Birthweight in Congenital Virilizing Adrenal Hyperplasia

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Qazi, Q. H., and Thompson, M. W. (1971). Archives of Disease in Childhood, 46, 350. Birthweight in congenital virilizing adrenal hyperplasia. Birthweights of 28 male and 38 female patients with congenital virilizing adrenal hyperplasia were compared with those of 27 male and 26 female unaffected sibs and newborns in Ontario. Only female patients were found to be significantly heavier than the female controls and female sibs. The mean birthweight of salt-losers was not significantly different from that of non-salt-losers. The female patients with more severe genital abnormalities did not have higher birthweight than those with less severe genital abnormalities.

The deficiency of C-21-hydroxylation in adrenocorticosteroid synthesis is the commonest cause of congenital virilizing adrenal hyperplasia (CVAH). Abnormal secretory activity in CVAH begins about the third month of gestation before the differentiation of external genitalia (Bierich, 1963). The masculinization of the female external genitalia noted at birth in almost all cases is thought to result from the androgenic action of excessive amounts of adrenal androgens produced by the fetal cortex. The rapid somatic growth and increased musculature in untreated patients are considered to be due to the protein anabolic effect of adrenal androgens which continue to be secreted postnatally in large amounts (Wilkins, 1965).

In the course of study of CVAH in Ontario, information regarding birthweight of patients with CVAH, their unaffected sibs, and newborn infants in the province of Ontario was obtained and the data were compared. This report summarizes our observations.

**Patients and Methods**

Seventy patients with C-21-hydroxylase deficiency were diagnosed at the Hospital for Sick Children, Toronto, in the period 1945–1967. In addition to characteristic clinical manifestations in the salt-losing and non-salt-losing patients, the diagnostic investigations included (1) estimation of urinary 17-ketosteroids and pregnanetriol before and after cortisone or dexamethasone administration; (2) x-rays for bone age; (3) intravenous pyelograms; and (4) sex chromatin study in female pseudohermaphrodites. In 3 cases the diagnosis was made at necropsy.

Most of the patients were referred to the hospital within a few weeks after birth. Information regarding birthweight and gestational age was obtained from the medical records, and was later verified in most instances by personal interviews with the parents. Data were available on 69 of 70 patients. Similar information was obtained about normal sibs of the patients, and about 3 affected sibs who had died earlier.

For birthweight analysis, only those children who were born after 37 weeks' gestation were used. The following 3 groups were compared: (1) affected children (28 male and 36 female); (2) normal sibs of the children in (1) above (27 male and 26 female); (3) children born in Ontario in 1960, used as population controls (71,817 male and 68,246 female).

The mean year of birth of patients in the present study was 1959, but since the 1959 Vital Statistics Report of Ontario did not include birthweight, 1960 was chosen as the nearest year.

**Results**

There were significantly more females with CVAH who had a birthweight of 3500 g or higher than expected (Table I). The mean birthweight of female patients was also significantly higher than that of their female sibs (Table II). Comparison of birthweight of male patients with male controls and their male sibs did not show similar differences. The mean birthweight of salt-losers was not
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significantly different from that of the non-salt-losers (Table III).

**TABLE I**

*Observed versus Expected Frequencies of Birthweights in Male and Female Patients with CVAH*

<table>
<thead>
<tr>
<th>Weight Groups (g)</th>
<th>Male Patients</th>
<th>Female Patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>2000-3499</td>
<td>11.39%</td>
<td>15.1%</td>
</tr>
<tr>
<td>3500-4999</td>
<td>17.50%</td>
<td>12.9%</td>
</tr>
</tbody>
</table>

For males, \( \chi^2 = 2.416, \text{D.F.}, 1, P > 0.10. \)
For females, \( \chi^2 = 11.149, \text{D.F.}, 1, P < 0.005. \)

**TABLE II**

*Birthweights of Affected Children and their Normal Sibs*

<table>
<thead>
<tr>
<th>Subjects</th>
<th>No.</th>
<th>Birthweight (g)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male patients</td>
<td>28</td>
<td>3,594</td>
<td>346</td>
</tr>
<tr>
<td>Male sibs</td>
<td>27</td>
<td>3,493</td>
<td>519</td>
</tr>
<tr>
<td>Female patients</td>
<td>38</td>
<td>3,593</td>
<td>424</td>
</tr>
<tr>
<td>Female sibs</td>
<td>26</td>
<td>3,094</td>
<td>408</td>
</tr>
</tbody>
</table>

**TABLE III**

*Birthweights of Patients with Salt-losing and Non-salt-losing Forms of CVAH*

<table>
<thead>
<tr>
<th>Subjects</th>
<th>No.</th>
<th>Birthweight (g)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Salt-losing males</td>
<td>20</td>
<td>3,651</td>
<td>350</td>
</tr>
<tr>
<td>Non-salt-losing males</td>
<td>8</td>
<td>3,452</td>
<td>300</td>
</tr>
<tr>
<td>Salt-losing females</td>
<td>23</td>
<td>3,662</td>
<td>477</td>
</tr>
<tr>
<td>Non-salt-losing females</td>
<td>15</td>
<td>3,488</td>
<td>314</td>
</tr>
</tbody>
</table>

The genital abnormalities of female patients were classified in 5 types according to Prader (1954). None of the patients had the rare type V abnormality. The mean birthweight of 28 patients with types III and IV (severe) abnormalities was not significantly different from that of 10 patients with types I and II (milder) abnormalities (Table IV).

**TABLE IV**

*Birthweights in Female Patients with Types I and II and Types III and IV Genital Abnormalities*

<table>
<thead>
<tr>
<th>Genital Types</th>
<th>No.</th>
<th>Birthweight (g)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>I + II</td>
<td>10</td>
<td>3,542</td>
<td>274</td>
</tr>
<tr>
<td>III + IV</td>
<td>28</td>
<td>3,560</td>
<td>470</td>
</tr>
</tbody>
</table>

Because of a more severe disturbance of corticosteroid synthesis, greatly increased production of androgens would be expected to occur in salt losers during fetal life. However, the mean birthweight of salt losers was not significantly higher than that of non-salt-losers. Similarly, an attempt
to correlate birthweight to the severity of genital defects in female patients did not show the mean birthweight of patients with the more severe genital abnormalities to be higher than that of patients with the milder abnormalities. The absence of significant correlation between the birthweight and severity of genital abnormalities may be due to a rather low ratio of anabolic to androgenic effects of the androgens produced by the fetal adrenal cortex (Suchowsky and Junkman, 1965).

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REFERENCES


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