

Bhalotra, Sonia R.; Rawlings, Samantha

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Sonia Bhalotra
Sam Rawlings

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Sonia Bhalotra

*University of Bristol (Economics and CMPO)
and IZA*

Sam Rawlings

University of Bristol (Economics and CMPO)

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IZA

P.O. Box 7240
53072 Bonn
Germany

Phone: +49-228-3894-0
Fax: +49-228-3894-180
E-mail: iza@iza.org

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ABSTRACT

Intergenerational Persistence in Health in Developing Countries: The Penalty of Gender Inequality?

This paper is motivated to investigate the often neglected payoff to investments in the health of girls and women in terms of next generation outcomes. This paper investigates the intergenerational persistence of health across time and region as well as across the distribution of maternal health. It uses comparable micro-data on as many as 2.24 million children born of about 0.6 million mothers in 38 developing countries in the 31 year period, 1970-2000. Mother's health is indicated by her height, BMI and anemia status. Child health is indicated by mortality risk and anthropometric failure. We find a positive relationship between maternal and child health across indicators and highlight non-linearities in these relationships. The results suggest that both contemporary and childhood health of the mother matter and that the benefits to the next generation are likely to be persistent. Averaging across the sample, persistence shows a considerable decline over time. Disaggregation shows that the decline is only significant in Latin America. Persistence has remained largely constant in Asia and has risen in Africa. The paper provides the first cross-country estimates of the intergenerational persistence in health and the first estimates of trends.

JEL Classification: I10, J11, O57

Keywords: intergenerational persistence, mobility, health, developing countries, cohort trends, inequality

Corresponding author:

Sonia Bhalotra
Department of Economics
University of Bristol
8 Woodland Road
Bristol BS8 1TN
United Kingdom
E-mail: s.bhalotra@bristol.ac.uk

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1. Introduction

There is growing evidence of the long term and often multiplicative benefits of investing in children (Case et al., 2005; Currie, 2009) and recognition that investments made during critical periods of development draw larger returns or, conversely, that failure to invest in early childhood can lead to irreversible damage (Cunha and Heckman, 2007; Almond and Currie, 2010). The origins of life *in utero* constitute one such critical period, when growth is rapid and neurological and physiological development is particularly sensitive to the environment. Maternal stress and nutritional deprivation tend to stimulate permanent changes in tissue structure and function that help the foetus survive but that are associated with abnormal structure, function, and disease in adult life. A later life penalty from foetal adaptation has been noted in terms of earlier onset of later life morbidities (Barker and Levy, 1994), reduced stature and increased risk of obesity (Deaton, 2007; Fung, 2009) and cognitive outcomes and earnings (Almond, 2006).

Mother's health is an important element of the foetal environment, exerting direct effects and also mediating the impact of shocks to the external environment. A vast cross-disciplinary literature documents the impact on intrauterine growth retardation of poor maternal health during pregnancy, measured as smoking, alcohol consumption, stress, infection or nutritional deprivation (Kramer 1987, Almond 2006, Camacho 2008, Cogswell et al. 2003). In fact it is not only the contemporaneous health of the mother or health during pregnancy that counts but rather the stock of her health (Barker, 1997), in the accumulation of which her own early life health environment plays a critical role. Of particular interest, a girl's exposure to adverse conditions can induce developmental changes that have intergenerational impacts (Barker and Osmond 1987; Barker 1992; Drake and Walker 2004; Gluckman and Hanson 2005). There is some evidence of this in the economics literature (Almond and Chay, 2006; Fung and Wei, 2009; Bhalotra, 2010a; Bhalotra and Venkataramani, 2010). Identified mechanisms include the following. Mothers who are themselves low birth weight tend to show reduced weight gain (Hackman et al., 1983) and increased risk of developing hypertension during pregnancy (Klebanoff et al., 1999), factors that predispose them to having low birth weight babies (Cawley et al., 1954; Currie and Moretti, 2007). Girls who are born small and do not exhibit catch up growth, failing to attain the height predicted by their genetic potential tend to have reduced uterine and ovarian size which also predicts lower birth weight of their offspring (Ibanez et al. 2000). Overall, evidence is emerging to suggest that the benefits of investments in foetal and early childhood health extend not only into adulthood but, through mothers, into the next generation.

The primacy of mother's health is pertinent in developing countries many of which are characterized by systematic under-investment in girls and women. The relative neglect of girls and women has been strikingly brought to light by Amartya Sen's work (e.g. Sen 1990, 1992). Further work has

documented the variety of forms this takes (Harriss 1990; Bhalotra and Attfield 1998; deRose et al., 2000; Bharadwaj and Nelson 2010), showing also that girls and women bear the brunt of adverse shocks (Behrman and Deolalikar 1990, Dercon and Krishnan 2000, Bhalotra 2010b). This neglect is not confined to societies characterized by son preference but, rather, is widespread (Hill and Upchurch 1995, Preston, 1976; Baird et al. forthcoming). The contribution of this paper is to investigate the implications of health deprivation amongst girls and women for next generation outcomes for both boys and girls. In this respect it provides the first systematic empirical investigation of the argument in Osmani and Sen (2003) that society at large suffers- for generations- from the unequal treatment of girls and women.

While childhood mortality rates in developing countries have declined over the period, they are still higher than can be explained by costs of nutrition or treatment, and income growth alone appears unlikely to make a substantial contribution to further reductions (Deaton, 2006). Mother's education and public health provision have been identified as having potential in this direction (Deaton, 2006), but there is relatively limited emphasis – or evidence- in the literature on the role of mother's health.

Two features of the analysis in this paper make it well suited to the objective of assessing next generation impacts of under-investment in the health of girls and women. First, it uses comparable micro-data on as many as 2.24 million children born of about 0.6 million mothers born in 38 developing countries in the 30 year period, 1970-2000. This allows us to draw general patterns, to make inter-regional comparisons and to investigate trends. We also investigate how the transmission coefficients change across the distribution of maternal health, so as to estimate directly the long run penalties of deprived health. Second, we use a number of indicators of mother's health and child health that together describe short and long term impacts. For the mother we investigate indicators of anemia and her body mass index around the time of birth alongside her (adult) height. Height relative to a country (or ethnic or sub-region) mean reflects the cumulative impact of her health environment from conception to adulthood (Cole, 2000; Deaton, 2007; Gunnell, 2002; Martorell and Habicht, 1986; Fogel, 2004; Schultz, 2009; Steckel, 2009; Strauss and Thomas, 2008), so capturing underinvestment in girls in early life. For the child, we use neonatal, infant and under-5 mortality risk, low birth weight, stunting and the age-gender adjusted z-score of height. To the extent that adult height summarises the history of life-course shocks sustained by the mother and low birth weight is portentous of adult health (e.g. Black and Devereux, 2010), the correlations described here are of relevance to understanding the emergence of "health dynasties".

There is no previous comparable cross-country evidence on levels of or trends in the intergenerational transmission of health and even single country studies are scarce (Solon 2002; Black and Devereux, 2010, pp. 63-65). It is particularly difficult to compare estimates of health persistence from

across individual studies because the measures of health used vary so much (e.g. birth weight, height, BMI, mental health, self-reported health status). It is even more difficult to find consistent time series data with which trends can be compared across regions.¹

The focus on developing countries is relevant not only because of the gender gap in investments but also because the absolute levels of health and resources tend to be low and the returns to health “mobility” are likely to be much higher than in richer environments. The three decades across which we analyse trends in the transmission coefficient constitute a period of momentous change and turbulence in the economic and health environment. For example, parts of Latin America and Asia experienced periods of unprecedented growth while large parts of Africa exhibited negative growth. The health environment was directly altered by new technologies and nationwide interventions including the Expanded Immunization Programme initiated in 1974 and rolled out over the next two decades. Many countries saw gains in life expectancy of greater than twenty years during 1960-2000 (Maddison, 2004). However, by the early 1990s, HIV-AIDS was reversing gains in Africa (Soares, 2006).

What do we find? Indicators of both the contemporary health of the mother and of her health stock are significant predictors of the survival and growth of her births in most countries in the sample. The average effects identified are large relative to the sample means of the dependent variables. In the linear models, a standard deviation decrease in mother's height or BMI raises the risk of poor child health by between 5 and 10 per cent, depending on the measure. These tendencies are widespread, being statistically significant in 20 to 29 of the 38 countries. Intergenerational persistence varies considerably across the three continents and within continent, by country, reflecting long standing differences in underlying levels and inequality in health. While the intergenerational persistence of health has weakened over time in Latin America, in Asia and Africa it has shown no tendency to fade in the last thirty years.

How do pairs of the alternative measures of health compare? Controlling for common unobservable trends and individual characteristics including the education of the mother and father and mother's age at birth, we find that children of shorter mothers are significantly more likely to be low birth weight, to experience neonatal, infant and under-5 mortality and, conditional upon survival (to the interview date) to be stunted. Low maternal BMI (<18.5) is a risk factor for low birth weight and stunting amongst children, indeed, these risks appear to fall monotonically with BMI. Low BMI appears to have no impact on mortality, which is instead elevated amongst mothers with high BMI (>24.9). Low BMI is widely regarded as indicating (current) nutritional deprivation; what is less recognised is that high BMI

¹ In contrast, education is typically measured in years, income is often measured as lifecycle-adjusted family income and a number of longitudinal or cohort data sets for richer countries contain education and income data.

(obesity) may arise from poor net nutrition in early life (Fung, 2009; Luo et al., 2006; Venkataramani, 2010a). We find that mothers with anemia are more likely to bear low birth weight children and more likely to lose them to early life mortality. The impact of anemia on stunting given survival depends upon its severity.

Most previous studies of mobility present linear estimates. But the mechanisms underlying health transmission suggest that it may be important to allow for nonlinearities. Also, all but one of our measures of child health indicates poor health. Estimating the persistence coefficient in different regions of the (mother's) health distribution, we find that increases in the risks of mortality and stunting associated with having a short mother are systematically larger than the decrease in risk derived from having a tall mother, and marginal effects are increasing in distance from mean height in both directions. For BMI we identify the expected non-linearity, with adverse effects on child health being associated with levels that are both too low and too high, using conventionally determined cut-offs. Overall, the estimates confirm that there is substantial intergenerational persistence at the low end of the (mother's) health distribution, suggesting that this may be where returns to investments in women's health are largest.

Given the finding that persistence is stronger at the low end, we may expect average persistence to decline with improvements in maternal health over time. Alternatively, positive trends in public health programmes that effectively target children at most risk will tend to weaken persistence. Using the pooled sample of 31 cohorts in 1970-2000 we find an erosion of 20-30% per decade in the regression coefficient relating child mortality to mother's height. Even after adjusting this for the rise in the relative inequality of mother's height and child mortality, we find average declines of 10-30%. This looks fairly impressive, especially in view of the record of limited income and education mobility. However, disaggregating by continent, we find that these trends are only statistically significant in Latin America. There is largely no trend in Asia except that the relationship between under-5 mortality and height has grown tighter. What is striking is that persistence, especially for neonatal mortality, has increased in Africa.

Given the bi-causal relationship between health and socio-economic status, these findings are relevant to understanding the intergenerational persistence of earnings. The once popular view was that cross-sectional income inequality was driven by stochastic elements like luck and ability (e.g. Roy, 1950). Influenced by estimates of intergenerational persistence in earnings in the US, economists have increasingly adopted from sociology the view that family background and the extent of intergenerational mobility are important determinants of inequality (Becker and Tomes, 1979; Atkinson, 1981; Bowles and Gintis, 2002). However the literature has not made much progress in identifying the precise mechanisms that drive intergenerational mobility (Currie, 2009). The focus has been upon education as the element of

human capital that determines earnings (Grawe and Mulligan, 2002; Behrman et al., 2001) but the evidence increasingly points towards at least equal importance of health (Case et al., 2005; Eriksson et al., 2005; Currie, 2009). This paper contributes to that evidence.

2. Methodology

The significance of the mother's early life environment (proxied by her height) in predicting the health of her babies underlines that the intergenerational transmission of health involves not only genomic but also non-genomic mechanisms. The dramatic variation in the transmission coefficient across countries and, in Latin America, across time would also seem to indicate non-genomic mechanisms.² While some variation is consistent with genetic mechanisms if, for example, the degree of assortative mating varies across region and time (Chadwick and Solon, 2002), we may expect that any such variation is small. A third indication of the importance of non-genomic elements arises from studies that show that mother's size has a stronger influence on offspring size than father's size. This suggests the importance of the *in utero* environment that the mother provides over and above genetic influences that flow from both parents (Walton and Hammond (1938), Morton (1955), Venkataramani 2010b, Bhalotra; in progress). For a given birth size, mother's health will further tend to influence post-birth health and survival given the greater significance of mother's health in relation to breastfeeding and early childhood care.

This said, our objective is to describe patterns of health mobility, distinguishing the transmission of weak and strong health. Like a lot of the mobility literature, we do not attempt to identify a structural parameter or to disentangle the non-genomic from the genomic element of the relationship between maternal and child health. This is inherently difficult to the extent that traits that appear to be genetically inherited may reflect a perpetuation of the consequences of environmentally driven foetal programming through several generations (Drake and Walker, 2004). For example, the evidence suggests that each of birth weight, adult height and obesity (at least amongst women) are heritable and that each is at the same time a barometer of the scars of environmental deprivation. Moreover, genomic and non-genomic mechanisms most likely interact (Ridley, 2003; Cunha and Heckman, 2007; van den Berg et al., 2008), invalidating the commonly used specifications for human capital production in which they appear additively. In their recent survey of intergenerational mobility, Black and Devereux (2010) argue that there have been relatively few attempts to estimate causal effects and that the estimates, as they exist, raise as many questions as they answer; also see Holmlund, Lindahl, and Plug (2008).

Baseline Model

² In other words, *differences* in health persistence between societies are likely to reflect differences in resources and health provision even if the *level* of health persistence is in part determined by genetic transmission.

Pooling the individual data from across countries, we estimate the following equation

$$C_{imjt} = a + \beta H_m + X'_{im} \lambda + \gamma_{jt} + \varepsilon_{imjt} \quad (1)$$

The dependent variable C_{imjt} is a binary variable indicating poor health of child i born to mother m in country j in year t and H_m indicates the health of the mother. Since the dependent variable is an inverse measure of health, we expect $\beta < 0$ for mother's height and BMI and $\beta > 0$ for anemia. The country-year fixed effects, γ_{jt} control for all relevant time-varying unobservables. They will, importantly, absorb average height differences between countries (Ruff, 2002), country-specific nonlinear trends in child health created by medical progress or changes in institutions for delivery of preventive health care (Cutler et al., 2006), and the impact of shocks like AIDS, civil war or famine that strike in the child's birth year. The estimation is on within-country variation across individual families.³ As there are ethnic differences in height, we investigate including fixed effects for ethnicity. Estimates are obtained with and without controls, X , for individual characteristics including birth month, gender and birth order of the child, education of the mother and her partner, her age at birth, religion and urban status, some of which pick up the socioeconomic position of the household. We use the linear probability estimator, having confirmed that predictions lie predominantly within the $[0,1]$ interval. Standard errors are robust to arbitrary forms of heteroskedasticity and to non-independence within countries, including correlation across years. Below we discuss extensions that allow β to vary with the (relative) level of mother's height, the birth year of the child and country. We also estimated a specification that allows more general slope heterogeneity at the mother level by including the demeaned covariates X_m interacted with H_m ; for the child-specific elements of X_{im} , we use the mother-level averages. We expect that the uninteracted coefficient on H_m is an estimate of the average partial effect (or, in the case below, where we use height dummies, the average treatment effect); see Wooldridge 2005.

Asymmetry and non-linearity

Nonparametric estimates of the intergenerational correlation of health indicate that it is stronger at both tails and stronger for height below the mean than for height above the mean. We therefore estimate these more general equations:

$$C_{imjt} = a + \beta H_m + \rho H_m^2 + X'_{im} \lambda + \gamma_{jt} + \varepsilon_{imjt} \quad (2)$$

$$C_{imjt} = a + \beta_1 tallhalf_m + \beta_2 tall1_m + \beta_3 tall2_m + \beta_4 shorthalf_m$$

³ Mothers are of different cohorts and there is therefore variation in their early life conditions.

$$+ \beta_5 short1_m + \beta_6 short2_m + X_{im}^* \lambda + \gamma_{jt} + \varepsilon_{imjt} \quad (3)$$

Equation (2) allows a single turning point to be estimated from the data. Equation (3) imposes the thresholds at which the slope of the relationship changes but allows for six different coefficients. In equations (1) and (2), maternal height is in metres. In equation (3) it is replaced with six dummy variables indicating the mother's place in the country-specific height distribution. These indicate whether her height is between 0.5 and 1, 1 and 2, or greater than 2 standard deviations above (*tall*) or below (*short*) mean height in her country. We similarly investigate asymmetry in the effects of BMI and anemia. We include indicator variables for low BMI (< 18.5) and high BMI (> 24.5), treating normal BMI, lying between these limits, as the omitted category. For anemia, we include indicators for 'none' and 'severe', casting 'mild' and 'moderate' as the omitted categories.

Trends in intergenerational persistence

For the analysis of trends we use only mother's height and childhood mortality since the other indicators are only available for recent births. We initially introduce an interaction of maternal height with a linear trend in equation (1). To allow for a more flexible trend, we estimate (1) for each of the six five-year cohorts of children born during 1970-2000 to obtain β_c . These estimates are standardized by the ratio of the within-cohort standard deviation of mother and child health to get:

$$\theta_c = \beta_c * \left(\frac{\sigma_m^c}{\sigma_i^c} \right)$$

θ_c is interpreted as the predicted standard deviation change in mortality risk for each standard deviation change in maternal height; it is bounded between 0 and 1. This adjustment factors out the cross-sectional dispersion of the health indicator in the two generations. Even if β is declining, θ may not decline if inequality in child health is falling faster than inequality in mother's health. In a third specification, we allow for a completely flexible relation over time. We estimate equation (1) for each *year*, plot the estimated β_t against year and fit a lowess curve to the scatter. We also standardise these estimates to obtain θ_t .

Spatial differences in intergenerational persistence

To investigate spatial differences in intergenerational persistence, equation (1) is estimated by continent and country to obtain β_j , and these are standardized using the continent or country specific ratio of the standard deviations of maternal and child health to get θ_j . As there are large differences in the level of child health across countries, we also present β_j normalised by the mean level of child health in the sample:

$$\tau_j = \left(\frac{\beta_j}{C_j} \right)$$

where C_i is the mean of the dependent variable.

3. Measures of Health

Child health

Improvements in life expectancy in developing countries arise primarily from improvements in child survival. As many as 30% of all deaths in developing countries are amongst children, compared with less than 1% in richer countries (Cutler et al., 2006). It is estimated that malnutrition is the underlying cause of about half of all infant deaths (Murray and Lopez, 1997). Poor nutrition increases vulnerability to infectious disease to which children under the age of five are especially vulnerable because their immune systems are not fully developed (Scrimshaw et al., 1968). Nutritional deprivation leaves a permanent scar even amongst those who survive through to adulthood (Elo and Preston, 1992; van den Berg et al., 2006). Low birth weight and stunting are early indicators of this scar. This paper therefore indicates child health by childhood mortality, measured as death up until the age of 1 (neonatal), 12 (infant) or 60 (under 5) months, low birth weight, and stunting (low height). We also investigate one measure of child health across the distribution, the z-score of child height adjusted for country, age-in-months and gender. Low birth weight, neonatal and infant mortality are often used as proxies for foetal nutrition (e.g. Barker and Osmond, 1986; Drake and Walker, 2004). Stunting and under-5 mortality, which involve longer postnatal exposure periods, incorporate more of the impact of the extrinsic environment. Indeed, infant and under-5 mortality rates are often used as indicators of the infectious disease environment (e.g. Deaton, 2007; Reidpath and Allotey, 2003). Stunting is an easily measured effect of the malnutrition-infection complex that predicts generalized functional impairment on a wide range of biological, behavioral and social dimensions in children and adults in developing countries (Martorell, 1995). Child height is commonly thought of as a good stock measure of net nutrition and height up to the age of three is strongly linked to adult stature (Martorell, 1999).

Maternal health

Malnutrition of women and girls is widespread in developing countries. There is the scourge of poverty and infection and, further, unequal distribution within households, with a tendency towards systematic under-investment in females (section 1). We use three measures of mother's nutrition- height (adjusted for country fixed effects), body mass index (BMI) and anemia status. The first is a stock measure and the latter two are flow measures. It is useful to obtain results on the same data for these alternative indicators. These measures are more objective than self-reported health status (e.g. Schultz, 2009). Previous studies, mostly set in richer countries, have tended to use birth weight to indicate maternal health (e.g. Drake and

Walker, 2004; Currie and Moretti, 2007). Mother's birth weight is not available in our data but it is correlated with final height (Emanuel et al., 1992; Costa, 1998).

An important aspect of wellbeing and of the bearing and rearing of children in developing countries is that mothers are often fatigued. BMI and anemia capture this. They are both correlated with disease, malaria, and poor nutrition (WHO, 2008) and strong predictors of low birth weight and preterm delivery (Neggars and Goldenberg, 2003; Levy et al., 2005). BMI is the most widely accepted measure of adult nutrition. Anemia reflects both dietary deficiencies and poor iron utilization associated with infection. Maternal anemia lowers offspring iron stores, possibly well into the first year of life (Allen, 2000). Potential mechanisms for the impact of the mother's height on the health of her offspring were discussed in section 1. It is estimated that in developed countries, environmental factors explain about 20% of variation in height, but this figure is likely to be higher in developing countries due to higher levels of environmental stress (Silventoinen, 2003) which tend to limit attainment of genetic potential (Mironov, 1999). There are permanent differences in height across regions that are hard to reconcile with nutritional differences and may arise from natural selection based on millennia of different climates and resources (Ruff, 2002). For example, African women are the tallest in the developing world but they grow up in very poor conditions (Deaton, 2007). Nevertheless, cohort variation in height within sub-Saharan Africa is sensitive to indicators of the early life nutritional environment (Akachi and Canning, 2007). We have confirmed that this is also the case in the current sample, where birth year GDP, the regional infant mortality rate and the average education of mothers in the cohort influences the height of adult women (Bhalotra 2009).

4. Data

The microdata are compiled from 77 Demographic and Health Surveys (DHS) for 38 developing countries. Multiple country surveys are pooled across rounds. The estimation sample contains comparable information on as many as 2.24 million children born to 600,000 mothers during 1970-2000 derived from surveys dated 1991-2006. Details are in Tables A.1, A.2 and Figure A.12. The data offer the following advantages. (1) They are of unprecedented scope in this literature; previous epidemiological research often relies upon small samples and no previous work by economists has analysed cross-country microdata on this scale or attempted to describe trends. (2) Biological mothers and children are linked through fertility records, avoiding the problem common in cross-sectional surveys of the presence of non-biological children which, in Africa, is a major issue. (3) A number of indicators of health are available, allowing us to analyse indicators of the contemporary and permanent health of the mother and of the survival and (prenatal and postnatal) growth of the child. (4) Height (mother and child) and the BMI and anemia status

of the mother are measured by surveyors using state of the art tools. This avoids the bias in commonly used subjective assessments of health. (5) Surveys were conducted across countries using very similar questionnaires and measurement tools. (6) The surveys provide complete retrospective fertility histories for women aged 15 to 49 years at the time of the survey in which the dates of all births and any deaths are recorded. Interview dates span 1986-2006, whilst births occur across 1952-2006. These histories are exploited to construct cohort data on mortality across three decades, as in (Bhalotra, 2010b), permitting analysis of trends. (7) Retrospective records that include all births eliminate sample selection problems in the mortality and birth weight measures since inclusion in the sample is not conditional on survival to the survey date. (8) An advantage of cross-country microdata is that they can be merged with country-level panel data (by country and birth year of the child) to look at how mobility varies with country-level income and institutions. We merge in GDP and immunization rates (a measure of public health). The appendix describes the countries in the sample, the sources and definitions of all variables analyzed, and explains how we deal with censoring, potential selectivity, missing data and outliers. It also presents descriptive statistics and graphs.⁴

5. Results

Nonparametric relationships between alternative pairs of indicators of maternal and child health are estimated using lowess predictions of the binary dependent variables (Figure 1).⁵ They consistently suggest a positive intergenerational correlation of health. The rest of this section discusses the parametric estimates presented in Tables 1-3. The estimated coefficients are significant and of the expected sign in every specification. To the extent that non-genomic mechanisms are at play (sections 1 and 2), these results suggest that the benefits of investing in the health of girls extend to their births.

Mother's height

Increases in mother's height lower the risk of poor child health for all measures of child health. They also raise the *z*-score of child health, a measure that is not weighted at the low end (Table A5). The average effects identified are large relative to the sample means of the dependent variables. For instance, a one standard deviation (s.d.) decrease in mother's height is associated with an elevation of low birth weight risk of 7.4% of the mean rate in the sample and with an elevation of neonatal mortality risk of 9.3% of the sample rate. A quadratic in height is significant for mortality and stunting risk, indicating attenuation of the effect of mother's height on child health as height increases. The probability of low birth weight, in

⁴ The appendix is available online at <http://www.efm.bris.ac.uk/ecsr/b/bhalotra.htm>. Henceforth Tables A.n indicate tables in this appendix.

⁵ Country-specific non-parametric plots are in the appendix and a discussion of the non-parametric plots is in the working paper version, Bhalotra and Rawlings (2010a).

contrast, appears to be well specified by the linear model. The specification that includes indicators for the mother's place in the height distribution of her country shows that the intergenerational correlation of health tends to be strongest at the tails of the height distribution and especially so at the low end of the distribution. For example, the increase in neonatal mortality risk from having a mother with height between 1 and 2 s.d. below the mean (*short1*) is more than twice the decrease in risk from having a mother between 1 and 2 s.d. above the mean (*tall1*). Moving from height more than two s.d. below the mean to height 1-2 s.d. below the mean reduces risks of low birth weight and neonatal mortality by 14% and 25% of their sample means. In contrast, moving up on the other side of the mean, from height 1-2 s.d. above mean to height more than 2 s.d. above mean yields gains of 5% and 0% of the mean respectively; see Table 1c. Asymmetry is apparent for all measures of ill-health but while it is significant for mortality and stunting, it cannot be rejected in the case of birth weight. No significant asymmetry is apparent when the z-score of child height is regressed upon the range of levels of mother's height.

The reported estimates are conditional upon characteristics X_{im} . Excluding these raises β by 50% in the specification with height in meters. The estimated coefficients on height are insensitive to controlling for ethnicity or sub-region fixed effects and to incorporating slope heterogeneity at the mother level. While there is some evidence of heterogeneity across mothers in the intergenerational correlation of health within country, this would appear to be independent of her relative height. The baseline specification therefore identifies the average partial (or treatment) effect. These additional estimates are reported in Bhalotra and Rawlings (2010b).

Mother's body mass index

In a linear specification, increases in maternal BMI lower the risk of low birth weight and stunting, raise the z-score of child height and *raise* mortality risk. For example, a one s.d. decrease in BMI is estimated to raise the risk of low birth weight by 10.8% of the mean and to lower the risk of neonatal mortality by 13.1% of the mean (Table 2b). We also estimate a model in which we distinguish *low* (< 18.5) and *high* (> 24.9) BMI from the intervening normal range. The risks of low birth weight and stunting fall monotonically with BMI but with different (implicit) curvature in the two cases. Low BMI raises the risk of low birth weight by almost twice as much as high BMI lowers it. On the other hand, gains in BMI only significantly reduce stunting risk when BMI crosses into the *high* range. When the dependent variable is the z-score of child height rather than stunting, high BMI has a positive influence on height that is significantly larger than the negative influence of low BMI. The positive association of child mortality and maternal BMI is driven by high BMI. This is striking, especially if viewed in relation to the emerging evidence that nutritional deprivation in early childhood can lead to obesity amongst women in adulthood

(section 1). However it must also be viewed alongside our evidence that increases in BMI, even into the obesity range, lower anthropometric failure.

Mother's anemia status

Anemia significantly raises the risk of low birth weight and mortality risk, and lowers the z-score of child height (Table 2). It only significantly lowers stunting risk when it is classed as severe. The risks of low birth weight and neonatal mortality are lower by 5.7% and 16.9% of their mean rates amongst mothers who do not have anemia relative to mothers who do. Distinguishing mild or moderate from severe anemia shows that its marginal effects increase with its severity.

Trends in intergenerational persistence: Overall and by region

Linear trends suggest no significant change over time in the influence of maternal height on neonatal mortality risk but there is a significant weakening of its influence on infant and under-5 mortality with a lowering of the intergenerational correlation by 0.018 and 0.026 per decade, respectively (appendix). A nonparametric fit to a scatter of the estimated intergenerational coefficient from cross-sections for each year against year is in Figure 2. For the earlier years in our sample, the average intergenerational correlation rises but, after about 1975, it declines monotonically. A linear trend fitted to the scatter of β_t shows significant declines in persistence over time at rates that are substantially larger than suggested by the linear specification. The rates of decline are 0.011, 0.036 and 0.043 per decade for neonatal, infant and under-5 mortality respectively (Table 3), which imply an erosion of about 31%, 35% and 31% per decade in the intergenerational coefficients. The correlation or the coefficient multiplied by the relative change in inequality of mother's height and child mortality declines by 10%, 33% and 30%, the decline for neonatal mortality being insignificantly different from zero. Estimates of β_c for five-year cohorts present a broadly similar picture to the estimates β_t ; details are in Bhalotra and Rawlings (2010a).

We repeat this exercise for each region-year; see Table 3 and Fig. A.4, A.5. Latin America shows a consistent improvement in health mobility over time through the three decades with rates of decline twice as large as the average rates reported above. Africa and Asia exhibit a tendency for intergenerational persistence to rise at the start of the period. Decline sets in by the early 1980s in Asia but only later, at the start of the 1990s, in Africa. The regression coefficients for Africa show a positive trend that is significant in the case of neonatal mortality and, once adjusted for the evolution of the ratio of standard deviations of height and mortality, for under-5 mortality. In Asia the relationship is stable over time, except that the relationship between under-5 mortality and height has grown significantly tighter. The average height of mothers grew by 0.16 mm, 0.07 mm, and 0.40 mm p.a. in Africa, Asia, and Latin America. So, younger cohorts of women in Latin America gained height over older cohorts many times faster than in the other

regions. Since the impact of mother's height on child mortality risk is decreasing in her height, this makes sense of the greater weakening of this tie in Latin America. Africa has suffered HIV/AIDS more than the other regions. Consistent with the mobility trend, economic growth picked up in Asia in the early 1980s and in Africa in the early 1990s. Latin America grew through the period albeit with a slowdown in the 1980s. It also experienced a greater expansion over the period of nationwide health interventions than the other two regions.

How do these results compare with previous estimates of intergenerational mobility? As discussed, there are no previous estimates of trends for measures of health. The few available studies of trends in intergenerational persistence in earnings and education suggest limited mobility in Europe and North America with the exception of the Nordic countries (Black and Devereux, 2010). Using data for 42 countries, 29 of which are developing or transition countries. Hertz et al. (2007) show that, while the slope of the relationship between parent and child years of education showed significant attenuation, the correlation remained more or less constant across half a century. Mobility measured by the correlation is weaker than that measured by the regression coefficient because the variance of parent's education has risen relative to the variance of children's education over time in many countries. Similarly, intergenerational income coefficients are often adjusted downwards because the variance of income tends to increase with age and the income of parents is often recorded at a more advanced age than that of their adult offspring (e.g. Blanden et al., 2004). The health data in our sample exhibit a similar tendency. Inequality measured by the standard deviation of mother's health σ_m^c fell slightly over the period but inequality in child mortality σ_i^c fell much more, in line with secular declines in mortality. This led to a rise in the within-cohort ratio σ_m^c/σ_i^c and the corresponding within-year ratio that is applied to adjust β_t . Adjusting for the trend in the relative variance of the mother and child health indicators therefore attenuates the trend in β . The trend in the correlation is further dampened by the data becoming less scattered around the regression line. This is illustrated by Figure A.11 which shows the slope and fit of the relationship for a random sample of data from India. The slope declines significantly between cohorts 25 years apart but the scatter fits more closely around the regression line for the younger cohort.

Spatial differences in intergenerational persistence

Continent-specific estimates are in Table A.7. For mortality, the marginal effect of height is largest in Asia and smallest in Latin America. Indicators of mother's health are worst in Asia and incomes are highest in

the LA countries.⁶ The ordering of continents by the size of the marginal effect of mother's health is reversed when we consider birth weight and stunting. A possible explanation for why, in Latin America, the effects of poor maternal health are largely felt on the anthropometric indices of children and not mortality is that there is less survival selection than in Asia and Africa. Estimates of equation (1) that include interaction terms between maternal height and 38 country dummies show that β_j is significantly different across countries for all measures of child health. The number of countries with a significant β and, amongst these, the range of β , θ , and τ are summarised in Table A.8 and Figure A.5. Country-specific estimates obtained by estimating the model for each country are in Tables A.9 – A.13. The range of θ_j is less than the range of β_j since σ_m^j is similar across countries, whilst σ_i^j is increasing in the size of β_j (see Tables A.9 – A.13). Thus σ_m^e/σ_i^e is decreasing with β_p , and this compresses the distribution of θ_j . The range of τ_j is greater than the range of β_j because the coefficient on maternal height increases with average poor child health. Figure A.6 shows plots of the β_j against country-specific averages of income, education, immunization rates and the levels of maternal and child health. Improvements in the environment weaken the tie between maternal height and child mortality (also see Bhalotra and Rawlings, 2010b). However, low birth weight and stunting risk show no significant association with the levels of maternal or child health, GDP immunization rates or mother's education.

6. Conclusions

This paper documents widespread intergenerational transmission of health from mothers to children using data on as many as 2.24 million births in 38 developing countries across three decades. We find that short stature and anemia of the mother each raise the likelihood that her births are low birth weight, suffer early life mortality and exhibit stunted growth. Low maternal BMI is a risk factor in low birth weight and stunting while high BMI is a risk factor for childhood mortality. Mother's health, by alternative indicators, is thus shown to impact size at birth, survival and child growth conditional upon survival. The results suggest that both contemporary and childhood health of the mother matter for the health of the next generation. BMI and anemia indicate health around the time of birth while adult height is a marker for health in childhood (Case and Paxson, 2010). Thus, under-investment in the health of girls and women contributes to explaining child mortality and intergenerational cycles of growth failure amongst survivors.

It is estimated that in at least two thirds of cases, low birth weight in developing countries signifies intrauterine growth retardation (ACC/SCN 2000: box 1). It also produces post-natal growth retardation to

⁶ These results contrast with Hertz et al. (2007) who found the strongest levels of persistence in education in Latin America. However, health trends may be different and the samples analysed are different. We have 28 African countries, 7 LA and 3 Asian, whilst they have 4, 7, and 10, respectively.

the extent that low birth weight babies are more prone to infection which, in turn, limits nutrient assimilation. More recent evidence indicates that low birth weight has a further range of impacts in adult life, including hypertension, insulin resistance, type 2 diabetes, cardio-vascular disease, obstructive lung disease, renal damage and some forms of cancer, and that these appear to be independent of SES and behaviours like smoking and drinking (Barker, 1998). Indeed, they are more likely to express in individuals who are growing up in conditions of resource plenty relative to their mothers. This may explain why, alongside infectious diseases, a range of chronic non-communicable diseases have taken their hold in developing countries (Yajnik, 2001, Osmani and Sen 2003).

Girls receive lower investments in both health and education in developing countries. To the extent that our estimates pick up non-genomic traits or, more generally, if genomic traits are more likely to express when resource conditions are poor (Turkheimer et al. 2003, van den Berg et al. 2008), our findings suggest that this may be a costly neglect. There is evidence from developing countries of the importance of health in raising educational attainment (Glewwe and Miguel, 2008) and productivity (Schultz, 2005) or, conversely, of poor health in generating poverty traps (Dasgupta and Ray, 1986). Together with evidence of the impact of education and income on health in poor households (Cutler et al., 2006; Bhalotra, 2010b) and of assortative matching in marriage (Pencavel, 1998), this implies that intergenerational persistence in health may explain some of the intergenerational persistence in education, earnings and economic inequality.

In the three decades analysed, there have been substantial advances in the diffusion of medical technology but at the same time, the majority of mothers continue to have little education and the effective provision of broad based public services remains poor. Economic growth has been erratic and unevenly distributed. We find a significant improvement in “health mobility” on average but disaggregation by continent presents a less rosy picture. Latin America exhibits a consistent improvement in health mobility through 1970-2000 with rates of decline twice as large as the average rates. In Africa and Asia, intergenerational persistence increased at the start of the period. Decline set in by the early 1980s in Asia but only later, at the start of the 1990s, in Africa. Overall, persistence increased in Africa, especially for neonatal mortality. It was more or less constant in Asia, with some indication of a tightening of the relationship for under-5 mortality.

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Table 1a: Intergenerational Persistence: Maternal stature effects on child health indicators

	Neonatal Mortality	Infant Mortality	Under-5 Mortality	Low BW	Stunted
A: Linear Height					
β	-0.066** (0.008)	-0.103** (0.010)	-0.139** (0.013)	-0.218** (0.031)	-0.098** (0.007)
θ	-0.021	-0.024	-0.027	-0.037	-0.043
B: Non-Linear Height: Quadratic					
Mother's Height	-0.835** (0.205)	-1.007** (0.203)	-1.227** (0.236)	-1.234 (1.005)	-1.239** (0.244)
Mother's Height ²	0.246** (0.064)	0.289** (0.064)	0.348** (0.075)	0.323 (0.315)	0.363** (0.077)
C: Non-Linear Height: Deviations from country mean					
Tall (0.5-1 s.d.)	-0.002** (0.001)	-0.005** (0.001)	-0.006** (0.001)	-0.011** (0.003)	-0.002** (0.001)
Tall (1-2 s.d.)	-0.004** (0.001)	-0.006** (0.001)	-0.008** (0.001)	-0.019** (0.003)	-0.004** (0.001)
Tall (> 2 s.d.)	-0.004 (0.002)	-0.009** (0.003)	-0.012** (0.003)	-0.029** (0.010)	-0.004 (0.003)
Short (0.5-1 s.d.)	0.004** (0.001)	0.005** (0.001)	0.007** (0.001)	0.013** (0.003)	0.007** (0.001)
Short (1-2 s.d.)	0.009** (0.001)	0.013** (0.001)	0.017** (0.001)	0.022** (0.005)	0.013** (0.002)
Short (> 2 s.d.)	0.021** (0.004)	0.030** (0.005)	0.039** (0.006)	0.051** (0.016)	0.035** (0.007)
Observations	2295740	2233720	1894526	324097	285440
mean (dep. var.)	0.049	0.097	0.145	0.202	0.025
s.d.(dep. var.)	0.216	0.296	0.352	0.402	0.157
mean (height)	1.559	1.559	1.559	1.569	1.566
s.d. (height)	0.069	0.069	0.069	0.068	0.069
$p > \chi^2$: (tallhalf/shorthalf)	0.000	0.000	0.000	0.000	0.000
$p > \chi^2$: (tall1/short1)	0.000	0.000	0.000	0.000	0.000
$p > \chi^2$: (tall2/short2)	0.000	0.000	0.000	0.014	0.000

Robust standard errors in brackets, allowing for clustering within country. * significant at 5%, ** significant at 1%. Mortality samples are adjusted to allow full exposure to the relevant risk. Low BW is 1(low birth weight). Birth weight and height are available only for recent births. Controls include country-year fixed effects, child gender, birth order and birth month, mother's religion, age at birth, education, partner's education and country-year fixed effects. In Panels A and B, mother's height is in metres. θ is β standardized by the ratio of the standard deviation of mother's height to the standard deviation of child health. Tall (0.5-1 s.d.), Tall (1-2 s.d.) and Tall (> 2s.d.) are indicators for being between 0.5 and 1, between 1 and 2, and greater than 2 standard deviations above mean height, respectively. The 'short' dummies are similarly defined. Means of Tall (0.5-1 s.d.), Tall (1-2 s.d.) and Tall (> 2 s.d.) are 0.15, 0.10, and 0.01, respectively. Means of Short (0.5-1 s.d.), Short (1-2 s.d.) and Short (> 2 s.d.) are 0.16, 0.10, and 0.01, respectively. The $p > \chi^2$ statistic gives the significance level at which we are unable to reject the null that the coefficients for tallhalf/shorthalf, tall1/short1, or tall2/short2 are significantly different in magnitude from one another.

Table 1b: Changes in predicted child health associated with a 1 s.d. increase in mother's height

	(1)	(2)		(3)		(4)		(5)	
	Mean	s.d.		Decrease in Risk		(3)/(1)		(3)/(2)	
		Individual	Country	Individual	Country	Individual	Country	Individual	Country
Low BW	0.202	0.400	0.080	0.015	0.007	0.074	0.032	0.038	0.082
Neonatal	0.049	0.216	0.030	0.005	0.002	0.093	0.040	0.021	0.066
Infant	0.097	0.296	0.050	0.007	0.003	0.073	0.032	0.024	0.062
Under-5	0.145	0.352	0.080	0.010	0.004	0.066	0.029	0.027	0.052
Stunted	0.025	0.157	0.010	0.007	0.003	0.270	0.118	0.043	0.294

Table indicates the changes associated with a 1 s.d. increase in mother's height at the country (individual level) which is 3.0cm (6.9cm). 'Low BW' refers to low birth weight. 'Individual' and 'Country' refer to which level the s.d. are calculated at. Means are the same at the country and individual level.

Table 1c: Changes in predicted child health associated with changes in mother's height across the height distribution

	Low BW	Neonatal	Infant	Under-5	Stunted
Change in Risk of Poor Child Health					
Tall (0.5-1 s.d.)	-0.011	-0.002	-0.005	-0.006	-0.002
Tall (1-2 s.d.)	-0.019	-0.004	-0.006	-0.008	-0.004
Tall (>2 s.d.)	-0.029	-0.004	-0.009	-0.012	-0.004
Short (0.5-1 s.d.)	0.013	0.004	0.005	0.007	0.007
Short (1-2 s.d.)	0.022	0.009	0.013	0.017	0.013
Short (>2 s.d.)	0.051	0.021	0.030	0.039	0.035
Change in Risk/Mean					
Tall (0.5-1 s.d.)	-0.054	-0.041	-0.052	-0.041	-0.080
Tall (1-2 s.d.)	-0.094	-0.082	-0.062	-0.055	-0.160
Tall (> 2 s.d.)	-0.144	-0.082	-0.093	-0.083	-0.160
Short (0.5-1 s.d.)	0.064	0.082	0.052	0.048	0.280
Short (1-2 s.d.)	0.109	0.184	0.134	0.117	0.520
Short (>2 s.d.)	0.252	0.429	0.309	0.269	1.400
Change in Risk/s.d.					
Tall (0.5-1 s.d.)	-0.138	-0.067	-0.100	-0.075	-0.200
Tall (1-2 s.d.)	-0.238	-0.133	-0.120	-0.100	-0.400
Tall (> 2 s.d.)	-0.363	-0.133	-0.180	-0.150	-0.400
Short (0.5-1 s.d.)	0.163	0.133	0.100	0.088	0.700
Short (1-2 s.d.)	0.275	0.300	0.260	0.213	1.300
Short (>2 s.d.)	0.638	0.700	0.600	0.488	3.500

Table indicates the changes associated with switching each tall/short indicator in equation (3) from 0 to 1. All changes are relative to the omitted category, which is height of mother within half a standard deviation of mean height in her country. Means and s.d. are calculated on the country-year panel. See Table 1a for sample rates, estimated coefficients, standard errors and significance levels.

Table 2: Alternative Measures of Mother's Health: BMI and anemia

	Mother's BMI				Mother's Anemia Status			
	Neonatal	Infant	Low BW	Stunted	Neonatal	Infant	Low BW	Stunted
A: Linear Effects								
BMI ($\beta * 10$)	0.005** (0.001)	0.008** (0.001)	-0.054** (0.007)	-0.004** (0.001)				
Anemia ($\beta * 10$)					0.054* (0.022)	0.129** (0.031)	0.106 (0.061)	-0.031 (0.028)
B: Non-Linear Effects								
Low BMI ($\beta * 10$)	0.003 (0.011)	0.010 (0.019)	0.461** (0.055)	0.009 (0.014)				
High BMI ($\beta * 10$)	0.044** (0.010)	0.052** (0.013)	-0.234** (0.040)	-0.028* (0.012)				
No Anemia ($\beta * 10$)					-0.049* (0.020)	-0.130** (0.030)	-0.094 (0.061)	0.032 (0.027)
Severe Anemia ($\beta * 10$)					0.216 (0.289)	-0.044 (0.198)	0.484 (0.386)	0.081 (0.124)
Observations	226458	167843	210821	201717	19610	15895	17542	17096
mean (dep. var)	0.025	0.050	0.205	0.024	0.031	0.062	0.19	0.025
s.d. (dep. var)	0.158	0.217	0.404	0.153	0.173	0.241	0.392	0.155
mean (bmi)	22.573	22.64	22.473	22.65				
s.d. (bmi)	3.786	3.844	3.721	3.784				
mean (anemia)					0.463	0.466	0.480	0.460
s.d. (anemia)					0.499	0.499	0.500	0.498
$p > \chi^2$: (lowbmi/highbmi)	0.004	0.075	0.000	0.019				
$p > \chi^2$: (no anemia/severe anemia)					0.383	0.662	0.155	0.722

Coefficients and standard errors are multiplied by 10 for ease of reading. Robust standard errors in brackets, allowing for clustering within country. * significant at 5%, ** significant at 1%. Mother's BMI is the ratio of weight in kilograms to the square of height in meters (kg/m^2). Anemia is a dummy variable indicating whether the mother had either mild, moderate, or severe anemia at the time of the survey. Controls include country-year fixed effects, child gender, birth order and birth month, mother's religion, age at birth, education, and partner's education. There are no estimates for under-5 mortality, since BMI and anemia are only collected for the 5 years preceding the survey, and for the mortality estimates we omit children who are not fully exposed to mortality risk. 'Low BW' refers to low birth weight.

Table 2b: Changes in predicted child health associated with a 1 s.d. increase in mother's body mass index, BMI

(A): Mean and s.d. from individual-level data					
	(1)	(2)	(3)	(4)	(5)
	Mean	s.d.	Change in Risk	(3)/(1)	(3)/(2)
Low BW	0.205	0.404	-0.022	-0.108	-0.055
Neonatal	0.025	0.158	0.003	0.131	0.021
Infant	0.05	0.217	0.002	0.041	0.009
Stunted	0.024	0.153	-0.002	-0.068	-0.011
(B): Mean and s.d. from country-level data					
	Mean	s.d.	Change in Risk	(3)/(1)	(3)/(2)
Low BW	0.197	0.139	-0.010	-0.052	-0.073
Neonatal	0.028	0.014	0.002	0.054	0.108
Infant	0.056	0.026	0.001	0.017	0.036
Stunted	0.024	0.013	-0.001	-0.031	-0.058

Using the coefficients in Table 2a, we compute changes associated with a 1 s.d. increase in maternal BMI at the country (individual level) which is 1.888 (4.092). Means and standard deviations at both the individual and the country-level are calculated from the sample of last-born children.

Table 2c: Changes in predicted child health associated with the mother not being anemic

(A): Mean and s.d. from individual-level data					
	(1)	(2)	(3)	(4)	(5)
	Mean	s.d.	Change in Risk	(3)/(1)	(3)/(2)
Low BW	0.185	0.388	-0.0106	-0.057	-0.027
Neonatal	0.032	0.177	-0.0054	-0.169	-0.031
Infant	0.063	0.242	-0.0129	-0.205	-0.053
Stunted	0.025	0.157	0.0031	0.124	0.020
(B): Mean and s.d. from country-level data					
	Mean	s.d.	Change in Risk	(3)/(1)	(3)/(2)
Low BW	0.197	0.139	-0.011	-0.054	-0.076
Neonatal	0.028	0.014	-0.005	-0.193	-0.386
Infant	0.056	0.026	-0.013	-0.230	-0.496
Stunted	0.024	0.013	0.003	0.129	0.238

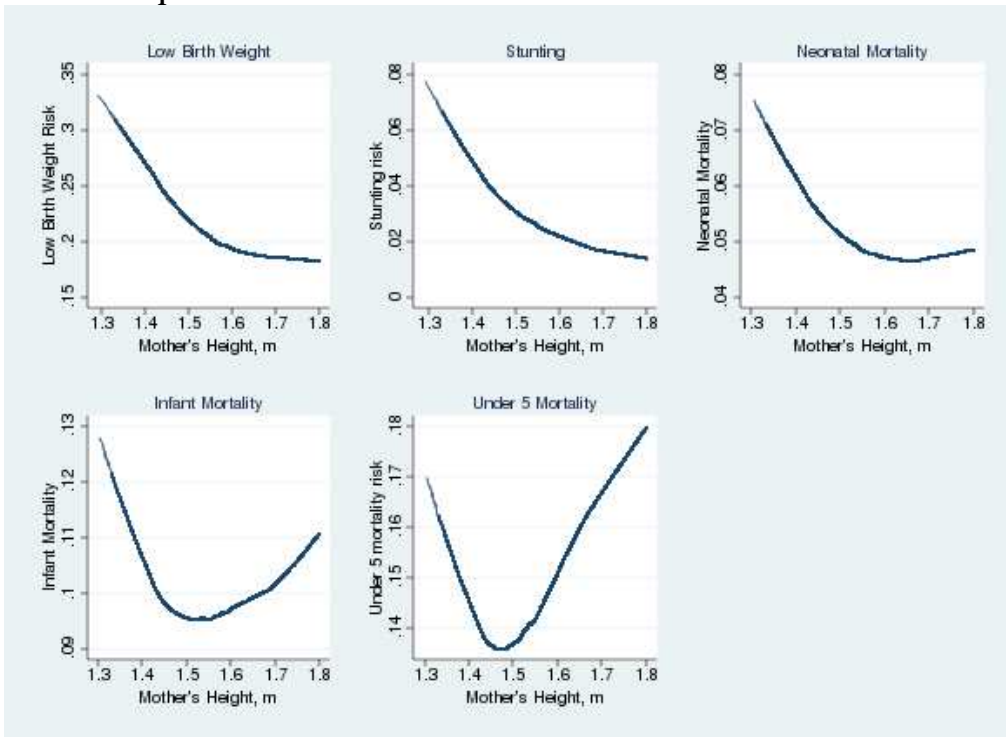
See notes to Table 2b.

Table 3: Linear change per decade in β_t and θ_t

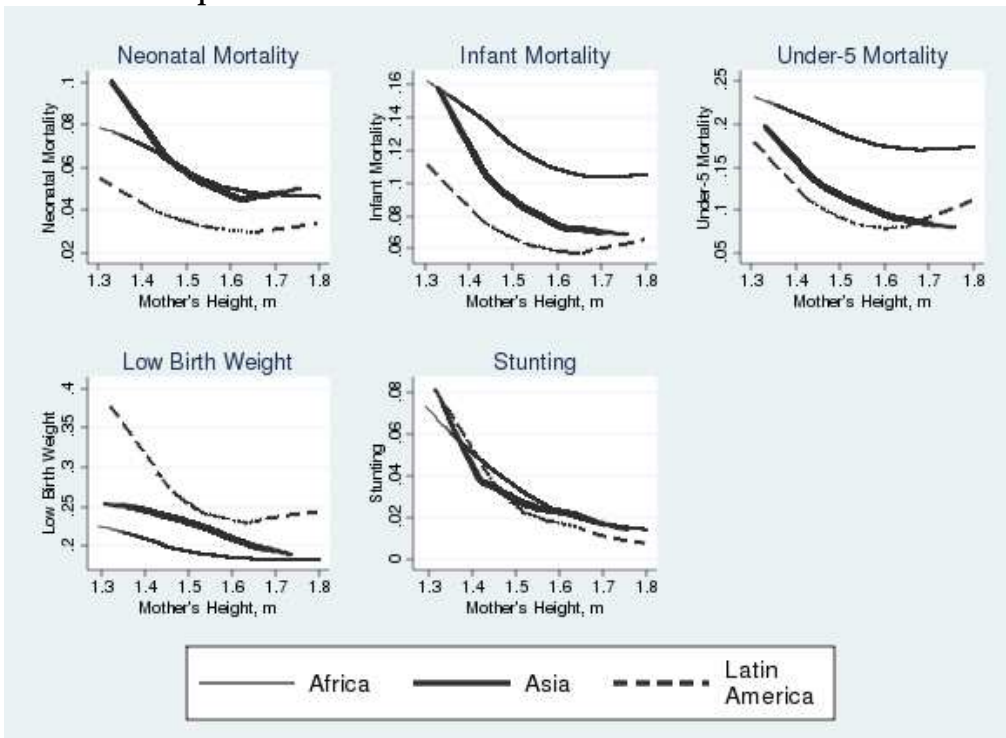
	Neonatal	Infant	Under-5
β_t , change per decade			
Pooled Sample	-0.011**	-0.036**	-0.043**
Africa	0.010*	0.006	0.014
Asia	-0.005	0.002	0.009
Latin America	-0.012	-0.039**	-0.070**
θ_t , change per decade			
Pooled Sample	-0.002	-0.008**	-0.008**
Africa	0.004**	0.002	0.003*
Asia	0.001	0.003	0.005*
Latin America	-0.002	-0.007**	-0.011*
σ_m^c/σ_i^c , change per decade			
<i>Pooled Sample</i>	<i>0.028**</i>	<i>0.021**</i>	<i>0.019**</i>
<i>Africa</i>	<i>0.021**</i>	<i>0.014**</i>	<i>0.011**</i>
<i>Asia</i>	<i>0.026**</i>	<i>0.024**</i>	<i>0.021**</i>
<i>Latin America</i>	<i>0.053**</i>	<i>0.052**</i>	<i>0.051**</i>

* significant at 5%, ** significant at 1%. These are linear trends fitted to β_t and θ_t . Scatters and lowess fits are in Figures 2, A4 and A5. The regressions are run by year and include controls for mother and child characteristics.

Figure 1: (a) Child health indicators against mother's height: Lowess predictions
Pooled sample

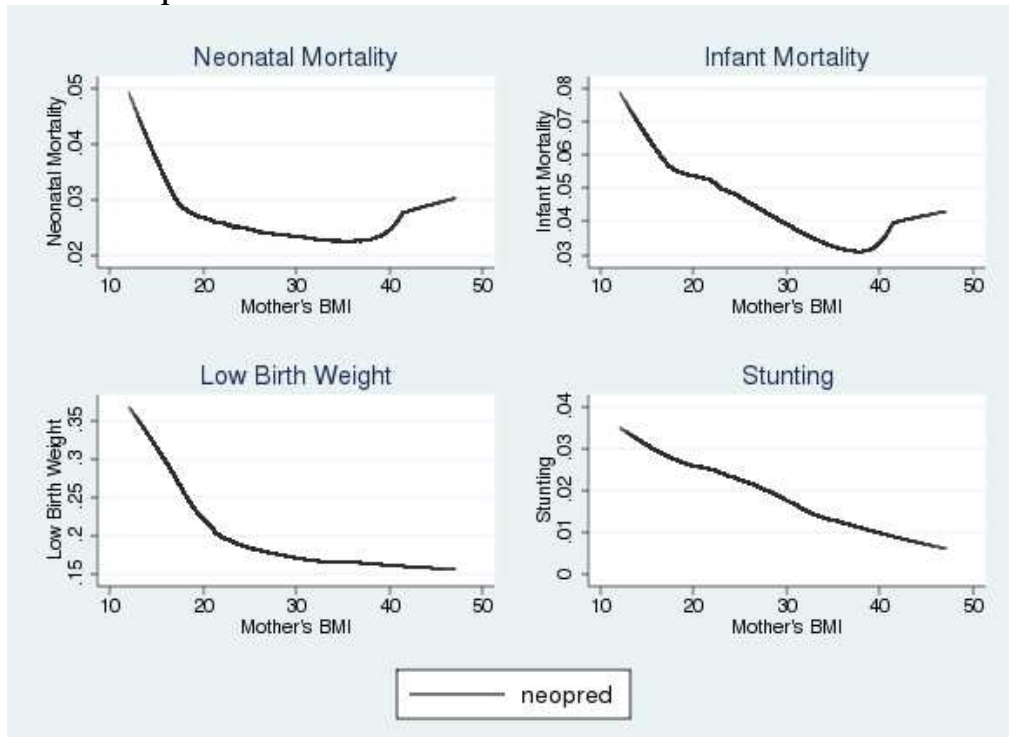


Continent samples

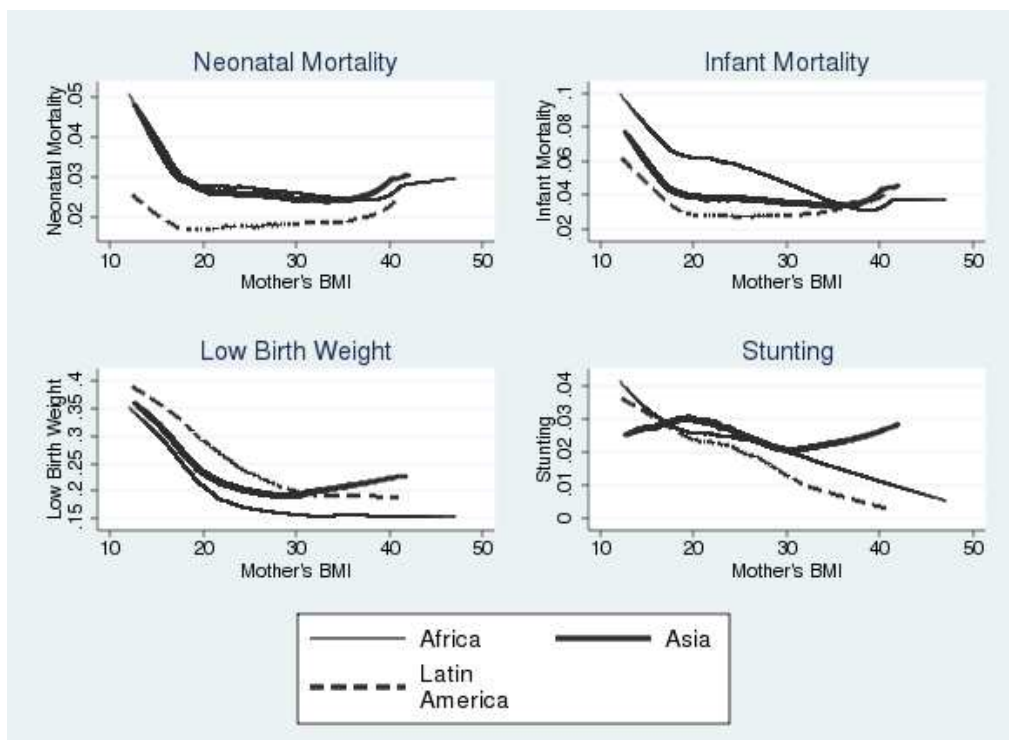


The lowess regressions are run on a random sample of 20% of observations (by country), since otherwise they are too computationally intensive. The 95th percentile of height is 1.675m, the 99th percentile is 1.724m, and the uppermost height observation in the sample is 1.9m.

(b) Child health indicators against mother's BMI: Lowess predictions
Pooled sample



Continent samples



(c) Country-averaged prevalence of poor child health against maternal anemia prevalence

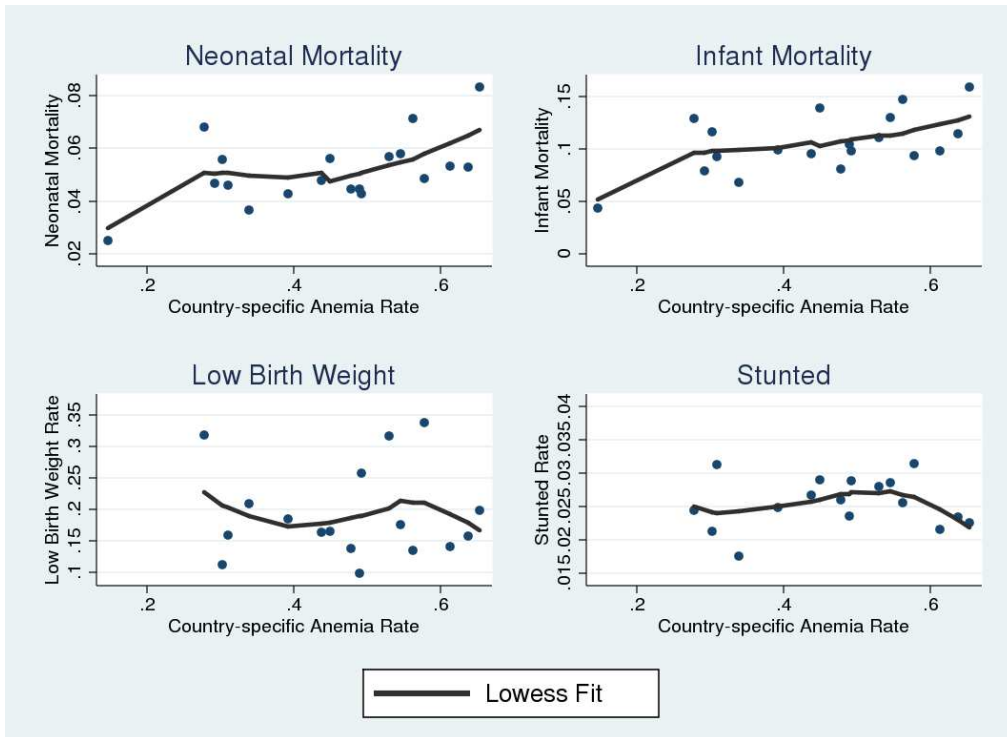
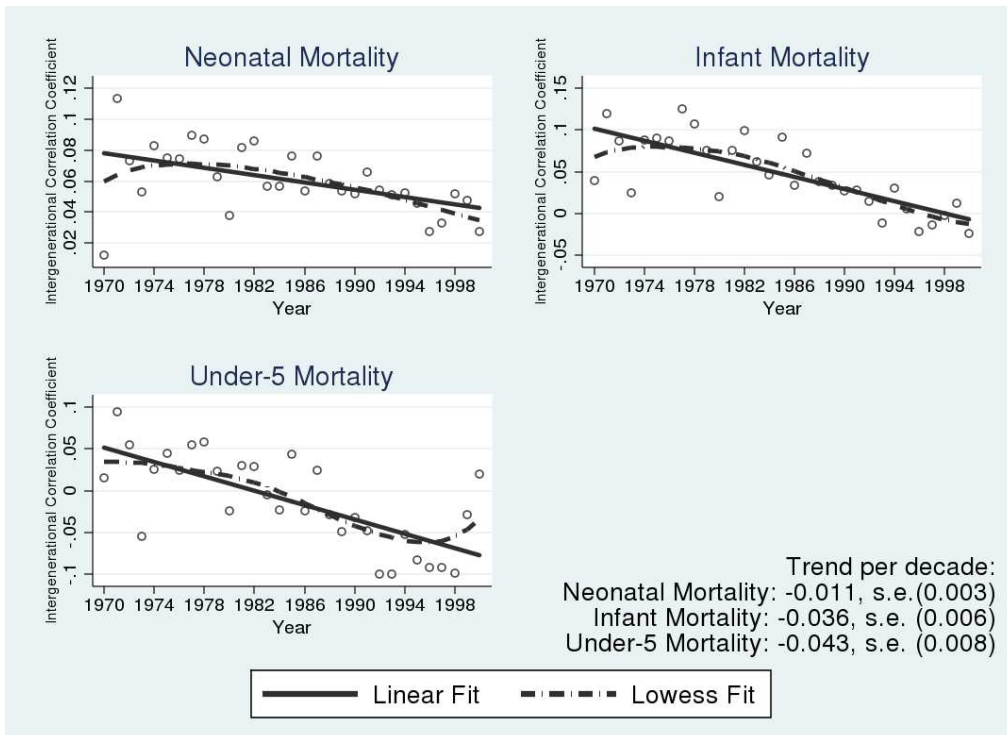
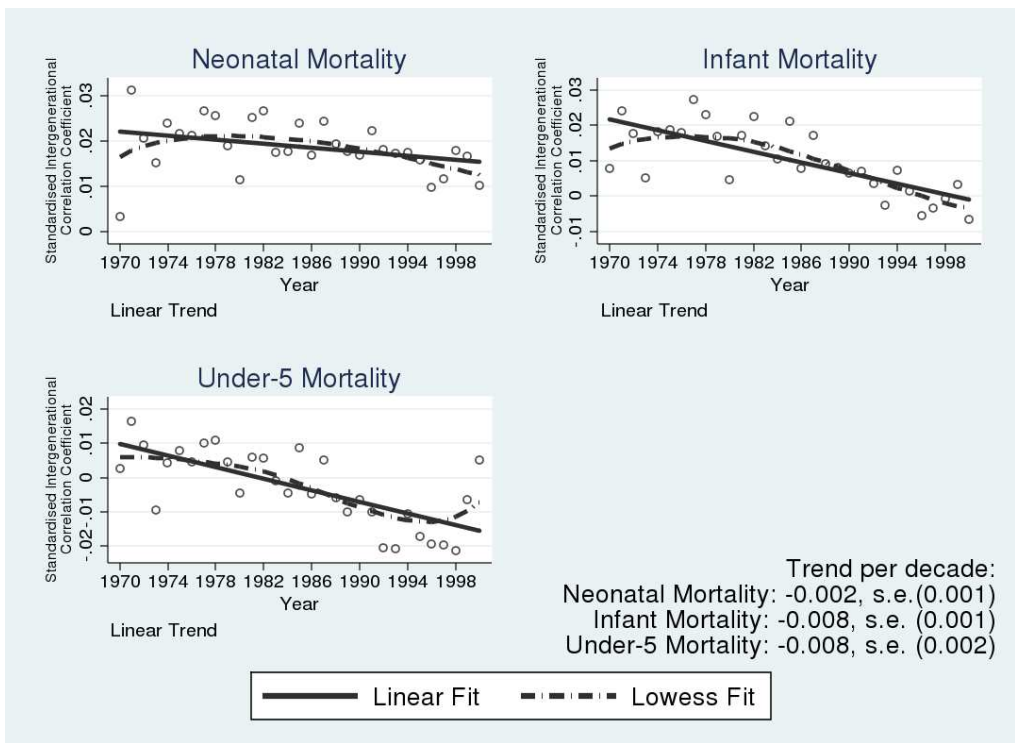


Figure 2: Trends in intergenerational persistence: β_t (regression slope)



Trends in intergenerational persistence: θ_t (correlation coefficient)



Data Appendix

Sources

The DHS data are obtained from www.measuredhs.com. Data on log GDP per capita in constant 2000 prices (chain series; henceforth GDP) for 1970-2000 is obtained from the Penn World Tables (Summers et al. 2009). Immunization rates for measles and DPT (diphtheria and tetanus toxoid with pertussis) are available for 1985-2000 from the World Development Indicators for all countries in our sample other than Namibia. These data measure the percentage of children aged 12-23 months who have been vaccinated

Mother's height

Once fully attained, height is time-invariant, at least until shrinking sets in, and this is likely to be after age 49. As a result, mother's height at the time of the survey can be matched to outcomes for births as long as 30 years before the date of the survey, which facilitates analysis of trends. The fact that height is a measure of permanent health also avoids a problem that marks the literature on intergenerational earnings persistence. This is that estimates often use current rather than permanent income and are therefore sensitive to the life cycle pattern of income. (Zimmerman, 1992). Mother's (and children's) heights are measured to the nearest millimeter at the time of interview by trained surveyors. This avoids the potential biases associated with data on self-reported height (Thomas and Strauss, 1998); the use of self-reported height has, for example, been shown to bias downwards estimates of the heritability of height (MacGregor et al., 2006). We deal with censoring, selectivity, missing data and outliers as follows.

Women in developing countries often continue to grow until about the age of 21 (Deaton, 2007). To allow for full growth, we restrict the sample to women who are at least 21 at the time of the survey. Some surveys interview only ever-married women.⁷ To the extent that shorter women marry younger (Bhalotra, 2008), this creates a selectivity problem. This is limited by eliminating women older than 21. We investigated instead, a cut-off at age 25 and at the (country-specific) 95th percentile of age at marriage. The results are robust to these variations. Height is missing for 21.5% of mothers. We impute these heights using prediction-matching⁸ on the assumption that the missing data are missing at random. To investigate this we estimated the model, pooling the actual and imputed data and including as a regressor an indicator for whether height was imputed. The coefficient on this regressor is small and insignificant for all measures of child health other than under-5 mortality for which it is significant and very small (Table A.4). To remove outliers, we exclude women with height more than three standard deviations away from the country mean, losing about 0.42% of the available sample.

⁷ Table A1 in Bhalotra and Umana-Aponte (2010) identifies these surveys.

⁸ This is done using the `-uvis-` command in Stata (Royston, 2004). For reproducibility of results, we set the random number seed to an arbitrary number, 1001. For robustness, we use the bootstrap method, which does not rely on the assumption that β is normally distributed. For some countries, mother's height is measured in some survey rounds and not in others. In such cases, we drop rounds in which height is not recorded, so the imputation is for surveys in which at least some heights are measured. We drop Nigeria from the analysis since it had a high occurrence of infeasible maternal heights, suggesting a problem with the measurement procedure.

Mother's BMI and anemia status

Mother's BMI (ratio of weight in kilograms to the square of height in meters (kg/m^2)) is measured by surveyors rather than self-reported. It is collected for women with births in the three years preceding the survey, excluding women who are pregnant or less than three months postpartum. To remove outliers, we exclude women whose BMI is greater than 3 standard deviations away from the country mean. Data on anemia are collected by a finger prick blood test and are available for 20 of the 34 surveys taken after 1999, in 19 countries. Testing is voluntary and 43.0% of the potential sample volunteer.⁹ The data are recorded as a categorical variable indicating no, mild, moderate, or severe anemia. The BMI and anemia analyses are run restricting the sample to the most recent birth of each mother.

Childhood mortality indicators

Mortality information is available for all births to each mother from retrospective accounts, as a result of which it spans 31 birth cohorts, 1970-2000. The rest of this subsection describes how we deal with potential issues of selectivity and recall bias raised by the retrospective nature of the data and the more general issues of age heaping and censoring. An upper limit to the age of the mother at the time of the survey implies fewer births for the earlier years in the sample and that these births are disproportionately from young mothers (Bumpass et al., 1978; Bhalotra, 2009). We therefore limit the sample to children born after 1970 and control for maternal age at birth. We drop children born after the year 2000 because the data are relatively thin and come disproportionately from a few countries. Naturally, the sample only includes mothers who are alive at the time of interview and this sample is likely to be selectively healthy. We might expect that in a linear model in which individual child health is conditional upon mother's health, the mother-child correlation of health will be unbiased but in practice this will seldom hold because our measures of mother's health are partial and no one will not fully capture her risk of early mortality. Given that we find a stronger relationship of mother-child health indicators for mothers in weaker health, our estimates are likely to under-estimate the strength of this relationship. Mortality (and birth weight) are reported by the mother but interviewers conduct a number of probes to ensure the quality of birth history data (ORC Macro, 2006). Recall bias is a common concern with retrospective data but the DHS design is sensitive to this. Also, births and deaths are, by their nature, less likely to be remembered with error than, for example, income or expenditure. To account for the heaping of age at 'round' ages, we define mortality to include deaths in the first, twelfth and sixtieth month, although the convention is to measure neonatal, infant and under-5 mortality as death before the age of 1, 12, and 60 months respectively.¹⁰ To ensure that children are fully exposed to the relevant mortality risk, the samples are

⁹ We recognise that the volunteers may not be a random sample of women but we control for age and education of mothers which are likely to determine selection into testing. There is no *a priori* reason that this self-selection is correlated with child health, although this cannot be ruled out.

¹⁰ The results for earnings transition matrices presented in O'Neill et al. (2007) suggest that measurement error in the dependent variable will lead us to under-estimate persistence. If there is measurement error in the indicators of both child and

adjusted to exclude children younger than the cut-off age at the time of interview (e.g. children less than or equal to 12 months of age at interview are excluded from the sample used to analyse infant mortality).

Indicators of growth failure in childhood

Observations on birth weight and child anthropometric outcomes are only available for recent births and, at irregular intervals, span 1986-2000. Child height is measured by surveyors for surviving births in the three, four or five years preceding the survey, the range depending upon the survey (see Table A.2 for survey years for each country). Mothers are asked to provide subjective recall data on birth weight for births in the five years preceding the survey, whether or not they survived.¹¹

We classify a child as stunted if their height is less than 2 standard deviations below the mean height in their country for their age and gender. This avoids the assumption that well nourished children in different populations have the same potential and the same trajectories of growth. The long-standing WHO convention has been to define stunting in developing countries with reference to the mean and s.d. of a sample of healthy American children but this has recently been called into question (Klasen and Moradi, 2000; De Onis et al., 2004). An unavoidable problem with using child height as the dependent variable is that the data are subject to survival selection. If taller children are systematically more likely to survive then this causes the stunting threshold to be higher than in the absence of mortality, leading to a 'healthier' group of stunted children. Outliers in child height are removed by setting boundaries at 30 cm and 120 cm, which results in a loss of about 750 observations.

Birth weight data are scarce in developing countries because most births occur at home and there is no systematic registration of births. Measured birth weight (in grams) is only available for around 25% of 'eligible' observations. Mothers' subjective assessment of birth weight, which classifies the child's birth weight as either above, at, or below average, is available for 60.54% of births in the 5 years prior to each survey. We use this to define low (i.e. below average) birth weight. Previous work has investigated the quality of these data- an encouraging result is that the mean birth weight for babies classified by mothers as small was less than 2500 grams in all except 7 of 62 countries (Blanc and Wardlaw, 2005).

mother's health then the impact on mobility estimates depends upon the correlation between the measurement errors. If it is low, as is likely in this setting where the indicators are different, then again we will tend to under-estimate persistence.

11 Lesotho, Honduras and Comoros have small sample sizes for child height and birth weight, and are excluded from these samples. Sample sizes for height (birth weight) were as follows: Comoros - 894 (998), Lesotho - 263 (670), Honduras - 15 (15)

Table A1: Countries in Sample, by Continent

Country	Country Code	Country	Country Code
African Countries			
Benin	BE	Madagascar	MD
Burkina Faso	BF	Malawi	MW
CAR	CR	Mali	ML
Cameroon	CM	Morocco	MO
Chad	CH	Mozambique	MZ
Comoros	CO	Namibia	NB
Cote d'Ivoire	CI	Niger	NG
Egypt	EG	Rwanda	RW
Ethiopia	ET	Senegal	SE
Gabon	GB	Tanzania	TZ
Ghana	GH	Togo	TO
Guinea	GU	Uganda	UG
Kenya	KE	Zambia	ZB
Lesotho	LE	Zimbabwe	ZW
Asian Countries			
Cambodia	CD	Turkey	TK
India	IN		
Latin American Countries			
Brazil	BR	Honduras	HO
Colombia	CB	Nicaragua	NC
Dominican Republic	DR	Peru	PE
Haiti	HA		

Table A2: Country Characteristics

Country	% GDP growth, 1970-2000	Potential Birth Year Sample	Interview Dates	Means of Child Health				
				Neonatal Mortality	Infant Mortality	Under-5 Mortality	Low Birth Weight	Stunting
Benin	0.103	1960-2001	1996, 2001	0.053	0.117	0.206	0.158	0.023
Brazil	1.367*	1959-1996	1996	0.033	0.067	0.079	0.244	0.017
Burkina Faso	0.955*	1955-2003	1992-1993, 1998-1999, 2003	0.053	0.132	0.23	0.176	0.027
CAR	-0.927*	1958-1995	1994-1995	0.048	0.1	0.135	0.141	0.021
Cambodia	-2.275*	1965-2000	2000	0.051	0.097	0.153	0.164	0.024
Cameroon	0.682	1960-2004	1998, 2004	0.059	0.107	0.17	0.163	0.03
Chad	-0.044	1960-2004	1996-1997, 2004	0.021	0.134	0.22	0.318	0.021
Colombia	1.781*	1958-2005	1995, 2000, 2004-2005	0.061	0.034	0.041	0.201	0.024
Comoros	-0.312	1961-1996	1996	0.034	0.11	0.15		
Cote d'Ivoire	0.123	1956-1999	1994, 1998-1999	0.059	0.106	0.165	0.163	0.024
Dominican Republic	2.277*	1952-1996	1991, 1996	0.046	0.061	0.082	0.2	0.021
Egypt	3.126*	1954-2005	1992-1993, 1995-1996, 2000, 2005	0.068	0.095	0.127	0.159	0.028
Ethiopia	1.018*	1955-1997	1992, 1997	0.034	0.13	0.201	0.318	0.021
Gabon	-1.722*	1962-2000	2000-2001	0.044	0.068	0.101	0.116	0.02
Ghana	0.188	1957-2003	1993-1994, 1998-1999	0.071	0.082	0.137	0.137	0.025
Guinea	-0.098	1961-2005	1999, 2005	0.057	0.149	0.237	0.134	0.027
Haiti	-0.12	1957-2000	1994-1995, 2000	0.025	0.114	0.178	0.317	0.028
Honduras	0.471*	1969-2006	2005-2006	0.063	0.044	0.057		
India	2.831*	1961-2000	1998-2000, 2005-2006	0.035	0.095	0.125	0.237	0.024

Table A2: Country Characteristics

Country	% GDP growth, 1970-2000	Potential Birth Year Sample	Interview Dates	Neonatal Mortality	Infant Mortality	Under-5 Mortality	Low Birth Weight	Stunting
Kenya	0.338*	1954-2003	1993, 2003	0.047	0.076	0.108	0.151	0.02
Lesotho	2.889*	1967-2005	2004-2005	0.043	0.08	0.094		
Madagascar	-1.860*	1962-2004	1997, 2003-2004	0.056	0.099	0.158	0.258	0.029
Malawi	1.210*	1954-2005	1992, 2000, 2004-2005	0.083	0.141	0.231	0.165	0.027
Mali	1.557*	1960-2001	1995-1996, 2001	0.047	0.163	0.281	0.198	0.02
Morocco	1.659*	1953-2004	1992, 2003-2004	0.067	0.08	0.104	0.246	0.028
Mozambique	-0.27	1960-2004	1997, 2003-2004	0.039	0.16	0.227	0.185	0.038
Namibia	-0.139	1957-1992	1992	0.029	0.07	0.106	0.17	0.026
Nicaragua	-2.984*	1961-2001	1997-1998, 2001	0.062	0.061	0.079	0.299	0.025
Niger	-0.931*	1954-1998	1992, 1998	0.037	0.157	0.335	0.364	0.021
Peru	-0.964*	1956-2000	1991-1992, 2000	0.058	0.07	0.102	0.208	0.016
Rwanda	-0.688	1963-2005	2000, 2005	0.056	0.118	0.198	0.112	0.018
Senegal	-0.091	1954-2005	1992-1993, 2005	0.049	0.096	0.17	0.337	0.032
Tanzania	0.353	1953-2005	1991-1992, 1996, 2004-2005	0.045	0.107	0.165	0.098	0.02
Togo	-1.965*	1958-1998	1998	0.05	0.096	0.166	0.183	0.02
Turkey	2.068*	1957-1998	1993, 1998	0.047	0.092	0.113	0.258	0.03
Uganda	-0.492	1959-2001	1995, 2000-2001	0.043	0.102	0.175	0.185	0.022
Zambia	-1.861*	1954-2002	1992, 1996-1997, 2001-2002	0.041	0.104	0.17	0.121	0.021
Zimbabwe	-0.057	1957-1999	1994, 1999	0.028	0.06	0.085	0.164	0.024

item * indicates growth was significant at 5% level. %`Low BW' refers to Low Birth Weight.

Table A3: Correlations amongst indicators of child and maternal health (proportions)

	Height		Neonatal		Infant		Under-5		Low Birth Weight		Stunted	
	Tall	Short	0	1	0	1	0	1	0	1	0	1
Neonatal	0.04	0.05	1	1	0	0.51	0	0.36	0.03	0.06		
Infant	0.09	0.1	0.05	1	1	1	0	0.7	0.06	0.1		
Under-5	0.12	0.14	0.09	1	0.04	1	1	1	0.08	0.12		
Low BW	0.18	0.21	0.2	0.34	0.2	0.28	0.2	0.26	1	1	0.2	0.27
Stunting	0.19	0.28							0.02	0.03	1	1

We use the smallest possible sample when calculating these proportions. For stunting and birth weight, this is the stunting sample. For birth weight and mortality, this is the birth weight sample. There are no figures for the correlation of stunting and mortality since any children who do not survive are not present in the stunting sample by construction. All children who die in the first month of life are naturally classified as having died in the first year and first 5-years of life. Similarly, children who survive the first 5 years of life necessarily survive their first year and month. ‘Tall’ refers to a mother's height $> \text{mean} + (0.5 \text{ s.d.})$, ‘Short’ is defined as a mother's height $< \text{mean} - (0.5 \text{ s.d.})$. We perform tests of the significance of differences in means within each column, and find that all means are significantly different at the 1% level. ‘Low BW’ refers to low birth weight.

Table A4: Investigating Missing Data on Heights

	Low Birth Weight	Stunted	Neonatal Mortality	Infant Mortality	Under-5 Mortality
Mother's Height	-0.217** (0.031)	-0.098** (0.007)	-0.066** (0.008)	-0.103** (0.010)	-0.139** (0.013)
Mother's Height is Missing	-0.006 (0.005)	-0.001 (0.003)	0.002 (0.001)	-0.000 (0.002)	-0.006* (0.003)
Observations	324097	285440	2295740	2233720	1894526

Robust standard errors in brackets. * significant at 5% ** significant at 1%. Height is missing is an indicator variable for height having been imputed.

Table A5: Alternative measure of child health: Z-scores of Height

A: Linear			
	Height	BMI	Anemia
Height	3.182*** (0.168)		
BMI		0.022*** (0.001)	
Anemia			-0.036* (0.017)
B: Non-Linear			
Tall (0.5-1 s.d.)	0.161** (0.011)		
Tall (1-2 s.d.)	0.293** (0.022)		
Tall (> 2 s.d.)	0.446** (0.044)		
Short (0.5-1 s.d.)	-0.172** (0.011)		
Short (1-2 s.d.)	-0.307** (0.015)		
Short (> 2 s.d.)	-0.504** (0.031)		
Low BMI		-0.085** (0.013)	
High BMI		0.142** (0.010)	
No Anemia			0.034* (0.016)
Severe Anemia			-0.102** (0.061)
Observations	283624	215193	20621
$p > \chi^2$: (tallhalf/shorthalf)	0.000		
$p > \chi^2$: (tall1/short1)	0.000		
$p > \chi^2$: (tall2/short2)	0.000		
$p > \chi^2$: (lowbmi/highbmi)		0.000	
$p > \chi^2$: (no anemia/severe anemia)			0.074

The dependent variable is the z-score of child height. The specifications correspond to those in Table 1 and 2. Robust standard errors in brackets, allowing for clustering within country. * significant at 5% sym** significant at 1%. Mother's height is in metres. Mother's BMI is the ratio of weight in kilograms to the square of the height in meters (kg/m^2). Anemia is a dummy variable indicating whether the mother had mild, moderate, or severe anemia at the time of the survey. Estimator is OLS. We control for child gender, birth order and birth month, mother's religion, age at birth, education, and partner's education. Estimates include country-year fixed effects. Means and s.d. of variables are in Tables 1 and 2. The $p > \chi^2$ statistic gives the significance level at which we are unable to reject the null that the coefficients for tallhalf/shorthalf, tall1/short1, or tall2/short2 are significantly different in magnitude from one another. Average Z-score for women who are greater than 2 standard deviations above the mean is 0.533, average z-score for women who are less than 2 standard deviations below the mean is -0.566.

Table A6: Linear Trends in Intergenerational Persistence: Overall and by Continent

	Neonatal Mortality	Infant Mortality	Under-5 Mortality
Pooled Sample			
Trend	0.001 (0.000)	0.002** (0.001)	0.003** (0.001)
Observations	2295740	2233720	1894526
Africa			
Trend	0.000 (0.000)	0.001 (0.001)	0.001 (0.001)
Observations	1325293	1285169	1062923
Asia			
Trend	0.002 (0.001)	0.003 (0.001)	0.003 (0.003)
Observations	514309	504395	454733
Latin America			
Trend	0.002 (0.001)	0.005** (0.001)	0.003** (0.001)
Observations	456138	444156	376870

Estimates are of the linear baseline model in panel A of Table 1, augmented with an interaction term between mother's height and a global trend. We now replace country-year fixed effects with country and year fixed effects and country-specific trends. 'Trend' reports the coefficient on the interaction term, an estimate of the rate of decline of the mother-child correlation of health. Robust standard errors in brackets, allowing for clustering within country. * significant at 5% ** significant at 1%. Means and s.d. for the pooled sample are in Table 1. Continent-specific means and s.d. are in Table A.5. See notes to Table 1.

Table A7: Intergenerational Persistence: Continent-Specific Estimates

	Neonatal Mortality		
	Africa	Asia	Latin America
Height (β)	-0.068** (0.007)	-0.093* (0.011)	-0.030* (0.011)
θ	-0.019	-0.023	-0.011
τ	-1.308	-1.691	-0.882
Observations	1325293	514309	456138
mean (neonatal)	0.052	0.055	0.034
s.d. (neonatal)	0.222	0.228	0.18

	Infant Mortality		
	Africa	Asia	Latin America
Height (β)	-0.108** (0.010)	-0.129** (0.009)	-0.057* (0.018)
θ	-0.022	-0.026	-0.015
τ	-0.964	-1.483	-0.877
Observations	1285169	504395	444156
mean (infant)	0.112	0.087	0.065
s.d. (infant)	0.316	0.282	0.246

	Under-5 Mortality		
	Africa	Asia	Latin America
Height (β)	-0.142** (0.012)	-0.170** (0.013)	-0.088* (0.029)
θ	-0.023	-0.031	-0.020
τ	-0.793	-1.504	-0.989
Observations	1062923	454733	376870
mean (under-5)	0.179	0.113	0.089
s.d. (under-5)	0.383	0.317	0.284
mean (height)	1.585	1.519	1.527
s.d. (height)	0.063	0.057	0.064

Table A7: Continued

	Low Birth Weight		
	Africa	Asia	Latin America
Height (β)	-0.182** (0.029)	-0.083** (0.028)	-0.424** (0.075)
θ	-0.029	-0.011	-0.064
τ	-0.973	-0.367	-1.717
Observations	228381	35404	60312
mean(low birth weight)	0.187	0.226	0.247
s.d. (low birth weight)	0.39	0.418	0.431
mean (height)	1.587	1.52	1.534
s.d. (height)	0.063	0.057	0.065
Stunted			
	Africa	Asia	Latin America
Height (β)	-0.091** (0.007)	-0.079* (0.015)	-0.122** (0.023)
θ	-0.037	-0.027	-0.051
τ	-3.640	-2.821	-5.083
Observations	188705	29090	67645
mean (stunting)	0.025	0.028	0.024
s.d. (stunting)	0.157	0.164	0.153
mean (height)	1.586	1.523	1.53
s.d. (height)	0.063	0.057	0.064

The linear baseline model in panel A of Table 1a is estimated by continent. β are estimated coefficients, We normalise the marginal effect of height (τ) by dividing it by the continent-specific mean of dependant variable. We standardize it (θ) by multiplying it by the continent-specific ratio of the standard deviation of maternal height to the standard deviation of the dependant variable. Robust standard errors in brackets, allowing for clustering within country. * significant at 5%, ** significant at 1%. See notes to Table 1a.

Table A.8: Country Differences in Intergenerational Persistence: Range and Significance

Health Measure	Country: Lowest		Country: Highest		Number of Significant Coefficients
Marginal Effect, beta					
Neonatal Mortality	Zambia	-0.029	Comoros	-0.182	28 / 38
Infant Mortality	Zimbabwe	-0.071	Mozambique	-0.212	27 / 38
Under-5 Mortality	Turkey	-0.065	Niger	-0.283	29 / 38
Low Birth Weight	Peru	-0.091	Morocco	-0.694	20 / 35
Stunting Incidence	Kenya	-0.062	Ethiopia	-0.243	25 / 35
Normalized Marginal Effect, tau					
Neonatal Mortality	Zambia	-0.009	Comoros	-0.044	
Infant Mortality	Guinea	-0.014	Mozambique	-0.035	
Under-5 Mortality	Turkey	-0.012	Nicaragua	-0.040	
Low Birth Weight	Chad	-0.655	Colombia	-2.910	
Stunting Incidence	Kenya	-0.022	Brazil	-0.104	
Standardized Marginal Effect, theta					
Neonatal Mortality	Zambia	-0.69	Honduras	-3.111	
Infant Mortality	Guinea	-0.541	Honduras	-2.426	
Under-5 Mortality	Chad	-0.462	Honduras	-2.528	
Low Birth Weight	Peru	-0.019	Colombia	-0.086	
Stunting Incidence	Kenya	-2.385	Brazil	-11.511	

We normalize the marginal effect of height by dividing it by the country-specific mean of child health. We standardize it by multiplying it by the country-specific ratio of the standard deviation of maternal height to the standard deviation of child health. The range of coefficients only includes coefficients which were significantly different from zero. 'Low BW' refers to low birth weight.

Table A9: Country-specific Estimates of Persistence: Low Birth Weight

	BE	BR	BF	CR	CD	CM	CH	CB	CO	CI	DR	EG
Height (β)	0.028 (0.075)	-0.444** (0.105)	-0.200** (0.053)	-0.136 (0.117)	-0.006 (0.073)	-0.170 (0.092)	-0.212* (0.089)	-0.574** (0.103)	-0.327** (0.091)	-0.346** (0.081)	-0.235** (0.039)	-0.270** (0.075)
θ	0.005	-0.065	-0.032	-0.025	-0.001	-0.029	-0.028	-0.087	-0.052	-0.051	-0.036	-0.035
τ	0.177	-1.755	-1.190	-0.855	-0.041	-1.090	-0.656	-2.914	-1.946	-1.640	-1.460	-0.833
Observations	6451	4263	14454	2058	7932	3884	7135	4509	4566	7105	29877	10345
mean (dep. var.)	0.158	0.253	0.168	0.159	0.147	0.156	0.323	0.197	0.168	0.211	0.161	0.324
s.d. (dep. var.)	0.365	0.435	0.374	0.366	0.355	0.363	0.468	0.398	0.374	0.408	0.368	0.468
mean (height)	1.583	1.555	1.617	1.588	1.523	1.604	1.627	1.544	1.594	1.564	1.573	1.572
s.d. (height)	0.06	0.064	0.059	0.068	0.054	0.061	0.061	0.06	0.06	0.06	0.056	0.06
	ET	GB	GH	GU	HA	HO	IN	KE	LE	MD	MW	ML
Height (β)	0.029 (0.099)	-0.166* (0.072)	-0.301** (0.078)	-0.251** (0.081)	-0.091 (0.050)	-0.244** (0.056)	-0.041 (0.111)	-0.287** (0.052)	-0.110 (0.057)	-0.174 (0.090)	-0.342** (0.079)	-0.300** (0.109)
θ	0.006	-0.028	-0.054	-0.033	-0.012	-0.042	-0.005	-0.045	-0.016	-0.023	-0.052	-0.050
τ	0.257	-1.153	-2.213	-0.789	-0.368	-1.638	-0.164	-1.783	-0.489	-0.702	-1.819	-1.667
Observations	3181	6436	5306	9161	24479	10607	4672	15712	14640	7284	7017	3146
mean (dep. var.)	0.113	0.144	0.136	0.318	0.247	0.149	0.25	0.161	0.225	0.248	0.188	0.18
s.d. (dep. var.)	0.316	0.351	0.343	0.466	0.431	0.356	0.433	0.367	0.417	0.432	0.391	0.384
mean (height)	1.585	1.589	1.588	1.582	1.514	1.598	1.536	1.56	1.617	1.574	1.559	1.61
s.d. (height)	0.061	0.059	0.061	0.061	0.056	0.062	0.058	0.057	0.06	0.056	0.06	0.064
	MO	MZ	NB	NC	NG	PE	RW	SE	TZ	TO	TK	
Height (β)	-0.694** (0.076)	-0.403** (0.083)	-0.332** (0.051)	-0.227** (0.059)	0.119 (0.100)	-0.091* (0.043)	-0.046 (0.113)	-0.238 (0.144)	-0.106 (0.061)	0.003 (0.044)	-0.132 (0.089)	
θ	-0.086	-0.050	-0.045	-0.044	0.015	-0.019	-0.007	-0.030	-0.017	0.001	-0.021	
τ	-2.298	-1.165	-1.566	-2.027	0.350	-0.910	-0.238	-0.919	-0.596	0.025	-0.772	
Observations	11640	9286	23634	8276	5916	13895	3543	2993	10210	15598	4886	
mean (dep. var.)	0.302	0.346	0.212	0.112	0.34	0.1	0.193	0.259	0.178	0.122	0.171	
s.d. (dep. var.)	0.459	0.476	0.408	0.316	0.474	0.3	0.395	0.438	0.383	0.327	0.376	
mean (height)	1.533	1.606	1.501	1.58	1.625	1.562	1.589	1.561	1.585	1.58	1.599	
s.d. (height)	0.057	0.059	0.055	0.061	0.06	0.062	0.06	0.055	0.063	0.06	0.061	

Robust standard errors in brackets. * significant at 5%, ** significant at 1%. See notes to Table 1a. Every equation includes a linear trend. Country name acronyms in the column heads are expanded in Appendix Table 1. We normalise the coefficient of height (β) by dividing it by the country-specific mean of dependant variable. We standardize it (θ) by multiplying it by the country-specific ratio of the standard deviation of maternal height to the standard deviation of the dependant variable.

Table A.10: Country-specific Estimates of Persistence: Stunting

	BE	BR	BF	CR	CD	CM	CH	CB	CO	CI	DR	EG
Height (β)	-0.100** (0.035)	-0.224** (0.041)	-0.105** (0.026)	-0.152 (0.078)	-0.043 (0.051)	-0.118* (0.057)	-0.097** (0.033)	-0.217** (0.030)	-0.107* (0.042)	-0.114** (0.034)	-0.086** (0.018)	-0.049 (0.031)
θ	-0.038	-0.103	-0.037	-0.060	-0.014	-0.044	-0.042	-0.084	-0.042	-0.043	-0.028	-0.020
τ	-4.000	-11.200	-3.750	-5.067	-1.792	-4.069	-4.619	-9.042	-4.458	-4.385	-2.966	-2.042
Observations	5573	3598	11619	1861	3493	2250	5684	9573	3970	5841	27093	8380
mean (dep. var.)	0.025	0.020	0.028	0.030	0.024	0.029	0.021	0.024	0.024	0.026	0.029	0.024
s.d. (dep. var.)	0.156	0.139	0.166	0.170	0.153	0.167	0.142	0.153	0.154	0.158	0.167	0.153
mean (height)	1.583	1.555	1.617	1.588	1.526	1.603	1.627	1.545	1.594	1.564	1.573	1.572
s.d. (height)	0.060	0.064	0.059	0.067	0.051	0.062	0.061	0.059	0.060	0.060	0.055	0.061
	ET	GB	GH	GU	HA	HO	IN	KE	LE	MD	MW	ML
Height (β)	-0.243** (0.060)	-0.109** (0.038)	-0.031 (0.036)	-0.131** (0.033)	-0.071** (0.022)	-0.110** (0.026)	-0.130** (0.047)	-0.062* (0.029)	-0.035 (0.024)	-0.047 (0.040)	-0.130** (0.036)	-0.037 (0.051)
θ	-0.095	-0.039	-0.012	-0.049	-0.024	-0.046	-0.047	-0.022	-0.014	-0.015	-0.047	-0.016
τ	-10.125	-4.037	-1.240	-4.679	-2.536	-4.783	-4.815	-2.385	-1.591	-1.516	-4.643	-1.682
Observations	2714	5555	4040	7692	20463	9051	4041	12624	12011	6581	5776	2292
mean (dep. var.)	0.024	0.027	0.025	0.028	0.028	0.023	0.027	0.026	0.022	0.031	0.028	0.022
s.d. (dep. var.)	0.154	0.163	0.156	0.164	0.165	0.148	0.162	0.159	0.147	0.172	0.166	0.148
mean (height)	1.583	1.589	1.589	1.582	1.514	1.597	1.535	1.560	1.617	1.573	1.559	1.609
s.d. (height)	0.060	0.059	0.061	0.061	0.056	0.062	0.058	0.056	0.059	0.056	0.060	0.065
	MO	MZ	NB	NC	NG	PE	RW	SE	TZ	TO	TK	
Height (β)	-0.061 (0.034)	-0.150** (0.029)	-0.097** (0.016)	-0.117** (0.031)	-0.110* (0.044)	-0.094** (0.023)	0.012 (0.045)	-0.137** (0.050)	-0.140** (0.030)	-0.101** (0.025)	0.004 (0.045)	
θ	-0.020	-0.063	-0.036	-0.049	-0.038	-0.038	0.005	-0.044	-0.056	-0.040	0.002	
τ	-2.033	-7.500	-4.619	-5.318	-3.548	-3.917	0.522	-4.567	-5.385	-4.391	0.174	
Observations	10156	7219	30785	6372	4295	11654	3260	5134	8211	12559	4020	
mean (dep. var.)	0.03	0.02	0.021	0.022	0.031	0.024	0.023	0.03	0.026	0.023	0.023	
s.d. (dep. var.)	0.17	0.14	0.144	0.146	0.174	0.153	0.149	0.17	0.158	0.15	0.151	
mean (height)	1.533	1.607	1.501	1.581	1.624	1.562	1.589	1.556	1.585	1.581	1.599	
s.d. (height)	0.057	0.059	0.054	0.061	0.06	0.062	0.059	0.055	0.063	0.06	0.06	

Robust standard errors in brackets. * significant at 5%, ** significant at 1%. See notes to Table 1a. Every equation includes a linear trend. Country name acronyms in the column heads are expanded in Appendix Table 1. We normalise the coefficient of height (β) by dividing it by the country-specific mean of dependant variable. We standardize it (θ) by multiplying it by the country-specific ratio of the standard deviation of maternal height to the standard deviation of the dependant variable.

Table A.11: Country-specific Estimates of Persistence: Neonatal Mortality

	BE	BR	BF	CR	CD	CM	CH	CB	CO	CI	DR	EG
Height (β)	-0.085** (0.020)	-0.042* (0.019)	-0.091** (0.014)	-0.048 (0.028)	-0.053* (0.021)	-0.049** (0.018)	-0.032 (0.019)	0.005 (0.008)	-0.182** (0.047)	-0.131** (0.024)	-0.017 (0.017)	-0.051** (0.009)
θ	-0.023	-0.014	-0.023	-0.015	-0.013	-0.014	-0.009	0.002	-0.044	-0.033	-0.006	-0.013
τ	-1.574	-1.200	-1.625	-0.960	-0.981	-1.065	-0.561	0.238	-2.984	-2.259	-0.515	-1.063
Observations	35216	23716	73621	14310	39881	36063	39581	98780	7341	27770	31350	183100
mean (dep. var.)	0.054	0.035	0.056	0.050	0.054	0.046	0.057	0.021	0.061	0.058	0.033	0.048
s.d. (dep. var.)	0.225	0.184	0.230	0.218	0.226	0.210	0.231	0.145	0.239	0.234	0.177	0.213
mean (height)	1.584	1.552	1.615	1.583	1.525	1.602	1.626	1.543	1.549	1.591	1.561	1.575
s.d. (height)	0.061	0.063	0.059	0.066	0.054	0.062	0.062	0.060	0.058	0.059	0.061	0.056
	ET	GB	GH	GU	HA	HO	IN	KE	LE	MD	MW	ML
Height (β)	-0.122** (0.015)	0.007 (0.024)	-0.049** (0.018)	-0.078** (0.021)	0.000 (0.020)	-0.084** (0.015)	-0.101** (0.006)	-0.048** (0.013)	-0.044 (0.035)	-0.051** (0.018)	-0.063** (0.015)	-0.078** (0.017)
θ	-0.030	0.002	-0.014	-0.018	0.000	-0.032	-0.025	-0.016	-0.012	-0.015	-0.016	-0.017
τ	-1.906	0.219	-1.089	-1.083	0.000	-3.111	-1.804	-1.371	-0.880	-1.275	-1.125	-0.940
Observations	71459	14679	37114	41874	36565	38312	441338	60662	11434	36567	77813	78973
mean (dep. var.)	0.064	0.032	0.045	0.072	0.054	0.027	0.056	0.035	0.050	0.040	0.056	0.083
s.d. (dep. var.)	0.245	0.177	0.208	0.258	0.227	0.161	0.230	0.185	0.218	0.197	0.229	0.275
mean (height)	1.573	1.585	1.589	1.587	1.579	1.514	1.516	1.595	1.574	1.539	1.561	1.616
s.d. (height)	0.061	0.061	0.060	0.061	0.061	0.061	0.056	0.062	0.061	0.058	0.057	0.060
	MO	MZ	NB	NC	NG	PE	RW	SE	TZ	TO	TK	UG
Height (β)	-0.064** (0.017)	-0.108** (0.018)	0.014 (0.029)	-0.060** (0.012)	-0.083** (0.019)	-0.038** (0.009)	-0.132** (0.017)	-0.008 (0.016)	-0.070** (0.013)	-0.067** (0.023)	-0.024 (0.020)	-0.046** (0.016)
θ	-0.017	-0.026	0.005	-0.020	-0.021	-0.010	-0.036	-0.002	-0.021	-0.019	-0.007	-0.015
τ	-1.362	-1.714	0.326	-2.069	-1.431	-0.950	-2.400	-0.160	-1.556	-1.396	-0.533	-1.122
Observations	47932	53648	11211	65311	46700	162104	48743	47204	68652	24303	33090	41175
mean (dep. var.)	0.047	0.063	0.043	0.029	0.058	0.040	0.055	0.050	0.045	0.048	0.045	0.041
s.d. (dep. var.)	0.212	0.242	0.202	0.168	0.234	0.196	0.228	0.217	0.207	0.215	0.207	0.198
mean (height)	1.575	1.561	1.610	1.533	1.605	1.498	1.581	1.623	1.560	1.589	1.553	1.585
s.d. (height)	0.057	0.059	0.065	0.057	0.059	0.054	0.062	0.062	0.062	0.061	0.059	0.063
	ZB	ZW										
Height (β)	-0.029* (0.014)	-0.039* (0.017)										
θ	-0.009	-0.014										
τ	-0.690	-1.345										
Observations	61007	27141										
mean (dep. var.)	0.042	0.029										
s.d. (dep. var.)	0.201	0.167										
mean (height)	1.580	1.599										
s.d. (height)	0.061	0.061										

Robust standard errors in brackets. * significant at 5%, ** significant at 1%. See notes to Table 1a. Every equation includes a linear trend. Country name acronyms in the column heads are expanded in Appendix Table 1. We normalise the coefficient of height (τ) by dividing it by the country-specific mean of dependant variable. We standardize it (θ) by multiplying it by the country-specific ratio of the standard deviation of maternal height to the standard deviation of the dependant variable.

Table A.12: Country-specific Estimates of Persistence: Infant Mortality

	BE	BR	BF	CR	CD	CM	CH	CB	CO	CI	DR	EG
Height (β)	-0.162** (0.029)	-0.043 (0.027)	-0.117** (0.021)	-0.108** (0.040)	-0.117** (0.029)	-0.089** (0.025)	-0.084** (0.028)	-0.001 (0.010)	-0.138* (0.063)	-0.170** (0.032)	-0.032 (0.024)	-0.102** (0.013)
θ	-0.031	-0.010	-0.021	-0.023	-0.021	-0.019	-0.016	0.000	-0.026	-0.033	-0.008	-0.019
τ	-1.361	-0.573	-0.907	-1.009	-1.125	-0.947	-0.651	-0.029	-1.255	-1.650	-0.508	-1.041
Observations	34104	22997	71846	13646	38351	35524	38487	97312	6999	26574	30198	178259
mean (dep. var.)	0.119	0.075	0.129	0.107	0.104	0.094	0.129	0.035	0.11	0.103	0.063	0.098
s.d. (dep. var.)	0.324	0.263	0.335	0.309	0.305	0.292	0.335	0.185	0.313	0.304	0.243	0.297
mean (height)	1.584	1.551	1.615	1.582	1.525	1.602	1.626	1.543	1.55	1.591	1.561	1.575
s.d. (height)	0.061	0.063	0.059	0.066	0.054	0.062	0.062	0.06	0.059	0.059	0.061	0.056
	ET	GB	GH	GU	HA	HO	IN	KE	IE	MD	MW	ML
Height (β)	-0.173** (0.021)	-0.025 (0.034)	-0.085** (0.025)	-0.080** (0.028)	-0.045 (0.028)	-0.114** (0.019)	-0.134** (0.008)	0.000 (0.018)	-0.050 (0.042)	-0.110** (0.026)	-0.108** (0.022)	-0.108** (0.023)
θ	-0.032	-0.006	-0.018	-0.014	-0.009	-0.033	-0.027	0.000	-0.011	-0.022	-0.018	-0.018
τ	-1.362	-0.385	-0.988	-0.541	-0.413	-2.426	-1.576	0.000	-0.602	-1.170	-0.777	-0.663
Observations	69886	14106	35963	41059	34999	38312	434073	58882	11434	35697	75165	75563
mean (dep. var.)	0.127	0.065	0.086	0.148	0.109	0.047	0.085	0.076	0.083	0.094	0.139	0.163
s.d. (dep. var.)	0.333	0.246	0.281	0.355	0.312	0.211	0.279	0.264	0.276	0.291	0.346	0.369
mean (height)	1.573	1.585	1.589	1.587	1.579	1.514	1.516	1.595	1.574	1.539	1.561	1.616
s.d. (height)	0.061	0.061	0.06	0.061	0.061	0.061	0.056	0.062	0.061	0.058	0.057	0.06
	MO	MZ	NB	NC	NG	PE	RW	SE	TZ	TO	TK	UG
Height (β)	-0.109** (0.022)	-0.212** (0.026)	-0.029 (0.039)	-0.118** (0.017)	-0.182** (0.029)	-0.071** (0.013)	-0.160** (0.025)	-0.023 (0.023)	-0.099** (0.020)	-0.076* (0.032)	-0.050 (0.027)	-0.018 (0.024)
θ	-0.023	-0.035	-0.007	-0.028	-0.030	-0.014	-0.031	-0.005	-0.020	-0.016	-0.010	-0.004
τ	-1.346	-1.442	-0.387	-1.903	-1.238	-0.922	-1.391	-0.235	-0.934	-0.768	-0.556	-0.184
Observations	47109	52561	10606	64006	44532	156332	47192	46342	66383	23111	31971	39091
mean (dep. var.)	0.081	0.147	0.075	0.062	0.147	0.077	0.115	0.098	0.106	0.099	0.09	0.098
s.d. (dep. var.)	0.273	0.354	0.263	0.241	0.354	0.267	0.32	0.298	0.308	0.299	0.286	0.298
mean (height)	1.575	1.561	1.61	1.533	1.605	1.498	1.581	1.623	1.56	1.589	1.553	1.586
s.d. (height)	0.057	0.059	0.065	0.057	0.059	0.054	0.062	0.062	0.062	0.061	0.059	0.063
	ZB	ZW										
Height (β)	-0.119** (0.021)	-0.072** (0.024)										
θ	-0.024	-0.019										
τ	-1.123	-1.200										
Observations	58965	26083										
mean (dep. var.)	0.106	0.06										
s.d. (dep. var.)	0.308	0.237										
mean (height)	1.58	1.599										
s.d. (height)	0.061	0.061										

Robust standard errors in brackets. * significant at 5%, ** significant at 1%. See notes to Table 1a. Every equation includes a linear trend. Country name acronyms in the column heads are expanded in Appendix Table 1. We normalise the coefficient of height (τ) by dividing it by the country-specific mean of dependant variable. We standardize it (θ) by multiplying it by the country-specific ratio of the standard deviation of maternal height to the standard deviation of the dependant variable.

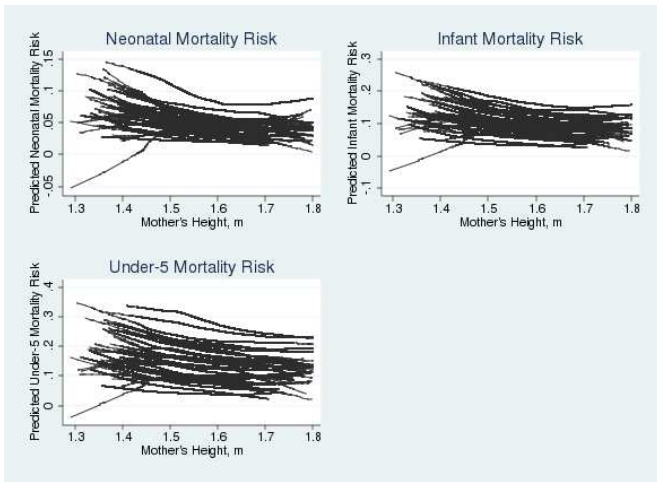
Table A.13: Country-specific Estimates of Persistence: Under-5 Mortality

	BE	BR	BF	CR	CD	CM	CH	CB	CO	CI	DR	EG
Height (β)	-0.183** (0.041)	-0.045 (0.033)	-0.137** (0.029)	-0.135* (0.056)	-0.137** (0.036)	-0.124** (0.033)	-0.098** (0.037)	-0.008 (0.012)	-0.157* (0.079)	-0.215** (0.043)	-0.016 (0.031)	-0.127** (0.015)
θ	-0.027	-0.010	-0.019	-0.023	-0.021	-0.022	-0.015	-0.002	-0.026	-0.035	-0.003	-0.022
τ	-0.880	-0.506	-0.620	-0.799	-0.979	-0.838	-0.462	-0.182	-1.047	-1.344	-0.186	-0.977
Observations	26657	19375	58808	10540	31648	30582	32322	87703	5533	20801	24238	153139
mean (dep. var.)	0.208	0.089	0.221	0.169	0.140	0.148	0.212	0.044	0.150	0.160	0.086	0.130
s.d. (dep. var.)	0.406	0.285	0.415	0.374	0.347	0.355	0.409	0.204	0.357	0.366	0.281	0.336
mean (height)	1.584	1.551	1.615	1.581	1.525	1.602	1.626	1.543	1.550	1.591	1.560	1.575
s.d. (height)	0.061	0.062	0.059	0.065	0.054	0.062	0.062	0.060	0.060	0.059	0.061	0.057
	ET	GB	GH	GU	HA	HO	IN	KE	LE	MD	MW	ML
Height (β)	-0.218** (0.027)	0.022 (0.045)	-0.109** (0.034)	-0.116** (0.035)	-0.085* (0.039)	-0.153** (0.021)	-0.179** (0.009)	0.002 (0.024)	-0.043 (0.048)	-0.159** (0.035)	-0.127** (0.029)	-0.159** (0.031)
θ	-0.033	0.005	-0.019	-0.017	-0.013	-0.039	-0.032	0.000	-0.009	-0.026	-0.017	-0.021
τ	-1.085	0.224	-0.757	-0.492	-0.489	-2.508	-1.613	0.019	-0.439	-1.067	-0.567	-0.566
Observations	61012	11436	29115	36274	27451	38260	396147	47888	10688	29639	61909	59899
mean (dep. var.)	0.201	0.098	0.144	0.236	0.174	0.061	0.111	0.108	0.098	0.149	0.224	0.281
s.d. (dep. var.)	0.401	0.298	0.351	0.424	0.379	0.240	0.314	0.311	0.297	0.356	0.417	0.449
mean (height)	1.573	1.585	1.589	1.587	1.578	1.514	1.516	1.594	1.574	1.540	1.562	1.616
s.d. (height)	0.061	0.061	0.060	0.061	0.060	0.061	0.056	0.061	0.061	0.058	0.057	0.060
	MO	MZ	NB	NC	NG	PE	RW	SE	TZ	TO	TK	UG
Height (β)	-0.131** (0.027)	-0.223** (0.033)	0.006 (0.054)	-0.188** (0.021)	-0.283** (0.042)	-0.121** (0.017)	-0.178** (0.033)	-0.081** (0.030)	-0.183** (0.026)	-0.060 (0.044)	-0.065* (0.032)	-0.061 (0.035)
θ	-0.024	-0.032	0.001	-0.040	-0.036	-0.021	-0.028	-0.013	-0.031	-0.010	-0.012	-0.010
τ	-1.248	-1.037	0.053	-2.321	-0.913	-1.071	-0.908	-0.466	-1.130	-0.349	-0.586	-0.361
Observations	40527	43777	8030	53395	34541	126448	40291	41112	54556	18108	26938	29548
mean (dep. var.)	0.105	0.215	0.113	0.081	0.310	0.113	0.196	0.174	0.162	0.172	0.111	0.169
s.d. (dep. var.)	0.307	0.410	0.317	0.273	0.463	0.317	0.397	0.379	0.369	0.377	0.314	0.375
mean (height)	1.576	1.561	1.610	1.533	1.605	1.497	1.582	1.623	1.560	1.589	1.552	1.586
s.d. (height)	0.057	0.059	0.065	0.058	0.059	0.054	0.062	0.062	0.062	0.062	0.060	0.063
	ZB	ZW										
Height (β)	-0.174** (0.030)	-0.091** (0.032)										
θ	-0.028	-0.020										
τ	-1.000	-1.083										
Observations	45366	20825										
mean (dep. var.)	0.174	0.084										
s.d. (dep. var.)	0.379	0.277										
mean (height)	1.580	1.599										
s.d. (height)	0.061	0.061										

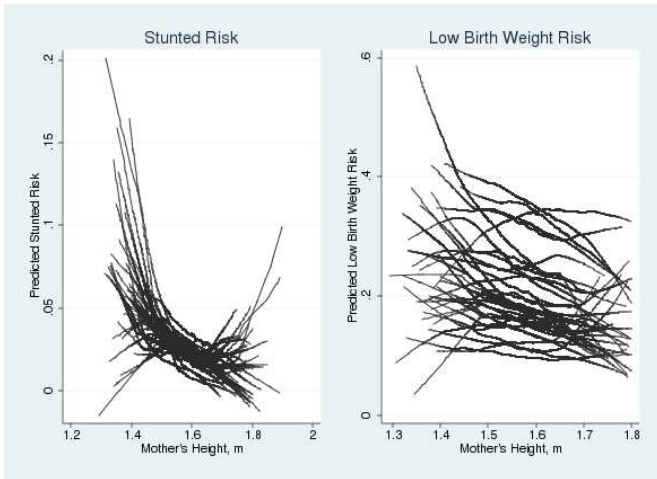
Robust standard errors in brackets. * significant at 5%, ** significant at 1%. See notes to Table 1a. Every equation includes a linear trend. Country name acronyms in the column heads are expanded in Appendix Table 1. We normalise the coefficient of height (τ) by dividing it by the country-specific mean of dependant variable. We standardize it (θ) by multiplying it by the country-specific ratio of the standard deviation of maternal height to the standard deviation of the dependant variable.

Figure A.1: Alternative Indicators of Child Health against Mother's Health: Country-specific Lowess Predictions

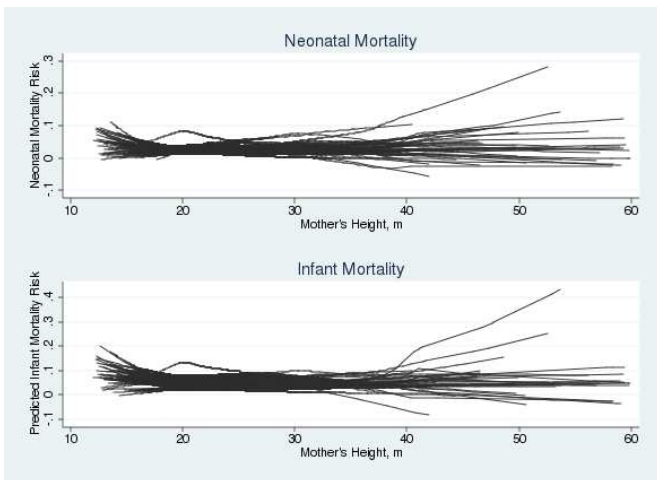
a) Height: Mortality Indicators



b) Height: Low Birth Weight and Stunting



c) BMI: Mortality Indicators



d) BMI: Low Birth Weight and Stunted

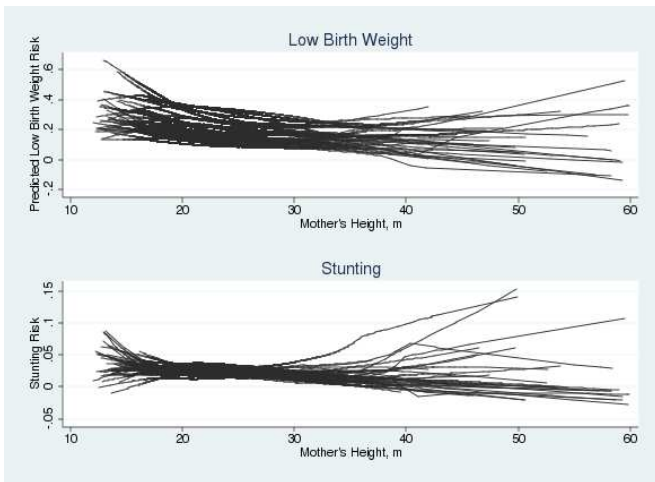


Figure A3: Trends in Intergenerational Persistence across Cohorts: β_c

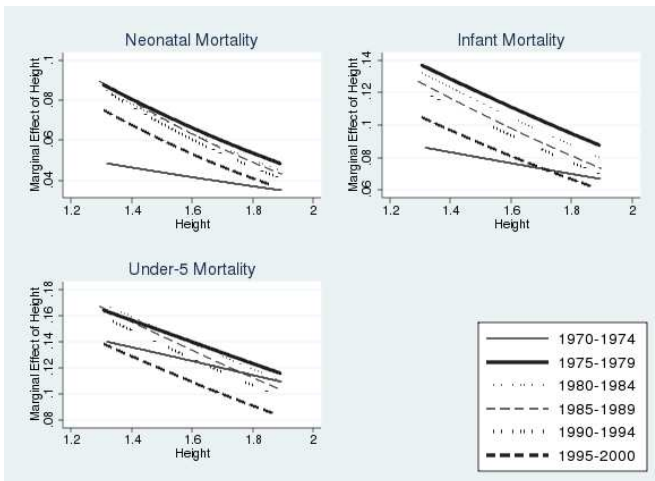
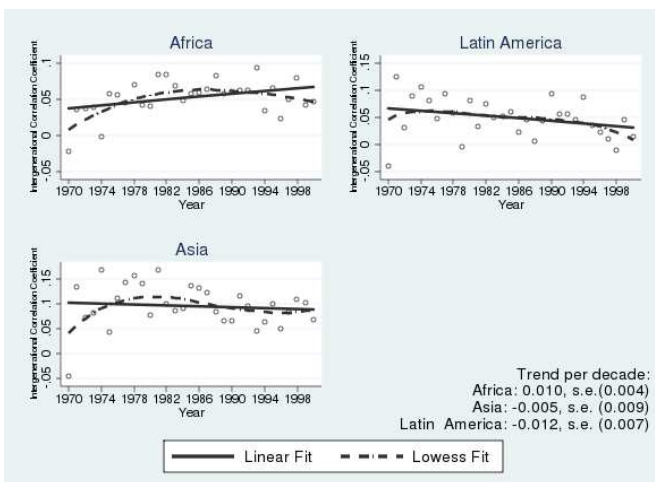


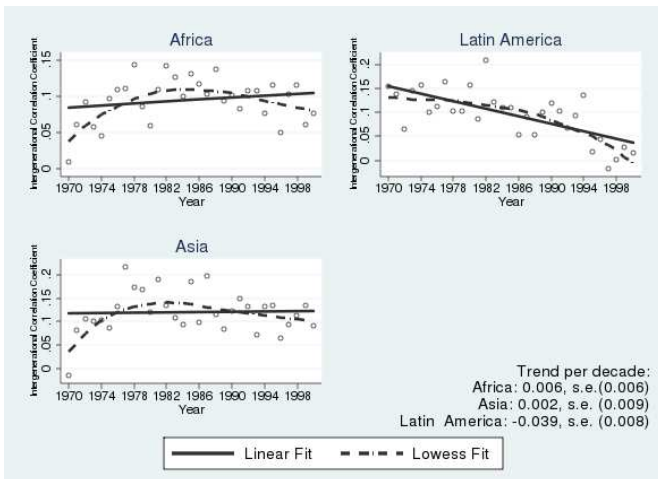
Figure plots the probit marginal effect of height over the height distribution, for each cohort

Figure A.4: Trends in Intergenerational Persistence across Years, β_t , by continent

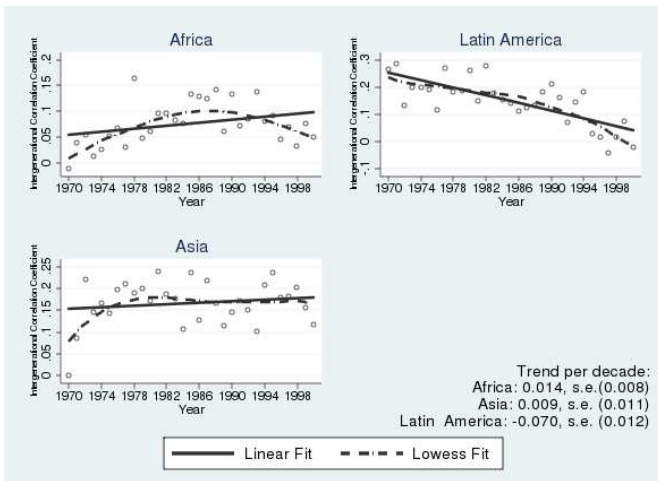
a) Neonatal Mortality



b) Infant Mortality



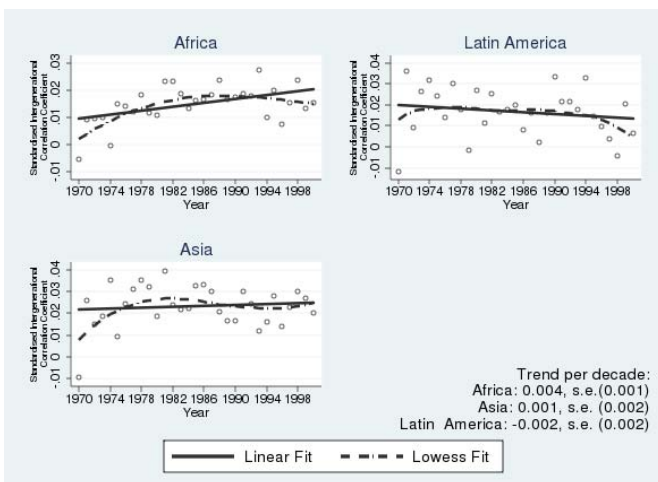
c) Under-5 Mortality



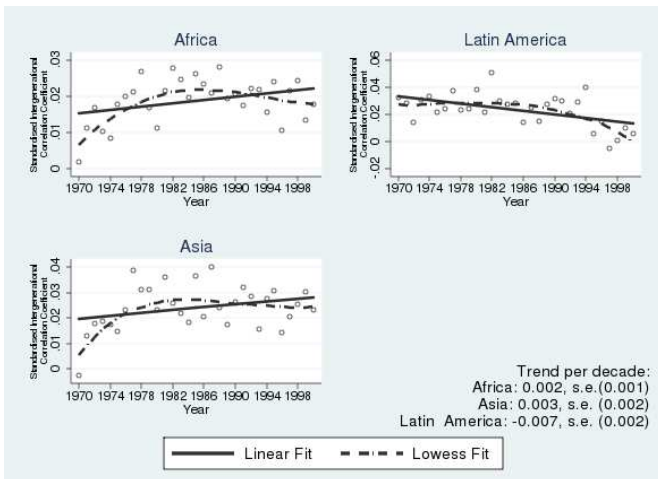
Plot of year-specific β from a regression of child health against mother's height, with child and household controls.

Figure A.5: Trends in Standardised Intergenerational Persistence across Years, θ_t , by continent

a) Neonatal Mortality



b) Infant Mortality



c) Under-5 Mortality

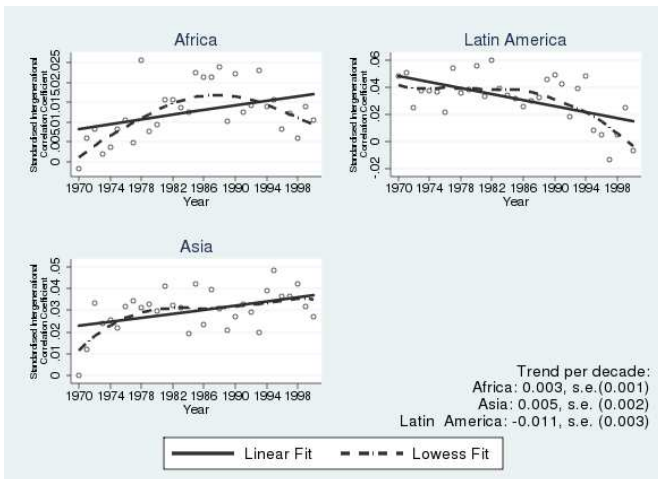


Figure A.6: Density Plots of β , θ , and η : Countries with significant β

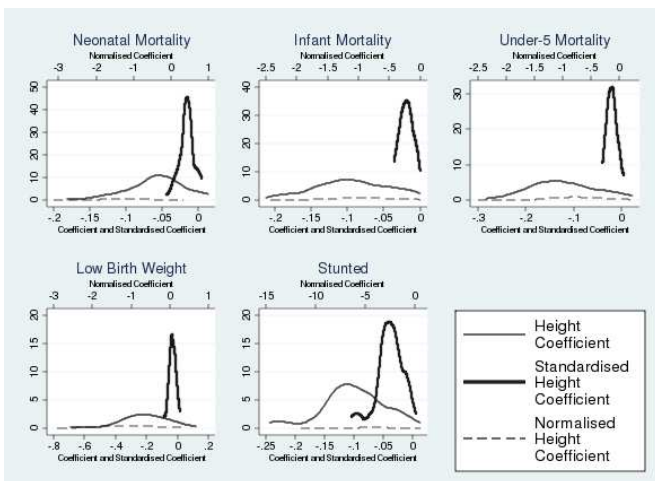
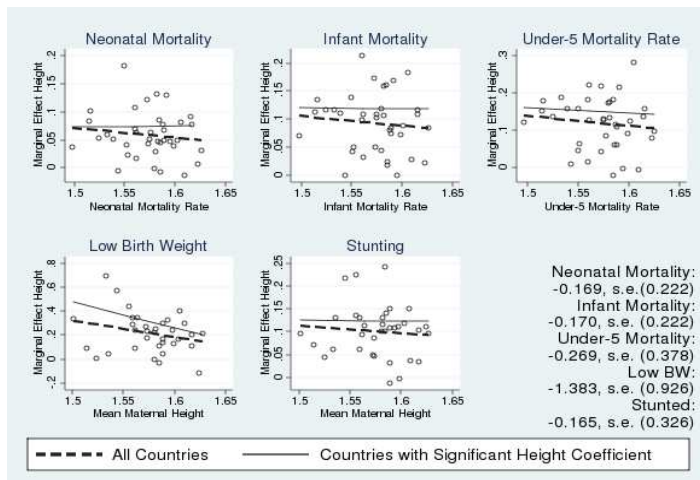
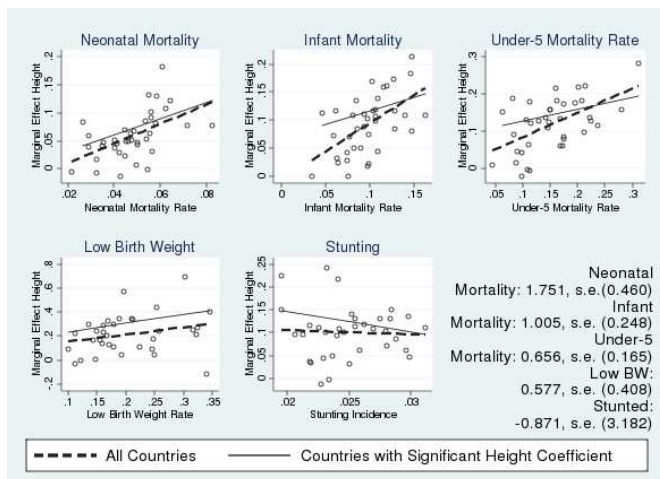


Figure A.7: Country Variation in Intergenerational Persistence

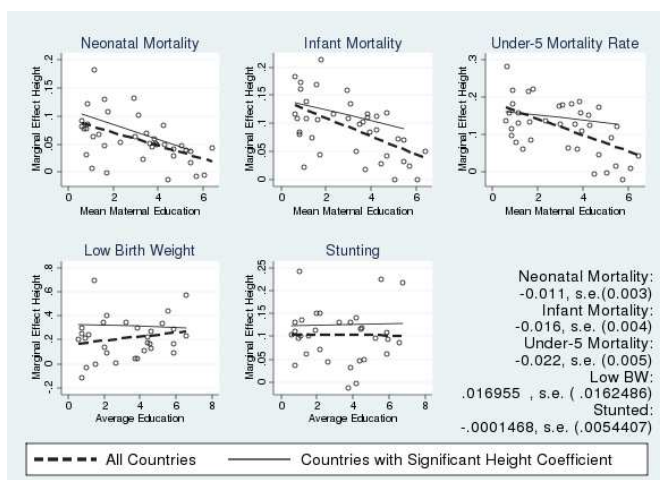
a) Against Mother Height



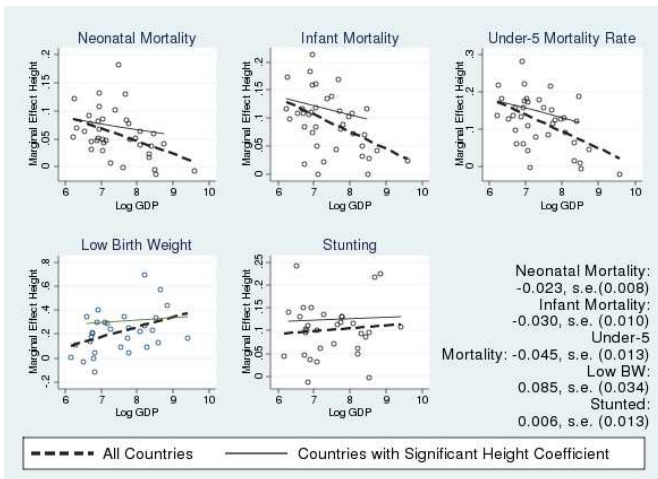
b) Against Level of Child Health



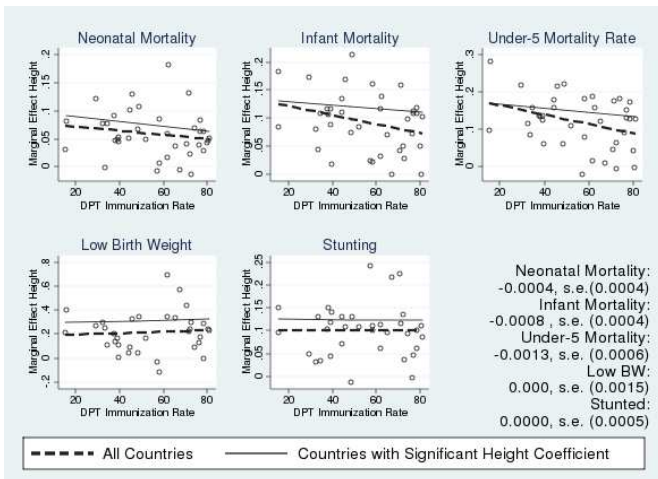
c) Average Education of Mothers



d) Against GDP



e) Against DPT Immunization Rates



f) Against Measles Immunization Rates

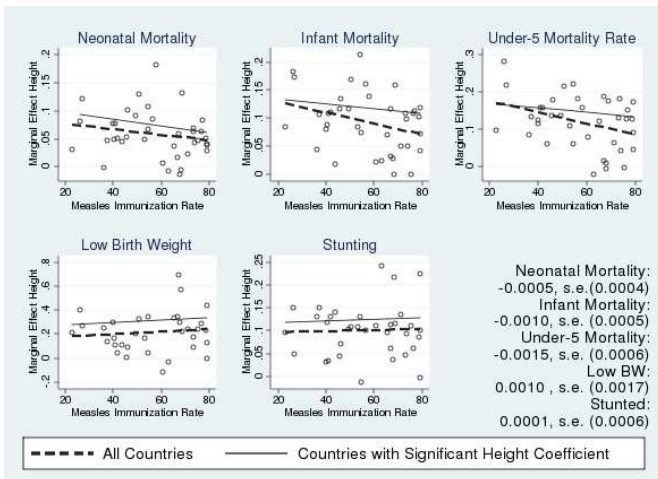
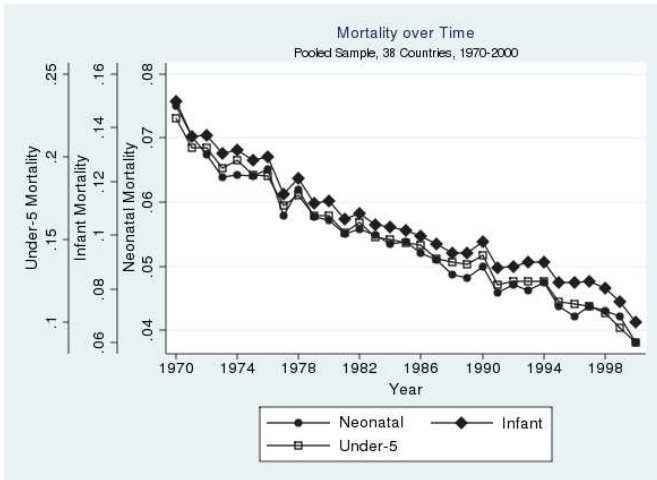
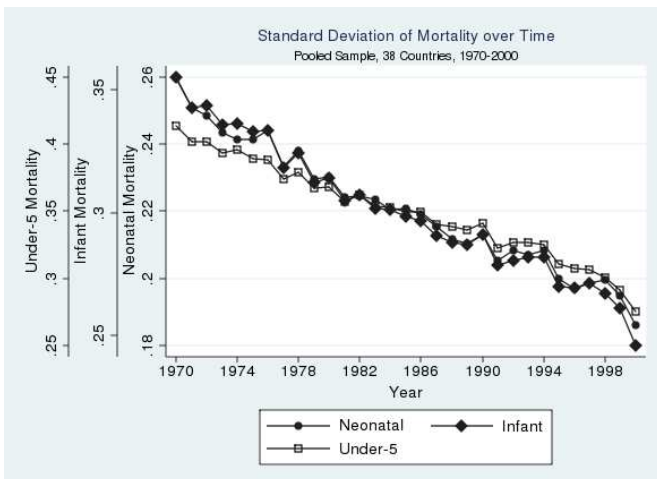


Figure A.8: Trends in Health and GDP: Pooled Sample

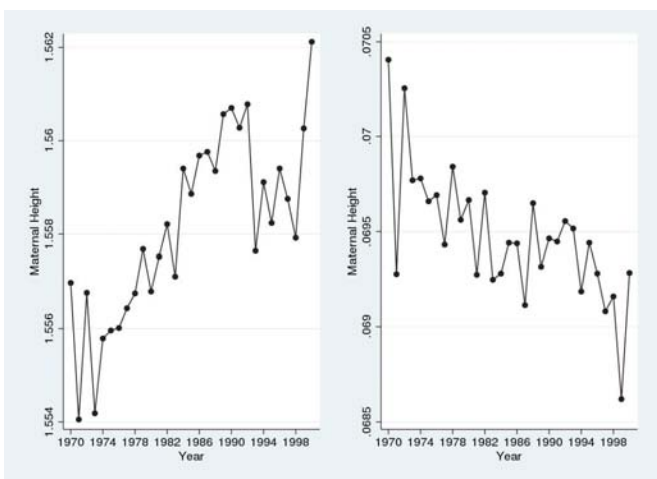
a) Mean Mortality: 38 Countries



b) Standard Deviation of Mortality Rates: 38 Countries



c) Mother's Height: Mean and s.d.



d) Log GDP: Mean and s.d., 38 Countries

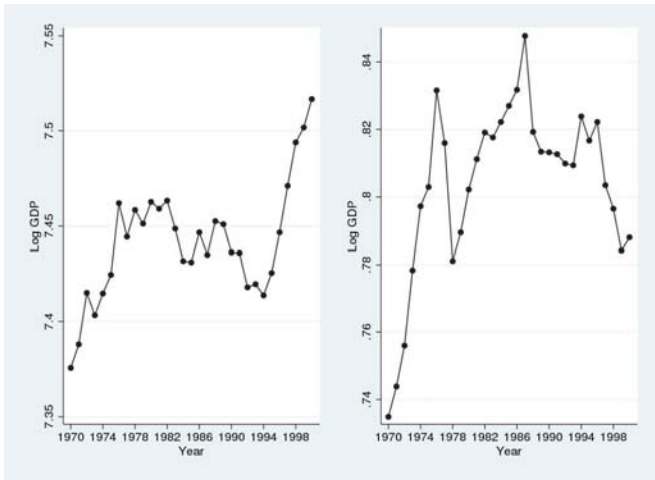
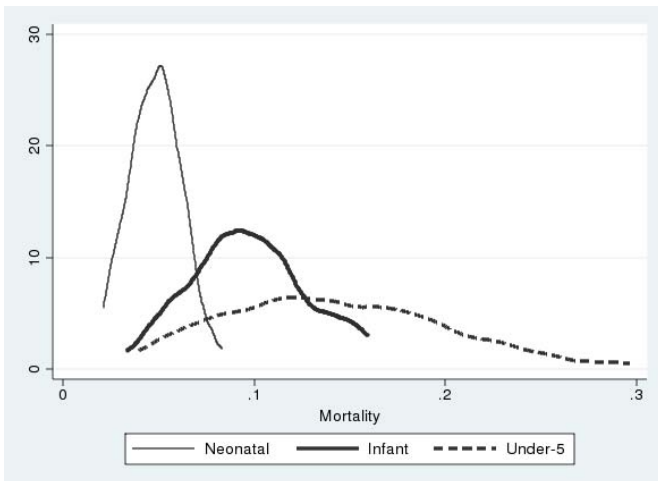
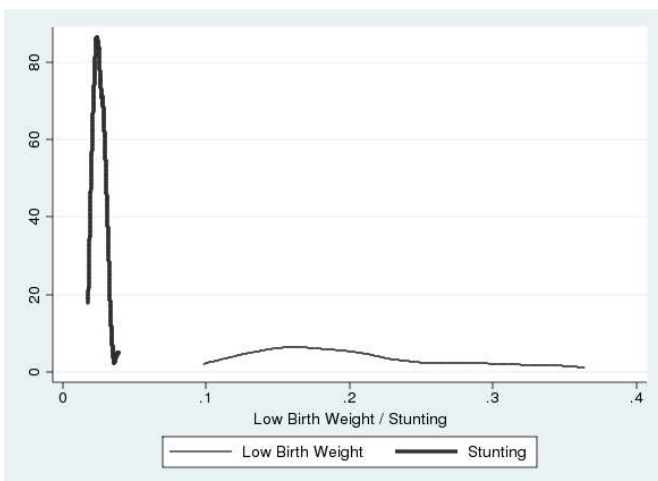


Figure A.9: Density plots of Country-Average Health

a) Mortality



b) Stunting and Birth Weight



c) Mother Health

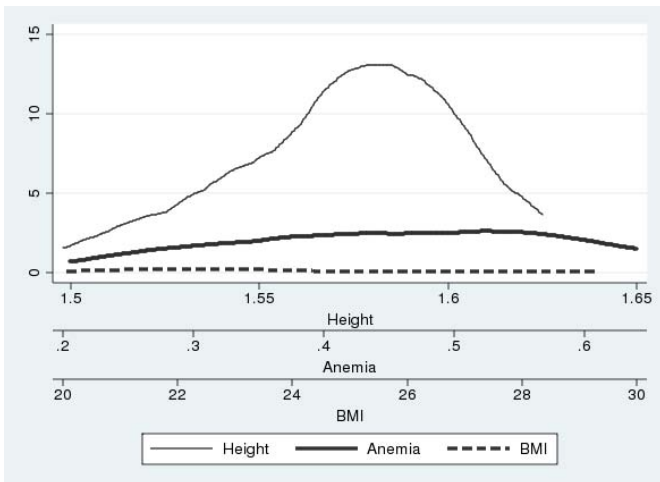
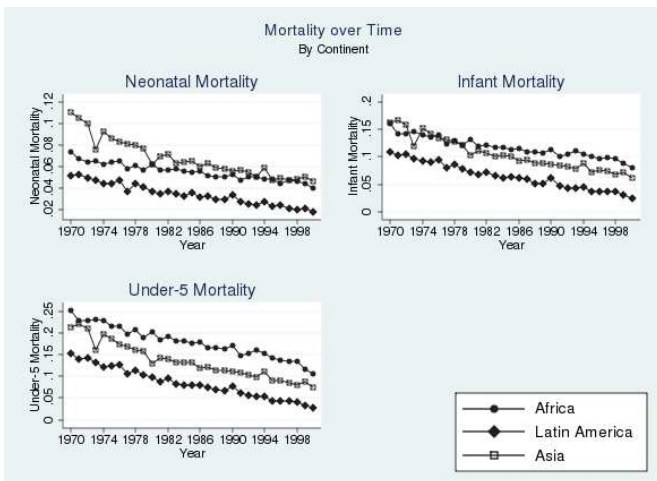


Figure A10 : Trends in Health and GDP: by Continent

a) Mortality Rates



b) Mother Height and Log GDP

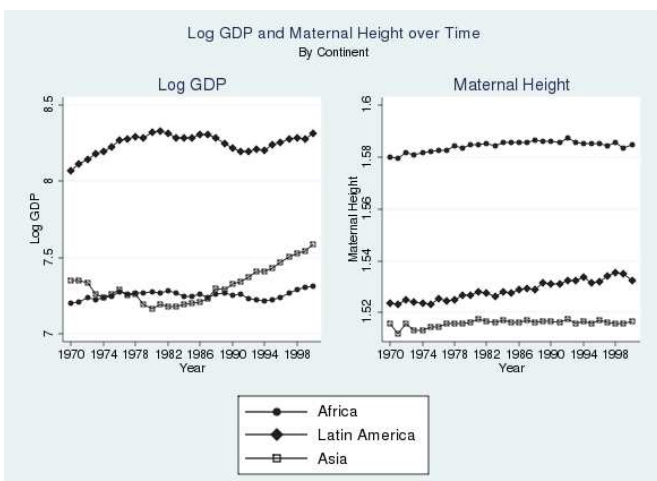
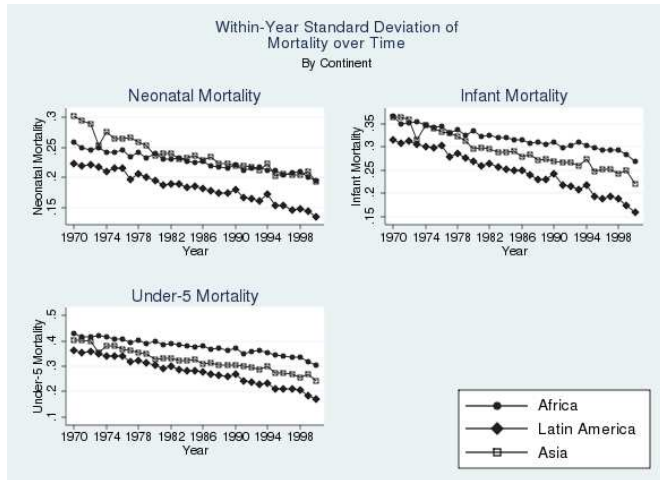


Figure A11: Trends in inequality in Health and GDP: by Continent

a) Mortality Rates



b) Within-Continent Standard Deviation of Height and GDP

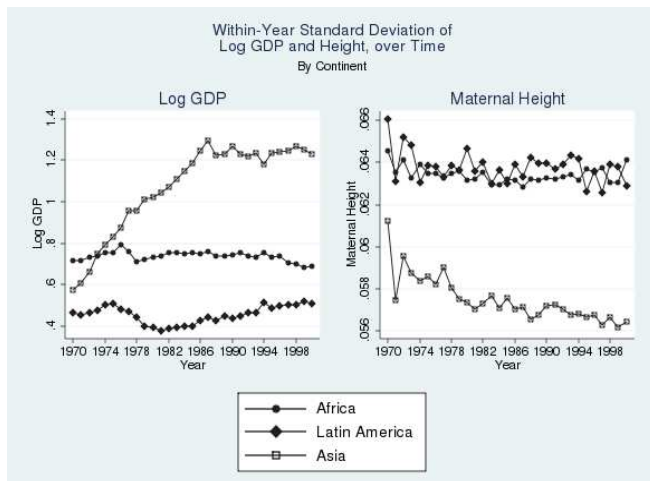


Figure A12: Changes over Time for the Case of India: Infant Mortality and Child Height

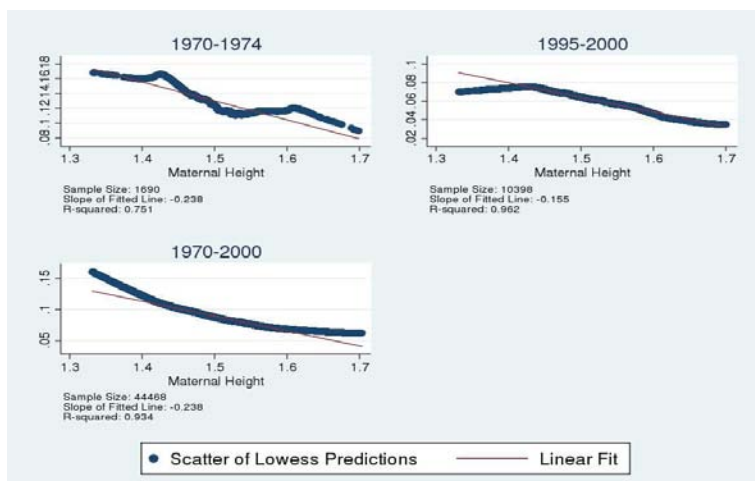


Figure A13: Map Displaying the 38 Countries in Sample

Map Displaying the 38 Countries in Sample

