Original Contribution

Associations of Birth Order With Early Growth and Adolescent Height, Body Composition, and Blood Pressure: Prospective Birth Cohort From Brazil

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Birth weight has been inversely associated with later blood pressure. Firstborns tend to have lower birth weight than their later-born peers, but the long-term consequences remain unclear. The study objective was to investigate differences between firstborn and later-born individuals in early growth patterns, body composition, and blood pressure in Brazilian adolescents. The authors studied 453 adolescents aged 13.3 years from the prospective 1993 Pelotas Birth Cohort. Anthropometry, blood pressure, physical activity by accelerometry, and body composition by deuterium were measured. Firstborns (n = 143) had significantly lower birth weight than later borns (n = 310). At 4 years, firstborns had significantly greater weight and height, indicating a substantial overshoot in catch-up growth. In adolescence, firstborns had significantly greater height and blood pressure and a lower activity level. The difference in systolic blood pressure could be attributed to variability in early growth and that in diastolic blood pressure to reduced physical activity. The magnitude of increased blood pressure is clinically significant; hence, birth order is an important developmental predictor of cardiovascular risk in this population. Firstborns may be more sensitive to environmental factors that promote catch-up growth, and this information could potentially be used in nutritional management to prevent catch-up “overshoot.”

birth order; blood pressure; body composition; growth; motor activity

Abbreviations: ALSPAC, Avon Longitudinal Study of Pregnancy and Childhood; CI, confidence interval; SD, standard deviation; WHO, World Health Organization.

Small size at birth has been associated in many studies with increased blood pressure in adolescence or adulthood (1). Initial research emphasized maternal and fetal malnutrition as a potential important mechanism (2), a hypothesis supported by experimental animal studies (3). However, associations with blood pressure hold across the whole range of birth weight, suggesting that a wide range of factors may be relevant.

Firstborn infants tend to have lower birth weight than later-born infants (4–7). In data from 3 Norwegian cities between 1860 and 1984, there was a substantial increase (approximately 200 g) in birth weight between the first and second pregnancies, followed by a much smaller increase (approximately 30 g) with each succeeding pregnancy (8). Physiologic studies have suggested possible anatomic explanations for the reduced birth weight of firstborns. During a mother’s first pregnancy, structural changes take place in the uterine spiral arteries, increasing blood flow with beneficial effects for fetal growth (9). These changes do not completely disappear following the pregnancy, such that subsequent offspring are from the start of pregnancy exposed to reduced vascular resistance and hence greater uterine blood flow compared with firstborns, promoting fetal growth (9).

The longer-term implications of these birth order associations remain unclear, in part because of inconsistent findings in previous studies and in part because of the possibility that both social and biologic mechanisms may be relevant. In some populations, firstborns remain shorter than later borns in adulthood (10), whereas other studies indicate that firstborns become significantly taller than later borns (11–13). Tanner (11) suggested that firstborn children may benefit from being...
the only child during their early life, resulting in improved nutrition and greater final size. Given the reduced birth weight of firstborns, this implies a tendency for infant catch-up growth to resolve early growth deficits.

Both low birth weight and rapid weight gain, particularly after the age of 2 years, are independent factors for cardiovascular risk (1, 14, 15), and recently we found that firstborns do indeed have elevated cardiovascular risk in Brazilian young adult men (13). We therefore investigated a second Brazilian birth cohort in order to determine in greater detail associations between birth order and subsequent phenotype. We investigated the association between birth order and 1) birth size, 2) postnatal growth rate, 3) adolescent body size, body composition, and physical activity level, and 4) adolescent blood pressure in the prospective 1993 Pelotas Birth Cohort Study. We tested the hypothesis that firstborns and later borns differ in each of these outcomes, and that these associations are independent of family size.

MATERIALS AND METHODS

Subjects

The 1993 Pelotas Birth Cohort recruited 5,249 individuals (16). Data on early growth at 6, 12, and 48 months were collected in a subsample of 1,272. For this study, we randomly selected 13% of those born in each calendar month of the year. This identified 655 individuals at 1 month of age, of whom 453 with full data at previous time points were successfully located and studied in adolescence. These individuals underwent measurements of body composition by deuterium dilution, physical activity by accelerometer, and blood pressure at 13.3 years. Birth order, based on the number of pregnancies, was obtained by maternal questionnaire at the time of recruitment. Ethics approval was obtained from the Federal University of Pelotas Medical School Ethics Committee.

Anthropometry

Birth weight and length were measured at the hospital by the research team. Weight and length or height at 6, 12, and 48 months were measured at the cohort participant’s household. At the 13.3-year visit, weight and height were again measured.

Body composition and pubertal stage

Body composition in adolescence was measured by using deuterium (17). Briefly, each adolescent was given a drink containing approximately 0.05 g of 99.9% deuterium oxide (\(\text{D}_2\text{O}\)) per kg of body weight. Saliva samples were obtained predose and 4 hours postdose by using absorbent salivettes at least 30 minutes after the last ingestion of food or drink and then stored frozen at \(-30^\circ\text{C}\), as described in detail elsewhere (18). The samples were shipped to the United Kingdom for analysis in duplicate with mass spectrometry, by use of the equilibration method (Delta plus XP; Thermofisher Scientific, Bremen, Germany) (19). For calculation of total body water, it was assumed that deuterium oxide dilution space overestimated total body water by a factor of 1.044 (17). Correction was made for dilution of the dose by water intake during the 4-hour equilibration period (17). Values for total body fluid were converted to lean mass (used here synonymously with fat-free mass), by using new reference data for the hydration of lean tissue (20). Fat mass was calculated as the difference of lean mass and weight. Both fat mass and lean mass were then adjusted for height to give the fat mass index and lean mass index, both expressed in the same kg/m\(^2\) units as body mass index (21, 22). Pubertal stage was assessed by Tanner staging, by use of line drawings. The 2 scores, each ranging from 1 to 5, were summed and the summed score analyzed.

Physical activity

Adolescent physical activity was assessed by using GT1M accelerometers (ActiGraph, Pensacola, Florida). The accelerometer was presented during the initial home interview and placed on the left side of the waist. In addition, an instruction sheet for the accelerometers including a diary was left at the participant’s home at the time of the interview. Participants were instructed to record if they did not wear the monitor for any period >1 hour during the day. Subjects wore the monitors from Wednesday to Monday and were encouraged to wear them 24 hours per day, except when showering, bathing, or swimming.

Blood pressure

Blood pressure was measured while the participant was seated, by using an HEM-629 digital portable wrist monitor (Omron Healthcare, Inc., Bannockburn, Illinois) in combination with a standard cuff size at the beginning and end of the interview (60 minutes apart). This monitor has been validated against a mercury sphygmomanometer in Brazilian adolescents (23). The mean value was used in analyses.

Statistics

Weight and height standard deviation (SD) scores for early growth data were calculated by using World Health Organization (WHO) reference curves. Preliminary analyses indicated differences between firstborns and later borns but not between secondborns and third-borns, as was the case in a similar prior analysis (13). The analyses therefore compared firstborns with all later borns. Preliminary analyses also considered whether the results changed if mothers aged <18 years or those with a high birth order of ≥4 were excluded. These factors did not alter the findings significantly, and therefore no such exclusions were applied to the full analyses.

Crude differences between firstborn and later-born individuals were assessed by chi-square tests, independent-sample t tests, or Mann-Whitney tests. To take into account other variables, we used regression analysis with a succession of models. Following unadjusted analyses, model 1 adjusted for maternal factors (age, height, body mass index, educational level, family income), as well as offspring sex. Model 2 further adjusted for birth weight z score, weight and height SD scores at 4 years, and the physical activity level at 13.3 years. Conditional growth between birth and 4 years was calculated as recommended by Keijzer-Veen et al. (24), by calculating

residuals for the regression of size at 48 months on size at birth for each of weight and length. As preliminary results showed firstborns to have higher maternal height than later borns, we considered the possible interaction between maternal height and birth order for predicting adolescent height, both with and without adjustment for birth size.

In order to separate potential birth order associations from those potentially arising from family size, we reran the analyses separately for firstborns who did or did not have a sibling and for siblings separately for firstborns who did or did not have a sibling. To control for the possibility of mortality confounding, we reran the analyses for those potentially arising from family size, we reran the analyses separately for firstborns who did or did not have a sibling and for siblings separately for firstborns who did or did not have a sibling. To control for the possibility of mortality confounding, we reran the analyses separately for firstborns who did or did not have a sibling and for siblings separately for firstborns who did or did not have a sibling.

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RESULTS

Table 1 shows that the subsample included in this analysis (n = 453) is comparable to the remainder of the cohort studied at 13.3 years (n = 5,249, including 1,843 firstborns) in terms of birth order, gender, family income, and maternal age. The subsample was significantly larger in body size at birth. However, this difference was relatively small and unlikely to indicate that our findings cannot be generalized to the whole cohort. Approximately one third (n = 143, 31.6%) of the subjects included in our analyses were firstborns, while 310 were later borns, and 52.3% of the sample were males. Of the firstborns, 47 had siblings by 4 years. Mean height at 13.3 years was 158 cm, mean body mass index was 20.3 kg/m², mean systolic blood pressure was 111.0 mm Hg, and mean diastolic blood pressure was 68.6 mm Hg.

The average weight SD score was below the WHO reference value at birth (mean: −0.18, 95% confidence interval (CI): −0.29, −0.08), but subjects were able to catch up and reach an SD score of 0.31 (95% CI: 0.20, 0.42) at 4 years of age. In terms of the height SD score, individuals were born below the WHO reference (mean: −0.36, 95% CI: −0.47, −0.25) but moved toward the reference mean, reaching −0.12 (95% CI: −0.23, −0.01) at 4 years of age.

In Table 2, we present the distribution of the maternal variables and the child's sex according to birth order. Subjects born to older mothers were less likely to be firstborns (P < 0.001). Also, the mean maternal age was 23.2 years among firstborns and 27.7 years among the others (P < 0.001). The likelihood of being a firstborn was not statistically associated with offspring sex, family income, or maternal height. However, the mothers of firstborns were 1.4 (95% CI: 0.1, 2.8) cm taller than mothers of later borns (P = 0.04), had completed on average 1.4 (95% CI: 0.7, 2.1) more years of education (P < 0.001), and had 0.8 (95% CI: −1.6, −0.1) kg/m² lower body mass index (P = 0.03). Tertile analysis showed that the association between maternal height and offspring height appeared to vary by birth order. Compared with those in the lowest tertile for maternal height, those in the second and third tertiles had 5.4 (95% CI: 2.7, 8.5) and 8.1 (95% CI: 5.2, 11.1) cm, respectively, if the child was first-born and 3.5 (95% CI: 1.4, 5.7) and 6.3 (95% CI: 4.3, 8.7) cm, respectively, if the child was a later born. These birth-order differences remained similar, although the absolute values were slightly reduced, if birth weight was held constant. However, the interaction between maternal height and birth order in relation to adolescent height was not statistically significant (P = 0.53).

Firstborns were born significantly smaller than later borns (Δ in birth weight SD scores = −0.31, 95% CI: −0.54, −0.08) (P = 0.008) and presented significantly greater catch-up than the WHO reference (0.31, 95% CI: 0.20, 0.42) at 4 years of age. In terms of height (Figure 2), firstborns were born shorter than later borns, although not significantly
so, but showed substantial catch-up growth within the first year of life, such that they were already significantly taller than later borns by 1 year of age, as well as above the reference mean. At 4 years of age, firstborns were 0.41 SD score (95% CI: 0.17, 0.64) taller than later borns ($P < 0.001$).

Table 3 shows that, in the unadjusted analyses, firstborns remained taller at 13.3 years than later borns, as well as presenting significantly higher blood pressure values. No significant crude differences were observed in terms of body mass index, sum of skinfolds, physical activity, or body composition indicators. Firstborns were also significantly more advanced in terms of pubertal stage: firstborns: 7.8 (95% CI: 7.5, 8.1); later borns: 7.4 (95% CI: 7.2, 7.6) ($P = 0.05$).

Table 2. Description of the Covariates (Sex, Maternal Age, Socioeconomic Status) According to Birth Order (Firstborns vs. Others) Among Brazilian Adolescents in the 1993 Pelotas Birth Cohort

<table>
<thead>
<tr>
<th>Variable</th>
<th>Firstborns</th>
<th>Others</th>
</tr>
</thead>
<tbody>
<tr>
<td>%</td>
<td>Mean (SD)</td>
<td>%</td>
</tr>
<tr>
<td><strong>Sex</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>58.7</td>
<td>49.4</td>
</tr>
<tr>
<td>Female</td>
<td>41.3</td>
<td>50.7</td>
</tr>
<tr>
<td>Maternal age, years</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;20</td>
<td>29.4</td>
<td>5.8</td>
</tr>
<tr>
<td>20–34</td>
<td>65.7</td>
<td>81.0</td>
</tr>
<tr>
<td>≥35</td>
<td>4.9</td>
<td>13.2</td>
</tr>
<tr>
<td>Maternal age, years</td>
<td>23.2 (5.9)</td>
<td>27.7 (5.8)</td>
</tr>
<tr>
<td>Quartile of family income</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 (poorest)</td>
<td>22.1</td>
<td>29.4</td>
</tr>
<tr>
<td>2</td>
<td>25.0</td>
<td>24.1</td>
</tr>
<tr>
<td>3</td>
<td>25.7</td>
<td>26.8</td>
</tr>
<tr>
<td>4 (wealthiest)</td>
<td>27.1</td>
<td>19.8</td>
</tr>
<tr>
<td>Family income, Brazilian real⁶</td>
<td>4.0 (4.0)</td>
<td>3.7 (4.1)</td>
</tr>
<tr>
<td>Quartile of maternal height</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 (shortest)</td>
<td>24.1</td>
<td>30.3</td>
</tr>
<tr>
<td>2</td>
<td>27.7</td>
<td>28.3</td>
</tr>
<tr>
<td>3</td>
<td>24.1</td>
<td>24.8</td>
</tr>
<tr>
<td>4 (tallest)</td>
<td>24.1</td>
<td>16.6</td>
</tr>
<tr>
<td>Maternal height, cm</td>
<td>161.0 (7.3)</td>
<td>159.6 (6.5)</td>
</tr>
<tr>
<td>Maternal schooling, years</td>
<td></td>
<td></td>
</tr>
<tr>
<td>0–4</td>
<td>14.7</td>
<td>33.3</td>
</tr>
<tr>
<td>5–8</td>
<td>51.8</td>
<td>46.6</td>
</tr>
<tr>
<td>≥9</td>
<td>33.6</td>
<td>20.1</td>
</tr>
<tr>
<td>Maternal schooling, years</td>
<td>7.6 (3.3)</td>
<td>6.2 (3.6)</td>
</tr>
<tr>
<td>Maternal BMI, kg/m²</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;18.5</td>
<td>10.7</td>
<td>8.8</td>
</tr>
<tr>
<td>18.5–24.9</td>
<td>71.4</td>
<td>65.3</td>
</tr>
<tr>
<td>25.0–29.9</td>
<td>14.3</td>
<td>19.2</td>
</tr>
<tr>
<td>≥30.0</td>
<td>3.6</td>
<td>6.7</td>
</tr>
<tr>
<td>Maternal BMI, kg/m²</td>
<td>22.2 (3.8)</td>
<td>23.1 (3.9)</td>
</tr>
</tbody>
</table>

Abbreviations: BMI, body mass index; SD, standard deviation.

⁴ Chi-square test.

⁵ t test.

⁶ 1.0 Brazilian real = 0.63000 US dollar.

⁷ Mann-Whitney test.
significant (adolescents in the 1993 Pelotas Birth Cohort. NS, nonsignificant.

birth and at 6, 12, and 48 months in Brazilian firstborn and later-born
Organization Child Growth Standards and 95% confidence interval at

Figure 2. Height standard deviation (SD) score of the World Health
Organization Child Growth Standards and 95% confidence interval at

for conditional weight or length at 4 years were made, the
increment in height for firstborns disappeared ($\Delta = 0$ cm, 95% CI: $-1.2, 1.2$ ($P = 0.99$), whereas the difference in
blood pressure remained significant (diastolic blood pressure:
$\Delta = 1.9$ mm Hg, 95% CI: 0.4, 3.4 ($P = 0.01$) or borderline
significant (systolic blood pressure: $\Delta = 1.9$ mm Hg, 95%
CI: $-0.3, 4.1$ ($P = 0.08$). If birth weight SD scores were
added to the model with size at 4 years, the birth order associ-
ation with systolic blood pressure lost further significance
although not magnitude ($\Delta = 1.9$ mm Hg, 95% CI: $-0.6, 4.3$
($P = 0.14$), whereas that for diastolic blood pressure remained
significant ($\Delta = 1.9$ mm Hg, 95% CI: 0.2, 3.5 ($P = 0.03$).
Only in the final model 2 (Table 4), when physical activity
was also included, did the birth order difference in diastolic
blood pressure lose significance. There were no significant
differences between birth order groups in lean mass or fat
mass in crude or adjusted models. Our analyses therefore in-
dicate that birth order affects blood pressure through the
impact of early growth and physical activity, although dif-
ferently so for the 2 blood pressure components.

When the analyses for height and blood pressure were
repeated separately for firstborns with or without siblings at
4 years, the difference between firstborns and later borns
was slightly reduced in those with siblings in terms of height
(with sibling: $\Delta = 2.0$ cm, 95% CI: $-0.5, 4.6$ ($P = 0.12$); no
sibling: $\Delta = 2.6$ cm, 95% CI: 0.7, 4.4 ($P = 0.007$)) and
diastolic blood pressure (with sibling: $\Delta = 2.5$ mm Hg,
95% CI: $-1.0, 6.0$ ($P = 0.16$); no sibling: $\Delta = 3.6$ mm Hg,
95% CI: 1.0, 6.1 ($P = 0.006$)) but much reduced for systolic
blood pressure (with sibling: $\Delta = 0.7$ mm Hg, 95% CI: $-1.5$,
2.8 ($P = 0.54$); no sibling: $\Delta = 2.3$ mm Hg, 95% CI: 0.8, 3.0
($P = 0.003$)). At 4 years, firstborns with no sibling were also
taller (height SD score = 0.2, 95% CI: 0.0, 0.4) than those
with siblings (height SD score = 0.1, 95% CI: $-0.3, 0.4$),
although the difference was not significant. However, none of
the interactions (between birth order and sibling presence in
relation to height, systolic blood pressure, or diastolic blood
pressure) was significant.

DISCUSSION

The smaller birth size in firstborns shown here has been
observed in not only humans but also other mammals such
as mice (25), sheep (26), and seals (27). However, this early
growth deficit was more than compensated for such that, by
6 months, the firstborns had become heavier and taller than
later borns, with a height difference of 2.6 cm persisting at
13.3 years. In turn, these early slow-fast growth patterns were
associated with greater blood pressure in firstborns, but not
with differences in body mass index or fatness.

Our analyses indicate that these birth order associations
are not due to confounding by maternal factors or the sex of
the offspring. The mothers of firstborns were 1.5 cm taller.
Hence, it is possible that minor differences in growth po-
tential might have become magnified during the window for
catch-up growth. Nevertheless, birth order differences in height
and blood pressure remained at 2–3 mm Hg, after adjustment
for sex and maternal height. In general, differences of such
magnitude are clinically important (28).

With further adjustment for early growth patterns and phys-
ical activity, the associations of birth order with height prac-
tically disappeared. This is not surprising as the literature
strongly suggests that adolescent and adult height deficits are
largely explained by growth patterns in early childhood (29).
Associations between birth order and blood pressure were re-
duced by about 0.5 mm Hg after such adjustment, and the
confidence interval of these differences included the null
value. This indicates that these factors partly contribute to the
mechanisms whereby birth order is associated with later blood
pressure. Early growth appeared more important for systolic
blood pressure, and activity patterns seemed more important
for diastolic blood pressure. Further work is required to
understand these contrasting findings in more detail. We
previously observed reduced activity levels in firstborns in this cohort at 11 years and suggested that the number of siblings might explain this association (30).

Several other studies have shown firstborns to have a tendency to overcompensate in catch-up growth and to become taller by adolescence and adulthood (11–13). Whether the smaller birth size of small-for-gestational-age infants tracks into later life is strongly dependent on the magnitude of early catch-up during the first 6–12 months (31), in other words, a “critical window.” This might account for the findings of Hermanussen et al. (10), who found that firstborns remained shorter than their peers in adulthood. The opportunity for catch-up may have been diminished in this population studied in the aftermath of World War II, causing early deficits to track into adulthood.

Elsewhere, early catch-up growth has been associated with increased adiposity in firstborns. Firstborns had greater central adiposity by 5 years in the Avon Longitudinal Study of

### Table 3. Outcomes at 13.3 Years According to Birth Order Among Brazilian Adolescents in the 1993 Pelotas Birth Cohort

<table>
<thead>
<tr>
<th>Variable</th>
<th>Mean (SD)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Firstborns (n = 143)</td>
<td>Others (n = 310)</td>
</tr>
<tr>
<td>Height, cm</td>
<td>159.6 (7.8)</td>
<td>157.0 (8.5)</td>
</tr>
<tr>
<td>BMI, kg/m²</td>
<td>20.3 (3.3)</td>
<td>20.3 (4.0)</td>
</tr>
<tr>
<td>Sum of skinfolds, mm</td>
<td>25.4 (12.0)</td>
<td>25.4 (13.7)</td>
</tr>
<tr>
<td>Accelerometry, 1,000 counts</td>
<td>384 (142)</td>
<td>401 (149)</td>
</tr>
<tr>
<td>Accelerometry, minutes/week MVPA</td>
<td>485 (218)</td>
<td>515 (218)</td>
</tr>
<tr>
<td>Systolic blood pressure, mm Hg</td>
<td>113.4 (13.9)</td>
<td>109.8 (14.0)</td>
</tr>
<tr>
<td>Diastolic blood pressure, mm Hg</td>
<td>70.7 (12.0)</td>
<td>67.7 (10.8)</td>
</tr>
<tr>
<td>Lean mass, kg</td>
<td>39.3 (7.4)</td>
<td>38.6 (7.0)</td>
</tr>
<tr>
<td>Fat mass, kg</td>
<td>11.9 (7.1)</td>
<td>11.8 (7.9)</td>
</tr>
<tr>
<td>Lean mass index, kg/m²</td>
<td>15.4 (1.9)</td>
<td>15.6 (1.9)</td>
</tr>
<tr>
<td>Fat mass index, kg/m²</td>
<td>4.7 (2.7)</td>
<td>4.8 (3.0)</td>
</tr>
</tbody>
</table>

Abbreviations: BMI, body mass index; MVPA, moderate-to-vigorous physical activity; SD, standard deviation.

<sup>a</sup> t test.

### Table 4. Adjusted Analyses for Outcomes at 13.3 Years According to Birth Order Among Brazilian Adolescents in the 1993 Pelotas Birth Cohort<sup>a</sup>

<table>
<thead>
<tr>
<th>Variable</th>
<th>Unadjusted Model 1&lt;sup&gt;b&lt;/sup&gt;</th>
<th>Model 2&lt;sup&gt;c&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Firstborns Coefficient 95% CI</td>
<td>Firstborns Coefficient 95% CI</td>
</tr>
<tr>
<td>Height, cm</td>
<td>2.6 1.0, 4.2&lt;sup&gt;0.01&lt;/sup&gt;</td>
<td>2.1 0.5, 3.8 0.01</td>
</tr>
<tr>
<td>BMI, kg/m²</td>
<td>−0.1 −0.8, 0.7 0.83&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.1 −0.7, 0.9 0.80&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Sum of skinfolds, mm</td>
<td>0.0 −2.6, 2.6 1.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.9 −1.8, 3.7 0.51&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Accelerometry, 1,000 counts</td>
<td>−17.3 48.8, 14.2 0.28&lt;sup&gt;a&lt;/sup&gt;</td>
<td>−22.6 55.3, 10.2 0.18&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Accelerometry, minutes/week MVPA</td>
<td>−30 −77, 17 0.21&lt;sup&gt;a&lt;/sup&gt;</td>
<td>−39 −88, 10 0.12&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Systolic blood pressure, mm Hg</td>
<td>3.5 1.2, 5.7 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.5 0.4, 4.7 0.02&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Diastolic blood pressure, mm Hg</td>
<td>2.0 0.7, 3.4 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.1 0.7, 3.6 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Lean mass, kg</td>
<td>0.77 −0.71, 2.26 0.31&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.10 −1.43, 1.63 0.90&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Fat mass, kg</td>
<td>0.06 −1.53, 1.66 0.94&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.57 −1.07, 2.21 0.50&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Abbreviations: BMI, body mass index; CI, confidence interval; MVPA, moderate-to-vigorous physical activity.

<sup>a</sup> Reference group: nonfirstborn.

<sup>b</sup> Model 1: adjusted for confounders (maternal age, maternal height, maternal BMI, smoking during pregnancy, family income, and sex).

<sup>c</sup> Model 2: adjusted for model 1 variables plus z score of birth weight, conditional weight and length at 48 months, and physical activity (counts) at 13.3 years.

<sup>d</sup> Not adjusted for physical activity (counts) at 13.3 years.

Pregnancy and Childhood (ALSPAC) cohort (7) and increased weight and body mass index in teenage girls from Poland (32). Birth order enhanced associations of socioeconomic status with central adiposity in young adult males from the Philippines (33). In adults, increased total and central adiposity was observed in firstborn Bengali women (12) and Brazilian men (13). In our study, however, birth order differences in adiposity were not apparent. This is consistent with several recent studies in developing countries, where infant catch-up appears beneficial for later height and lean mass but much weaker in association with adiposity (34, 35), whereas in industrialized populations, rapid growth is associated with later adiposity (36, 37) and obesity (38, 39). Such differences may also be due to variation in the duration of catch-up growth.

Other studies have suggested family size associations with growth. Among Da-an boys of Taiwan, boys without sisters were ~2.5 cm taller and ~3.5 kg heavier than those with 1 or 2 sisters (40). A study of Cairo children showed that boys in smaller family sizes were taller and heavier than those of larger family sizes, but with no specific birth-order association (41). In the ALSPAC cohort, family size was negatively related to height, so that compared with children lacking siblings, those with 4 siblings had 0.9 cm lower birth length and 3.1 cm reduced height at age 10 years (42). In our study, firstborns without a sibling achieved greater height by 4 years and had the highest values for height and blood pressure at 13.3 years. However, these differences were not significant, and the association with family size appears much less important than that with early growth patterns for explaining the birth-order difference in height and systolic blood pressure.

The strengths of the study include the prospective nature of the early growth data, the relatively large sample size, and the objective measurements of body composition and physical activity level. The main limitations comprise the definition of birth order according to previous pregnancies and the lack of information about siblings. However, our definition of "firstborn" would tend to a conservative estimate of any differences, because some individuals may have been classified as secondborn despite the previous pregnancy’s not persisting long enough to affect uterine physiology. A second limitation is that we did not adjust our study for multiple comparisons. However, we believe this approach is acceptable, as previously published studies have reported associations of birth order with all the outcomes we considered here.

Our findings are important in understanding the pathways whereby early growth patterns are associated with later degenerative disease. There is little reason to assume that maternal nutritional status itself varies substantially across successive pregnancies in this population, where undernutrition is practically nonexistent. Many components of maternal phenotype (e.g., height, uterine volume) are relatively consistent across the reproductive career and reflect early maternal growth and development. Other components of phenotype may change modestly across pregnancies, such as body weight and adipose tissue distribution (43, 44). The primary shift in fetal growth pattern occurs between the first and second pregnancies (8) and appears attributable to changes in uterine vascular function (9) rather than to drastic changes in maternal nutritional status. The reduced birth size of firstborns therefore appears due to reduced access to maternal resources rather than the inadequacy of those resources.

Our findings therefore do not support the predictive adaptive response hypothesis (45), which attributes associations between early growth and later metabolic phenotype to proactive “anticipation” of future breeding conditions. It is difficult to see how or why firstborns should generate different predictions of future environmental conditions than their later-born peers, given a common maternal phenotype. Instead, our findings are consistent with the maternal capital hypothesis (46), and they indicate that firstborns have reduced access to the maternal nutritional supply during pregnancy, resulting in their smaller size at birth. Firstborns can undergo rapid compensatory growth during infancy but at a cost of long-term associations with blood pressure. The greater sensitivity of firstborns to factors promoting catch-up could be addressed in nutritional management to avoid catch-up “overshoot.”

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