Parental Responses to Social Insurance for Children: Evidence from CHIP

Job Market Paper

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December 24, 2020

Abstract

This paper estimates the parental responses to a large expansion of children's insurance following the roll-out of the Children's Health Insurance Program (CHIP) in the US. Although CHIP does not cover children in utero, pregnant mothers exposed to the roll-out of CHIP invest more in utero, and birth weight improves for children with in-utero exposure to CHIP. The investments are consistent with a 14.3% reduction in the present bias of mothers exposed to the roll-out. Because mothers highly value the child's birth weight, they value the exposure to CHIP as much as expansions of own insurance due to the benefit on birth weight. In the long run, in-utero investments increase college enrollment and predict higher earnings and tax payments that lower the net cost of the program by 8.4%. The private and social benefits of the investment responses provide strong motivations for outreach efforts engaging parents in children's insurance programs.

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1 Introduction

A growing empirical literature has identified substantial benefits of early-life investments on the health, skill, and economic success of children over the life course. These benefits motivate social insurance to target children in low-income families, where private investments may be inadequate. While targeting children, the insurance relies on the behavior of parents to enroll and to invest in the child. Yet, parental behavior remains under-studied in the literature on children's insurance. In particular, there exists little evidence on how parents respond to children's insurance, and how their investments could impact child outcomes and the effectiveness of insurance.

This paper provides evidence on these open questions exploiting the roll-out of the Children's Health Insurance Program (CHIP) in the US. The program doubled the share of children eligible for public insurance from 15% to 30%, but did not expand insurance for pregnant mothers or their children in utero. As such, CHIP is unlikely to impact the birth outcomes of children. Nonetheless, birth outcomes may improve if pregnant mothers responded to the roll-out by increasing private investments in the child. Therefore, I examine mother's in-utero investments and the child's birth weight in the roll-out of CHIP to understand parental responses to social insurance for children.

I have three main findings from the analysis. First, exposure to the roll-out significantly increased mother's investments in the child, and birth weight already improved for children with in-utero exposure to CHIP. Second, the exposure increased investments by reducing the present bias of mothers. Due to the large benefits of investments on birth weight, mother values the exposure as much as expansions of her own insurance. Third, investments increased the college enrollment rate for children, and predict higher earnings and tax payments that could lower the social cost of insurance by 8.4% in the long run.

Conceptually, expanding insurance for children could decrease parental investments in utero due to the incentive effect on ex-ante investments. By investing more in the less healthy child, CHIP reduces the gap in child outcomes by child health. The insurance may lower parents' incentive to invest in utero in child health. Empirically, however, the roll-out of CHIP led to a substantial "crowd-in" of investments, suggesting the existence of additional mechanisms besides the incentive effect on investments. I motivate and show support for a behavioral effect on the present bias of mothers as the leading mechanism.

Empirically, I estimate parental responses exploiting the roll-out as an exogenous shock on mother's exposure to children's insurance. The roll-out increased the income limit of children's insurance across states. Within states, mothers exposed to CHIP early in the pregnancy had greater duration of exposure to the expansion. I capture the variations and

construct mother's exposure as a weighted average of income limits before and after CHIP, with the weights equal to the share of pregnancy exposed to each limit. In the regressions, I exploit the within-state variation in the duration of exposure across cohorts to estimate parental responses to children's insurance.

I find that exposure to the roll-out significantly increased mother's investments in the child, resulting in earlier onset of pre-natal visits and less smoking in pregnancy. Specifically, the exposure reduced late care onset past the first trimester by 5.6%, reduced very late onset in the third trimester by 11.2%, and reduced smoking by 5.3%. The investments increased birth weight by 8.5 grams, and decreased low birth weight by 3.6%. These effects were fully concentrated among children of single mothers exposed to CHIP since the first trimester. By contrast, exposure had little effect on the investments of married mothers whose children had low predicted eligibility for CHIP.

To understand mechanisms, I examine whether the exposure to children's insurance increased mother's uptake of own insurance, or increased her incomes in pregnancy. Drawing data from the Survey of Income and Program Participation, I find that the exposure had no effect on mother's insurance coverage, cash transfer incomes, borrowing from future incomes, or health spending in pregnancy. Moreover, mothers who invested more in utero also had substantially higher uptake of CHIP for their newborns, but these mothers did not indicate higher expected education for the child. This suggests that the investments were not motivated by the long-run effects of investments on education.

After ruling out mother's insurance, income, and the long-run effects of investments as mechanisms, I explore two behavioral mechanisms whereby the exposure may have adjusted mother's perception of investments. First, the exposure may have increased mother's utility weight on child outcomes relative to own consumption in pregnancy, increasing her altruism for the child. Second, the exposure may have shifted mother's inter-temporal weights towards utility in future periods, resulting in less present bias and more forward thinking in pregnancy.

I investigate the behavioral mechanisms using a dynamic model of in-utero investments. In the model, mother chooses pre-natal visits and smoking each trimester to invest in the child's birth weight. Because investments occur over multiple periods, mothers with lower valuation of future utility may delay investments and end up investing less in the child. By reducing present bias, the exposure could shift investments to early trimesters in pregnancy. It could further increase investments by increasing the altruism for the child. Exploiting the exposure to CHIP as a shifter of mother preferences, I match investments across exposure to quantify the behavioral effects. I find that the exposure to the roll-out lowered present bias by 14.3%, with little effect on altruism.

Turning to welfare, I quantify mother's valuation of the exposure from her investment responses. Mother values the exposure because it reduces her behavioral biases and results in better outcomes for the child. In the roll-out of CHIP, the exposure increased mother utility by \$0.31 due to the benefit of investments on birth weight. Compared to the program's spending on outreach, mother values the exposure at 76% of the spending, which is comparable to her valuation of own insurance from Medicaid (Hendren and Sprung-Keyser, 2020). For children's insurance, this result suggests that outreach efforts engaging parents can increase parental utility as effectively as expansions of parents' own insurance.

In the long run, investments increased college enrollment by 8.7% for children of single mothers. This effect could lower the social cost of insurance by increasing the tax payments of children. However, since mothers did not internalize the long-run effect on education, the fiscal externality is under-stated in mother's valuation of the exposure. To quantify the externality, I predict an increase in earnings by \$1,101.1 for children of single mothers based on the effect on college enrollment. The resulting increase in tax payments recoups 8.4% of the program cost in childhood. Thus, parental investments can have large impacts on the social cost of insurance through the long-run effects on child outcomes.

These findings suggest that parental investments can powerfully impact the effectiveness of children's insurance. Positive investment responses not only improve child outcomes before the program onset, but also lower the social cost of insurance through the long-run effects in adulthood. Investigations into the mechanism of the responses reveal that parents may suffer from behavioral biases that limit their investments in the child. By reducing the biases, the information of insurance can "nudge" investments and increase parental utility through the benefits on children. These results provide strong motivations for outreach efforts engaging parents in children's insurance programs.

This paper contributes to several strands of literature. First, although numerous studies have documented the beneficial impacts of insurance programs for children (Currie and Gruber 1996a; Currie and Gruber 1996b; Goodman-Bacon 2018), less is known about the parental responses to these programs, or whether parental investments could improve child outcomes over and above the direct effects of insurance. This paper examines inutero investments in the roll-out of insurance to isolate parental responses from program investments in the child. Hence, it contributes to the literature on children' insurance by establishing parental investments as a critical pathway to the benefits on children. Therefore, effective designs of children's insurance should encourage positive investment responses from parents. The role of parental investments has received similar attention in the literature of early-childhood interventions (Heckman and Mosso, 2014), where

programs improving parenting skills through information and preference change are shown to have larger impacts on child outcomes than simple transfer programs.

This paper also contributes to a growing literature that evaluates the welfare of social insurance programs through the lens of beneficiaries' valuation of the insurance (Finkelstein and Hendren 2020; Finkelstein et al. 2019a; Finkelstein et al. 2019b). Here, I quantify mother's valuation of her exposure to CHIP by first estimating the behavioral effects of exposure implied by her investment responses. In doing so, I also contribute to the literature on structural behavioral models, in particular models of dynamic inconsistencies and self control (Laibson 1997; O'Donoghue and Rabin 1999; DellaVigna and Malmendier 2006; Duflo et al. 2011; Sadoff et al. 2020), with new evidence from parental investments.

Lastly, this paper contributes to the literature applying behavioral insights to the design of social policies. To combat present bias, for instance, policymakers may consider taxes and subsidies to help individuals internalize their biases (Herrnstein *et al.* 1993; Gruber and Köszegi 2001; O'Donoghue and Rabin 2006). Alternatively, information nudges (Thaler and Sunstein, 2008) can achieve behavioral changes by adjusting the behavioral biases. In the roll-out of CHIP, the information of children's insurance reduced mother's present bias. The behavioral effect then increased mother utility as effectively as subsidized expansions of mother's own insurance.

The rest of the paper proceeds as follows. I introduce the Children's Health Insurance Program in Section 2, motivate the investment responses to CHIP exposure in Section 3, and describe the data in Section 4. Section 5 presents empirical evidence on the investment responses and explores mechanisms. Section 6 estimates the long-run effects on education outcomes. Section 7 estimates the behavioral effects and evaluates welfare using a structural model of in-utero investments. Section 8 concludes.

2 Children's Health Insurance Program

The Children's Health Insurance Program, or CHIP, was created by the Balanced Budget Act (BBA) of 1997 under a new Title XXI of the Social Security Act. Title XXI offered states the option to enroll children ineligible for Medicaid either through an expansion of the existing Medicaid program or by establishing a separate insurance program for children. States opting for either type of expansion were eligible for federal funding totaling \$40 billion in the first ten years of Title XXI (FY1998-FY2007). The funding should finance insurance expansions for children (age 0 to 18), not for adult parents.

Nearly all states expanded insurance for children between 1997 and 2000. Table 1 lists

¹The only exception is Tennessee, where the Medicaid program discrolled a large number of enrollees

the timing of CHIP onset by states, along with changes in the income limit of insurance for different age groups of children.² Of the 50 states that expanded insurance for children, 11 states also expanded insurance for adult parents using state funding.³ I focus on the 39 states expanding insurance exclusively for children to understand the parental responses to social insurance for children.

Overall, CHIP increased the income limit of Medicaid/CHIP by 80% of the federal poverty level (henceforth FPL) during the roll-out. The expansion significantly increased the *expected* insurance eligibility for children. In Figure 1, I predict insurance eligibility across birth cohorts applying the income limits of insurance (Table 1) known at the time of pregnancy. Children born in Jan. 1997 can expect 15% of childhood, or 2.9 childhood years, eligible for Medicaid/CHIP. Eligibility increased for children born in 1997-2000 during the roll-out. By Dec. 2000, all states (except Tennessee) had expanded insurance for children. Compared to the Jan-1997 birth cohort, the Dec-2000 birth cohort can expect 30% of childhood, or 5.7 childhood years, eligible for public insurance.

3 Conceptual Framework

To conceptualize the investment responses, I consider a two-period model where parental investments occur in utero (t=0) and in childhood (t=1). In each period, parents receive an exogenous income Y_t , and divide the income between own consumption c_t – which generates utility $u(c_t)$ – and costly investment v_t in the child. In-utero investments affect the child's health status at birth. Children born with lower health, such as those with disability and special health service needs, require larger medical expenses in childhood. CHIP lowers the medical expenses for parents investing in the less healthy child. I analyze the effect of this insurance on investments in childhood and in utero below.

in 2002. Before the disenrollment, individuals with income up to 400% FPL were eligible for the state's Medicaid program.

²I collect the income limit and the program onset date of Medicaid/CHIP from program fact sheets available at https://www.medicaid.gov/CHIP. For instance, the fact sheet for the state of New York is available at https://www.medicaid.gov/sites/default/files/CHIP/Downloads/NY/NYCurrentFactsheet.pdf. I track subsequent expansions from 2001 to 2013 combining the program fact sheets and the Trends of CHIP/Medicaid Eligibility charts published by the Kaiser Family Foundation available at https://www.kff.org/medicaid/state-indicator/medicaidchip-upper-income-eligibility-limits-for-children/?currentTimeframe=0&sortModel=%7B%22co1Id%22:%22Location%22,%22sort%22:%22asc%22%7D.

³States expanding insurance for children as well as adult members of the family are marked with an asterisk in the "infant" column in Table 1.

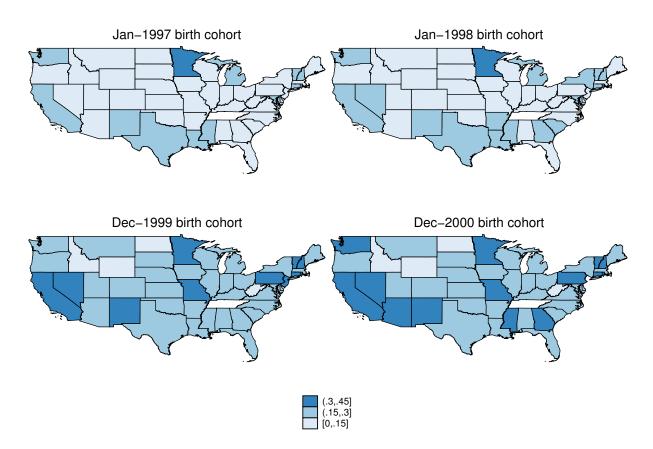
⁴I follow the standard "simulated eligibility" approach pioneered by Currie and Gruber (1996a) and Currie and Gruber (1996b) to calculate the expected eligibility. I show details of the calculation in Appendix A.

Table 1: CHIP onset and the income limit for children's insurance

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Notes. States marked with * in the "infant" column expanded insurance for children as well as adult members of the family. See footnote 2 in the main text for the sources of the information.

Figure 1: Expected insurance eligibility for birth cohorts, 1997-2000



Notes. The map shows the expected insurance eligibility for birth cohorts between Jan. 1997 and Dec. 2000. The expected eligibility calculates the fraction of childhood years (age 0-18) in which the child is eligible for Medicaid/CHIP based on program rules known at the time of pregnancy. Appendix A details the calculation.

3.1 Childhood Investments

Let superscript h = 0, 1 denote the child's health status at birth, with h = 1 indicating a healthy child. A less healthy child incurs out-of-pocket medical expenses OOPC in childhood. The medical expenses limit non-health investments v_1^h which are inputs to the child's education outcome $s(v_1^h)$. Given health status h, parents choose investment v_1^h to maximize the following utility

$$U_1^h(v_1^h) = u(c_1^h) + \Gamma(s(v_1^h)) + \delta V^h(s(v_1^h)), \tag{1}$$

where $c_1^h = Y_1 - v_1^h - OOPC \cdot 1\{h = 0\}$ is consumption after health and non-health investments, and $\Gamma(\cdot)$ is parents' utility from the child's education $s(v_1^h)$. $V^h(s(v_1^h))$ is the utility from the child's adult outcomes given education $s(v_1^h)$ and health h. δ is the discount factor.

Due to the medical expenses OOPC, non-health investment is smaller in the less healthy child.⁵ By reducing the medical expenses, CHIP increases v_1^0 and narrows the education gap $\Delta\Gamma = \Gamma(s(v_1^1)) - \Gamma(s(v_1^0))$ by child health. The reduction in the education gap results in smaller gaps in adult outcomes $\Delta V = V^1(s(v_1^1)) - V^0(s(v_1^0))$ and in consumption $\Delta u = u(c_1^1) - u(c_1^0)$ by child health.

3.2 In-Utero Investments

In t = 0, parents choose in-utero investments to maximize the sum of utility in utero and in childhood as follows

$$U_0(v_0) = u(c_0) + w(v_0) + \delta \rho(v_0) \tilde{U}_1^1 + \delta (1 - \rho(v_0)) \tilde{U}_1^0, \tag{2}$$

where $c_0 = Y_0 - v_0$ is consumption after investments, and $w(v_0)$ is the utility from the child's birth weight. $\rho(v_0)$ is the probability of giving birth to a healthy child, which increases with in-utero investment v_0 at decreasing rates: $\rho' > 0$ and $\rho'' < 0$. $\tilde{U}_1^h = \operatorname{argmax}_{v_1^h} U_1^h(v_1^h)$ is the maximized utility in t = 1 given health h.

Optimal in-utero investment satisfies the following first-order condition

$$\delta \rho'(v_0) \Delta \tilde{U}_1 + w'(v_0) = u'(c_0). \tag{3}$$

The first term $\delta \rho'(v_0) \Delta \tilde{U}_1 = \delta \rho'(v_0) (\Delta u + \Delta \Gamma + \delta \Delta V)$ is the marginal benefit of investment on the future utility of parents. This benefit is larger if the utility gap by child health, $\Delta u + \Delta \Gamma + \delta \Delta V$, is greater in childhood. $w'(v_0)$ is the marginal benefit on the utility from

⁵I show proofs in Appendix B.

birth weight. Optimal investment balances the marginal benefits with the marginal cost $u'(c_0)$.

CHIP affects the trade-off by lowering the benefit of in-utero investment on future utility. Because CHIP reduces the utility gap by child health, it lowers the marginal benefit $\delta \rho'(v_0) \Delta \tilde{U}_1$ in equation 3. Thus, parents have less incentive to invest in utero in child health. Since CHIP did not affect the cost of in-utero investments but lowered the marginal benefits, equation 3 predicts smaller in-utero investments in response to CHIP.

3.3 Investment Crowd-In

Equation 3 is consistent with the standard prediction that public insurance can "crowd-out" private investments. However, empirical evidence from the roll-out overwhelmingly shows an increase in in-utero investments. The crowd-in response suggests the existence of additional mechanisms that increase the marginal benefit of investments instead of decreasing it.⁶ Here, I consider two behavioral mechanisms in which the exposure to the roll-out may have adjusted parents' perception of investment benefits. First, the exposure may have increased parents' utility weight on child outcomes relative to own consumption, increasing the altruism of parents. Second, the exposure may have shifted inter-temporal weights towards utility in future periods, decreasing the present bias of parents.

Altruism. I model altruism as the weight on child outcomes in parents' utility. Exposure to the roll-out could increase altruism by raising parents' awareness of the child's well-being. In equation 3, increasing the child's weight to $\alpha > 1$ revises the first-order condition as follows

$$\delta \rho'(v_0) \Delta \tilde{U}_1(\alpha) + \alpha w'(v_0) = u'(c_0), \tag{4}$$

where the utility gap $\tilde{U}_1(\alpha) = \Delta u + \alpha (\Delta \Gamma + \delta \Delta V)$ now depends on the gap in child outcomes $\Delta \Gamma + \delta \Delta V$ multiplied by the altruism parameter α . Higher altruism thus increases parents' perceived benefit of investments. When parents care sufficiently more about child outcomes, the perceived benefit could increase despite smaller gaps in $\Delta \Gamma + \delta \Delta V$, resulting in a crowd-in of in-utero investments.

Present Bias. The exposure could further affect investments through the inter-temporal preferences of parents. Because in-utero investments have immediate costs but delayed benefits on children, parents who are overly sensitive to their immediate utility tend to under-invest in utero. By shifting inter-temporal weights towards future utility, the

⁶I focus on marginal effects assuming that the income and the resources of pregnant mothers did not increase in the roll-out. I show empirical support for the assumption in Section 5.5.

exposure could reduce parents' parent bias and increase investments in utero. More generally, present bias has been shown to affect health investments such as smoking (Gruber and Köszegi, 2001), preventive care (Fang and Wang, 2015), and food choice (Sadoff *et al.*, 2020). Here, I illustrate the implications for parental investments using the $\beta - \delta$ representation of present bias (Laibson, 1997).

Present bias affects in-utero investments according to the following condition

$$\beta \,\delta \,\rho'(v_0) \,\Delta \tilde{U}_1(\alpha,\beta) + \alpha \,w'(v_0) = u'(c_0). \tag{5}$$

Compared to equation 4, present-biased parents value the future benefits of investments less due to the bias term β < 1, and invest less in utero. Exposure to the roll-out could shift parents' inter-temporal weights towards future benefits for children, resulting in less present bias (higher β) and greater tolerance of investments in utero.

In equation 5, increasing either α or β predicts greater in-utero investment v_0 . However, the behavioral effects have different implications for the timing of investments across trimesters within pregnancy. Because present bias generates cost avoidance in each trimester, a behavioral effect lowering present bias further predicts shifts of in-utero investments towards early stages of pregnancy. Absent investment increases by the end of pregnancy, an increase in the early onset of investments is consistent with a reduction in present bias as the leading mechanism. I exploit this distinction to interpret the empirical results in Section 5.5 and to structurally estimate the behavioral effects in Section 7.

4 Data

I use the universe of birth certificate records in the US to study the investment responses to the roll-out of CHIP. The birth certificate shows the child's birth weight, mother's demographics, and in-utero investments such as pre-natal visits and smoking. I focus on the 39 states that expanded insurance exclusively for children during the roll-out, and focus on mothers between age 21 and 40 at the time of delivery. I therefore exclude teen pregnancies and pregnancies above age 40 from the analysis.⁷

Table 2 summarizes insurance eligibility, in-utero investments, and birth weight for children born in 1997-2001. The average income limit of Medicaid/CHIP is 175% FPL over this period. While exposed to the same income limits, children of single mothers are twice as likely to be eligible (44% compared to 23% on average) based on predicted

 $^{^7}$ 17% of the births in 1997-2001 were given by women below age 20. I exclude this group because states can extend maternity coverage for CHIP enrollees up to age 20. 82% of the births were given by women between age 21-40, and the remaining 1% were by women above age 40.

eligibility. Single mothers are less likely to start pre-natal care in the first trimester, more likely to delay care till the third trimester, and have higher rates of smoking in pregnancy. Birth weight is lower by 100 grams for children of single mothers.

Table 2: Summary statistics, children born in 1997-2001

	All Mothers		Single Mo	thers
	Observations	Mean	Observations	Mean
Medicaid/CHIP				
income limit (100% FPL)	12,094,302	1.75	2,978,094	1.74
eligibility	12,094,302	0.23	2,978,094	0.44
care started in 1st trimester (%)	12,117,214	84.99	2,966,116	72.67
care started in 3rd trimester (%)	12,117,214	5.44	2,966,116	11.29
# doctor visits	12,000,696	11.72	2,929,395	10.79
≥ 5 cigarettes daily (%)	9,727,610	8.13	2,365,340	15.49
≥ 15 cigarettes daily (%)	9,727,610	2.96	2,365,340	5.55
birth weight (grams)	12,407,979	3339.43	3,067,358	3234.10
low birth weight (% <2,500 grams)	12,407,979	7.10	3,067,358	9.64

Notes. Table summarizes insurance eligibility, in-utero investments, and birth weight for cohorts born between Jan. 1997 and Dec. 2001 in the 39 states expanding insurance exclusively for children during the roll-out of CHIP. I predict children's eligibility for Medicaid/CHIP based on the income limits of insurance known at the time of pregnancy. Appendix A details the calculation.

5 In-utero Investments

5.1 Empirical Strategies

To estimate the investment responses, I exploit the roll-out of CHIP as an exogenous shock on mother's exposure to children's insurance. The roll-out increased the income limit of children's insurance across states. Within states, mothers exposed to CHIP early in the pregnancy had greater duration of exposure to the expansion. I measure mother's exposure to children's insurance using the average income limit of Medicaid/CHIP in pregnancy, and construct in-utero exposure, *eliginc*, as follows

$$eliginc_{st} = \begin{cases} inc_s^{pre} & \text{if } j \leq -10, \\ \frac{|j|}{9} \cdot inc_s^{pre} + \left(1 - \frac{|j|}{9}\right) \cdot inc_s^{post} & \text{if } -9 \leq j \leq -1, \\ inc_s^{post} & \text{if } j \geq 0, \end{cases}$$

$$(6)$$

where $j = t - T_s$ is the gap between the pregnancy onset time t and the CHIP onset time T_s in state s. inc_s^{pre} and inc_s^{post} denote the income limit of children's insurance before and

after CHIP, respectively.8

In equation 6, in-utero exposure is a weighted average of income limits inc_s^{pre} and inc_s^{post} , with the weights equal to the share of pregnancy exposed to each limit. Mothers gaining exposure to CHIP in month |j| of the pregnancy have an weight $\frac{|j|}{9}$ associated with the pre-CHIP limit inc_s^{pre} . Mothers starting pregnancy more than 9 months before CHIP $(j \le -10)$ are exposed only to inc_s^{pre} , whereas mothers starting pregnancy after CHIP $(j \ge 0)$ are fully exposed to the CHIP limit inc_s^{post} . Thus, in-utero exposure increases within states for mothers with earlier exposure and hence greater duration of exposure to CHIP. In the empirical analysis, I exploit the variation in exposure timing across mothers starting pregnancy between 16 months before and 4 months after CHIP $(-16 \le j \le 4)$.

I estimate parental responses to children's insurance using the following specification

$$y_{itc} = \beta_1 \cdot eliginc_{ts(c)} \cdot single + \beta_2 \cdot eliginc_{ts(c)} + \beta_c \cdot single + \alpha_c + \tau_t + \alpha_{s(c)} \cdot \tau_{v(t)} + \epsilon_{itc},$$
 (7)

where y_{itc} is the investment of mother i starting pregnancy in time t. $eliginc_{ts(c)}$ is her in-utero exposure to children's insurance. I include county fixed effects α_c and control for time-varying factors by state-year in $\alpha_{s(c)} \cdot \tau_{y(t)}$. Since children of single mothers are twice as likely to be eligible for insurance (Table 2), I examine whether investments differ by single motherhood indicated by single. Based on the eligibility differences, we should expect larger responses from single mothers captured in β_1 than from married mothers captured in β_2 . This difference allows me to further control for time-varying factors by state-year-month in the following specification

$$y_{itc} = \beta \cdot eliginc_{ts(c)} \cdot single + \beta_c \cdot single + \alpha_c + \tau_t + \alpha_{s(c)} \cdot \tau_t + \epsilon_{itc}, \tag{8}$$

where $\alpha_{s(c)} \cdot \tau_t$ absorbs the main effect of exposure $eliginc_{ts(c)}$ and fully controls for unobserved differences across states and over time. β estimates the investment responses of single mothers.

Both equation 7 and 8 control for differences by single motherhood in local areas (counties) with $\beta_c \cdot single$. Within counties and mother groups, I exploit the variation in exposure timing across cohorts to estimate the investment responses. The empirical strategy requires that mothers do not selectively enter pregnancy or single motherhood in response to CHIP, and that confounding policy changes across states do not explain the investments. I rule out fertility responses in Appendix Table I1, and fully control for time-varying factors across states in equation 8. I illustrate the identifying variation showing

⁸Using the program language, I refer to income limits in 100% federal poverty levels (FPL). For instance, an income limit of "200% FPL" in the CHIP program implies that $inc_s^{post} = 2$.

sharp changes in investments across exposure timing in the event study. I incorporate additional heterogeneity across mother groups using simulated eligibility in Section 5.4, finding similar results for single mothers.

5.2 Birth Weight

I first examine the effect of in-utero exposure on the child's birth weight. Because CHIP did not expand insurance for pregnant mothers, the roll-out is unlikely to impact the birth weight of children. An increase in the birth weight, however, indicates the existence of private investment responses to the roll-out. In Table 3, I estimate the effect on birth weight using equation 7 and 8.

Exposure to the roll-out significantly increased the child's birth weight. In column 1, gaining a 100% FPL exposure increased birth weight by 10.6 grams, or by 0.32% above the mean. In column 3, the exposure decreased the probability of low birth weight (<2,500 grams) by 0.32 percentage points, or by 4.5% below the mean. Both effects are fully concentrated among children of single mothers. Moreover, replacing the main effect of *eliginc* with a full set of state-year-month fixed effects in column 2 and 4 yields very similar results for children of single mothers. These estimates imply that the roll-out of CHIP, which expanded income limits from 122% FPL to 202% FPL in 1997-2000, increased birth weight by $10.6 \cdot 80\% = 8.5$ grams, and decreased low birth weight by $4.5\% \cdot 80\% = 3.6\%$ for children of single mothers.

Table 3: Effects of exposure on birth weight

	(1)	(2)	(3)	(4)
	birth weig	ght (grams)	low birth	weight (%)
eliginc·single	10.66***	10.61***	-0.32***	-0.32***
eliginc	(2.51) -1.76 (4.09)	(2.51)	(0.10) 0.005 (0.14)	(0.10)
y mean	3342.57		7.0	08%
R^2 N	0.02 4,31	0.02 5,394	0.01 4,31	0.01 5,394

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table shows the effect of in-utero exposure on birth weight (grams) and low birth weight (<2,500 grams). I estimate separate effects for single and married mothers using equation 7 in column 1 and 3, and estimate effects on single mothers using equation 8 in column 2 and 4. Robust standard errors clustered at the level of states in the parenthesis.

To illustrate the identifying variation, I estimate the following event study specification

$$y_{itc} = \sum_{\substack{j=-16\\j\neq-10}}^{4} \beta_{j} \cdot eliginc_{ts(c)} \cdot single \cdot 1\{t - T_{s(c)} = j\} + \gamma \cdot eliginc_{ts(c)}$$

$$+ \beta_{c} \cdot single + \alpha_{c} + \tau_{t} + \alpha_{s(c)} \cdot \tau_{y(t)} + \epsilon_{itc}, \qquad (9)$$

where I expand the term $\beta_1 \cdot eliginc_{ts(c)} \cdot single$ in equation 7 by the timing of exposure indexed by j, and estimate separate effects across cohorts with β_j . I normalize the effect on children conceived 10 months before CHIP to zero ($\beta_{-10} = 0$). These children and earlier cohorts ($j \le -10$) were born before the CHIP onset, and hence were not exposed to CHIP in utero. In-utero exposure then increased with j for later cohorts.

Figure 2 plots estimates of β_j for birth weight in panel (a). Birth weight did not increase for children never exposed to CHIP in utero ($j \le -10$). For children exposed to CHIP in the second and third trimester ($-9 \le j \le -4$), the exposure had no significant impact on birth weight. Exposure significantly increased birth weight for children exposed to CHIP in the first trimester ($-3 \le j \le -1$) and those fully exposed to CHIP in utero ($j \ge 0$). Among these children, gaining a 100% FPL exposure increased birth weight by 4.1 grams, and decreased low birth weight by 1.6% (Appendix Table I2). The overall effect on birth weight was concentrated among children of single mothers exposed to CHIP since the first trimester.

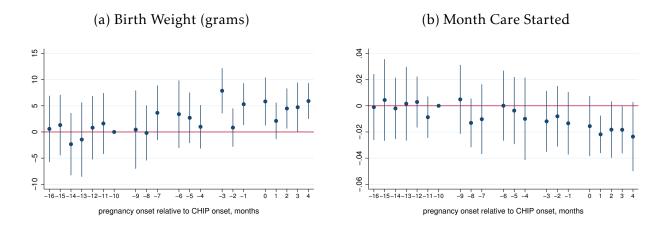
5.3 In-Utero Investments

The effect on birth weight suggests that exposure to the roll-out may have increased mother's private investments in the child. To detect the investment responses, I examine mother's pre-natal visits and smoking in pregnancy, and estimate the effects of exposure on the *timing* and the *level* of investments using equation 7 and 8. I find that exposure to the roll-out increased mother's early onset of pre-natal visits, reduced the delay in pre-natal visits, and reduced smoking in pregnancy.

Investment Timing. I examine responses in the timely onset of pre-natal visits in Table 4. Pre-natal care begins when the pregnant woman has the first pregnancy-related doctor visit. The Guidelines for Perinatal Care (Freeman and Poland, 1992) recommends that pregnant women have one doctor visit per month in the first two trimesters, and have 4

⁹To succinctly summarize the results across cohorts, I group children by the trimester of exposure and estimate effects for six exposure groups in Appendix Table I2.

Figure 2: Effects of exposure on birth weight and pre-natal care onset, event study



Notes. Figure plots the effect of in-utero exposure on birth weight in panel (a) and on the onset month of pre-natal visits in panel (b), for cohorts with different timing of exposure to CHIP. Mothers starting pregnancy more than 10 months before CHIP were not exposed to CHIP in pregnancy. In-utero exposure then increased for later cohorts. 95% confidence intervals are based on robust standard errors clustered by states.

visits per month towards the end of pregnancy. Nonetheless, around 15% of mothers delay the onset of care till the second or third trimester, and 5% start care in the third trimester. In Table 4, gaining a 100% FPL exposure decreased late care onset past the first trimester by 1.1 percentage points, or by 7% below the mean, and decreased third trimester onset by 14%. The exposure reduced the time to the first pre-natal visit by 0.05 months. These responses were fully concentrated among single mothers.

Panel (b) of Figure 2 plots the event study estimates for the timing of care onset from equation 9. The month of care onset did not decrease for mothers not exposed to CHIP in pregnancy, and decreased by a small and insignificant amount for mothers exposed to CHIP in the first trimester. The overall effect on the timing of care onset was concentrated among single mothers with full exposure to CHIP ($j \ge 0$). For these mothers, gaining a 100% FPL exposure reduced late care onset by 2.5%, and reduced very late onset in the third trimester by 5.5% (Appendix Table I3).

Investment Levels. I next examine if improvements in the early onset of visits led to greater number of visits by the end of pregnancy. In Table 5, gaining a 100% FPL exposure increased pre-natal care by 0.09 visits. This effect is small in magnitude. Compared to the sample average of 11.7 visits in pregnancy, the exposure increased pre-natal visits by less than 1%. Results by the timing of exposure in Appendix Table I4 show a similar null

Table 4: Effects of exposure on the timing of pre-natal visits

	(1)	(2)	(3)	(4)	(5)	(6)
	month ca	month care started		late onset (%) (2nd/3rd trimester)		onset (%)
eliginc · single	-0.046***	-0.046***	-1.09***	-1.10***	-0.75***	-0.75***
	(0.014)	(0.014)	(0.26)	(0.27)	(0.25)	(0.25)
eliginc	0.018		0.33		0.29	
	(0.015)		(0.32)		(0.17)	
y mean	2.	45	15.0	08%	5.4	8%
R^2	0.08	0.08	0.06	0.06	0.04	0.04
N	4,200	0,326	4,20	0,326	4,20	0,326

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table shows the effect of in-utero exposure on the month of care onset in column 1-2, late care onset past the first trimester in column 3-4, and very late onset in the third trimester in column 5-6. I estimate separate effects for single and married mothers using equation 7 in odd-numbered columns, and estimate effects on single mothers using equation 8 in even-numbered columns. Robust standard errors clustered at the level of states in the parenthesis.

effect on the number of visits, including for mothers with full exposure to CHIP ($j \ge 0$). Among these mothers, despite earlier onset of pre-natal visits, the number of pre-natal visits did not increase significantly with exposure.

Table 5: Effects of exposure on the number of pre-natal visits and smoking

	(1)	(2)	(3)	(4)	(5)	(6)
	# pre-na	tal visits		ng (%) ettes daily)		oking (%) ettes daily)
eliginc · single	0.091**	0.089*	-0.45**	-0.46**	-0.38***	-0.39***
	(0.045)	(0.045)	(0.18)	(0.18)	(0.13)	(0.13)
eliginc	-0.050		-0.073		0.058	
	(0.041)		(0.16)		(0.13)	
y mean	11.	.74	8.4	1%	3.1	2%
R^2	0.07	0.07	0.06	0.06	0.03	0.03
N	4,157	7,327	3,33	1,203	3,33	1,203

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table shows the effect of in-utero exposure on the number of pre-natal visits and smoking intensity. I estimate separate effects for single and married mothers using equation 7 in odd-numbered columns, and estimate effects on single mothers using equation 8 in even-numbered columns. Robust standard errors clustered at the level of states in the parenthesis.

The birth certificate also indicates the number of cigarettes consumed daily by the pregnant mother. Because the consumption is recalled as an average in pregnancy, the duration of the recall adds noises to the data and potentially attenuates the smoking responses. To limit the attenuation bias, I focus on the intensive margin and examine smoking more than 5 and 15 cigarettes daily in column 3-6 of Table 5. Gaining a 100%

FPL exposure reduced smoking more than 5 cigarettes daily by 5%, and reduced smoking more than 15 cigarettes daily by 13%. These effects were similarly concentrated among single mothers exposed to CHIP since the first trimester (Appendix Table I4).

5.4 Heterogeneity

Effects by States. In addition to examining investment responses by the timing of exposure, I further explore heterogeneity across states using the following specification

$$y_{itc} = \sum_{k} \beta_{k} \cdot eliginc_{ts(c)} \cdot single \cdot 1\{s = k\} + \gamma \cdot eliginc_{ts(c)}$$

$$+ \beta_{c} \cdot single + \alpha_{c} + \tau_{t} + \alpha_{s(c)} \cdot \tau_{y(t)} + \epsilon_{itc},$$

$$(10)$$

where β_k estimates the effect of CHIP exposure in state k.¹⁰ I plot estimates of β_k by the size of expansion across states in Appendix Figure J1. Exposure had the largest impact on birth weight among small expansion states increasing income limits by less than 70% FPL.¹¹ The expanded income limit (162% FPL) remained 40% FPL below the national average. In these states, a 100% FPL expansion would increase birth weight by 16 grams, or by 50% above the average effect across states. The reduction in smoking is also larger in the small expansion states.¹² By comparison, the majority of states (65% of births) expanded income limits by 75%-90% FPL, and the effects on birth weight and investments in this range are comparable to the average effects across states.¹³ Finally, a handful of states (10% of births) expanded income limits by more than 100% FPL.¹⁴ The effects on birth weight and investments tend to be smaller in the largest expansion states.

$$y_{it}^k = \beta_1^k \cdot eliginc_t^k \cdot single_i^k + \beta_2^k \cdot single_i^k + \tau_t^k + \epsilon_{it}^k, \tag{11}$$

where $eliginc_t^k$ differs in state k by the timing of exposure across cohort t. In practice, equation 10 and equation 11 give very similar estimates by states. I plot estimates of β_k from equation 10.

¹⁰The state-specific effects can also be estimated from the state-level regression

¹¹Around 25% of all births occurred in the small expansion states. The largest expansion state in this group is Maine, where income limit increased by 55% FPL. States such as Louisiana, South Dakota, Wyoming, Iowa, and Mississippi expanded income limits by only 22.58% FPL. I do not estimate state-specific effects for Minnesota where the income limit increased by a minimal 0.26% FPL in the roll-out.

 $^{^{12}}$ I examine smoking intensity by half packs (10s) of cigarettes daily. Specifically, I use integer 0, 1, and 2 to indicate non-smokers (<5 cigarettes daily), light smokers (≥5 and <15 cigarettes daily), and heavy smokers (≥15 cigarettes daily), respectively, so that intensity increases by half packs (10s) of cigarettes daily.

¹³This range includes states with the largest number of births such as California, Texas, New York, and Florida. In these populous states, birth weight increased by 13.6 grams per 100% FPL exposure, or by 30% above the average effect across states.

¹⁴These states are Rhode Island, Connecticut, New Hampshire, Pennsylvania, and Missouri.

Simulated Exposure. Complementing the main analysis, I incorporate additional heterogeneity in mother's exposure to children's insurance using the simulated eligibility strategy (Currie and Gruber 1996a; Currie and Gruber 1996b). Specifically, I apply the income limit of Medicaid/CHIP to calculate the probability of being eligible for insurance for a fixed sample of children. Because children of low-income mothers are more likely to be eligible, I calculate the probability by mother demographics – across race, education, and marital status – to simulate mother's exposure to children's insurance. ¹⁵ I then estimate investment responses to simulated exposure *eligCHIP* in the following specification

$$y_{its} = \beta \cdot eligCHIP_{d(i)ts} + \alpha_s \cdot \gamma_d + \alpha_s \cdot \tau_t + \epsilon_{its}, \tag{12}$$

By construction, exposure $eligCHIP_{d(i)ts}$ is larger for mother demographics with greater predicted eligibility for CHIP, for cohorts exposed to CHIP earlier in pregnancy, and in states expanding insurance to higher income limits. I control for time-varying factors across states with $\alpha_s : \tau_t$, and control for unobserved differences across states and mother

where exposure $eligCHIP_{d(i)ts}$ differs across state s, cohort t, and mother demographics d.

across states with $\alpha_s \cdot \tau_t$, and control for unobserved differences across states and mother demographics with $\alpha_s \cdot \gamma_d$. Within states and mother demographics, I exploit eligibility differences by the timing of exposure across cohorts to estimate the investment responses.

I find similar effects on birth weight and investments using the simulated exposure in Appendix Table I5. Increasing exposure by 10 percentage point eligibility increases birth weight by 3.2 grams, and the effect is concentrated among children of single mothers. For these children, the roll-out increased insurance eligibility from 0.33 to 0.50, increasing birth weight by 8.1 grams. This effect is comparable to the 8.5 gram increase implied by estimates from equation 8 in Table 3. Similarly, investment responses to eligibility increases are comparable to the effects of expanded income limits for single mothers, ¹⁷ supporting the results from equation 8.

¹⁵I show details of the simulation in Appendix A. Simulated exposure differs sharply by the marital status of mothers. Specifically, the roll-out of CHIP increased simulated exposure to 50% for single mothers, and to 16% for married mothers.

 $^{^{16}}$ Specifically, simulated exposure implies a birth weight gain of $(0.50-0.33)\times47.5=8.1$ grams. The roll-out expanded income limits by 80% FPL, implying an increase in birth weight by $10.6\times80\%=8.5$ grams applying estimates from equation 8 in Table 3.

¹⁷Specifically, increases in simulated exposure imply a 0.9 percentage point reduction in late care onset and a 0.2 percentage point reduction in heavy smoking for single mothers in the roll-out. These effects are comparable to the 0.9 percentage point reduction in late care onset and the 0.3 percentage point reduction in heavy smoking implied by estimates from equation 8.

5.5 Mechanism

The empirical results show compelling evidence of investment crowd-in following the roll-out of CHIP. To understand mechanisms, I first examine whether the exposure to children's insurance increased mother's take-up of own insurance and safety net benefits. I also examine whether mothers increased in-utero investments to improve long-run outcomes of the child. I find little support for these mechanisms. Instead, behavioral effects on mother's altruism and present bias could explain the investment responses.

Budget Constraint. I examine mother's insurance and income in pregnancy using the Survey of Income and Program Participation (SIPP). I find no evidence that the exposure to children's insurance increased mother's uptake of own insurance in pregnancy (Appendix Table I6). I also find no evidence that the exposure increased mother's cash transfer incomes or health spending in pregnancy (Appendix Table I7). Moreover, I find no evidence that mothers borrowed more from future incomes to invest more in pregnancy (Appendix Table I8). These results are inconsistent with a resource-based mechanism where the exposure increased in-utero investments by relaxing the budget constraints of mothers.

Long-Run Effects. Investments could also increase if mothers internalized the long-run effects of investments on child outcomes. In this case, mothers investing more during the roll-out should expect better outcomes for their children later in life. I test this implication examining mother's decision to enroll the child in CHIP and her expectation of the child's education attainment in Appendix Table I9. Mothers exposed to CHIP since the first trimester, in addition to investing more in utero, also had substantially higher uptake of CHIP for their newborns, but despite their investments, these mothers did not indicate higher expected education for the child. These results are inconsistent with a mechanism motivated by the long-run effects of investments on child outcomes.

Altruism. I next examine the two behavioral mechanisms which I take to the data in the structural analysis. Exposure to the roll-out may increase mother's altruism for the child by increasing her awareness of the child's well-being. The increase in altruism tends to increase investments in all trimesters, and predicts greater number of pre-natal visits and less smoking in pregnancy. Empirically, however, the exposure significantly shifted the timing of pre-natal visits across trimesters with smaller and weaker effects on the number of visits by the end of pregnancy. The effects on investment timing suggest the existence of mechanisms in addition to the effects on altruism.

Present Bias. To explain responses in the investment timing, I explore behavioral effects of the exposure on the time preferences of mothers. Specifically, exposure to the roll-out may have shifted mother's intertemporal weights towards utility in future periods, reducing mother's sensitivity to immediate costs of investments. The reduction in present bias predicts less delay in costly investments and more self control over additive consumption such as smoking, consistent with earlier onset of visits and reduced smoking in pregnancy. I formally investigate the behavioral mechanisms using a dynamic model of in-utero investments in Section 7, where I exploit the investment responses to quantify the behavioral effects of the exposure.

6 Education Outcomes

I next estimate the long-run effects of in-utero exposure on children's education outcomes using the American Community Survey (ACS). I find that cohorts with greater exposure to CHIP in utero are more likely to graduate high school and enroll in college. In high school, they are more likely to attend the grade appropriate for their age. These effects are concentrated among children of single mothers.

6.1 Empirical Strategy

I estimate the effects of in-utero exposure on education outcomes using the following specification

$$y_{ibqt} = \beta_0 \cdot eliginc_{ibq}^{utero} \cdot single + \beta_1 \cdot eliginc_{ibq}^{utero} + \beta \cdot X_{ibqt}$$

$$+ \gamma_b + \psi_q + \tau_t + \gamma_b \cdot \psi_{y(q)} + \gamma_b \cdot \tau_t + \beta_b \cdot single + \epsilon_{ibqt},$$
 (13)

where y_{ibqt} is the education of child i born in year-quarter q and state b surveyed in year t. $eliginc_{ibq}^{utero}$ is the in-utero insurance exposure of child i, calculated from a 3-quarter average of income limits during pregnancy. single indicates children of single mothers. Since in-utero responses to CHIP are concentrated among single mothers, the long-run effects on education tend to be larger for children of single mothers (β_0) than for children of married mothers (β_1). In X_{ibqt} , I include childhood exposure $eliginc_{ibqt}^{child}$, constructed as the average income limit from birth till year t, and child age. 18

I control for unobserved differences by single motherhood across states in $\beta_b \cdot single$. Within states, I exploit differences in the timing of exposure across cohorts to estimate

¹⁸I include separate controls of childhood exposure for children of single and married mothers in X_{ibqt} .

the effect of CHIP exposure on education. In addition, I use state-year fixed effects to account for state-specific trends during the roll-out and in the long-run follow-up. ¹⁹ In the robustness check, I further include *single*-year fixed effects to absorb the long-term trends of single motherhood. I estimate equation 13 for children conceived 7 quarters before till 2 quarters after CHIP to exploit the roll-out of CHIP. ²⁰

I define two measures of single motherhood based on mother's marital status. The first measure requires that the mother has never married by the time of the survey. Under this measure, the child was born to single mothers who remained single throughout the parenthood. The second measure only requires that the mother is unmarried in survey year t. Ex ante, one might expect larger effects of CHIP exposure on children raised continuously in single-parent households.

Equation 13 may yield biased estimates of the effect of CHIP exposure, if mothers adjusted the probability of single parenthood in response to CHIP. Empirically, I examine and rule out the effects of CHIP exposure on the marital status of mothers in Appendix Table I10, and rule out fertility responses to the roll-out of CHIP in Appendix Table I1. These results suggest that differences in the timing of exposure are plausibly exogenous for children of single mothers. To make explicit the variation by the timing of exposure, I plot event study estimates by the quarter of exposure on education outcomes below.

6.2 Effects on Education

I study the education outcomes of children expected to enter Grade 9 or above in 2010-2018. Following the academic progress of these children through high school and into college, I estimate the effect of CHIP exposure on their grade-for-age status, high school graduation, and college enrollment applying equation 13. I focus on college enrollment first.

College Enrollment. Table 6 estimates the effects on college enrollment. Column 3-4 estimates the preferred specification in equation 13. Gaining a 100% FPL exposure to CHIP increases college enrollment by 2.91 percentage points for children of never married mothers, or by 18.44% above the mean. For children of single mothers in general, the exposure increases college enrollment by 1.71 percentage points. Controlling for the

¹⁹Specifically, $\gamma_b \cdot \psi_{y(q)}$ controls for state-specific trends in the roll-out of CHIP, and $\gamma_b \cdot \tau_t$ controls for trends in the long-run follow-up.

²⁰These children are born between the second quarter of 1996 and the first quarter of 2002. The average child – born in the fourth quarter of 1998 – expects to enter Grade 1 in 2004-2005 and expects to graduate high school (Grade 12) in 2016-2017. I track their education outcomes into college using ACS 2010-2018. I detail the sample construction in Appendix C.

long-run trends of single motherhood in column 5-6 yields similar estimates. Removing year-specific effects for both states and single motherhood in column 1-2 also yields similar estimates for children of single mothers. Overall, the long-run impacts of in-utero exposure on college enrollment are not sensitive to the choice of controls in the specification.

To illustrate the identifying variation, Appendix Figure J2 plots the effects on college enrollment by the timing of exposure across cohorts. Children conceived more than 4 quarters before CHIP were not exposed to CHIP in utero, and college enrollment did not increase for these children. College enrollment increased by a small and insignificant amount for children exposed to CHIP in the first and second trimester. The overall effect on college enrollment is concentrated among the full exposure cohort, where a 100% FPL exposure to CHIP increased college enrollment by 1.14 percentage points for children of never married mothers, and by 0.69 percentage points for children of single mothers (Appendix Table I11).

Table 6: Effects of CHIP exposure on college enrollment

	(1)	(2)	(3)	(4)	(5)	(6)
eliginc ^{utero} · single	2.72***	1.58***	2.91***	1.71***	2.60***	1.86***
	(0.60)	(0.41)	(0.58)	(0.40)	(0.99)	(0.70)
eliginc ^{utero}	-1.09*	-1.26**	-0.76	-0.94	-0.74	-1.06
	(0.57)	(0.59)	(0.90)	(0.93)	(0.93)	(0.99)
single						
never married	Y		Y		Y	
state-year FE			Y	Y	Y	Y
single-year FE					Y	Y
y mean	15.7	78%	15.7	78%	15.7	78%
R^2	0.34	0.34	0.35	0.35	0.35	0.35
N	385	,065	385	,063	385	,063

^{***} *p* < 0.01 ** *p* < 0.05 * *p* < 0.10

Notes. Table estimates the effect of CHIP exposure on college enrollment. *single* indicates children of single mothers. In odd-numbered columns, single mothers have never married, whereas in even-numbered columns they are unmarried in the survey year. I examine the robustness of the results with different controls in the specification. The preferred specification in column 3-4 controls for state-year fixed effects for the roll-out of CHIP and for the long-run follow-up. The full specification in column 5-6 further controls for long-run trends in single motherhood with *single*-year fixed effects. I remove both sets of controls in column 1-2. Regressions are weighted by the ACS sampling weights. Robust standard errors clustered at the level of states in the parenthesis.

High School. I also examine the effect of CHIP exposure on children's academic progress through high school, focusing on whether the child attends the grade appropriate for her age (the "grade-for-age" status), and whether the child graduates high school. Appendix Table I12 shows the estimates. In-utero exposure significantly improves grade-for-age.

For children of never married mothers, gaining a 100% FPL exposure in utero lowers late grade entry by 2.36 percentage points, or by 20% below the mean. For children of single mothers, the exposure lowers late grade entry by 1.56 percentage points.

In-utero exposure further improves high school graduation rates for children of single mothers. Gaining a 100% FPL exposure increases high school graduation by 1.71 percentage points for children of never married mothers, or by 5.94% above the mean. Compared to the effect on college enrollment, the increase in high school graduation is smaller in magnitude, suggesting that CHIP exposure increased college enrollment among high school graduates who would not have attended college absent the exposure. Across cohorts, the effects on high school graduation is concentrated among children gaining full exposure to CHIP in utero (Appendix Table I11).

Effect Magnitude. Focusing on college enrollment, I compare the effect of gaining inutero exposure to CHIP with the effect of gaining Medicaid insurance to understand the magnitude of the results. Exploiting Medicaid expansions that simultaneously expanded insurance for pregnant mothers and infants, Miller and Wherry (2019) finds that a ten percentage point increase in Medicaid eligibility in utero and in the first year of life increases college enrollment by 0.35 percentage points.²¹ This effect implies that the rollout of CHIP would increase college enrollment by 0.60 percentage points.²² In practice, the exposure to CHIP increased college enrollment by 1.13 percentage points for children of single mothers, or by 0.29 percentage points for children on average.²³ Therefore, in-utero exposure to CHIP increased college enrollment by around half the direct effect of insurance to pregnant mothers and infants.

6.3 Discussion

Empirical results from Section 5 and 6 show that in-utero exposure to the roll-out of CHIP significantly improved the birth weight and the college enrollment of children. Specifically, the exposure decreased low birth weight by 3.6% and increased college enrollment by 1.13 percentage points for children of single mothers. To understand the magnitude of the effect on birth weight, I compare the effect of CHIP exposure with the direct effects of transfer programs for single mothers. For instance, Hoynes *et al.* (2015) estimates that

²¹Similarly, Levine and Schanzenbach (2009) finds that CHIP eligibility in the first year of life improves Reading scores in Grade 4, providing one pathway for the longer-term effect on attainment.

 $^{^{22}}$ Specifically, CHIP increased children's insurance eligibility by 17 percentage points, leading to an increase in college enrollment by 17*0.035 = 0.60 percentage points.

²³Specifically, CHIP expanded income limits by 80% FPL, increasing college enrollment by 0.8*1.41 = 1.13 percentage points for children of single mothers, or by 25.46%*1.13 = 0.29 percentage points on average.

single mothers receiving a \$1,000 payment of Earned Income Tax Credit (EITC) have lower rates of low birth weight by 6.5%. By comparison, in-utero exposure to the roll-out of CHIP resulted in around half the reduction in low birth weight. Therefore, for both birth weight and college enrollment, in-utero exposure to CHIP improved child outcomes by around half the direct effects of transfers to low-income families and children.

7 Behavioral Mechanisms

I next investigate the behavioral mechanisms using a dynamic model of in-utero investments. In the model, mothers choose smoking and pre-natal visits in each trimester to invest in the birth weight of the child. Because investments occur over multiple trimesters, mothers may delay the onset of investments and end up under-investing in the child. The exposure to CHIP shifted the timing and the level of investments, which I exploit to estimate the behavioral effects of the exposure. I then evaluate welfare based on the model estimates.

7.1 Setting

Pre-Natal Visits. Let v_{it} be the monthly number of visits chosen by mother i in trimester t, where t = 1, 2, 3. Following the medical guideline, I allow the maximum number of visits each month to increase from 1 visit in the first trimester to 2 visits in the second trimester, and finally to 4 visits in the third trimester. The higher frequency of visits in the third trimester matches the guideline recommendation of weekly visits for near-term mothers. In the model, mothers can take a maximum of 21 visits during pregnancy. Empirically, mothers on average take 11 visits during pregnancy.

The out-of-pocket cost of a pre-natal visits is c_{it} . I allow the cost to vary by mother age a_i and a health shock ξ_{it} . The health shock is assumed to be i.i.d. N(0,1), and determines whether the mother pays zero out-of-pocket cost or pays a positive amount, in which case the cost is a quadratic function of mother age. I match the probability of paying positive costs and the level of the costs with empirical moments observed in the Medical Expenditure Panel Survey (MEPS). I show details of the cost calibration in Appendix D.

Mother's utility per visit has a permanent component η_i and a transitory component v_{it} . η_i is a random coefficient reflecting mother's taste for visits. I assume that mother's tastes belong to one of the three types in the population, and the distribution of η_i across types depends on mother characteristics X_i . v_{it} , drawn from i.i.d. $N(0, \sigma_v^2)$, is the transitory taste shock in trimester t. To account for investment costs, I subtract the out-of-pocket cost

per visit c_{it} from the utility, and absorb any non-monetary cost inside the taste type η_i . I therefore measure the utility from pre-natal visits v_{it} in dollar terms using the following equation

$$\vartheta(v_{it}; v_{it}, \xi_{it}, a_i) = 3v_{it} [\eta_i + v_{it} - c_{it}(a_i, \xi_{it})]$$
(14)

Smoking. I discretize mother's smoking decision s_{it} based on cut-points at 5 and 15 cigarettes daily. Mothers consuming fewer than 5 cigarettes daily are non-smokers with $s_{it} = 0$. Mothers consuming 5 to 15 cigarettes, or on average half a pack daily, are light smokers with $s_{it} = 1$. Mothers consuming over 15 cigarettes daily are heavy smokers with $s_{it} = 2$. The utility from an additional level of smoking – roughly a half pack daily – depends on mother i's taste for smoking ζ_i and a transitory taste shock ω_{it} . ζ_i is drawn from the type distribution of mother i, and ω_{it} is drawn from i.i.d. $N(0, \sigma_s^2)$. Since the price of tobacco does not vary by mother characteristics, I absorb the money cost of smoking inside the taste ζ_i . This gives the utility from smoking s_{it} in dollar terms as

$$\psi(s_{it}; \omega_{it}) = s_{it} \left(\zeta_i + \omega_{it} \right) \tag{15}$$

Birth Weight. The child is born at the end of the third trimester. Birth weight, b_i , depends on mother's pre-natal visits and smoking in pregnancy. I specify a flexible birth weight production function as follows

$$log(b_i) = \phi_i + \phi_1 \cdot V_i + \phi_2 \cdot V_i^2 + \phi_3 \cdot smoke_i + \phi_4 \cdot heavy_i + \Phi(V_i, V_i^2, smoke_i, heavy_i) + \sigma \xi_{i3},$$
(16)

where $V_i = 3 \sum_{t=1}^{3} v_{it}$ is the total number of visits of mother *i*. I use the quadratic form to allow for decreasing marginal benefits of visits given by $\phi_1 + 2\phi_2 \cdot V_i$.

 $smoke_i$ and $heavy_i$ are binary variables derived from the average smoking intensity $\bar{s}_i = \frac{1}{3} \sum_{t=1}^3 s_{it}$ in pregnancy. I let $smoke_i = 1$ for mothers consuming more than 5 cigarettes daily $(\bar{s}_i \geq 5)$, and let $heavy_i = 1$ for heavy smokers consuming more than 15 cigarettes daily $(\bar{s}_i \geq 15)$. The effect of smoking on birth weight is captured by ϕ_3 , and the effect of heavy smoking is captured by $\phi_3 + \phi_4$.

I further include flexible interaction terms between pre-natal visits and smoking in $\Phi(V_i, V_i^2, smoke_i, heavy_i)$. The interactions allow the marginal benefit of visits to vary by smoking intensity. Moreover, birth weight depends on an endowment effect ϕ_i drawn from the type distribution of mother i. Larger ϕ_i increases the return of investments on birth weight. Finally, the health shock ξ_{i3} affects birth weight through the parameter σ .

Altruism. Mother's utility from the child's birth weight is given by $L(b_i) = \alpha b_i^{\theta}$, where θ governs the marginal utility of birth weight, and α is mother's utility weight on birth weight. I use α to measure mother's altruism for the child. Larger α increases mother's utility from birth weight relative to her own utility $\vartheta(v_{it}; v_{it}, \xi_{it}, a_i)$ and $\psi(s_{it}; \omega_{it})$ from investments.

Present Bias. I model present bias using the β - δ representation. I fix $\delta = 1$ for quarterly discounting in the long run. In the short run, present-biased mothers choose investments in t = 3 to maximize the following utility

$$U_{3}(v_{i3}, s_{i3}; \varepsilon_{i3}, \mathcal{I}_{i3}) = \vartheta(v_{i3}; v_{i3}, \xi_{i3}, a_{i}) + \psi(s_{i3}; \omega_{i3}) + \beta \mathbb{E}[L(b_{i})|\mathcal{I}_{i3}, v_{i3}, s_{i3}],$$
(17)

where $\varepsilon_{i3} = (v_{i3}, \xi_{i3}, \omega_{i3})$ is the vector of transitory shocks in t = 3. $\mathcal{I}_{i3} = (3\sum_{t=1}^{2} v_{it}, \sum_{t=1}^{2} s_{it}, X_i)$ is the state vector summarizing previous investments and mother characteristics X_i . Let (v_{i3}^*, s_{i3}^*) denote investments in t = 3. The long-run utility implied by the investments is

$$\mathcal{U}_{3}\left(v_{i3}^{*},s_{i3}^{*};\epsilon_{i3},\mathcal{I}_{i3}\right) = \vartheta(v_{i3}^{*};v_{i3},\xi_{i3},a_{i}) + \psi(s_{i3}^{*};\omega_{i3}) + \mathbb{E}\left[L(b_{i})|\mathcal{I}_{i3},v_{i3}^{*},s_{i3}^{*}\right]. \tag{18}$$

Compared to the long-run utility (equation 18), present-biased mothers under-value the utility from birth weight in equation 17, discounting $\mathbb{E}[L(b_i)|\mathcal{I}_{i3}, v_{i3}, s_{i3}]$ by $\beta < 1$.

In t=2, present-biased mothers discount future utility U_3 by $\beta < 1$, and choose investments to maximize the following utility

$$U_2(v_{i2}, s_{i2}; \varepsilon_{i2}, \mathcal{I}_{i2}) = \vartheta(v_{i2}; v_{i2}, \xi_{i2}, a_i) + \psi(s_{i2}; \omega_{i2}) + \beta \mathbb{E}[\mathcal{U}_3 | \mathcal{I}_{i2}, v_{i2}, s_{i2}],$$
 (19)

where \mathcal{U}_3 is the long-run utility implied by investments in t=3 (equation 18). Maximizing equation 19 yields investments (v_{i2}^*, s_{i2}^*) in t=2. Recursively, the long-run utility implied by (v_{i2}^*, s_{i2}^*) is discounted by β to determine investments in $t=1.^{24}$ I solve for the investment profile $\left(v_{it}^*, s_{it}^*\right)_{t=3,2,1}$ of mother i from equation 17 to 20, discounting the long-run utility in the next period by β to determine investments in the current period. I then match the profiles with empirical patterns in the data to estimate the model parameters.

$$U_1(v_{i1}, s_{i1}; \varepsilon_{i1}, \mathcal{I}_{i1}) = \vartheta(v_{i1}; v_{i1}, \xi_{i1}, a_i) + \psi(s_{i1}; \omega_{i1}) + \beta \mathbb{E}[\mathcal{U}_2 | \mathcal{I}_{i1}, v_{i1}, s_{i1}], \tag{20}$$

where U_2 is the long-run utility implied by investments in t = 2.

²⁴Specifically, investments in t = 1 maximize the utility

7.2 Estimation

I estimate the model parameters using the method of simulated moments (MSM). To quantify the behavioral effects of the exposure, I allow altruism α and present bias β to vary by mother's exposure to CHIP, and match investments across CHIP exposure to estimate the effects on preferences. I detail the construction of the exposure measure and the moment conditions identifying the model parameters below.

CHIP Exposure. Building on the definition of in-utero exposure *eliginc* (equation 6), I construct CHIP exposure $\Delta \ell_{js} = eliginc_{js} - inc_s^{pre}$ as the change in *eliginc* due to the onset of CHIP. Intuitively, $\Delta \ell_{js}$ is larger in states with greater expansion of insurance, and for mothers gaining earlier exposure to CHIP. Specifically, for mothers starting pregnancy within 9 months before CHIP $(-9 \le j \le -1)$, $\Delta \ell_{js} = (1 + \frac{j}{9}) \cdot \Delta inc_s$ increases with j for cohorts with earlier exposure to CHIP, and increases with expansion size $\Delta inc_s = inc_s^{post} - inc_s^{pre}$. Cohorts gaining full exposure to CHIP $(j \ge 0)$ have $\Delta \ell_{js} = \Delta inc_s$, and cohorts never exposed to CHIP in utero $(j \le -10)$ have $\Delta \ell_{js} = 0$.

I calculate $\Delta \ell_{js}$ for 324,400 non-college educated single mothers starting pregnancy within one year before CHIP ($-12 \le j \le -1$). These mothers lived in states where the income limit was between 110% and 130% FPL before CHIP, and the expansion was between 20% and 90% FPL after CHIP. This restriction allows me to estimate the behavioral effects of CHIP exposure starting from a common baseline exposure level across states. ²⁵ I discretize $\Delta \ell_{js}$ into 5 nodes, corresponding to 0%, 10%, 30%, 50%, and 70% FPL exposure to CHIP. I assign mothers with $j \le -10$ to the zero exposure node, and assign mothers with $j \ge -9$ to the nearest positive node to construct CHIP exposure $\Delta \ell_i$ for mother i.

Behavioral Effects. To model the behavioral effects, I allow mother's altruism α_i to differ across CHIP exposure $\Delta \ell_i$ according to the following equation

$$\alpha_i = \alpha_0 + \alpha_1 \Delta \ell_i, \tag{21}$$

where the intercept α_0 is the baseline altruism for mothers unexposed to CHIP ($\Delta \ell_i = 0$). The slope $\alpha_1 = \frac{\alpha_i - \alpha_0}{\Delta \ell_i}$ gives the change in altruism for a unit increase in CHIP exposure. Starting from the baseline altruism α_0 , the slope α_1 determines the altruism for mothers with different exposure to CHIP. In practice, I normalize $\alpha_0 = 1.26$ Similarly, I specify a

²⁵91% of single mothers lived in states where the income limit before CHIP was between 110% and 130% FPL. I show details of the sample construction in Appendix E.

²⁶This is because mother's utility weight for her own investments is already estimated in the taste types η_i and ζ_i . Different scaling of α_0 does not affect the relative utility weights between the mother and the child.

behavioral equation for present bias

$$\beta_i = \beta_0 + \beta_1 \, \Delta \ell_i \,, \tag{22}$$

where β_0 is the baseline present bias for $\Delta \ell_i = 0$, and $\beta_1 = \frac{\beta_i - \beta_0}{\Delta \ell_i}$ is the change in present bias for a unit increase in CHIP exposure. I therefore estimate parameters $(\alpha_1, \beta_0, \beta_1)$ to quantify the behavioral effects of CHIP exposure.

Moment Conditions. Based on equation 21 and 22, I quantify the behavioral effects using CHIP exposure as a shifter of mother preferences. An increase in the altruism tends to increase mother's overall investments in the child, whereas an effect on present bias further impacts the timing of investments across trimesters of pregnancy. I therefore match the level and the timing of investments across CHIP exposure to quantify the behavioral effects

- 1. percent of mothers starting pre-natal visits in the first, second, and the third trimester at each exposure level in $\Delta \ell_i$,
- 2. number of pre-natal visits at each exposure level in $\Delta \ell_i$,
- 3. percent of smokers at each exposure level in $\Delta \ell_i$.

To estimate the birth weight production function in equation 16, I construct moment conditions exploiting the exposure levels in $\Delta \ell_i$ as instruments.²⁷ Specifically, I match the reduced-form relationship between birth weight and $\Delta \ell_i$ in addition to the first-stage relationship between investments and $\Delta \ell_i$. The relative shifts in birth weight and investments across different $\Delta \ell_i$ inform the production of birth weight from investments. I also match the empirical patterns between birth weight and investments in a separate set of moment conditions.

I further include a large number of auxiliary moment conditions to capture additional investment responses and heterogeneity across mothers. Compared to the moment conditions identifying the behavioral effects and the production of birth weight, the auxiliary moments receive lower weights in the estimation. In total, I employ 272 moment conditions to estimate the model parameters. I detail the full list of moment conditions and the estimation procedure in Appendix F.

Mother Types. I model the distribution of mother types using a multinomial Probit. Given mother characteristics X_i , the multivariate Probit generates the probability distribution

²⁷The production function in equation 16 requires three instruments for three endogenous investments: pre-natal visits V_i and smoking indicators $smoke_i$ and $heavy_i$. I use indicators for the five exposure levels in $\Delta \ell_i$ as instruments in the moment conditions.

of mother i across three potential types in the population. Specifically, the probability of being type k = 0, 1, 2 for mother i is the follows

$$P_i^k = P^k(X_i; \pi^k) = \frac{F(\pi^k \cdot X_i)}{\sum_{n=0,1,2} F(\pi^n \cdot X_i)},$$
 (23)

where F is the cumulative distribution function of a standard normal. X_i includes mother age, whether the mother had fetal death in previous pregnancies, has any maternal risk factor on the birth certificate, and the smoking rate in the mother's county of residence. I summarize mother characteristics in Appendix E. I illustrate the identification of mother types from mother characteristics with estimation results. As is standard in multinomial Probit, I normalize the coefficients for one of the types (type 1) to zero: $\pi^1 = 0$.

7.3 Results

Behavioral Effects. The top panel of Table 7 estimates the behavioral effects of CHIP exposure on altruism and present bias. The exposure had little effect on the altruism of mothers. The estimated slope parameter α_1 is small and indistinguishable from zero. Therefore the investments are not driven by higher altruism for the child. By contrast, CHIP exposure significantly lowered the present bias of mothers. Gaining a 100% FPL exposure increases mother's short-term patience by $\beta_1 = 0.13$. Compared to the baseline patience ($\beta_0 = 0.75$), the roll-out – which expanded income limits by 80% FPL – increased short-term patience to 0.85 (= $0.75 + 80\% \cdot 0.13$) for mothers with full exposure to CHIP in pregnancy, lowering their present bias by 14.3% (= $80\% \cdot 0.13/0.75$).

I illustrate the identification of the behavioral effects comparing simulated investments with empirical counterparts in the data. In Figure 3, the share of mothers staring care in the first trimester increased with CHIP exposure (panel a), and the share starting care in the third trimester decreased with the exposure (panel b). The simulation matches these patterns in the data. Despite earlier onset of visits, the number of visits increased little with CHIP exposure (panel c), and smoking decreased with exposure in panel (d). The simulation matches these patterns in the data. Thus, the early onset of visits and the reduction in smoking are primarily explained by the behavioral effect on present bias.

Mother Types. The middle section of Table 7 summarizes mother's taste types and the birth weight endowment. The majority of mothers (53.16%) are type 2 mothers who derive utility from pre-natal visits and dis-utility from smoking. Another large share of mothers (37.31%), the type 1 mothers, derive significant disutility from smoking and have

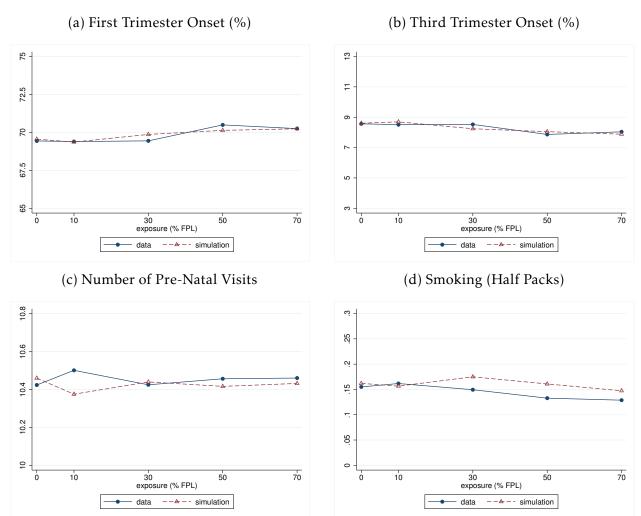
Table 7: Estimated model parameters

Behavioral Effects			
α_0 :	1	α_1 :	0.001
	_		(0.006)
β_0 :	0.75	eta_1 :	0.13
	(0.071)		(0.040)
Mother Types	Type 0	Type 1	Type 2
Taste for visits η_i	-0.017	-0.38	3.76
	(0.067)	(0.27)	(0.32)
Taste for cigar. ζ_i	38.77	-947.77	-17.69
	(8.30)	(<0.001)	(1.85)
Endowment ϕ_i	3.56	2.06	2.00
	(0.40)	(0.16)	(0.15)
Share (%)	9.53	37.31	53.16
Birth Weight Produ	ıction		
V_i	1.54	$V_i \cdot smoke$	0.61
	(0.010)		(0.022)
V_i^2	-0.093	$V_i \cdot heavy$	4.72
•	(<0.001)	•	(3.20)
$smoke\ (\bar{s}_i \geq 5)$	-10.54	$V_i^2 \cdot smoke$	0.025
	(0.022)	ı	(0.001)
$heavy \ (\bar{s}_i \ge 15)$	-4.95	$V_i^2 \cdot heavy$	-0.77
• ,	(7.19)	,	(0.32)
Birth Weight Valua	tion		
θ :	0.53		
	(0.011)		

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table shows estimates of the model parameters. The top panel shows estimates of the behavioral effects in equation 21 and 22. The middle panel shows estimates of mother types – her tastes for visits, smoking, and the birth weight endowment – and the share of each type in the population. The last panel shows estimates of the birth weight production function (equation 16) and mother's marginal utility from birth weight captured by θ . Standard errors of the estimated parameters in the parenthesis.

Figure 3: Investment responses to CHIP exposure, model fit



Notes. Figure compares simulated investments with observed investments for different exposure levels in $\Delta \ell_i$. I focus on the timing of care onset in panel (a) and panel (b), the number of pre-natal visits in panel (c), and smoking intensity in half packs (10 cigarettes) daily in panel (d). I simulate investments for ten million non-college educated single mothers and plot simulated investments by exposure levels in dotted lines. I plot the empirical counterparts in solid lines.

near-zero probability of smoking in pregnancy. Their taste for visits is not statistically different from zero. The remaining share of mothers (9.53%) are type 0 mothers who have strongly prefer smoking, almost always smoke in pregnancy, and derive zero utility from visits. The birth weight endowment is significantly larger for type 0 mothers.

To illustrate the identification of mother types, I show in Appendix Figure J3 that simulated investments recover differences by mother characteristics exploited in the multinomial Probit. Specifically, simulated number of visits matches the difference by mother risk factors in panel (a), and simulated smoking intensity closely matches empirical counterparts across county smoking rates in panel (b). Moreover, birth weight is lower for mothers with fetal death in previous pregnancies, and the simulation matches this difference in the data (Appendix Figure J4).

Birth Weight Production. The lower panel of Table 7 estimates the birth weight production function in equation 16. Pre-natal visits significantly increase birth weight, but the marginal effect diminishes with larger number of visits. By contrast, smoking has significant detrimental impacts on birth weight. The relative magnitude between the two effects suggests that the negative impacts of smoking are comparable to missing 7 pre-natal visits in pregnancy. The interaction terms between visits and smoking imply that the optimal number of visits is around 9 for non-smoking mothers, and around 15 for smokers.

The endowment ϕ_i impacts birth weight by increasing the return to investments. Highendowment mothers have higher returns to investments and may invest more in response to the exposure. To illustrate this point, I simulate investments and birth weight across mother types in Appendix Figure J5. Consistent with higher return to investments, type 0 mothers significantly reduced smoking (panel c) despite having strong tastes for smoking, and further increased the number of visits by a small amount despite having near-zero tastes for visits (panel b). By contrast, the investment responses from type 1 and type 2 mothers are very modest. As a result, the overall increase in birth weight is highly concentrated among children of type 0 mothers (panel d).

7.4 Welfare

I next use the model estimates to quantify the welfare impacts of the exposure. Specifically, I reveal mother's valuation of the exposure from her investments, and ask how her valuation compares with the social cost of the exposure and with expansions of own insurance based on the marginal value of public funds (Finkelstein and Hendren, 2020).

Utility. I use mother's long-run utility to evaluate the welfare impacts of investments. Under the welfare metric, optimal investments maximizing the long-run utility generate welfare \mathcal{U}^{*} . By deviating investments away from the long-run optimal, present bias results in welfare loss $\frac{\mathcal{U}^{*}-\mathcal{U}^{i}}{\mathcal{U}^{*}}$ relative to the long-run utility \mathcal{U}^{*} , where \mathcal{U}^{i} represents welfare in the status quo (i=1) and in the absence of CHIP (i=0). Compared to the absence of CHIP, the status quo exposure to the roll-out lowered present bias and increased welfare by $\frac{\mathcal{U}^{1}-\mathcal{U}^{0}}{\mathcal{U}^{*}}$. I summarize this welfare effect in Table 8.

CHIP exposure increased mother's long-run utility by 0.063%. Across mother types, this effect is mainly concentrated among type 0 mothers, where long-run utility increased by 0.20%. To understand how investments contribute to the welfare effect, I calculate the utility from investments M^i and the utility from the child's birth weight C^i .²⁹ Investments decreased long-run utility more for type 0 and type 1 mothers, but the benefit on birth weight more than offset the costs. On net, the exposure increased utility the most for type 0 mothers due to the large benefit on birth weight.

MVPF. Since I measure mother utility in dollars, the utility difference $\mathcal{U}_i^1 - \mathcal{U}_i^0$ is also mother's willingness to pay (WTP) for the exposure. Specifically, mother i is willing to pay up to the utility difference for the exposure compared to receiving zero exposure to CHIP. Formally, I calculate the WTP as

$$WTP_{i} = \frac{U_{i}^{1} - U_{i}^{0}}{\Delta \ell_{i}} = \frac{C_{i}^{1} - C_{i}^{0}}{\Delta \ell_{i}} + \frac{M_{i}^{1} - M_{i}^{0}}{\Delta \ell_{i}},$$
(25)

where I normalize the utility difference $\mathcal{U}_i^1 - \mathcal{U}_i^0$ by the exposure $\Delta \ell_i$ to calculate the WTP for a 100% FPL exposure to CHIP. I similarly calculate the WTP for birth weight $\frac{C_i^1 - C_i^0}{\Delta \ell_i}$ and for investments $\frac{M_i^1 - M_i^0}{\Delta \ell_i}$.

I then compare mother's WTP with the social cost of the exposure to construct the marginal value of public funds (MVPF) for the exposure (Finkelstein and Hendren 2020; Hendren and Sprung-Keyser 2020). I measure the social cost using the spending on

$$\mathcal{U}(I) = \mathbb{E}\left[\sum_{t=1}^{3} \vartheta(v_t; \varepsilon_t, \mathcal{I}_t) + \sum_{t=1}^{3} \psi(s_t; \varepsilon_t, \mathcal{I}_t) + L(b^i; \varepsilon_3, \mathcal{I}_3)\right],\tag{24}$$

which sums over utility in each trimester applying the long-run discount factor $\delta = 1$. Optimal investments I^* maximizing equation 24 give the long-run utility $\mathcal{U}^* = \mathcal{U}(I^*)$.

²⁹Specifically, mother utility $\mathcal{U}^i = C^i + M^i$ is the sum of utility from birth weight and investments. I therefore decompose the welfare effect $\frac{\mathcal{U}^1 - \mathcal{U}^0}{\mathcal{U}^*}$ by the effect on birth weight $\frac{C^1 - C^0}{C^*}$ and on investments $\frac{M^1 - M^0}{M^*}$ in Table 8.

²⁸Formally, given investment profile $I = (v_t, s_t)_{t=1,2,3}$, the long-run utility is given by

Table 8: Effects of CHIP exposure on birth weight and investments

	All	Type 0	Type 1	Type 2
$\frac{\mathcal{U}^1-\mathcal{U}^0}{\mathcal{U}^*}$ (%)	0.063	0.20	0.066	0.037
	(<0.001)	(0.001)	(<0.001)	(<0.001)
$\frac{C^1-C^0}{\mathcal{U}^*}$ (%)	0.12	0.35	0.21	0.012
	(<0.001)	(0.001)	(<0.001)	(<0.001)
$\frac{M^1-M^0}{\mathcal{U}^*}$ (%)	-0.057	-0.15	-0.15	0.025
	(<0.001)	(0.001)	(<0.001)	(<0.001)
Share (%)	100	9.53	37.31	53.16

Notes. Table summarizes the welfare effect of CHIP exposure relative to the long-run utility \mathcal{U}^* . I calculate utility from the child's birth weight C^i and from mother investments M^i , where subscript i indicates utility in the status quo (i=1) and in the absence of CHIP (i=0). I then summarize welfare separately for birth weight and for investments using the equation $\frac{\mathcal{U}^1-\mathcal{U}^0}{\mathcal{U}^*}=\frac{C^1-C^0}{\mathcal{U}^*}+\frac{M^1-M^0}{\mathcal{U}^*}$. Standard errors of the welfare effects from ten million simulated individuals in the parenthesis.

program outreach during the roll-out. Since the goal of the outreach was to introduce the program to the public (Williams and Rosenbach, 2007), I assume an even distribution of outreach spending across households. This gives a cost of exposure $\Delta G = \$0.42$ per household.³⁰

I calculate the MVPF for CHIP exposure as follows

$$MVPF = \varphi \frac{WTP}{\Lambda G} = \varphi \frac{WTP^C}{\Lambda G} + \varphi \frac{WTP^M}{\Lambda G}, \qquad (26)$$

where $WTP = \frac{1}{N} \sum_{i} WTP_{i} \Delta inc_{s(i)}$. Because CHIP exposure is larger in states expanding insurance to higher income limits, I scale WTP_{i} by the size of expansion $\Delta inc_{s(i)} = inc_{s(i)}^{post} - inc_{s(i)}^{pre}$ to calculate the average WTP in the roll-out. To the extent that society may value the transfer to pregnant mothers more than the dollar costs, I allow for higher welfare weight $\varphi > 1$ for pregnant mothers.

Table 9 summarizes the MVPF. Mothers are willing to pay \$0.49 for the benefit of the exposure on birth weight, which more than offsets the program spending on outreach (\$0.41). Net of investment costs, mothers are willing to pay \$0.31 for the exposure, and hence value the exposure at $\frac{\$0.31}{\$0.41} = 76\%$ of the spending. To understand magnitude, I ask how mothers value the exposure relative to expansions of own insurance. For instance, according to recent Medicaid expansions in Oregon and Massachusetts, low-income adults

³⁰I detail the calculation of outreach costs and examine robustness in Appendix G.

value the expansion of own insurance between 55% and 116% of the program spending (Finkelstein *et al.* 2019a; Finkelstein *et al.* 2019b). By comparison, the MVPF of CHIP exposure falls in the middle range of these estimates.³¹ This suggests that, for each dollar of program spending, mothers value the exposure to children's insurance as much as expansions of own insurance.

For the insurance program, this result suggests that the information of insurance can effectively "nudge" investments by reducing the behavioral biases of parents. Parents value the information because they value the benefit on birth weight. Thus, in addition to improving child outcomes, outreach efforts encouraging parental investments in the child also improve parents' own utility as effectively as direct expansions of insurance.

Table 9: MVPF of CHIP Exposure

	WTP	WTP^C	WTP^{M}	MVPF
$\varphi = 1$	0.31	0.49	-0.18	0.76
	(0.001)	(0.002)	(0.001)	(0.002)
$\varphi = 2$	0.62	0.98	-0.36	1.51
	(0.002)	(0.003)	(0.003)	(0.005)
$\varphi = 3$	0.93	1.47	-0.54	2.27
	(0.003)	(0.005)	(0.004)	(0.007)

Notes. Table summarizes the marginal value of public funds (MVPF) of CHIP exposure. To calculate the MVPF, I divide mother's WTP by the cost of the exposure (\$0.41) measured by the spending on program outreach. I calculate separate WTP for birth weight WTP^C and for investments WTP^M , and summarize the MVPF varying mother's welfare weight φ in the table. Standard errors from ten million simulated individuals in the parenthesis.

7.5 Fiscal Externality

Because MVPF is based on mother's WTP for the exposure, social benefits not internalized by mother's investments are not captured in the welfare metric. One example of social benefit is the fiscal impact of investments on program costs, which may operate through the long-run effects on education and earnings. In the roll-out of CHIP, since mothers investing more in utero did not expect higher education for the child, the increase in college enrollment and the potential impacts on program costs are not internalized in

³¹See Appendix Table D.I of Hendren and Sprung-Keyser (2020) for a summary of these estimates.

mother's WTP for the exposure. Here, I quantify the fiscal externality predicting future tax payments from college enrollment.

Specifically, Table 6 estimates that the roll-out of CHIP increases college enrollment by $80\% \cdot 1.71 = 1.37$ percentage points for children of single mothers. Assuming that college students induced by the exposure attend college for two years, and applying an 11.3% return on earnings for each year of college enrollment (Zimmerman, 2014), I calculate that exposure to the roll-out increases earnings by $2 \cdot 11.3\% \cdot 0.0137 = 0.31\%$ for children of single mothers. Discounted to year 1997, the life-cycle earning benefit amounts to \$1,101.11 per child of single mother under a 2% annual discount rate.

Assuming that the earning benefit is subject to a 18.9% marginal tax rate (Hendren and Sprung-Keyser, 2020), exposure to the roll-out increases tax payments by $18.9\% \cdot \$1,101.11 = \208.11 per child of single mother, or by $\frac{\$208.11}{\$2,483.10} = 8.4\%$ of the initial program cost in childhood. Hence, parental responses to the roll-out could lower the net cost of the program by 8.4% in the long run. I examine robustness in Appendix H. Fiscal externality increases to 15.4% of the cost when students enroll in college for 4 years, and drops to 6.0% assuming a 2-year enrollment and a 3% annual discount rate. Compared to the fiscal externality of program investments in children, parental responses to the roll-out could further lower the program cost by over 6% in the long run. 32

8 Discussion and Conclusion

Social policies for children increasingly harness parental investments to augment the policy impacts on children. In K-12 and pre-school, for instance, parent-school partnerships and family-based interventions lower the informational and behavioral frictions facing parents, improving their investments and the education outcome of the child.³³ To effectively engage parents, policymakers need information on how parents invest in the child and how responses to policies might impact investments and child outcomes. In the context of social insurance for children, I provide evidence on these questions exploiting the roll-out of the Children's Health Insurance Program (CHIP) in 1997-2000.

I find that expanding insurance for children increases parental investments in the child. Pregnant mothers exposed to greater insurance expansions due to CHIP reduced smoking

³²Focusing on children born in the early 1980s, Brown *et al.* (2020) estimates that each childhood year of insurance eligibility increases tax payments in age 19-28 by \$178 in 2000 dollars. Exposure to the roll-out of CHIP predicts an increase in tax payments between \$12 and \$32 by age 28, further reducing the net cost of the program by 6.7% to 18.0% in the long run.

³³See Bergman (2019) for a survey of parental engagements in K-12, and see Brooks-Gunn *et al.* (2000) for a survey in pre-school.

and improved the early onset of pre-natal visits. Using a dynamic model of in-utero investments, I show that CHIP exposure increased investments by lowering the present bias of mothers, and increased mother utility due to the benefit on birth weight. This result suggests that mothers highly value child outcomes, but nonetheless under-invest due to behavioral biases limiting their investments in the child.

The behavioral mechanism has several implications for policy. First, parents may exhibit short-term biases due to unawareness of future investment opportunities for the child. Informing parents of program eligibility through outreach and reminders can foster forward thinking and increase investments, with potentially larger effects on more present-biased parents (Mayer *et al.*, 2019). Second, effective engagement strategies can feature low-cost, light-touch interventions that "nudge" investments without significant financial incentives to parents. Because parents highly value child outcomes, the intervention could improve parental utility as effectively as expansions of parents' own insurance.

Finally, combating the behavioral biases also generates social benefits that lower the fiscal cost of insurance. This is because fetal and early-life environments have persistent impacts on later-life outcomes (Almond and Currie 2011; Almond *et al.* 2018), implying large social externality from parental investments. Investment responses to the roll-out of CHIP, for instance, can lower the net cost of the program by 8.4% through the increase in earnings and tax payments. These benefits provide strong motivations for outreach efforts engaging parents in children's insurance programs.

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Appendix

A Insurance Eligibility

I simulate children's insurance eligibility using the sample of all women between age 21 and 40 in the 2000 decennial census. In state s and time t, child i in age band b is eligible for Medicaid/CHIP if the family income $finc_i$ is below the income limit inc_{st}^b . I parameterize the income limit in terms of the share of children eligible for insurance as follows

$$eligCHIP_{st}^b = \frac{1}{N} \sum_{i} 1\{finc_i \le inc_{st}^b\}, \tag{A1}$$

where N = 1,948,731 is the number of women in age 21-40 in the 2000 census. That is, I assume that each woman in the sample had a child in age band b, and calculate the share of women whose children would be eligible for Medicaid/CHIP given the income limit inc_{st}^b . Thus, $eligCHIP_{st}^b$ captures insurance eligibility in a fixed population of mothers and their incomes. Differences in $eligCHIP_{st}^b$ are due to exogenous variation in program rules that shifted income limits across state, time, and age.

I calculate $eligCHIP_{st}^b$ for three age bands: infants (age 0), small children (age 1-5) and older children (6+). Changes in the income limits across states by age bands are listed in Table 1. I average out age groups to calculate simulated eligibility $eligCHIP_{st}$ in childhood:

$$eligCHIP_{st} = \frac{1}{19}eligCHIP_{st}^{0} + \frac{5}{19}eligCHIP_{st}^{1-5} + \frac{13}{19}eligCHIP_{st}^{6+},$$
 (A2)

where instead of using b in the superscript, I make explicit the ages in the age band.

The income limits imply that children from low-income families are more likely to qualify for Medicaid/CHIP. To explicitly account for the eligibility differences by incomes, I calculate separate eligibility for mother demographic groups that differ by marital status, race, and education. Specifically, eligibility for children in age b and group g is given by

$$eligCHIP_{gst}^b = \frac{1}{N_g} \sum_{i \in \mathcal{G}} 1\{finc_i \le inc_{st}^b\}, \tag{A3}$$

where \mathcal{G} is the collection of children in demographic group g. $eligCHIP_{gst}^b$ is the share of children in age b and group g who would be eligible for Medicaid/CHIP in a fixed population of mothers and incomes. Within group g, differences in eligibility $eligCHIP_{gst}^b$ are due to exogenous variation in income limits across state, time, and age.

I then average out ages to calculate simulated eligibility $eligCHIP_{gst}$ in childhood

$$eligCHIP_{gst} = \frac{1}{19}eligCHIP_{gst}^{0} + \frac{5}{19}eligCHIP_{gst}^{1-5} + \frac{13}{19}eligCHIP_{gst}^{6+}.$$
 (A4)

Note that $eligCHIP_{gst}$ is specific to income limits known in time t. To calculate in-utero

exposure, I use the average eligibility in a 9-month pregnancy

$$eligCHIP_{g(i)st} = \frac{1}{9} \sum_{\tau=0}^{8} eligCHIP_{gs\tau}, \tag{A5}$$

where $eligCHIP_{gs\tau}$ is eligibility in month τ of the pregnancy, and $eligCHIP_{g(i)st}$ gives the in-utero exposure for mothers starting pregnancy in time t. I plot $eligCHIP_{g(i)st}$ across states and cohorts during the roll-out in Figure 1, and use it as the simulated exposure to children's insurance in Section 5.4. Eligibility increased substantially from 0.15 to 0.30 between 1997 and 2000 (or from 2.85 eligible years to 5.70), and increased much more for children of single mothers from 0.33 to 0.50 over the same period.

In most of the empirical analyses, I simply use the income limit of Medicaid/CHIP as the exposure measure (without calculating the implied share of children eligible), and use the average income limit in a 9-month pregnancy as mother's in-utero exposure to children's insurance. Specifically, in-utero exposure is given by

$$eliginc_{st} = \frac{1}{9} \sum_{\tau=0}^{8} inc_{s\tau}, \tag{A6}$$

where $inc_{s\tau} = \frac{1}{19}inc_{s\tau}^0 + \frac{5}{19}inc_{s\tau}^{1-5} + \frac{13}{19}inc_{s\tau}^{6+}$ is the income limit known in state s and month τ of pregnancy. $eliginc_{st}$ gives the in-utero exposure for mothers starting pregnancy in time t in state s.

B Detailed Proofs

I solve parents' investment problem starting from the childhood stage in t = 1. Given child health h = 0, 1, parents choose non-health investment v_1^h to maximize the utility

$$U_1^h(v_1^h) = u\left(Y_1 - v_1^h - OOPC \cdot 1\{h = 0\}\right) + \Gamma(s(v_1^h)) + \delta V^h(s(v_1^h)), \tag{B1}$$

where $u(Y_1 - v_1^h - OOPC \cdot 1\{h = 0\})$ is the utility from consumption after investing v_1^h . A less healthy child (h = 0) incurs medical expenses OOPC. Non-health investments improve education $s(v_1^h)$ and adult outcomes summarized in $V^h(s(v_1^h))$. Optimal investments v_1^{h*} satisfy the first order condition

$$u'(c_1^{h*}) = \Gamma' s'(v_1^{h*}) + \delta V' s'(v_1^{h*}), \tag{B2}$$

where $c_1^{h*}=Y_1-v_1^{h*}-OOPC\cdot 1\{h=0\}$ is the optimal consumption given child health h. Equation B2 states that optimal investment v_1^{h*} balances the marginal cost of investment $u'(c_1^{h*})$ with the marginal benefit on the child's education and adult outcomes $\Gamma's'(v_1^{h*})+\delta V's'(v_1^{h*})$. When CHIP reduces OOPC, a decrease in v_1^{h*} would lower the marginal cost on the left hand side but increase the marginal benefit on the right hand side assuming concavity for Γ , s, and V. Therefore, v_1^{h*} must increase when CHIP reduces OOPC. Since OOPC=0 for the healthy child, non-health investment is larger in the healthy child. This implies that the marginal benefit of investment is larger for the less healthy child, and from equation B2, so is the marginal cost of investment. Therefore, total investment is larger in the less healthy child, but non-health investment is smaller due to the medical expenses OOPC.

Let $\tilde{U}_1^h = U_1^h(v_1^{h*})$ denote the maximized utility in childhood given optimal investments v_1^{h*} . In the fetal stage (t=0), parents choose in-utero investment v_0 to maximize utility

$$U_0(v_0) = u(c_0) + w(v_0) + \delta \rho(v_0) \tilde{U}_1^1 + \delta (1 - \rho(v_0)) \tilde{U}_1^0, \tag{B3}$$

where consumption $c_0 = Y_0 - v_0$, and $w(v_0)$ is parents' utility from birth weight as a function of in-utero investment v_0 . Larger v_0 also increases the probability of giving birth to a healthy child, and the probability $\rho(v_0)$ is concave in v_0 . Optimal in-utero investment v_0^* satisfies

$$\delta \rho'(v_0^*) \Delta \tilde{U}_1 + w'(v_0^*) = u'(c_0^*), \tag{B4}$$

where $\Delta \tilde{U}_1 = u(c_1^*) - u(c_0^*) + \Gamma(s(v_1^{1*})) - \Gamma(s(v_1^{0*})) + \delta V^1(s(v_1^{1*})) - \delta V^0(s(v_1^{0*})) = \Delta u + \Delta \Gamma + \delta \Delta V$ is parents' utility gap in own consumption (Δu) , child's education $(\Delta \Gamma)$ and adult outcomes (ΔV) . Because reducing OOPC increases non-health investment in the less healthy child, CHIP narrows the gap in child outcomes $\Delta \Gamma + \Delta V$ by child health. From equation B2, the investment lowers the marginal benefit on the right hand side, implying higher consumption levels for parents of the less healthy child. Therefore, consumption gap Δu also decreases after CHIP. As a result, CHIP reduces the overall utility gap $\Delta \tilde{U}_1 = \Delta u + \Delta \Gamma + \delta \Delta V$, and hence reduces marginal benefit $\delta \rho'(v_0^*) \Delta \tilde{U}_1$ in equation B4. In

response, v_0 decreases after CHIP.

Altruism. I model altruism as the weight on child outcomes in parents' utility. Suppose that exposure to the roll-out increases the utility weight on child outcomes to $\alpha > 1$. Optimal investments in t = 1 now solve

$$u'(c_1^{h*})/\alpha = \Gamma' s'(v_1^{h*}) + \delta V' s'(v_1^{h*}), \tag{B5}$$

and in-utero investment v_0 solves

$$\delta \rho'(v_0^*) \Delta \tilde{U}_1(\alpha) + \alpha w'(v_0^*) = u'(c_0^*),$$
 (B6)

where the utility gap $\Delta \tilde{U}_1(\alpha) = \Delta u + \alpha (\Delta \Gamma + \delta \Delta V)$ places weight α on the gap in child outcomes.

When CHIP increases altruism α in addition to lowering OOPC, non-health investments increase for both health types in equation B5. The additional investments lower consumption c_1^{1*} for parents of the healthy child. For parents of a less healthy child, because OOPC also decreases after CHIP, the change in consumption is ambiguous. In the case that parental consumption is a normal good, decreasing OOPC increases consumption c_1^{0*} . This would imply that the consumption gap Δu narrows after CHIP. However, $\Delta\Gamma + \delta\Delta V$ depends on the size of investment responses and the marginal effects of investments ($\Gamma's' + \delta V's'$) by child health. These quantities are indeterminate from equation B5. In general, altruism could increase parents' perception of investment benefits by increasing their valuation of child outcomes in the utility gap $\Delta \tilde{U}_1(\alpha) = \Delta u + \alpha (\Delta\Gamma + \delta\Delta V)$, 34 potentially increasing in-utero investments based on equation B6.

Present Bias. I model present bias using the $\beta-\delta$ representation in Laibson (1997). Present-biased parents discount future utility to the current period by an additional factor $\beta < 1$. When investments have delayed payoffs, present bias generates over-sensitivity to immediate costs and predicts under-investments in the short run. Specifically, investments of present-biased parents satisfy the following condition in childhood

$$u'(c_1^{h*})/\alpha = \Gamma' s'(v_1^{h*}) + \beta \delta V' s'(v_1^{h*}), \tag{B7}$$

and the following condition in utero

$$\beta \,\delta \,\rho'(v_0^*) \,\Delta \tilde{U}_1(\alpha, \beta) + \alpha \,w'(v_0^*) = u'(c_0^*), \tag{B8}$$

where utility gap $\Delta \tilde{U}_1(\alpha, \beta) = \Delta u + \alpha (\Delta \Gamma + \delta \Delta V)$ depends on investments v_1^{h*} under present bias β and altruism α .

When CHIP increases α and β in addition to lowering *OOPC*, non-health investments increase even more for both health types. However, the effect on consumption is am-

³⁴For illustration, consider the special case where $u(c) = \log(c)$, and $\Gamma(s(v)) + \delta V(s(v)) = (1 + \delta k) \log(v)$, where k captures the return of education on adult outcomes. It is easy to solve that $\Delta u + \alpha \left(\Delta \Gamma + \delta \Delta V\right) = (1 + \alpha + \alpha \delta k) \log \left(\frac{Y_1}{Y_1 - OOPC}\right)$, which may increase with $\alpha > 1$ from the exposure to CHIP.

biguous for parents of a less healthy child, and the gap in child outcomes may change depending on the investment responses and the marginal effects of investments by child health. In general, the behavioral effects allow parents' perception of investment benefit $\beta \, \delta \, \rho'(v_0^*) \, \Delta \tilde{U}_1(\alpha, \beta)$ to increase with β and α from the exposure, ³⁵ potentially increasing in-utero investments based on equation B8.

³⁵Continuing the example in footnote 34, allowing for present bias, perceived investment benefit $\beta \delta \rho'(v_0^*) \Delta \tilde{U}_1(\alpha, \beta) = \rho'(v_0^*) \beta \delta (1 + \alpha + \alpha \beta \delta k) \log \left(\frac{Y_1}{Y_1 - OOPC}\right)$ could increase due to larger β and α from the exposure.

C Education Outcomes

Sample Construction. I estimate the effect of CHIP exposure on children conceived 7 quarters before till 2 quarters after CHIP in the American Community Survey (ACS). To focus on education outcomes from high school to college, I restrict the sample to children who are age-ready for high school (Grade 9) or above when surveyed in the ACS. Because the oldest cohort – those conceived 7 quarters before CHIP – are born in the second quarter of 1996, the earliest time for the children to be age-ready for Grade 9 is 2010-2011. I therefore use 2010-2018 waves of ACS to follow the academic progress of children from high school to college. During this period, the average child in the sample – born in the fourth quarter of 1998 – expects to enter Grade 1 in 2004-2005, and expects to complete high school (Grade 12) in 2016-2017. Appendix Table C1 summarizes the estimation sample.

Table C1: Sample summary, education outcomes (N = 385,063)

	mean	s.e.		mean	s.e.
grade-for-age (%)	88.21	0.052	child age	16.89	0.003
graduate high school (%)	28.79	0.073	birth year	1998.36	0.002
enroll in college (%)	15.78	0.059	survey year (t)	2015.25	0.003
-			Grade 1 entry year	2004.72	0.002
single motherhood					
never married (%)	8.64	0.045	eliginc ^{utero} (100% FPL)	1.58	0.001
unmarried in t (%)	25.46	0.070	eliginc ^{child} (100% FPL)	2.33	0.001

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table summarizes the estimation sample for education outcomes. I construct the sample from children conceived 7 quarters before till 2 quarters after CHIP, and focus on the education outcomes of children expected to attend high school or above in 2010-2018 waves of the American Community Survey (ACS). I determine the expected grade given child age based on the school entry age in the state, and identify mothers using the family interrelationships variables developed by Ruggles *et al.* (2020). Based on mother's marital status, I define single motherhood requiring that the mother has never married, or is unmarried in survey year *t*.

Variable Definition. I determine the child's grade-for-age status – whether the child attends the grade level expected of her age and graduates high school on time – based on the school entry age in the state. Specifically, children turning five before a cut-off month in a year can enter kindergarten in the same year, and those turning five after the cut-off enter in the next year. Since ACS provides the quarter of birth rather than the month, I assume that children born in the same quarter as the cut-off are born after the cut-off month, and hence adopt the more generous criterion in determining grade-for-age. For children expected to have finished Grade 12, grade-for-age indicates whether the child has obtained a regular high school diploma or is enrolled in college. I also separately examine high school graduation and college enrollment as distinct outcomes after Grade 12.

I identify mothers of school-age children in the ACS using the family interrelationships variables developed by Ruggles *et al.* (2020). Based on mother's marital status, I define

³⁶Similar definition is adopted in Kearney and Levine (2019). Following Deming and Dynarski (2008), I code school entry cut-offs for CHIP cohorts based on Appendix Table 1 of Bedard and Dhuey (2007).

single motherhood requiring that the mother has never married, or is unmarried in survey year t. 8.64% of the children live in single-mother households where the mother has never married, and 25.46% live in households where the mother is unmarried in year t. Over the sample period, in-utero exposure to CHIP averages 158% FPL, and increased from 122% FPL before CHIP to 202% FPL after. Childhood exposure averages 233% FPL, and increased from 201% FPL when the average child is age 0 to 263% FPL when the child reaches age 18.

D Calibration of Out-Of-Pocket Costs

I calibrate the out-of-pocket cost of pre-natal visits from the Medical Expenditure Panel Survey (MEPS) "Event Files." The Event Files contain information on the visits to physicians, hospitals and other health facilities by household members in a calendar year. For each visit, the Event Files record the reason for seeking medical care, the insurance coverage of the medical costs, and payments made by the individual and by the provider. From these records, I identify pregnant mothers as hospital patients whose reason of the visit was "to give birth to a child."

For pregnant mothers, I identify pre-natal visits if the visit was related to a pregnancy condition during a 9-month period prior to the birth event. I look up pregnancy-related conditions from the Clinical Classification Codes provided by MEPS. I further require that the mother consult directly with a medical doctor in an office setting during the visit. Appendix Table D1 summarizes the out-of-pocket cost per visit for low-educated single mothers who had a birth event in 1997-2001.

Table D1: Out-of-pocket costs (OOPC) per pre-natal visit, MEPS

	(1)	(2)	(3)
	OOPC	OOPC > 0	OOPC(>0)
age	0.80	0.10***	-15.32
	(1.05)	(0.024)	(9.79)
age ²	-0.011	-0.0016***	0.25
	(0.018)	(0.0004)	(0.16)
constant	-9.69	-1.41***	249.04
	(15.02)	(0.33)	(144.72)
y mean	3.42	0.14	24.26
N	1,750	1,750	251

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table predicts the out-of-pocket cost (OOPC) per prenatal visit using a quadratic function of age. Column 1 predicts the average cost per visit. Column 2 predicts the probability of incurring a positive out-of-pocket cost. Column 3 predicts the average cost conditional on incurring a positive cost. Sample includes pre-natal visits by non-college educated single mothers who had a birth event in a hospital between 1997 and 2001 in MEPS. All dollars adjusted to the 2000 level by the CPI-Urban price index.

Over 85% of the pre-natal visits had zero out-of-pocket costs (y mean in column 2). The probability of zero costs varied significantly with age. To approximate the empirical

distribution, I model the out-of-pocket cost as following a two-node distribution between zero costs and a positive amount depending on the mother's age. The mass at zero is determined by the regression coefficients in column 2. The level of the positive cost is determined by the regression coefficients in column 3. Although I predict out-of-pocket costs using a quadratic function of mother age, I discretize mother age into four 5-year age groups, and calibrate the cost per visit as follows

$$c_{it}(a_i, \xi_{it}) = \sum_{g=0}^{3} 1\{\xi_{it} \ge F^{-1}(1 - z_g)\} \cdot 1\{20 + 5g \le a_i \le 25 + 5g\} \cdot \left[249.04 - 15.32 \cdot (22.5 + 5g) + 0.25 \cdot (22.5 + 5g)^2\right], \tag{D1}$$

where g=0,1,2,3 indexes age groups 21-25, ..., 36-40. z_g is probability of positive out-of-pocket costs for group g. In the model, positive out-of-pocket costs occur when the health shock ξ_{it} is greater than $F^{-1}(1-z_g)$, where F is the cumulative distribution function of a standard normal.³⁷ Larger health shocks increase the probability of paying positive out-of-pocket costs. I use the median age in each age group and the regression coefficients in column 3 of Appendix Table D1 to calibrate the positive cost level.

³⁷Empirically, $z_0 = 0.105$, $z_1 = 0.128$, $z_2 = 0.295$, $z_3 = 0.369$.

E Structural Sample

E.1 Sample Definition

The structural model estimates the investment responses of non-college educated single mothers whose pregnancy onset was within one year prior to the CHIP onset. Compared to the analysis in Section 5, the structural model examines a more homogeneous group of single mothers (those without college education) and uses a shorter event window around CHIP. In particular, mothers starting pregnancy more than one year before CHIP and those starting pregnancy after CHIP are not included in the structural analysis.

I further restrict the sample to states with homogeneous income limits before CHIP, and exclude states with very small or very large expansion of income limits after CHIP. Specifically, states increasing income limits by less than 20% FPL (MN) and by over 90% FPL (CT, MO, NH, PA, RI) are excluded. These states account for 10% of the births by low-educated single mothers. Moreover, 5 states (MN, MI, NM, VT, WA) expanded insurance above 130% FPL prior to CHIP, and the remaining states (91% of the births by low-educated single mothers) have pre-CHIP income limits between 110% -130% FPL. I focus on the latter set of states and estimate the behavioral effects starting from a homogeneous exposure level (110%-130% FPL) prior to CHIP. In equation 21 and 22, the intercept (α_0 , β_0) corresponds to the altruism and present bias at this exposure level.

The final sample further excludes a small fraction of mothers giving birth before the 7th month of pregnancy, and those with missing birth weight or the onset time of pre-natal visits. I exclude these mothers because the structural model assumes that the child is born in the third trimester, relies on the timing of visits to estimate time preferences, and examines birth weight as the main outcome of the child. The final sample includes 324,400 low-educated single mothers. I summarize the sample in Table E1.

	Endogenous Variables			Mother C	Characteristics
	mean	s.e.		mean	s.e.
log birth weight	8.07	0.20	$\Delta \ell_i$ (% FPL)	24.79	24.07
care onset in			age group	0.69	0.90
first trimester (%)	69.68	45.96	prior fetal death (%)	28.01	44.90
second trimester (%)	21.94	41.39	any risk factor (%)	28.76	45.26
third trimester (%)	8.37	27.70	county smoking (%)	30.51	16.11
# pre-natal visits	10.46	4.27	trimester of exposure	1.51	1.11
≥ 5 cigar. daily (%)	10.88	31.14	missing smoking (%)	30.26	45.94

Table E1: Summary statistics, structural estimation sample

Notes. Table summarizes the sample of non-college educated single mothers in the structural analysis. The left panel summarizes birth weight and investments to be matched with predictions from the model. The right panel summarizes exogenous variables such as mother characteristics and CHIP exposure $\Delta \ell_i$. The last two variables – the trimester of exposure and the percent of mothers missing records of smoking – are exploited in the construction of moment conditions. Because not all states ask mothers about her smoking during pregnancy (affecting 30% of the estimation sample), I treat "missing" as a distinct level of smoking and calculate the share of smokers including all mothers in the denominator.

^{***} p < 0.01 ** p < 0.05 * p < 0.10

E.2 Mother Characteristics

The right panel of Table E1 summarizes mother characteristics exploited in the structural analysis. Mother's age group and CHIP exposure $\Delta \ell_i$ enter as state variables in the dynamic model. Mother's age (grouped into 5-year age bands between age 21 and 40) affects out-of-pocket costs and hence the utility from pre-natal visits ϑ . $\Delta \ell_i$ shifts altruism and present bias through equation 21 and 22. The next three characteristics – whether mother has fetal death in previous pregnancies, has any pregnancy risk factor, and the population smoking rate in her county of residence – determine the distribution of taste types η_i and ζ_i and the endowment type ϕ_i according to equation 23. Mothers have previous fetal deaths if they report lower live birth order than the total birth order for the current birth. Risk factor is an indicator set to 1 if the mother has at least one comorbidity indicated on the birth certificate. To construct the county smoking rate, I first calculate monthly smoking rates for non-college educated women (both pregnant and non-pregnant) in each county from the Behavioral Risk Factor Surveillance System (BRFSS). I then use the average smoking rate in the fifth quarter prior to CHIP (or three months before the start of the estimation sample) as the county smoking rate.

The last two characteristics, the trimester of exposure and the share of mothers with missing records of smoking, are exploited in the construction of moment conditions. I group mothers by the trimester of exposure in some of the moment conditions to examine investment responses specifically to the timing of exposure across cohorts. Because not all states ask mothers about her smoking during pregnancy (affecting 30% of the estimation sample), I treat "missing" as a distinct level of smoking, and calculate the share of smokers including all mothers in the denominator in the moment conditions.

F Moment Conditions and Estimation

Moment Conditions. I construct moment conditions to estimate three key equations in the structural model: the behavioral effects in equation 21 and 22, and the birth weight production in equation 16. To identify the behavioral effects, I exploit responses in the timing and the level of investments by exposure levels in $\Delta \ell_i$. I identify the birth weight production function using exposure levels in $\Delta \ell_i$ as instruments. Therefore, the model parameters can be estimated from the following set of moment conditions

- 1. 4×5 moments on the percent of mothers starting pre-natal visits in the first, second, and third trimester, and the percent without pre-natal visits, by exposure $\Delta \ell_i$,
- 2. 5 moments on the number of pre-natal visits, by exposure $\Delta \ell_i$,
- 3. 5 moments on the percent of mothers smoking more than 5 cigarettes daily, by exposure $\Delta \ell_i$,
- 4. 6×3 moments on log birth weight interacted with indicators of 6 levels of pre-natal visits and 3 levels of smoking,
- 5. 5 moments on log birth weight, by exposure $\Delta \ell_i$,

In addition, I include the following set of auxiliary moment conditions to capture additional investments and heterogeneity by mother age. These moment conditions are less weighted in the estimation.

- 6. 4×4 moments on the percent of mothers starting pre-natal visits in the first, second, and third trimester, and the percent without pre-natal visits, by cohort j,
- 7. 16×5 moments on the percent of mothers starting pre-natal visits in a given trimester and taking a given number of visits by the end of pregnancy, by exposure $\Delta \ell_i$,
- 8. 2×5 moments on the percent of mothers who are non-smokers (<5 cigarettes daily) or heavy smokers (≥ 15 cigarettes daily), by exposure $\Delta \ell_i$,
- 9. 4×5 moments on the percent of mothers with very low or very high number of visits (≤ 6 or ≥ 15) who also smoke less than 5 or over 15 cigarettes daily, by exposure $\Delta \ell_i$,
- 10. $3 \times 6 \times 3$ moments on the probability of birth weight falling below 2,500 grams and below two terciles (3,062 grams and 3,450 grams), interacted with 6 levels of pre-natal visits and 3 levels of smoking,
- 11. 3×5 moments on the probability of birth weight falling below 2500 grams and below two terciles (3,062 grams and 3,450 grams), by exposure $\Delta \ell_i$,
- 12. 6×4 moments on the number of pre-natal visits, by mother age a_i .

In total, I employ 272 moment conditions to estimate the model parameters.

Simulated Moment Conditions. Moment conditions that vary by exposure $\Delta \ell_i$ take the following form

$$\mathbb{E}\left[d_i^l | \Delta \ell_i = k\right] - D^l(\Theta; k) = 0, \tag{F1}$$

where d_i^l is the outcome of interest in moment condition l for individual i, and Θ is the model parameters. The outcome implied by the model for exposure k is $D^{l}(\Theta; k)$, obtained by simulating optimal investments given Θ . At true parameter values, simulated outcomes $D^l(\Theta; k)$ should match the sample counterpart $\mathbb{E}\left[d_i^l|\Delta\ell_i=k\right]$. I transform the conditional moment in equation F1 into an unconditional one as follows

$$\mathbb{E}\left[\left(d_i^l - D^l(\Theta; k)\right) \cdot 1\{\Delta \ell_i = k\}\right] = 0, \tag{F2}$$

and the sample counterpart of the moment condition is given by

$$m^{l} = \frac{1}{N} \sum_{i} \left(d_{i}^{l} - D^{l}(\Theta; k) \right) \cdot 1\{ \Delta \ell_{i} = k \}, \tag{F3}$$

where m^l is the moment residual given model parameter Θ .

I similarly construct the moment residuals for investments by mother age groups and for the birth weight production function.³⁸ Stacking up, $m = (m^l)$, l = 0, ..., 271 is a vector of moment residuals with the variance covariance matrix **S**.

Estimation. The method of simulated moment (MSM) searches for parameter values that best match the simulated outcomes $D^l(\Theta)$ with the sample counterparts. The estimated parameter $\hat{\Theta}$ minimizes moment residuals m according to the following objective function

$$\hat{\Theta} = \operatorname{argmin}_{\Theta} m(\Theta)' \mathbf{W} m(\Theta), \tag{F5}$$

where **W** is the weighting matrix. I choose a diagonal weighting matrix where the weights are inverse to the variance of moment residuals and are larger for the main identifying moments.³⁹ The estimate $\hat{\Theta}$ is asymptotically normal: $\sqrt{I}(\hat{\Theta} - \Theta_0) \sim N(0, \mathbf{V})$, and the variance-covariance matrix V equals

$$\mathbf{V} = (1+r)(\mathbf{D}'\mathbf{W}\mathbf{D})^{-1}\mathbf{D}'\mathbf{W}\mathbf{S}\mathbf{W}\mathbf{D}(\mathbf{D}'\mathbf{W}\mathbf{D})^{-1},\tag{F6}$$

$$m^{l} = \frac{1}{N} \sum_{i} \left[\log(b_{i}) \cdot 1\{V_{i} = p\} \cdot 1\{\bar{s}_{i} \ge q\} - D^{l}(\Theta) \right], \tag{F4}$$

where p = 3, 6, ..., 18, and q = 5, 15.

³⁸In particular, moment conditions for log birth weight interacted with visits and smoking inputs are the follows

³⁹Specifically, diagonal element $w_{ll} = \gamma_l \left[\frac{1}{N} \sum_i (d_i^l - D^l)^2 \cdot 1\{\Delta \ell_i = k\} \right]^{-1}$ for condition l in equation F3, where D^l is the sample statistic. I increase γ_l so that the 53 main moment conditions receive the largest weights in the estimation.

where $\mathbf{D} = \frac{\partial m}{\partial \Theta'}\Big|_{\Theta_0}$ is the Jacobian of moment residuals at the true parameter, and r is the ratio of observed to simulated number of individuals.

I simulate investments for 10 million individuals drawn from the data. For each simulated individual, I draw a vector of standard normal shocks that differ by trimesters. I transform the z-draws into taste and health shocks based on model parameters Θ . I then solve for optimal investments in state $\mathcal{I}_{it} = (3\sum_{\tau=1}^{t-1}v_{i\tau},\sum_{\tau=1}^{t-1}s_{i\tau},X_i)$ from equation 17 to 20. I use the decision rules to generate simulated profiles $D^l(\Theta)$, and calculate the fit with sample counterparts according to equation F5. The algorithm tries different parameter values to find the best fitting parameters $\hat{\Theta}$.

G Calculation of MVPF

Outreach Spending. The outreach spending in the roll-out of CHIP is based on program reports in year 2000. According to the Balanced Budget Act of 1997, the national budget for CHIP each year is capped at an amount chosen by the federal government, and states can spend no more than a fixed share of the allotment on program outreach and other administrative costs. Specifically, states can spend no more than 10% of the allotment on program administration, outreach, and additional health assistance and initiatives related to the program. In year 2000, the program allotment at the federal level is \$4.3 billion, and spending on program outreach constitutes a small share of total administrative costs subject to the 10% cap, ranging from 6% in Pennsylvania to 14% in California. Assuming that 10% of the administrative costs are outreach costs, total outreach spending is \$4.3 billion \cdot 10% \cdot 10% = \$43 million in 2000. Since the goal of the outreach is to inform the public of the program (Williams and Rosenbach, 2007), I assume an even distribution of the spending across the population. I therefore divide the total outreach spending by the number of US households (105 million) in 2000, and calculate the cost of exposure ΔG to be $\frac{\$43 \text{ million}}{105 \text{ million}} = \0.41 per household.

Robustness. I calculate alternative MVPFs when the outreach spending increases by a factor of $1+\omega$, where ω captures the additional administrative costs resulting from greater awareness of the program. For instance, program outreach may increase application to the program and the costs of processing the application. For states expanding insurance through the Medicaid program, information about CHIP may increase the take-up of Medicaid insurance among parents, and the woodwork effect may increase the overall administrative burden of insurance programs. I allow for these potential effects by setting $\omega = 0.5$, the upper bound of the marginal cost of public funds commonly applied in the literature. Appendix Table G1 calculates the MVPF of CHIP exposure using the adjusted cost ($\Delta G = \$0.62$ per household). The main findings for welfare remain unchanged under alternative cost calculations.

⁴⁰A detailed list of items subject to the 10% cap is available in the attachment of a letter from the Health Care Financing Administration, available at https://www.medicaid.gov/sites/default/files/Federal-Policy-Guidance/downloads/SMD120897b.pdf.

⁴¹A report prepared by the United States General Accounting Office (GAO) summarizes outreach spending based on state responses to a 2000 survey. The report is available at https://www.gao.gov/new.items/he00086.pdf.

⁴²Historical households tables are published by the Census Bureau at https://www.census.gov/data/tables/time-series/demo/families/households.html.

Table G1: MVPF of CHIP exposure, $\Delta G = \$0.62$

	WTP	WTP^{M}	WTP^C	MVPF
$\varphi = 1$	0.29	0.43	-0.13	0.46
	(0.003)	(0.005)	(0.004)	(0.004)
$\varphi = 2$	0.59	0.85	-0.27	0.94
	(0.005)	(0.010)	(0.007)	(0.009)
$\varphi = 3$	0.88	1.28	-0.40	1.40
	(0.008)	(0.015)	(0.011)	(0.013)

Notes. Table summarizes the marginal value of public funds (MVPF) of CHIP exposure, applying an alternative cost of exposure $\Delta G = \$0.62$. The new cost measure accounts for potential increases in administrative costs as a result of the program outreach, and adjusts the original outreach spending by a marginal cost of public funds of 50%. Standard errors from ten million simulated individuals in the parenthesis.

H Fiscal Externality

I calculate the fiscal externality of CHIP exposure in three steps. First, I calculate the cost of initial program investments based on spendings in the first 19 years of the program (FY1998-FY2016). Next, I predict the life-cycle increases in earnings based on the effect of CHIP exposure on college enrollment. Finally, I calculate the increase in tax payments and compare it with the initial program costs to quantify the fiscal externality of CHIP exposure. I detail the calculations below.

Program Costs. Because spendings of CHIP are capped at the federal level by the annual allotment, I divide the allotment by the number of children (age 0-18) to calculate the cost of program investment per child in a given year. I accumulate the investment costs through the childhood years of the 1998 birth cohort (which overlap with the first 19 years of CHIP since 1998), and discount the cumulative cost to the year before birth (1997) using a 2% annual discount rate. Appendix Table H1 lists the annual CHIP allotment (in 2000 dollars), number of children each year, and the cost per child discounted to 1997. In total, CHIP invested \$1,354.42 per child in the first 19 years of the program.

I then adjust the average cost to derive the cost per child of single mothers. In the National Health Interview Survey (NHIS), the take-up of CHIP is 30% in 1998-2016, and among single mothers, 55% enrolled their children in CHIP. The implied cost per enrolled child is $\frac{\$1,354,42}{30\%} = \$4,514.73$. Among single mothers, CHIP invested an average of $\$4,514.73 \cdot 55\% = \$2,483.10$ per child of single mothers.

Earning Benefits. I predict the increase in earnings from the effect of CHIP exposure on college enrollment. For children of single mothers, the roll-out of CHIP increases college enrollment by $80\% \cdot 1.71 = 1.37$ percentage points (Table 6). Following Hendren and Sprung-Keyser (2020), I assume that students induced by the exposure to attend college remain in college for two years. Applying a 11.3% return on earnings for each year of college enrollment (Zimmerman, 2014), I calculate that in-utero exposure increases earnings by $2 \cdot 11.3\% \cdot 0.0137 = 0.31\%$ for children of single mothers.

I then apply the 0.31% effect to the life-cycle earning profile for children of single mothers. I construct the profile from average labor incomes in age 19-64 in the 2014-2018 American Community Survey (ACS). All adjust the earning profile of an average individual to match earnings for children of single mothers using estimates from Lopoo and DeLeire (2014). Because the 0.31% effect on earnings is relative to children without college education, I calculate earnings for the latter group using the college enrollment rate among children of single mothers. Consistent with Zimmerman (2014), I assume that the return of college education on earnings begins from age 23 onward, and each

⁴³I assume a 0.5% wage growth rate to predict the earning profiles for children in adulthood. Results are very similar using static wages observed in 2014-2018.

⁴⁴Specifically, Lopoo and DeLeire (2014) finds that children of single parents have lower adult incomes by 27% compared to children of continuously married parents, or by 21% compared to the population average.

⁴⁵College enrollment is 45% among children of single mothers in the ACS. Assuming that students attend college for two years, the implied earning loss for children without college education is $9.09\% = 1 - \frac{1}{45\% \cdot (1+2\cdot 11.3\%) + 55\%}$ below the population average.

Table H1: Cost of program investments, by year

	allotment	# children	cost per child
FY	(billions)	(millions)	(discounted to 1997)
1998	4.56	75.37	62.59
1999	4.45	75.89	59.42
2000	4.30	76.42	55.90
2001	4.21	76.74	53.36
2002	3.07	76.95	38.08
2003	3.01	77.16	36.47
2004	2.92	77.37	34.59
2005	3.52	77.58	40.72
2006	3.43	77.90	38.72
2007	4.15	78.11	45.86
2008	4.02	78.22	43.39
2009	8.52	78.22	90.25
2010	9.93	78.22	103.03
2011	10.36	78.01	105.62
2012	11.31	77.79	113.29
2013	12.93	77.69	127.19
2014	13.99	77.69	134.78
2015	8.25	77.71	77.91
2016	10.07	77.69	93.25
Total	126.99	1,470.72	1,354.42

Notes. Table calculates the cost of program investments per child in FY1998-FY2016. Each year, CHIP spendings are capped by an allotment determined by the federal government. I divide the allotment by the number of children to determine the cost of investment per child, and discount the cost to year 1997 using a 2% annual discount rate. I adjust all dollars to 2000 levels using CPI-U. Between FY1998 and FY 2016, CHIP invested a total of \$1,354.42 per child. For the 1998 birth cohort, this coincides with the cost of program investments in childhood. I obtain CHIP allotments from the Federal Register, and obtain the number of children each year from the Federal Interagency Forum on Child and Family Statistics, available at https://www.childstats.gov/americaschildren/tables/pop1.asp.

year of college lowers annual earnings in age 19-22 by 12.80%. ⁴⁶ Discounted to 1997, the life-cycle earning benefit amounts to \$1,101.11 per child of single mothers.

Tax Payments. Following Hendren and Sprung-Keyser (2020), I assume that the earning benefits are subject to a marginal tax rate of 18.9%. The implied increase in tax payment, 18.9%.\$1,101.11 = \$208.11, amounts to $\frac{\$208.11}{\$2,483.10} = 8.38\%$ of the program costs in childhood. This suggests that the government can expect to recoup 8.4% of the program investments from parental responses to the roll-out.

Confidence Intervals and Robustness. I construct confidence intervals for the fiscal externality to account for uncertainties in the estimated effects on college enrollment and earnings. Specifically, I bootstrap estimates of the effect on college enrollment based on the estimate ($\hat{\beta} = 1.37\%$, s.e.=0.32%) in Table 6, and bootstrap estimates of the effect on earnings based on the estimate ($\hat{\beta} = 11.3\%$, s.e.=3.83%) in Zimmerman (2014).

Appendix Table H2 compares fiscal externality under different assumptions and shows the empirical 95% confidence intervals from 1,000 bootstrapped effects in the square brackets. Under a 3% annual discount rate and a 2-year enrollment in college, fiscal externality of CHIP exposure amounts to 6.0% of the program cost in childhood, and the government can rule out a fiscal externality less than 1% or above 15% at the 95% confidence level. Lower discount rate at 2% increases fiscal externality to 8.4% of the cost, and increases it to 15.4% if students attend college for 4 years. In these cases, the more optimistic forecast of the fiscal externality can exceed 20% of the program cost. The preferred estimate (8.4% assuming a 2% discount rate and a 2-year enrollment in college) is in the lower tail of estimates shown in Table H2.

⁴⁶Specifically, Zimmerman (2014) finds lower earnings among college attendees in the 4 years after high school. Scaled by the first-stage effect on the years of college, each additional year of college lowers annual earnings in age 19-22 by 12.08%.

⁴⁷The tax rate is based on CBO estimates of tax-and-transfer rates, which include state and federal individual income taxes, SNAP benefits, and subsidies on health insurance benefits. The CBO estimates are available at https://www.cbo.gov/sites/default/files/114th-congress-2015-2016/reports/50923-marginaltaxrates.pdf. Hendren and Sprung-Keyser (2020) deducts federal payroll taxes (13.9%) from the estimates and adds a small adjustment for state individual income taxes (2.6%). This is consistent with the view that payroll taxes are partly returned to workers as benefits and do not strictly increase government revenues. Ultimately, I apply the adjusted tax-and-benefit rates reported in Table G.I of Hendren and Sprung-Keyser (2020) for the calculation.

Table H2: Fiscal externality in percent of initial program costs, robustness

	r = 3%	<i>r</i> = 2%
2-year enrollment	6.02% [1.05%, 14.99%]	8.38% [1.49%, 20.81%]
4-year enrollment	11.02% [1.92%, 27.45%]	15.35% [2.72%, 38.10%]

Notes. Table calculates the fiscal externality of CHIP exposure in percent of initial program costs in childhood, by discount rate r and the duration of college enrollment. To account for the uncertainty in the effect on college enrollment and earnings, I follow Hendren and Sprung-Keyser (2020) and bootstrap 1,000 estimates from the asymptotic distribution of the estimates. I calculate the implied fiscal externality for each draw, and show the empirical 95% confidence interval in the square brackets.

Additional Tables

Table I1: Effect of exposure on fertility and single motherhood

	(1)	(2)
	fertility (%)	single mothers (%)
exposure in		
4-7 months of age	0	-0.16
	(0.005)	(0.17)
1-3 months of age	0	0
	_	-
3rd trimester	-0.001	0.12
	(0.005)	(0.22)
2nd trimester	0.008	0.16
	(0.009)	(0.30)
1st trimester	0.005	0.34
	(0.009)	(0.27)
0-4 months pre-utero	0.005	0.15
	(0.009)	(0.31)
y mean	0.66%	24.21%
R^2	0.92	0.97
N	808	808

*** p < 0.01 ** p < 0.05 * p < 0.10Notes. Table shows the effect of in-utero exposure on fertility rates and the share of single mothers among pregnant mothers. To calculate fertility rates, I combine the birth certificate records, which contain the universe of live births, with the fetal death records to arrive at the full count of mothers giving birth in each year-month. I use the gestation estimates in these records to infer the pregnancy onset time, and construct the fertility rate for women between age 21 and 40 by state-year-month. I calculate the share of pregnant mothers who are single based on her marital status at the time of delivery. I group mothers by the trimester of exposure to CHI, and estimate separate effects across six exposure groups in the table. Robust standard errors clustered at the level of states in the parenthesis.

Table I2: Effects of exposure on birth weight

	(1)	(2)
	birth weight (grams)	low birth weight (%)
$single \cdot eliginc \cdot exposure$ in		
4-7 months of age	-1.31	0.05
· ·	(1.66)	(0.09)
1-3 months of age	0	0
· ·	-	_
3rd trimester	0.70	0.02
	(1.70)	(0.07)
2nd trimester	1.66	-0.01
	(1.71)	(0.07)
1st trimester	4.07***	-0.12*
	(1.03)	(0.07)
0-4 months pre-utero	4.12***	-0.11**
•	(1.09)	(0.05)
y mean	3342.57	7.08%
R^2	0.02	0.01
N	4,315,394	4,315,394

**** p < 0.01 *** p < 0.05 * p < 0.10Notes. Table estimates the effect of in-utero exposure on birth weight (grams) and low birth weight (<2,500 grams). I group children by the trimester of exposure and estimate effects for six exposure groups. Robust standard errors clustered at the level of states in the parenthesis.

Table I3: Effects of exposure on the timing of pre-natal visits

	(1)	(2)	(3)
	()	late onset (%)	very late onset (%)
	month care started	(2nd/3rd trimester)	(3rd trimester)
$single \cdot eliginc \cdot exposure$ in			
4-7 months of age	0.002	0.10	-0.021
, and the second	(0.010)	(0.23)	(0.18)
1-3 months of age	0	0	0
- C	_	_	_
3rd trimester	-0.006	0.034	-0.10
	(0.009)	(0.24)	(0.10)
2nd trimester	-0.004	-0.10	-0.13
	(0.012)	(0.23)	(0.18)
1st trimester	-0.010	-0.25	-0.27
	(0.010)	(0.19)	(0.17)
0-4 months pre-utero	-0.019**	-0.38**	-0.30**
-	(0.008)	(0.16)	(0.13)
y mean	2.45	15.08%	5.48%
R^2	0.08	0.06	0.04
N	4,200,326	4,200,326	4,200,326

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table estimates the effects of in-utero exposure on the timing of pre-natal visits, focusing on the month of pre-natal care onset in column 1, late care onset past the first trimester in column 2, and very late onset in the third trimester in column 3. I group mothers by the trimester of exposure and estimate effects across six exposure groups. Effects on mothers whose children are already 1-3 months old at the onset of CHIP are normalized to zero. Robust standard errors clustered at the level of states in the parenthesis.

Table I4: Effects of exposure on the number of pre-natal visits and smoking

	(1)	(2) smoking (%)	(3) heavy smoking (%)
	# pre-natal visits	(≥5 cigarettes daily)	(≥15 cigarettes daily)
$single \cdot eliginc \cdot exposure$ in			
4-7 months of age	-0.014	0.12	0
G	(0.016)	(0.15)	(0.073)
1-3 months of age	0	0	0
_	_	-	_
3rd trimester	0.006	0.12	0.029
	(0.022)	(0.14)	(0.086)
2nd trimester	-0.003	-0.012	-0.092
	(0.024)	(0.10)	(0.064)
1st trimester	0.022	-0.11	-0.16**
	(0.021)	(0.097)	(0.071)
0-4 months pre-utero	0.035	-0.20**	-0.19***
-	(0.022)	(0.082)	(0.062)
y mean	11.74	8.41%	3.12%
R^2	0.07	0.07	0.03
N	4,157,327	3,331,203	3,331,203

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table estimates the effect of CHIP exposure on the number of pre-natal visits and smoking intensity. I group mothers by the trimester of exposure and estimate effects across six exposure groups. Robust standard errors clustered at the level of states in the parenthesis.

Table I5: Effects of simulated exposure on birth weight and investments

	(1)	(2)	(3)	(4)	(5)	(6)
		birth weight (grams)		late onset (%) (2nd/3rd trimester)		oking (%) ar. daily)
eligCHIP	32.09***	13.08	-0.048***	-0.027	-0.029***	-0.023***
	(11.63)	(12.84)	(0.018)	(0.018)	(0.008)	(0.006)
eligCHIP \cdot single		47.51***		-0.054***		-0.014**
		(11.01)		(0.013)		(0.006)
y mean	334	3.95	14.9	94%	3.1	3%
R^2	0.03	0.03	0.08	0.08	0.05	0.05
N	4,240	6,535	4,142	2,279	3,27	9,807

*** p < 0.01 ** p < 0.05 * p < 0.10Notes. Table estimates the effects of simulated exposure on birth weight (column 1-2), late care onset (column 3-4), and smoking (column 5-6). I simulate CHIP exposure based on the predicted eligibility for children applying income limits of Medicaid/CHIP known at the time of pregnancy. I show details of the simulation in Appendix A. I estimate separate effects by mother's marital status in even-numbered columns. Robust standard errors clustered at the level of states in the parenthesis.

Table I6: Effects of exposure on mother's insurance

	(1)	(2)	(3)	(4)	(5)	(6)
	any insu	rance (%)		insurance dicaid (%)		insurance ployer (%)
$eliginc \cdot treat \cdot post$	-1.72 (2.42)	-10.93 (6.96)	-1.13 (1.85)	-11.38 (7.90)	-3.87 (4.53)	-8.48 (7.06)
eliginc · treat	5.87 (8.59)	29.38 (17.68)	3.63 (5.45)	35.09 (21.44)	4.54 (10.52)	19.22 (20.76)
eliginc	-1.82 (1.29)	-1.62 (1.23)	0.039 (0.050)	-0.038 (0.49)	-3.14 (1.92)	-3.26 (1.94)
treat pregnant single	Y	Y Y	Y	Y Y	Y	Y Y
y mean	80.20%	80.20%	4.87%	4.87%	41.80%	41.80%
R^2 N	0.03 34,915	0.03 34,853	0.02 34,421	0.03 34,360	0.02 34,421	0.02 34,360

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table estimates the effect of CHIP exposure on mother's insurance status, using data from the Behavioral Risk Factor Surveillance System (BRFSS). I restrict the analysis to women in age 21-40 surveyed between 8 months before and 8 months after the onset of CHIP. I show results from the following difference-in-differences specification

 $y_{its} = \beta_0 \cdot eliginc_{st} \cdot treat \cdot post + \beta_1 \cdot eliginc_{st} \cdot treat + \beta_2 \cdot eliginc_{st} + \beta_s \cdot treat + \alpha_s + \tau_t + \alpha_s \cdot \tau_t + \epsilon_{its},$

where eliginc is the income limit of children's insurance in state s and year-month t. Specifically, $eliginc_{st} = inc_s^{post}$ before the onset of CHIP, and $eliginc_{st} = inc_s^{post}$ after the onset. treat indicates pregnant mothers in odd-numbered columns, and indicates single pregnant mothers in even-numbered columns. The specification controls for differences by mother groups across states (β_s -treat), and differences across states over time (α_s - τ_t). β_0 estimates the effect on (single) pregnant mothers for each 100% FPL exposure to the CHIP insurance. BRFSS sampling weights applied in the regressions. Robust standard errors clustered at the level of states in the parenthesis.

Table I7: Effects of exposure on cash benefits and health spending

	(1)	(2)	(3)	(4)
	means-tested cash transfer (\$)		out-of-pocket health spending (thousands \$)	
$eliginc \cdot treat \cdot post$	-5.58 (16.18)	58.86 (90.74)	0.056 (0.85)	0.34 (0.36)
eliginc · treat	33.64 (42.07)	-185.74	-2.75 (1.91)	-0.29 (1.25)
eliginc	5.47 (5.42)	6.02 (5.42)	0.36**	0.36**
treat	(3.12)	(3.12)	(0.10)	(0.10)
pregnant single	Y	Y Y	Y	Y Y
y mean	46.81	46.82	444.37	444.37
R^2	0.02	0.02	0.02	0.02
N	33,808	33,803	11,963	11,963

^{***} *p* < 0.01 ** *p* < 0.05 * *p* < 0.10

Notes. Table estimates the effect of CHIP exposure on monthly cash benefits from means-tested programs (column 1-2) and out-of-pocket health spending in the past 12 months (column 3-4), using data from the Survey of Income and Program Participation (SIPP). I restrict the analysis to women in age 21-40 surveyed between 8 months before and 8 months after the onset of CHIP. I show results from the following difference-in-differences specification

 $y_{its} = \beta_0 \cdot eliginc_{st} \cdot treat \cdot post + \beta_1 \cdot eliginc_{st} \cdot treat + \beta_2 \cdot eliginc_{st} + \beta_s \cdot treat + \alpha_s + \tau_t + \alpha_s \cdot \tau_t + \epsilon_{its}$

where eliginc is the income limit of children's insurance in state s and year-month t. Specifically, $eliginc_{st} = inc_s^{pre}$ before the onset of CHIP, and $eliginc_{st} = inc_s^{post}$ after the onset. treat indicates pregnant mothers in odd-numbered columns, and indicates single pregnant mothers in even-numbered columns. The specification controls for differences by mother groups across states ($\beta_s \cdot treat$), and differences across states over time ($\alpha_s \cdot \tau_t$). β_0 estimates the effects on (single) pregnant mothers for each 100% FPL exposure to the CHIP insurance.

Means-tested cash benefits are asked in the main survey for each month in a 4-month recall. I use benefit amounts reported for the most recent month (the fourth month). Out-of-pocket health spending includes payments made in the household towards the mother's healthcare utilization in the past 12 months, which include health insurance premiums net of reimbursements from third parties. Dollar amounts are in the 2000 levels with CPI-U. SIPP sampling weights applied in the regressions. Robust standard errors clustered at the level of states in the parenthesis.

Table I8: Effects of exposure on personal debts

	(1)	(2)	(3)	(4)	
	credit card + store bill debt (\$)			personal debt (thousands \$)	
$eliginc \cdot treat \cdot post$	-9.64	-5.93	2.00	-0.58	
	(8.81)	(13.81)	(2.41)	(1.45)	
eliginc · treat	26.92	19.61	-3.68	1.79	
	(18.64)	(25.90)	(5.33)	(2.93)	
eliginc	3.02	2.81	-0.15	-0.17	
	(2.63)	(2.57)	(0.48)	(0.47)	
treat					
pregnant	Y	Y	Y	Y	
single		Y		Y	
y mean	1,986.98	1,987.73	4.00	4.00	
R^2	0.01	0.01	0.01	0.01	
N	12,214	12,209	12,219	12,219	

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table estimates the effect of CHIP exposure on credit card and store bill debts in column 1-2, and total personal debt further including loans and other debts in column 3-4, using data from the Survey of Income and Program Participation (SIPP). I restrict the analysis to women in age 21-40 surveyed between 8 months before and 8 months after the onset of CHIP. I show results from the following difference-in-differences specification

 $y_{its} = \beta_0 \cdot eliginc_{st} \cdot treat \cdot post + \beta_1 \cdot eliginc_{st} \cdot treat + \beta_2 \cdot eliginc_{st} + \beta_s \cdot treat + \alpha_s + \tau_t + \alpha_s \cdot \tau_t + \epsilon_{its},$

where eliginc is the income limit of children's insurance in state s and yearmonth t. Specifically, $eliginc_{st} = inc_s^{pre}$ before the onset of CHIP, and $eliginc_{st} = inc_s^{post}$ after the onset. treat indicates pregnant mothers in odd-numbered columns, and indicates single pregnant mothers in even-numbered columns. The specification controls for differences by mother groups across states ($\beta_s \cdot treat$), and differences across states over time ($\alpha_s \cdot \tau_t$). β_0 estimates the effects on (single) pregnant mothers for each 100% FPL exposure to the CHIP insurance.

Table I9: Effects of exposure on program uptake and education expectation

	(1)	(2)	(3)	(4)
		Expected Education		
	Medicaid/CHIP	college degree	graduate school	attainment
	(%)	(%)	(%)	(1-5 scale)
$single \cdot eliginc \cdot exposure in$				
4-7 months of age	1.97	-4.26	7.55	12.35
, and the second	(12.71)	(7.93)	(11.41)	(23.98)
1-3 months of age	0	0	0	0
	_	_	_	_
3rd trimester	19.66	-0.63	-0.85	-0.89
	(14.15)	(2.69)	(2.10)	(4.86)
2nd trimester	13.61*	2.58	-2.64	4.53
	(7.64)	(2.01)	(3.51)	(6.55)
1st trimester	47.69***	3.77	-0.68	5.71
	(15.68)	(2.52)	(3.82)	(6.95)
0-4 months pre-utero	36.96**	-2.68	-0.75	-5.08
	(14.26)	(4.96)	(4.59)	(11.93)
y mean	19.41%	86.15%	26.93%	4.06
R^2	0.28	0.12	0.05	0.11
N	1,542	1019	1019	1019

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table estimates the effect of CHIP exposure on program uptake in column 1, and mother's expected education for the child in column 2-4, using data from the Survey of Income and Program Participation (SIPP). In SIPP, respondents are asked their enrollment in public insurance programs for each month in a 4-month recall. I focus on enrollment in the most recent month (the fourth month), and examine the child's enrollment in Medicaid/CHIP in the first year of the child's life (age 0) in column 1. Outcomes in column 2-4 are mother's stated expectation of the child's education attainment. The expectation is asked for all children in SIPP households in wave 6 (the middle wave) and wave 12 (the final wave) of the 1996-2000 survey panel, which spans the roll-out of CHIP. In column 4, education attainment is coded on a 1-5 scale, with the integers corresponding to no degree (less than high school), high school, some college, college degree, and graduate school, respectively. SIPP sampling weights applied in the regressions. Robust standard errors clustered at the level of states in the parenthesis.

Table I10: Effects of CHIP exposure on mother's marital status

	(1)	(2)
	never married (%)	unmarried in t (%)
eliginc ^{utero}	0.62	0.99
	(0.72)	(0.81)
eliginc ^{child}	0.99	0.33
	(0.87)	(1.22)
y mean	8.64%	25.46%
R^2	0.01	0.01
N	385,063	385,063

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table estimates the effects of CHIP exposure on single mother-hood, measured by the percentage of mothers who have never married in column 1, and by the percentage of mothers unmarried in the survey year t in column 2. Regressions are weighted by the ACS sampling weights. Robust standard errors clustered at the level of states in the parenthesis.

Table I11: Effects of CHIP exposure on high school graduation and college enrollment

	(1)	(2)	(3)	(4)
	graduate high school (%)		enroll in college (%)	
eliginc ^{utero} · single · exposure in				
7-12 months of age	-0.54	-0.14	-0.81	-0.31
-	(0.87)	(0.46)	(0.92)	(0.56)
1-6 months of age	0	0	0	0
	_	_	_	_
second - third trimester	0.39	-0.11	0.18	0.32
	(0.49)	(0.28)	(0.73)	(0.49)
first trimester in and pre utero	0.47	0.12	0.86**	0.59**
	(0.29)	(0.20)	(0.41)	(0.25)
4-9 months pre utero	0.59**	0.29	1.14**	0.69**
	(0.29)	(0.18)	(0.45)	(0.29)
single				
never married	Y		Y	
y mean	28.79%		15.78%	
R^2	0.64	0.64	0.35	0.35
N		385,063		5,063

^{***} p < 0.01 ** p < 0.05 * p < 0.10

Notes. Table shows the effects of CHIP exposure on high school graduation rates (column 1-2) and on college enrollment rates (column 3-4), for children of never married mothers and for children whose mothers were unmarried in the survey year. I group children by the timing of exposure and show separate effects for six exposure groups. Effects on children already 1-6 months old at the onset of CHIP are normalized to zero. Regressions are weighted by ACS sampling weights. Robust standard errors clustered at the level of states in the parenthesis.

Table I12: Effects of in-utero exposure to CHIP on grade progression in high school

	(1)	(2)	(3)	(4)	
	grade-fo	grade-for-age (%)		graduate HS (%)	
eliginc ^{utero} · single	2.36***	1.56***	1.71**	0.75**	
	(0.61)	(0.51)	(0.68)	(0.37)	
eliginc ^{utero}	0.028	-0.16	-0.75	-0.80	
	(1.01)	(0.99)	(0.66)	(0.69)	
treat					
never married	Y		Y		
y mean	88.21%		28.79%		
R^2	0.095	0.095	0.64	0.64	
N	385,063		385,063		

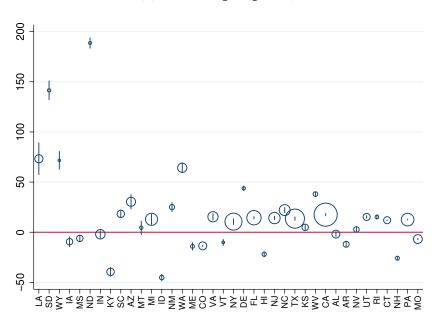
^{***} *p* < 0.01 ** *p* < 0.05 * *p* < 0.10

Notes. Table shows the effects of CHIP exposure on the grade-for-age status (whether the child attends the grade expected of her age) in column 1-2, and on high school graduation in column 3-4. *single* indicates children of single mothers. I estimate effects for children of never married mothers in odd-numbered columns, and for children of single mothers unmarried in the survey year in even-numbered columns. Regressions are weighted by the ACS sampling weights. Robust standard errors clustered at the level of states in the parenthesis.

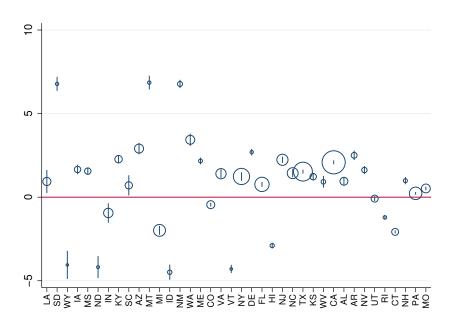
J Additional Figures

Figure J1: Effects of exposure across states

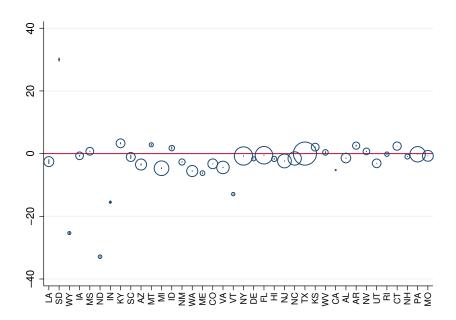
(a) Birth Weight (grams)



(b) First Trimester Care (%)

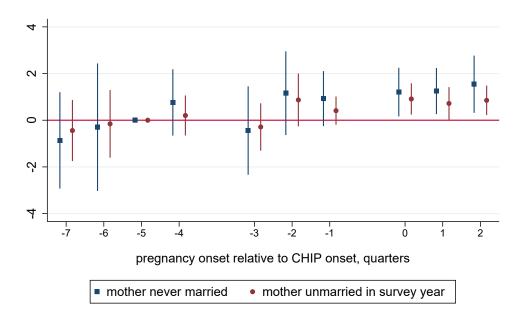


(c) Smoking Intensity (half packs)



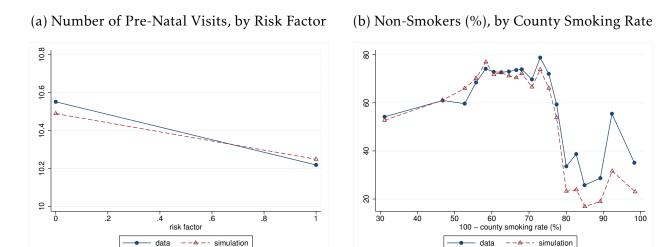
Notes. Figure plots state-specific effects of exposure on birth weight in panel (a), first trimester care in panel (b), and smoking intensity in panel (c). Smoking intensity is measured in half packs (10s) of cigarettes daily. I rank states by the size of expansion from small to large on the horizontal axis. On average, states expanded income limits by 80% FPL. Small expansion states from Louisiana (LA) to Maine (ME) expanded income limits between 22.5% FPL and 54.8% FPL. States beginning with Rhode Island (RI) expanded income limits by over 100% FPL. I do not estimate state-specific effects for Minnesota (MN) where the income limit increased by a minimal 0.26% FPL. I indicate the sample size in each state with the circle around the estimates. 95% confidence intervals are based on robust standard errors clustered at the level of states.

Figure J2: Effect of CHIP exposure on college enrollment (%), event study



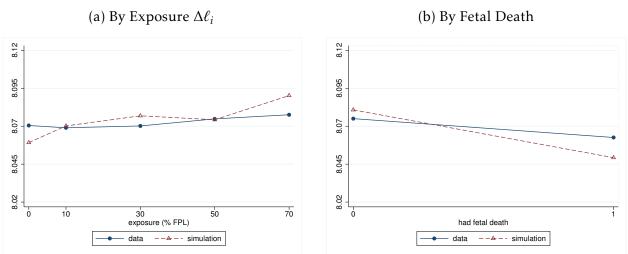
Notes. Figure plots the effect of CHIP exposure on college enrollment by the timing of exposure across cohorts. I show effects for children born to never married mothers and for children of single mothers in the survey year. The horizontal axis compares the pregnancy onset quarter with the CHIP onset quarter. Children conceived more than 4 quarters before CHIP are not exposed to CHIP in utero. In-utero exposure then increases for later cohorts. I normalize the effects on children conceived 5 quarters before CHIP to zero. 95% confidence intervals are based on robust standard errors clustered by states.

Figure J3: Simulated investments by mother characteristics



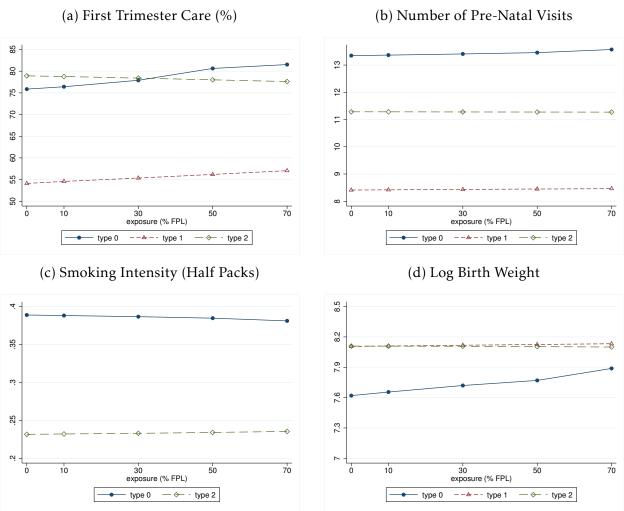
Notes. Figure compares simulated investments with empirical counterparts across mother characteristics used as predictors of mother types in the multinomial Probit. Panel (a) shows differences in pre-natal visits across mother risk factors, and panel (b) shows differences in smoking across county smoking rates in panel (b). I construct county smoking rates using the share of smokers among non-college educated women (pregnant and non-pregnant) in each county in the Behavioral Risk Factor Surveillance System (BRFSS). I simulate investments for ten million non-college educated single mothers and plot simulated investments in the dotted line. I plot empirical counterparts in the solid line.

Figure J4: Effect of CHIP exposure on birth weight



Notes. Figure compares simulated birth weight with empirical counterparts across CHIP exposure $\Delta \ell_i$ in panel (a), and by fetal death in previous pregnancies in panel (b). Mother had fetal death in previous pregnancies if the live birth order of the current birth is smaller than the total birth order. I simulate birth weight for ten million low-educated single mothers and plot simulated investments in the dotted line. I plot empirical counterparts in the solid line.

Figure J5: Effect of CHIP exposure on investments and birth weight, by mother types



Notes. Figure plots simulated investments and birth weight by CHIP exposure $\Delta \ell_i$ for different mother types. I simulate investments and birth weight for ten million non-college educated single mothers at each exposure level in $\Delta \ell_i$, and plot the results by exposure and mother types. In panel (c), since type 1 mothers never smoked in the simulation due to significant disutility from smoking, I plot simulated smoking intensity only for type 0 and type 2 mothers.

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