

Long-Term Causal Effects of Access to Institutional Delivery on Dementia Risk

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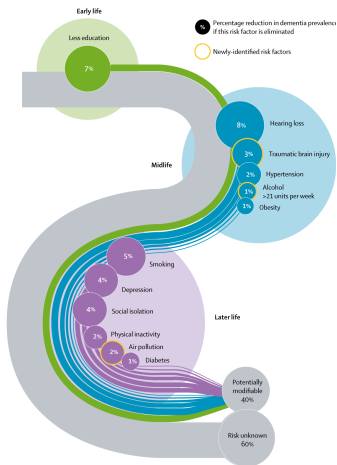


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Risk Factors for Dementia (Lancet Commission)

Risk factors for dementia

An update to the Lancet Commission on Dementia prevention, intervention, and care presents a life-course model showing that 12 potentially modifiable risk factors account for around 40% of worldwide dementias



- ▶ 40% of known risk factors potentially preventable.
- ▶ 60% either unknown or considered non-preventable.
- ▶ Known risk factors represent associations, not causal relationships.
- ▶ **Very early life period under-explored.**

Path Diagram

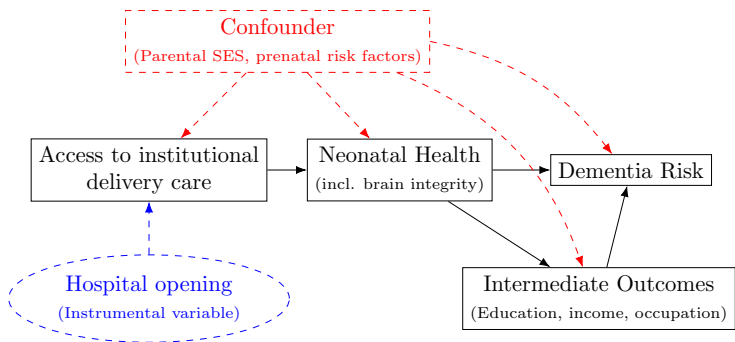


Figure: Path Diagram.

Reserve – Threshold Models

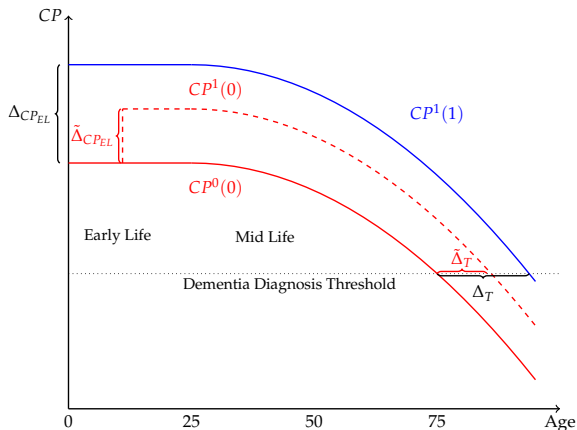
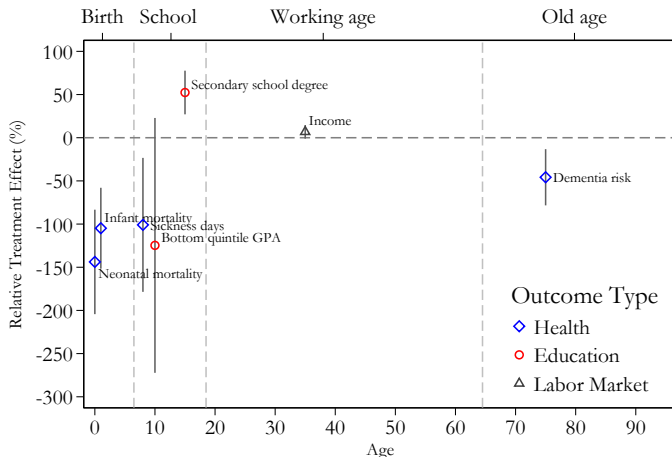


Figure: Cognitive Reserve – Potential Outcomes.

Treatment effects from being born in hospital estimated by instrumental variable



How to interpret the results – running LATE

- ▶ Estimated treatment effects from IV VERY large.
- ▶ Suggests that born in maternity ward eradicates infant mortality, the risk of getting dementia and doubles educational attainment. Note very reasonable...And also not in line with aggregated trends.
- ▶ Know that IV under **heterogenous treatment effects** estimates effect on **subpopulation of compliers**.
- ▶ Put effects into perspective by
 - a) using marginal treatment framework to explain the large effects stemming from **selection into treatment**,
 - b) present evidence that compliers were chosen through **sophisticated risk selection by midwives**,
 - c) realign large treatment effects with general aggregated trends,
 - d) Consequences for external validity and policy implications.

Transition from Home to Hospital Births

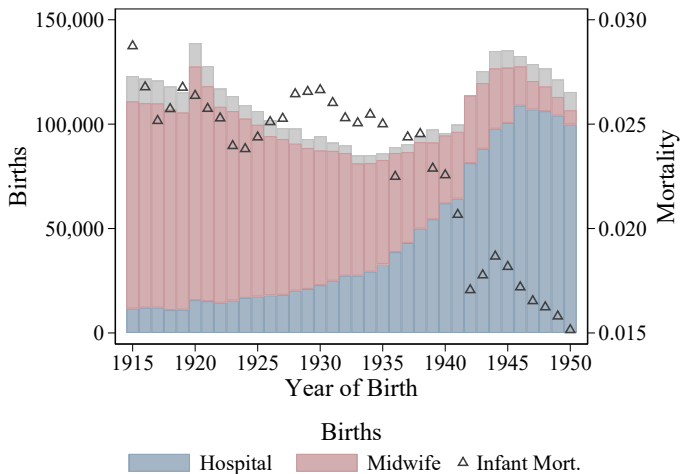


Figure: Live Births and Neonatal Mortality in Sweden, 1915–50.

Transition from Home to Hospital Births

- ▶ **Changing preferences** to give births in maternity wards.
- ▶ Transition **earlier** than in other Nordic countries.
- ▶ Swedish **midwives** generally well-educated but not for births with serious complications.
- ▶ Excess demand for giving birth in maternity wards ⇒ hospitals operated at capacity constraint.
- ▶ Supply shifts as *natural experiments*:
 - ▶ Establishment of **new hospitals**.
 - ▶ **Extensions** of existing institutions.

Data Sources

1. Swedish **Administrative Data** (SIP, individual level)
 - ▶ Censuses (1950, 1970)
 - ▶ Outcome: Dementia
 - ▶ Cause of Death Register
 - ▶ Inpatient Register
 - ▶ Mediators: Income, Education, Occupation
2. **Intervention Data** (expansions in the hospital sector)
 - ▶ 51 hospital openings or extensions.
 - ▶ Requirements for in sample:
 - ▶ Historical records on openings or extensions
 - ▶ Presence of a birthbook
 - ▶ Identification of **catchment area** around hospital (parishes which contributed to births at the hospital $\sim 30\text{--}40$ km).

Data Sources, Ctd.

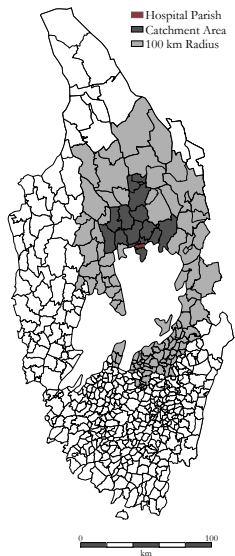
3. **School Performance Data** (individual level)

- ▶ Digitised from exam catalogues for subsample ($N = 10,965$)
- ▶ Includes performance years 1 and 4 (ages 7–11).
- ▶ Read/speak, write, math, religion, physical education.
- ▶ Sickness absence (pct of school days).

4. **Midwife Data** (universe of home births)

- ▶ Entire population 1928–38.
- ▶ Data on mothers, complications, procedures, parity...
- ▶ Available at **health district** level ($N = 400$).

Catchment Area Karlstad (1937)



1. Start with radius 60km (120 km in the 5 northern counties).
2. Data-driven approach to identify parishes within radius which contribute to maternity ward births.
3. Typical radius 30-40km.
4. Radius is in line with historical sources on travel distances, using quite sophisticated modes of transportation for 1930s (cars, sometimes even planes!).

Empirical Strategy: Instrumental Variables

First Stage:

$$H_i = \beta_0 + \beta_1 Z_i + \beta_2' x_i + \mu_c + \delta_t + \eta_i,$$

H_i hospital birth

Z_i born after an extension/opening

μ_c catchment area fixed effects

δ_t fixed effects birth year

Second stage:

$$Y_i = \gamma_0 + \gamma_1 \hat{H}_i + \gamma_2' x_i + \mu_c + \delta_t + \epsilon_i,$$

Empirical Models

1. Pooled Regression Discontinuity (before and after)
2. Two-way fixed effects (DiD) (with 2-year window around opening)
3. Difference-in-Discontinuity (with 2-year window around opening)
 - ▶ Linear regression models
 - ▶ Survival models (time of first diagnosis of dementia).
 - ▶ Control function
 - ▶ G-estimation

Local average treatment effect

- ▶ With our binary instrument and under the assumptions of
 - ▶ **Independence**
 - ▶ **Exclusion**
 - ▶ and **Monotonicity**...
- ▶ γ_1 identifies a LATE for those being *born in a hospital due to a supply side shift*.

$$\gamma_1^{LATE} = \mathbb{E} \left(Y_i^1 - Y_i^0 \mid H_i^1 > H_i^0 \right)$$

- ▶ **Not necessarily**

$$\gamma_1^{LATE} = \mathbb{E} \left(Y_i^1 - Y_i^0 \right)$$

$$\gamma_1^{LATE} = \mathbb{E} \left(Y_i^1 - Y_i^0 \mid H_i = 1 \right)$$

$$\gamma_1^{LATE} = \mathbb{E} \left(Y_i^1 - Y_i^0 \mid H_i = 0 \right)$$

First Stage

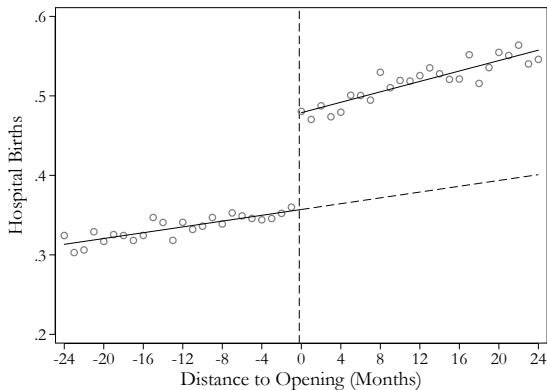


Figure: The First Stage: Hospital Birth

First Stage

Table: First Stage

	RDD	DiDisc
MATERNITY WARD OPENING/EXTENSION (P.P.) [95% CI]	12.42 [10.69,14.16]	12.51 [9.06,15.97]
Mean Dep. Var	37.37	37.37
F-Statistic	218.46	53.59

Notes: Robust standard errors are clustered at the level of the running variable.

Early Life and Mid-life Outcomes

Table: Instrumental Variable Estimates

	Neonatal mortality	Life-Cycle Earnings	Years of education	Secondary school
BORN IN MATERNITY WARD				
RDD	-0.041** (0.017)	11225.51 (16808.78)	1.48*** (0.39)	0.18*** (0.04)
Mean dep. var.	0.028	175,944.46	9.42	0.24
DiD	-0.045*** (0.009)	19457.92** (8280.75)	1.35*** (0.33)	0.17*** (0.05)
Mean dep. var.	0.028	175,944.46	9.42	0.24

Main Results

Table: Effects of Hospital Delivery on Dementia Risk

	RDD	DiDisc
PROPORTIONAL HAZARD MODEL		
ITT	0.876	0.883
[95% CI]	[0.774,0.990]	[0.798,0.976]
IV	0.339	0.462
	[0.130,0.885]	[0.218,0.979]
<i>N</i>	43,512	86,675
Cluster	24	48

Notes: Robust standard errors are clustered at the level of the running variable. Significance levels: * 0.10 ** 0.05 *** 0.01.

Untreated Outcomes Test

- ▶ Subpopulation of individuals who were **not** born in hospital
 - ▶ before a supply-side expansion: compliers (C) and never-takers (N)
 - ▶ after supply-side expansion only never-takers (N)
- ▶ any change in outcomes or in control variables $W^0 \in \{Y^0, X^0\}$ within the untreated group which coincides with the intervention Z must be a result of selection.
- ▶ Use two different data source:
 1. Midwife diaries on homes births (by construction conditioning on no treatment).
 2. Individual level data conditioning on **not** being treated.

Untreated Outcomes Test (UOT)

- ▶ Restrict analysis to **untreated**.
- ▶ **Before a supply-side expansion** the mean in the non-treated group is the weighted average between compliers and never-takers

$$\mathbb{E}(W^0|Z=0) = \frac{\mathbb{P}(C)}{\mathbb{P}(C) + \mathbb{P}(N)} \mathbb{E}(W^0|C) + \frac{\mathbb{P}(N)}{\mathbb{P}(C) + \mathbb{P}(N)} \mathbb{E}(W^0|N). \quad (1)$$

- ▶ **After the supply-side expansion** simply the mean of never-takers $\mathbb{E}(W^0|Z=1) = \mathbb{E}(W^0|N)$.
- ▶ We can quantify selection by

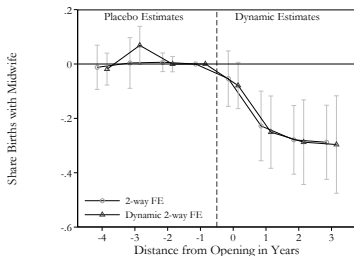
$$\begin{aligned} \Delta^0 &= \mathbb{E}(W^0|C) - \mathbb{E}(W^0|N) \\ &= (-1) \frac{\mathbb{P}(C) + \mathbb{P}(N)}{\mathbb{P}(C)} \left[\underbrace{\mathbb{E}(W^0|Z=1) - \mathbb{E}(W^0|Z=0)}_{\theta_0} \right], \end{aligned}$$

with θ_0 the reduced form estimand (ITT) of the supply-side expansion **based on untreated only**.

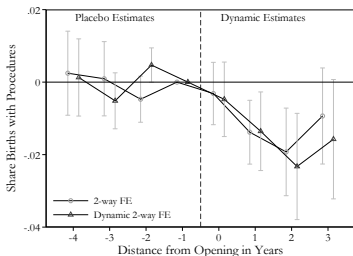
Untreated Outcomes Test (UOT)

- ▶ Testing whether there is **non-random uptake of treatment**.
- ▶ Δ^0 represents the mean difference between compliers and never-takers prior to treatment.
- ▶ Note that this difference cannot be due to the treatment itself, as all are untreated.
- ▶ We will use Δ^0 to investigate to which extent compliers are a selected group in terms of health risk and socio-economic status.
- ▶ Statistical inference is based on the delta method or bootstrapping.

Event Study Midwife Data (UOT)



(a)



(b)

Figure: Event Study: **(a)** Births with Midwife; **(b)** Midwife Birth with Procedure

UOT on Midwife Assisted Home Births

	(1)	(2)	(3)	(4)	(5)
PANEL A: DiD (2-WAY FE)					
	Births			Mother ill /	Twins
	with Midwife	with Procedures	with Complications	diseased	
Hospital Opening θ^0	-0.284*** (0.054)	-0.013*** (0.004)	-0.003** (0.001)	-0.006*** (0.002)	-0.001 (0.001)
<i>N</i>	791,755	622,930	622,930	622,930	622,930
Health Districts	397	413	413	413	413
UNTREATED OUTCOME TEST (SELECTION)					
$\mathbb{E}(W^0 N)$	0.513	0.029	0.003	0.011	0.012
Δ^0		0.055	0.011	0.027	0.003
Relative Risk		2.916	4.730	3.529	1.252
ROBUST DiD ESTIMATOR (DYNAMIC 2-WAY FE)					
Robust Effect	-0.232	-0.012	-0.002	-0.008	-0.000
SE Robust Effect	(0.061)	(0.004)	(0.001)	(0.007)	(0.002)

Notes: Table shows effects of the a hospital opening or extension in a given health district.

Source: Midwife Diaries. Own calculations.

Transition from Home to Hospital Births

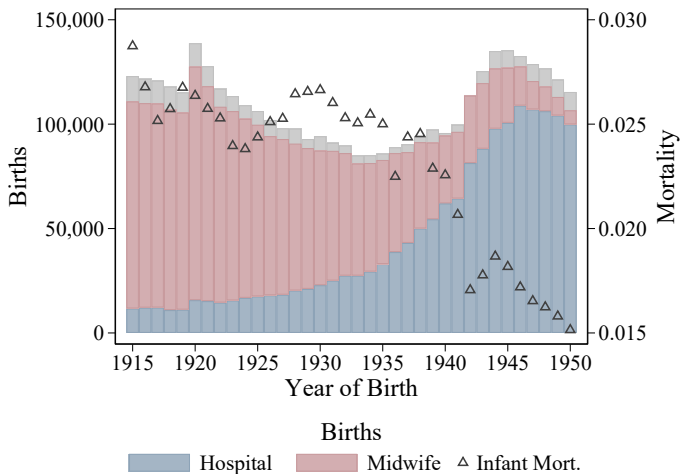


Figure: Live Births and Neonatal Mortality in Sweden, 1915–50.

Treated and Untreated Outcomes: Neonatal Mortality

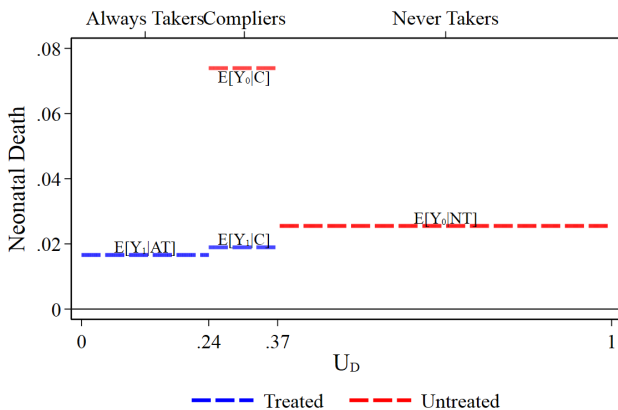


Figure: Treated and Untreated Outcomes by Group.

Treated and Untreated Outcomes: Neonatal Mortality

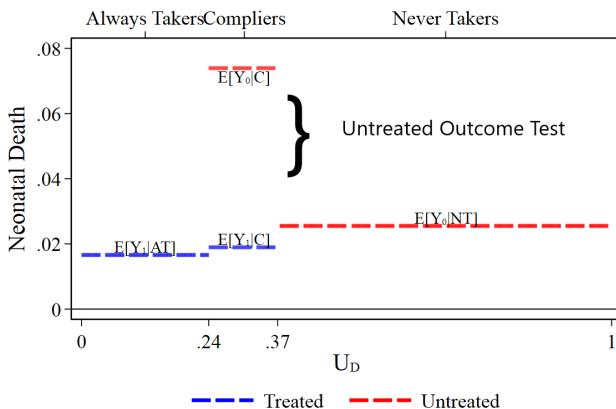
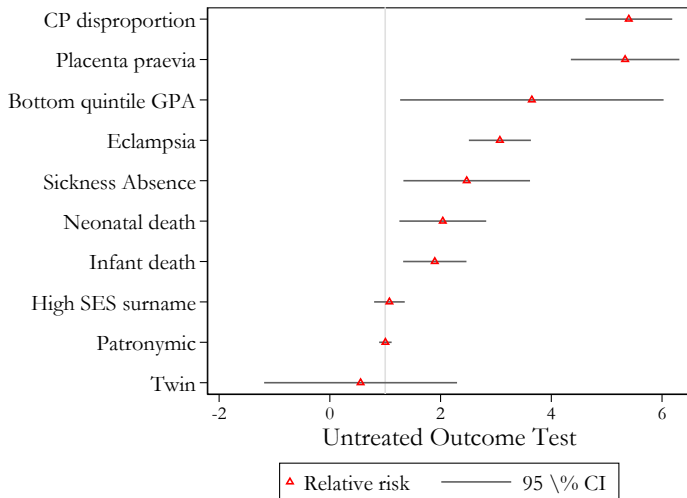


Figure: Treated and Untreated Outcomes by Group.

Selection: Summary



Is strong risk selection in line with historical sources?

▶ General

- ▶ By law (SFS 1919:798) the midwife had to contact a doctor in case of complications during child birth.
- ▶ The National healthcare committee (1928): hospitals should be for complicated cases.

▶ Related to specific complications

- ▶ Midwives not allowed to do surgery or use sharp instruments.
- ▶ Only doctors allowed to give medicine (pitocin 1927, ergometrin 1935, heparin 1937).
- ▶ Eclampsia could be identified by urine test by 1923.
- ▶ Hospitals had instruments to initiate contractions and from 1927 doctors could provide pitocin.

Can we do more? External validity?

- ▶ Until now only binary instrument.
- ▶ Estimates of IV restricted to compliers.
- ▶ How well do our IV estimates extrapolate?
- ▶ Important question, when decision whether extending a programme based on (self-)selection into treatment.
- ▶ Imposing more structure
 - ▶ Impose linearity in unobserved heterogeneity / *resistance* to being born in a hospital U_D
 - ▶ Continuous instrument - distance to next hospital from families place of residence.

Linear MTE: Neonatal mortality

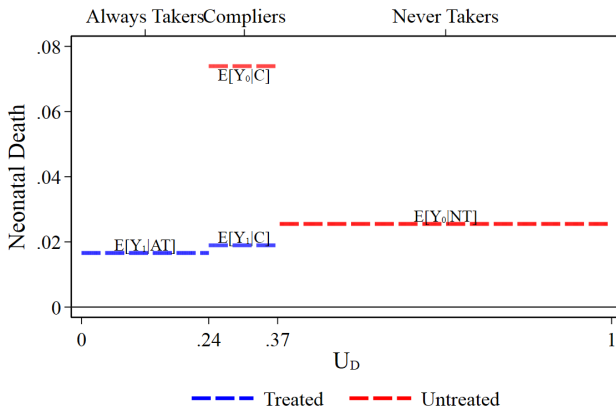


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Linear MTE: Neonatal mortality

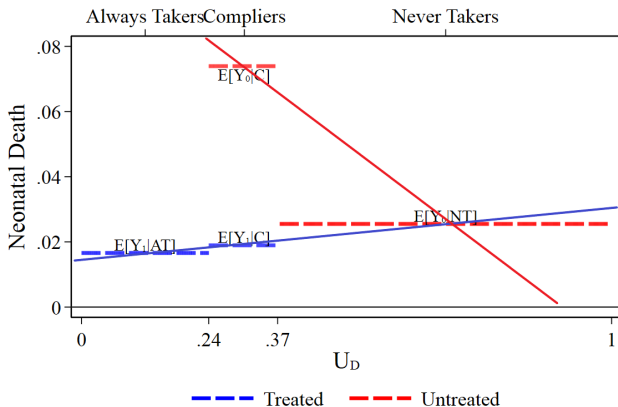


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Linear MTE: Neonatal mortality

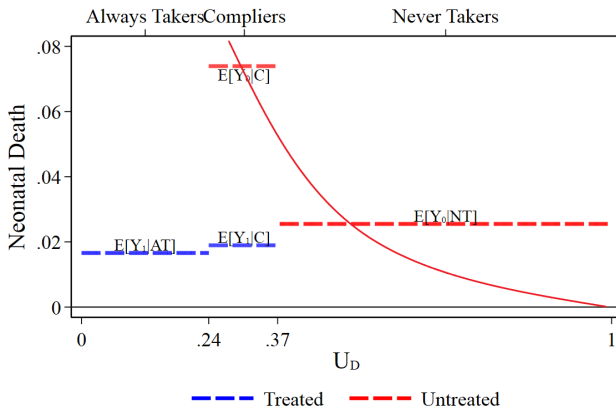
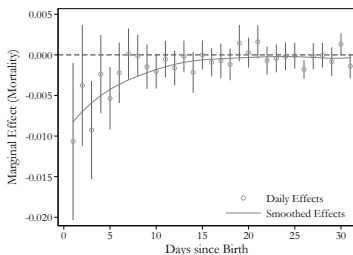
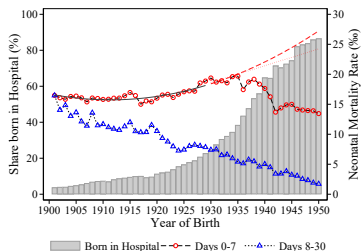


Figure: Treated and Untreated Outcomes by Group.

Reconciliation Effects and Trends



(a)



(b)

Figure: (a) Births with Midwife; (b) Midwife Birth with Procedure

Conclusions

- ▶ Critical births seem to have **shifted to hospitals**.
- ▶ **Better care** in case of (severe) **complications**.
- ▶ Find substantial decrease in the relative dementia risk of **10-15% from improved access** to services.
- ▶ Parts of the effect runs through **education** and **income**.
- ▶ Unexplored potential to address the **early-life origins** of dementia.
- ▶ Potential for large policy leverage in developing countries...
- ▶ However strong **self-selection** suggests $ATE \ll LATE$.