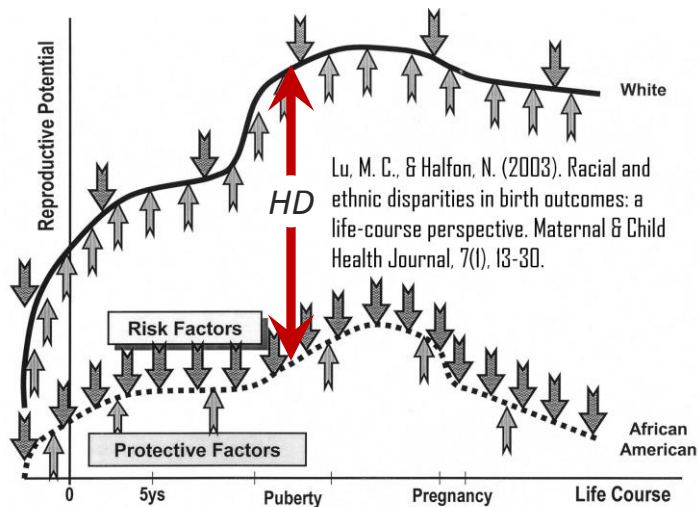
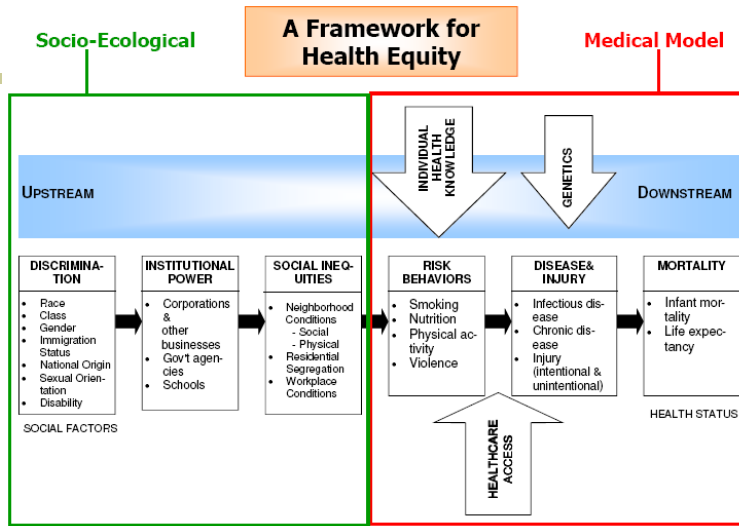


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Another intuitive model of HDs



- Adapted by ACPHD from the Bay Area Regional Health Inequities Initiative, Summer 2008

Tony Iton "Framework for Health Equity", referenced in the text: *Tackling Health Inequities Through Public Health Practice: Theory to Action*, Edited by Richard Hofrichter and Rajiv Bhatia. Based on a project by the National Association of County and City Health Officials. Oxford University Press, 2010 (page 380).
 Iton, A., & Shrimali, B. P. (2016). Power, Politics, and Health: A New Public Health Practice Targeting the Root Causes of Health Equity. *Maternal and Child Health Journal*, 20(8), 1753-1758. doi:10.1007/s10995-016-1980-6

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A testable model of HDs



Caveats:

1. Causality: can race 'cause' health?
2. Compare the comparable
3. Control for 'covariates'

VanderWeele T.J., & Hernán MA. (2012). Causal effects and natural laws: towards a conceptualization of causal counterfactuals for non-manipulable exposures with application to the effects of race and sex. In Berzuini C, Dawid P, & Bernardinelli L (Eds.), *Causality: Statistical Perspectives and Applications* (pp. 101–113). West Sussex, UK John Wiley & Sons.
 VanderWeele, T. J., & Robinson, W. R. (2014). On the causal interpretation of race in regressions adjusting for confounding and mediating variables. *Epidemiology*, 25(4), 473-484.

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Modeling options for HDs

1. Independent samples t-test
2. Anova
3. Regression
4. Instrumental Variable regression
5. SEM
6. Matching methods
7. +

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Modeling options for HDs

1. Independent samples T-test is a 2-group model

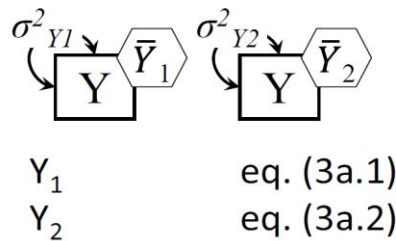


Figure 3a: The t-test model

Note: The independent samples t-test is testing the hypothesis: $\bar{Y}_1 = \bar{Y}_2$; this two-group setup allows for inclusion of group specific covariates; the two equations are simply one variable for each group, but across-group constraints are possible, like $\sigma^2_{Y1} = \sigma^2_{Y2}$.

Modeling options for HDs

1. Independent samples

T-test is a 2-group model

```
ttest y, by(binary)
pwmean y, over(binary) effects
```

2. Anova – similar for 2 groups

```
anova y binary
```

➤ Allows for covariates though

```
anova y binary c.x1 c.x2
```

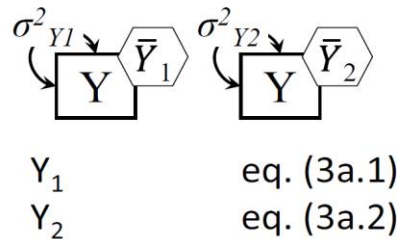


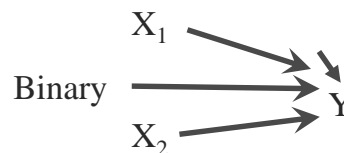
Figure 3a: The t-test model

Note: The independent samples t-test is testing the hypothesis: $\bar{Y}_1 = \bar{Y}_2$; this two-group setup allows for inclusion of group specific covariates; the two equations are simply one variable for each group, but across-group constraints are possible, like $\sigma^2_{Y1} = \sigma^2_{Y2}$.

Modeling options for HDs

3. Regression

```
reg y binary x1 x2
```

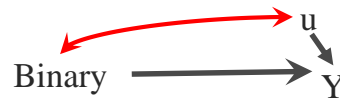


But: ? Can race cause health?

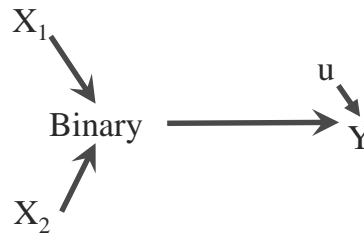
Modeling options for HDs

4. IV regression [Cameron]

➤ Binary **correlated** with u



➤ Need IV \rightarrow X but not IV \rightarrow Y



```
ivregress 2sls y
(binary = x1 x2), first
```

➤ Tests the IVs:

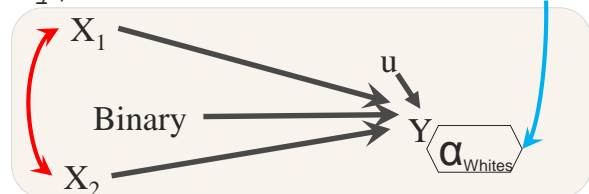
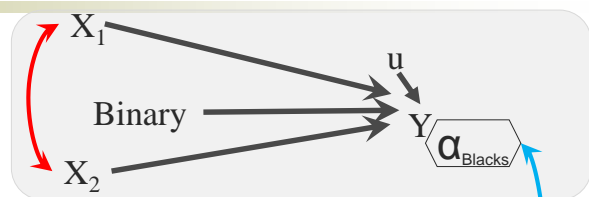
```
estat endogenous
```

Modeling options for HDs

5. SEM

2-group model, compare intercepts (needs centering)

```
sem y <- x1 x2, group(binary)
ginvariant(covex)
```



Test Δ

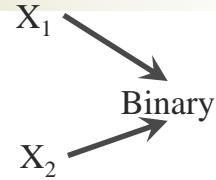
Can relax the equal Y variances assumption

Modeling options for HDs

6. Matching options

```
teffects psmatch (y)
(binary x1 x2 )
```

Stage 1

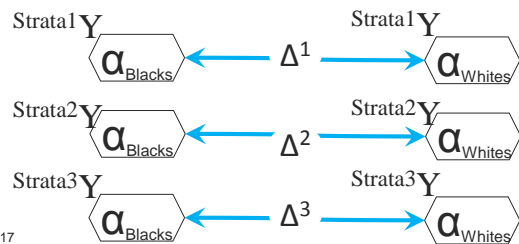


```
psmatch2 binary x1 x2 ,
outcome (y) common ate
```

Stage 2

Match in strata on $\widehat{\text{Binary}}$

Stage 3



```
attnd y binary x1 x2
```

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Modeling options for HDs

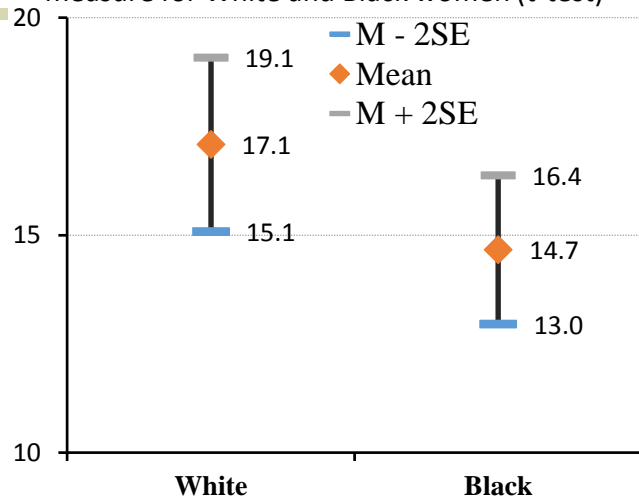
1. T-test

2. Anova – identical

➤ Allows for covariates though

```
anova y binary c.x1 c.x2
```

Means (+/- 2 SE) of the anxiety baseline measure for White and Black women (t-test)



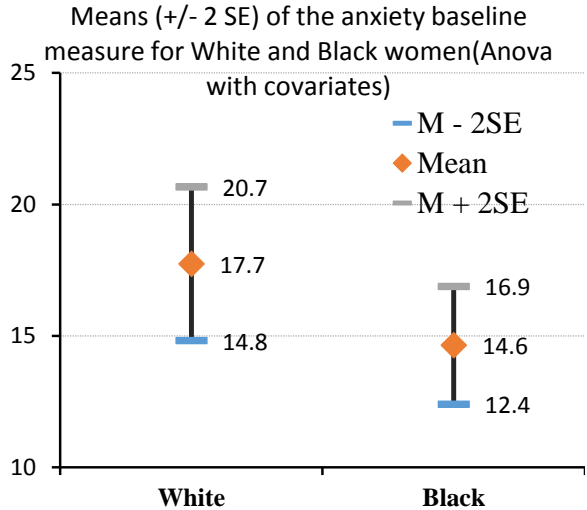
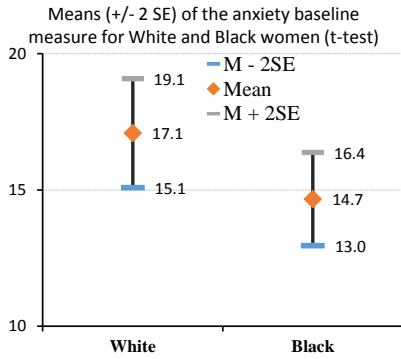
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Modeling options for HDs

T-test

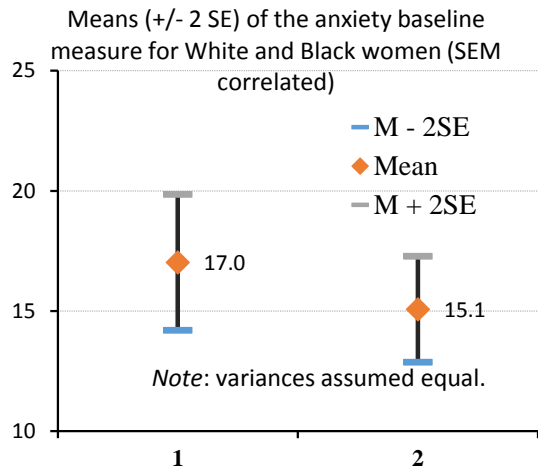
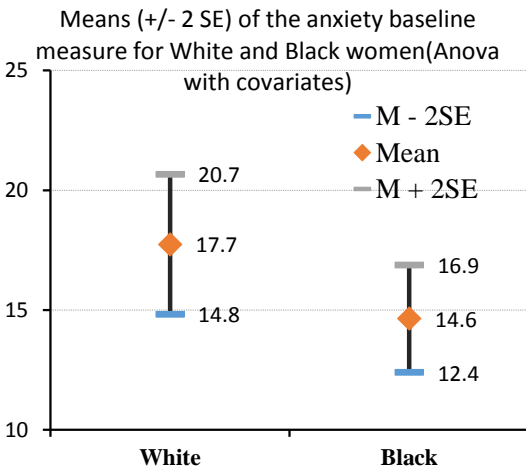
Anova – with covariates



Modeling options for HDs

Anova – with covariates

SEM with *correlates*



Modeling options for HDs

SEM with *covariates* [0=White; 1=Black]; variances freed

```
sem y <- x1 x2, group(binary) ginvariant(covex)
      |
      |      OIM
      |      Coef.  Std. Err.    z    P>|z|    [95% Conf. Interval]
-----+-----
Structural
y <-
  x1
  0 |   .0199704   2.16228    0.01  0.993   -4.21802   4.257961
  1 |   2.61248   1.673379    1.56  0.118   -0.6672832  5.892244
  x2
  0 |   .164855   .3766247    0.44  0.662   -0.5733159  .9030258
  1 |   .4905938   .3001652    1.63  0.102   -0.0977191  1.078907
  _cons
  0 |  13.43964   8.259071    1.63  0.104   -2.747845  29.62712
  1 |  3.422483   6.895766    0.50  0.620   -10.09297  16.93794
-----+-----
var(e.y)
  0 |   60.78133  13.59112           39.21351   94.21166
  1 |   84.67747  14.74046           60.19947   119.1086
-----means centered at .050 (X1 neighborhood disorder) and 22.91y -----
```

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Modeling options for HDs

teffects psmatch

```
. teffects psmatch (y) (binary x1 x2 )
```

```
Treatment-effects estimation      Number of obs      =      106
Estimator      : propensity-score matching      Matches: requested =      1
Outcome model  : matching                      min =      1
Treatment model: logit                        max =      1
```

```
      |
      |      AI Robust
      |      Coef.  Std. Err.    z    P>|z|    [95% Conf. Interval]
-----+-----
ATE
  binary
  (1 vs 0) |  -3.169811  2.228166   -1.42  0.155   -7.536936   1.197313
-----+-----
```

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Modeling options for HDs

psmatch2

```

psmatch2 binary x1 x2 , outcome (y) common ate Probit regression
Number of obs = 106

LR chi2(2) = 12.08
Prob > chi2 = 0.0024
Pseudo R2 = 0.0860
Log likelihood = -64.210562

-----+-----
      binary |      Coef.   Std. Err.      z    P>|z|     [95% Conf. Interval]
-----+-----
      x1 |   .6428318   .2054421    3.13   0.002   .2401726   1.045491
      x2 |   .0468275   .0366658    1.28   0.202  -.0250361   .1186911
      _cons |  -.7222502   .8205248   -0.88   0.379  -2.330449   .8859489

psmatch2: |   psmatch2: Common
Treatment |           support
assignment | Off suppo  On suppor |      Total
-----+-----
Untreated |           0         40 |         40
Treated   |          11         55 |         66
-----+-----
Total    |          11         95 |        106

```

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Modeling options for HDs

psmatch2

```

psmatch2 binary x1 x2 , outcome (y) common ate Probit regression
Number of obs = 106

-----+-----
Variable      Sample |   Treated   Controls   Difference      S.E.   T-stat
-----+-----
      y  Unmatched | 15.0757576   17.025   -1.94924242   1.8091337   -1.08
           ATT |      13.8   18.9272727   -5.12727273   2.19360704   -2.34
           ATU |      17.025   12.5   -4.525           .           .
           ATE |           .   .   -4.87368421           .           .
-----+-----

```

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Modeling options for HDs

attnd

```
attnd y binary x1 x2 , comsup boot reps(100) dots logit detail
```

Note: the common support option has been selected

The region of common support is [.30252502, .89642859]

The distribution of the pscore is

Pr(binary)

```
-----
Percentiles      Smallest
1%               .3173758       .302525
5%               .3990624       .3173758
10%              .4397769       .3197376      Obs               141
25%              .4943597       .3332007      Sum of Wgt.      141

50%              .6010811
75%              .7131706       Largest
90%              .7813184       .854728
95%              .8165104       .8644204      Mean              .6028369
99%              .8644204       .8964286      Std. Dev.         .1374292
                                Variance          .0188868
                                Skewness         -.0160976
                                Kurtosis         2.2178
```

The program is searching the nearest neighbor of each treated unit.

This operation may take a while.

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Modeling options for HDs

attnd

```
attnd y binary x1 x2 , comsup boot reps(100) dots logit detail
```

ATT estimation with Nearest Neighbor Matching method

(random draw version)

Analytical standard errors

```
-----
n. treat.   n. contr.   ATT   Std. Err.   t
-----
           85           23   -0.524   2.008   -0.261
-----
```

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Modeling options for HDs

ivregress

```
ivregress 2sls y (binary = x1 x2 ), first
```

First-stage regressions

		Number of obs =		106	
		F(2, 103) =		6.05	
		Prob > F =		0.0033	
		R-squared =		0.1051	
		Adj R-squared =		0.0878	
		Root MSE =		0.4652	

binary	Coef.	Std. Err.	t	P> t	[95% Conf. Interval]	
x1	.2176514	.0672757	3.24	0.002	.0842259	.3510769
x2	.0148822	.0124655	1.19	0.235	-.0098403	.0396046
_cons	.2805711	.281939	1.00	0.322	-.2785883	.8397305

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Modeling options for HDs

ivregress

```
ivregress 2sls y (binary = x1 x2 ), first
```

Instrumental variables (2SLS) regression

Number of obs	=	106
Wald chi2(1)	=	1.57
Prob > chi2	=	0.2098
R-squared	=	.
Root MSE	=	10.126

y	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
binary	7.847662	6.257221	1.25	0.210	-4.416267	20.11159
_cons	10.92504	4.018222	2.72	0.007	3.04947	18.80061

Instrumented: binary
Instruments: x1 x2

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Modeling options for HDs

ivregress

estat endogenous

Tests of endogeneity

H₀: variables are exogenous

Durbin (score) chi2(1) = 3.51188 (p = 0.0609)

Wu-Hausman F(1,103) = 3.52942 (p = 0.0631)

"If the test statistic is significant, the variables must be treated as endogenous" [URL](#)

estat overid

Tests of overidentifying restrictions:

Sargan (score) chi2(1) = .872447 (p = 0.3503)

Basmann chi2(1) = .854791 (p = 0.3552)

"A statistically significant test statistic indicates that the instruments may not be valid" [URL](#)

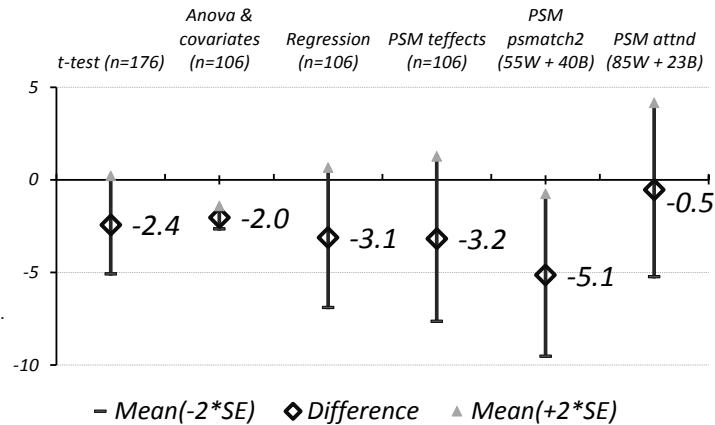
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Modeling options for HDs

Comparisons

Comparison of racial/ethnic (R/E) differences (92 White- 145 Black) in Anxiety baseline scores, by estimation method

All models used age and neighborhood stress as: covariates (regression); propensity predictors (PSM); instruments for R/E (IV).



Some methods are 'better' than others in identifying health disparities.

SEM is not shown, the test of HD is a Wald (or chi-squared test).

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[Conclusions – email for >: comanus@gmail.com]

- Some methods are ‘better’ than others in identifying health disparities.
- Comparisons encourage analysts to think about ‘the place’ of the variables in the analytical model.
- One general insight: we can trust estimates when that are somewhat consistent across methods.