#### Social disadvantage and infant mortality:

the birth weight paradox revisited

Bianca De Stavola

with Rhian Daniel, Richard Silverwood, Rachel Stuchbury, Emily Grundy

UK Causal Inference Meeting  $\cdot$  28 – 29 April 2014





#### Infant mortality (deaths <1 yr):

- negatively related to birth weight (BW)
- patterned by socio-economic conditions.



#### Infant mortality (deaths <1 yr):

- negatively related to birth weight (BW)
- patterned by socio-economic conditions.

#### Complication:

low BW babies in high-risk populations tend to have lower mortality rates than low BW babies in low-risk populations.

・ロト ・ 同ト ・ ヨト ・ ヨト … ヨ



#### Infant mortality (deaths <1 yr):

- negatively related to birth weight (BW)
- patterned by socio-economic conditions.

Complication:

low BW babies in high-risk populations tend to have lower mortality rates than low BW babies in low-risk populations.

First observed by Yerushalmy (1964, 1971) and interpreted as BW modifying the effect of many factors associated with infant mortality:

#### **BW** paradox





- Smoking known risk factor for low BW.
- Low BW babies born to smokers lower mortality than those of non-smokers:



Figure: Birth-weight-specific infant mortality curves, US, 1991 (Hernandez-Diaz, AJE 2006)



#### 1 Background

- 2 An alternative model
- 3 Questions and estimands
- 4 Preliminary results
- 5 Critique and Conclusions

イロン 不得 とくほ とくほう 一日



- BW is on the causal pathway from "Disadvantage" (*E*) to "Infant death", but there are unmeasured confounders *U*<sub>1</sub>.
- Comparing infant mortality rates at given values of BW leads to opening up a spurious path from E to "Infant death" (Hernandez-Diaz et al., 2006).
- Paradox explained if U<sub>1</sub> and U<sub>2</sub> act in opposite directions (Basso et al., 2006 & CONT)





- BW is on the causal pathway from "Disadvantage" (*E*) to "Infant death", but there are unmeasured confounders *U*<sub>1</sub>.
- Comparing infant mortality rates at given values of BW leads to opening up a spurious path from E to "Infant death" (Hernandez-Diaz et al., 2006).
- Paradox explained if U<sub>1</sub> and U<sub>2</sub> act in opposite directions (Basso et al., 2006 & COON)





- BW is on the causal pathway from "Disadvantage" (*E*) to "Infant death", but there are unmeasured confounders *U*<sub>1</sub>.
- Comparing infant mortality rates at given values of BW leads to opening up a spurious path from E to "Infant death" (Hernandez-Diaz et al., 2006).
- Paradox explained if U<sub>1</sub> and U<sub>2</sub> act in opposite directions (Basso et al., 2006 & COON)





- BW is on the causal pathway from "Disadvantage" (*E*) to "Infant death", but there are unmeasured confounders *U*<sub>1</sub>.
- Comparing infant mortality rates at given values of BW leads to opening up a spurious path from E to "Infant death" (Hernandez-Diaz et al., 2006).
- Paradox explained if U<sub>1</sub> and U<sub>2</sub> act in opposite directions (Basso et

al. , 2006 & 2000)





- It is only a proxy of intrauterine growth rate and time,
- neither intrauterine dimensions are usually available in large observational studies.
- Other pathways may link exposure to the infant mortality (hence the added arrows).





- It is only a proxy of intrauterine growth rate and time,
- neither intrauterine dimensions are usually available in large observational studies.
- Other pathways may link exposure to the infant mortality (hence the added arrows).





- It is only a proxy of intrauterine growth rate and time,
- neither intrauterine dimensions are usually available in large observational studies.
- Other pathways may link exposure to the infant mortality (hence the added arrows).





- It is only a proxy of intrauterine growth rate and time,
- neither intrauterine dimensions are usually available in large observational studies.
- Other pathways may link exposure to the infant mortality (hence the added arrows).

But how can we proceed without information on intrauterine growth?





Wilcox (1983,2001) suggested that there are two sub-populations of newborns:

- (a) predominant: mostly term babies,
- (b) compromised: mostly pre-term babies and small-for-gestational-age.



# Background Alternative Questions Preliminary results Conclusions Reformulated alternative model



- The model can be reformulated in terms of these classes.
- Assuming that the birth weight distribution for each sub-population is normal,
- and including predictors, we can estimate
  - Prob(class = compromised) using Latent Class Modelling.



イロン 不得 とくほ とくほう 一日

### Background Alternative Questions Preliminary results Conclusions Reformulated alternative model



#### The model can be reformulated in terms of these classes.

- Assuming that the birth weight distribution for each sub-population is normal,
- and including predictors, we can estimate Prob(class = compromised) using Latent Class Modelling.





- The model can be reformulated in terms of these classes.
- Assuming that the birth weight distribution for each sub-population is normal,
- and including predictors, we can estimate Prob(class = compromised) using Latent Class Modelling.





- The model can be reformulated in terms of these classes.
- Assuming that the birth weight distribution for each sub-population is normal,
- and including predictors, we can estimate
   Prob(class = compromised) using Latent Class Modelling.



イロン 不得 とくほ とくほう 一日



With this more general theoretical framework, we reconsider the two main questions. Is BW:

- 1 an effect modifier of the effect of "Disadvantage" on Infant mortality?
- 2 a mediator for the effect of "Disadvantage" on Infant mortality?



With this more general theoretical framework, we reconsider the two main questions. Is BW:

- 1 an effect modifier of the effect of "Disadvantage" on Infant mortality?
- 2 a mediator for the effect of "Disadvantage" on Infant mortality?

▲□▶ ▲□▶ ▲□▶ ▲□▶ □ のへの



- BW: potential mediator (*M*); "Disadvantage": exposure (*E*); Infant mortality: outcome (*Y*); "Intrauterine growth": intermediate confounder (*L*).
- **Replacing** *L* with  $\hat{L} = \Pr(L = 1)$  (1: compromised, 0: predominant),





- BW: potential mediator (*M*); "Disadvantage": exposure (*E*); Infant mortality: outcome (*Y*); "Intrauterine growth": intermediate confounder (*L*).
- **Replacing** L with  $\hat{L} = \Pr(L = 1)$  (1: compromised, 0: predominant),



(日) (同) (E) (E) (E)



#### We address the first question:

- by comparing Controlled Direct Effect of *E* on *Y* holding *M* at either 0 or 1.
- If these effects are similar there is no support for effect modification by M.





- We address the first question:
- by comparing Controlled Direct Effect of *E* on *Y* holding *M* at either 0 or 1.
- If these effects are similar there is no support for effect modification by M.



(注) (注) (注) (注) (○)



- We address the first question:
- by comparing Controlled Direct Effect of *E* on *Y* holding *M* at either 0 or 1.
- If these effects are similar there is no support for effect modification by M.



(注) (注) (注) (注) (○)

Bianca De Stavola/BW Paradox



- We address the second question:
- by estimating the Natural Direct and Indirect Effects of *E* on *Y*, where:
  - the indirect effect is made of (a)
  - and (b),
  - m and the direct effect is (c):





- We address the second question:
- by estimating the Natural Direct and Indirect Effects of E on Y, where:
  - the indirect effect is made of (a)
  - and (b)
  - and the direct effect is (c):





- We address the second question:
- by estimating the Natural Direct and Indirect Effects of E on Y, where:
  - the indirect effect is made of (a)
    and (b),
    and the direct effect is (c):





- We address the second question:
- by estimating the Natural Direct and Indirect Effects of E on Y, where:
  - the indirect effect is made of (a)
  - and (b),

and the direct effect is (c):





- We address the second question:
- by estimating the Natural Direct and Indirect Effects of E on Y, where:
  - the indirect effect is made of (a)
  - and (b),
  - and the direct effect is (c):





# Estimands (CDE(m) and PNDE, TNIE) are expressed as OR contrasts.

Assumptions:

No interference, consistency, conditional exchangeability, and, because of L, either:

- No *E*-*M* interaction: Model | (Robins and Greenland, 1992).
- No non-linearities in L:Model II (Petersen et al., 2006).

Estimation:

- via Monte Carlo G-computation (Daniel, et al, 2011),
- accounting for the estimation of Pr(L = 1) and clustering of children.



Estimands (CDE(m) and PNDE, TNIE) are expressed as OR contrasts.

Assumptions:

No interference, consistency, conditional exchangeability, and, because of L, either:

■ No *E*-*M* interaction: Model | (Robins and Greenland, 1992).

■ No non-linearities in L:Model II (Petersen et al. , 2006).

Estimation:

- via Monte Carlo G-computation (Daniel, et al, 2011),
- accounting for the estimation of Pr(L = 1) and clustering of children.

イロン 不得 とくほ とくほう 一日



Estimands (CDE(m) and PNDE, TNIE) are expressed as OR contrasts.

Assumptions:

No interference, consistency, conditional exchangeability, and, because of L, either:

■ No *E*-*M* interaction: Model | (Robins and Greenland, 1992).

■ No non-linearities in L:Model II (Petersen et al., 2006).

Estimation:

- via Monte Carlo G-computation (Daniel, et al, 2011),
- accounting for the estimation of Pr(L = 1) and clustering of children.

▲□▶ ▲□▶ ▲□▶ ▲□▶ □ のへの

#### Background Alternative Questions Preliminary results Conclusions The ONS Longitudinal Study (ONS LS)



- Record linkage study set up in 1974 (see http://celsius.lshtm.ac.uk/).
- Comprises linked census and event (and thus infant mortality<sup>1</sup>) records for 1% of the population of England and Wales (about 500,000 people at any one census).
- Includes BW of babies born to LS mothers (regularly since 1981, recorded at registration).
- Several indicator of social disadvantage: here we show results for maternal education
- Today: data restricted to births of white mothers (85%), with complete information on maternal education (loss of 3.8%).

(Data only available at a dedicated lab at the Office for National Statistics, all results vetted before release.)



- Record linkage study set up in 1974 (see http://celsius.lshtm.ac.uk/).
- Comprises linked census and event (and thus infant mortality<sup>1</sup>) records for 1% of the population of England and Wales (about 500,000 people at any one census).
- Includes BW of babies born to LS mothers (regularly since 1981, recorded at registration).
- Several indicator of social disadvantage: here we show results for maternal education
- Today: data restricted to births of white mothers (85%), with complete information on maternal education (loss of 3.8%).

(Data only available at a dedicated lab at the Office for National Statistics, all results vetted before release.)

## Background Alternative Questions Preliminary results Conclusions The study population



- 160,366 singleton live births in 1981-2011.
- *E*: 38% of mother with fewer that 5 O-levels ("Low education").
- M: 5.3% with birth weight < 2.5kg.
- Y: 0.54% (862) infant deaths.
- Mortality rates vary greatly by BW, moderately by sex, improving with calendar time:

## Background Alternative Questions Preliminary results Conclusions The study population



- 160,366 singleton live births in 1981-2011.
- **E**: 38% of mother with fewer that 5 O-levels ("Low education").
- *M*: 5.3% with birth weight<2.5kg.
- Y: 0.54% (862) infant deaths.
- Mortality rates vary greatly by BW, moderately by sex, improving with calendar time:



- 160,366 singleton live births in 1981-2011.
- **E**: 38% of mother with fewer that 5 O-levels ("Low education").
- *M*: 5.3% with birth weight<2.5kg.
- Y: 0.54% (862) infant deaths.
- Mortality rates vary greatly by BW, moderately by sex, improving with calendar time:



- 160,366 singleton live births in 1981-2011.
- **E**: 38% of mother with fewer that 5 O-levels ("Low education").
- *M*: 5.3% with birth weight<2.5kg.
- Y: 0.54% (862) infant deaths.
- Mortality rates vary greatly by BW, moderately by sex, improving with calendar time:

イロン 不得 とくほ とくほう 一日

### Background Alternative Questions Preliminary results Conclusions The study population



- 160,366 singleton live births in 1981-2011.
- E: 38% of mother with fewer that 5 O-levels ("Low education").
- M: 5.3% with birth weight<2.5kg.
- Y: 0.54% (862) infant deaths.
- Mortality rates vary greatly by BW, moderately by sex, improving with calendar time:



Background Alternative Questions Preliminary results Conclusions

Natural direct and indirect effects of low maternal education

VERY PRELIMINARY RESULTS- SEs not yet corrected

	Model I		Model II	
	ln OR	(SE)	ln OR	(SE)
CDE(0)	_	_	0.205	(0.076)
CDE(1)	-	-	0.206	(0.076)
PNDE TNIE	<mark>0.221</mark> 0.011	(0.082) (0.007)	<mark>0.227</mark> -0.012	(0.077) (0.005)

#### **TCE** 0.232 (0.082) 0.205 (0.076)

- Model I and II give similar results, despite the difference in assumptions.
- CDE(0) and CDE(1) from Model II are very similar: no evidence of effect modification.
- There is little support for a mediating effect of BW (also supported by sensitivity analyses).
- However problems of stability of the results.

Background Alternative Questions Preliminary results Conclusions

Natural direct and indirect effects of low maternal education

VERY PRELIMINARY RESULTS- SEs not yet corrected

	Model I		Model II	
	ln OR	(SE)	ln OR	(SE)
<b>CDE(0)</b>	_	_	0.205	(0.076)
<b>CDE(1)</b>	-	_	0.206	(0.076)
PNDE	0.221	(0.082)	0.227	(0.077)
TNIE	0.011	(0.007)	-0.012	(0.005)

**TCE** 0.232 (0.082) 0.205 (0.076)

- Model I and II give similar results, despite the difference in assumptions.
- CDE(0) and CDE(1) from Model II are very similar: no evidence of effect modification.
- There is little support for a mediating effect of BW (also supported by sensitivity analyses).
- However problems of stability of the results.

# Background Alternative Questions Preliminary results Conclusions Critique



#### What about unmeasured confounders?

- Results would still be biased.
- However, not if  $U_1$  and  $U_2$  influenced L directly.



イロン 不得 とくほ とくほう 一日

# Background Alternative Questions Preliminary results Conclusions Critique



#### What about unmeasured confounders?

- Results would still be biased.
- However, not if  $U_1$  and  $U_2$  influenced L directly.



▲□▶▲□▶▲□▶▲□▶ □ のQ@



- Approach may contribute to the debate about the BW paradox by representing the underlying biological process via a latent variable.
- Results depends on strong and partly unverifiable assumptions, although similarity of results from alternative parametric specifications are reassuring.
- Estimation of mediation effects and their SEs raises several problems. There are issues with:
  - estimation of the class probability,
  - correlations among the outcomes of siblings,
  - instability due to small number of events.
- These are being addressed by extending the Monte Carlo G-formula algorithm.



- Approach may contribute to the debate about the BW paradox by representing the underlying biological process via a latent variable.
- Results depends on strong and partly unverifiable assumptions, although similarity of results from alternative parametric specifications are reassuring.
- Estimation of mediation effects and their SEs raises several problems. There are issues with:
  - estimation of the class probability,
  - correlations among the outcomes of siblings,
  - instability due to small number of events.
- These are being addressed by extending the Monte Carlo G-formula algorithm.



- Approach may contribute to the debate about the BW paradox by representing the underlying biological process via a latent variable.
- Results depends on strong and partly unverifiable assumptions, although similarity of results from alternative parametric specifications are reassuring.
- Estimation of mediation effects and their SEs raises several problems. There are issues with:
  - estimation of the class probability,
  - correlations among the outcomes of siblings,
  - instability due to small number of events.
- These are being addressed by extending the Monte Carlo G-formula algorithm.



- Approach may contribute to the debate about the BW paradox by representing the underlying biological process via a latent variable.
- Results depends on strong and partly unverifiable assumptions, although similarity of results from alternative parametric specifications are reassuring.
- Estimation of mediation effects and their SEs raises several problems. There are issues with:
  - estimation of the class probability,
  - correlations among the outcomes of siblings,
  - instability due to small number of events.
- These are being addressed by extending the Monte Carlo G-formula algorithm.



This work is supported by the ESRC Pathways Node (Award ES/1025561/2) of the National Centre for Research Methodology.

The permission of the Office for National Statistics to use the Longitudinal Study is gratefully acknowledged, as is the help provided by staff of CeLSIUS.

CeLSIUS is supported by the ESRC Census of Population Programme (Award Ref: ES/K000365/1).

The authors alone are responsible for the interpretation of the data.

Census output is Crown copyright and is reproduced with the permission of the Controller of HMSO and the Queen's Printer for Scotland.

・ロト ・ 同ト ・ ヨト ・ ヨト … ヨ

#### Background Alternative Questions Preliminary results Conclusions References

- Basso O, Wilcox AJ, Weimberg CR. Birth Weight and Mortality: Causality or Confounding? AJE 2006;164:303-311.
- Basso O, Wilcox AJ. Intersecting Birth Weight-specific Mortality Curves: Solving the Riddle. AJE 2009;169:787-797
- Daniel RM, De Stavola BL, Cousens SN. gformula: Estimating causal effects in the presence of time-varying confounding or mediation using the g-computation formula. Stata J. 2011;11(4):479-517.
- Hernandez-Diaz S, Schisterman EF, Hernan MA. The birth weight "paradox" uncovered? AJE 2006;164(11):1115-2.
- Kramer MS, Zhang X, Platt RW. Analysing risks in adverse pregnancy outcomes. AJE 2014;179(3): 361-367.
- Petersen ML, Sinisi SE, van der Laan MJ. Estimation of direct causal effects. Epidemiology. 2006;17(3):276-284.
- Robins JM, Greenland S. Identifiability and exchangeability for direct and indirect effects. Epidemiology. 1992;3(2):143-155.
- Yerushalmy, J. Mother's cigarette smoking and survival of infant. AJOG 1964;88:505-518.
- Wilcox AJ, Russell I.Birthweight and perinatal mortality standardizing for birthweight is biased. AJE 1983; 118 (6):857-864.
- Wilcox AJ. On the importance and the unimportance of birth weight. International Journal of Epidemiology. 2001 Dec;30(6):1233-41.
- Yerushalmy, J. The relationship of parents cigarette smoking to outcome of pregnancy. Implications as to
  problem of infering causation from observed associations. AJE 1971;93(6):443-456.

イロン 不得 とくほ とくほう 一日





Bianca De Stavola/BW Paradox

(ignoring the confounders in these definitions; Vansteelandt, 2012)

The total causal effect (TCE):

$$TCE^{OR} = \frac{E[Y(1)]/\{1 - E[Y(1)]\}}{E[Y(0)]/\{1 - E[Y(0)]\}}$$

#### ■ The natural direct effect (NDE):

$$NDE^{OR} = \frac{E[Y(1, M(0))]/\{1 - E[Y(1, M(0))]\}}{E[Y(0, M(0))]/\{1 - E[Y(0, M(0))]\}}$$

#### The natural indirect effect (NIE):

$$NIE^{OR} = \frac{E[Y(1, M(1))] / \{1 - E[Y(1, M(1))]\}}{E[Y(1, M(0))] / \{[1 - E[Y(1, M(0))]]\}}$$

where Y(x) is the potential value of Y that would have occurred had X been set to x and Y(x, m) the potential value of Y that would have occurred had X been set to x and M to m

#### Bianca De Stavola/BW Paradox





	<b>Birth weight</b> $\geq$ 2.5 kg		Birth weight < 2.5 kg	
Mat Education	Low	High	Low	High
Births Deaths	92,704 220	59,141 222	4,393 225	4,128 195
Rates (x 1,000)	2.4	3.8	51.24	47.2
Sex-adjusted OR heterog test (p)	<b>1.58</b> (1.31, 1.91)		<b>0.92</b> (0.76, 1.12)	
Adjusted <sup>e</sup> OR heterog test (p)	<b>1.23</b> (1.01, 1.49)		<b>0.9</b> .036)	<b>2</b> (0.76, 1.12)

Bianca De Stavola/BW Paradox



	Variable	Class 1	Class 2
For $\mu$			
	Intercept	3.51	3.65
	sex	-	-
	year birth	-	+
	mat age	+	+
	birth order	-	+
For $\sigma$	Intercept	0.90	0.45
For $\pi$			
	sex	-	
	Mat educ	+	

About 10% of births predicted to be "compromised".



- There is another source of bias: conditioning on live birth.
- Still births are a form of competing event, reducing the denominator of possible infant deaths.
- Consider the composite outcome of Infant death or Still birth (Kramer *et al.*, 2014):

	Only Infant deaths Model I		Only Infant deaths & Still birth Model II	
	ln OR	(SE)	ln OR	(SE)
PNDE TNIE	<mark>0.221</mark> 0.011	(0.082) (0.007)	0.174 0.018	(0.067) (0.008)
TCE	0.232	(0.082)	0.192	(0.066)

イロン 不得 とくほ とくほう 一日