Fasting Serum Insulin Levels in Childhood

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There is little published information on insulin levels in normal children. Ehrlich and Bammers (1964) found the fasting plasma insulin levels of 6 normal children to be similar to adult levels. Slone, Soeldner, Steinke, and Crigler (1966) have reported low fasting serum insulin levels in 13 normal children between the ages of 3 and 9 years.

This paper describes the fasting immuno-reactive insulin and blood glucose concentrations in 86 European children who had been admitted to hospital for elective surgical procedures. Insulin and glucose levels in 9 premature infants and 12 young adults are also given.

Subjects and Methods

Children. The group of children consisted of 50 boys and 36 girls, and their ages ranged from 3 months to 14 years. All the children were well and had been leading relatively normal lives before admission. None was considered to need either medical investigation or medical treatment. The conditions for which the children were admitted to hospital are shown in Table I.

The distribution of the children’s weights expressed as centile weight (Tanner, 1958) is shown in Table II. The difference between the numbers of children in each centile group and the expected numbers is not statistically significant (p > 0·05). In most of the older children sexual development was assessed (Tanner, 1955). Almost all the children over the age of 11 years showed evidence of the onset of puberty.

Fasting blood specimens were collected before breakfast on the morning after admission to hospital, after a period of fasting which varied between 8 to 14 hours. Blood was obtained by finger-prick or heel-prick at the same time as blood was collected for a pre-operative haemoglobin estimation. Care was taken to avoid haemolysis.

Adults. Venous blood was collected from 12 young adults after an overnight fast.

Premature infants. Blood was collected by heel-prick from 9 infants with birthweights below 2500 g. The birthweights of 7 of these infants were compatible with gestational age. The blood was collected 3 hours after a feed, at the same time as blood was collected for Hb estimation. All the infants were less than 6 weeks old, were gaining weight, and were about to be discharged from the nursery.

Capillary blood was collected as this is a much simpler procedure in small children than a venepuncture, particularly if multiple samples are required. It has been found that, providing haemolysis is avoided, insulin levels in venous and capillary samples collected at the same time correlate well (Fig. 1). Providing insulin levels are not changing rapidly, the speed at which a capillary specimen is obtained does not seem to be important.

Serum, rather than heparinized plasma, was used for the insulin studies, as it is difficult to control the heparin content of capillary samples and high concentrations of heparin were found to interfere with the immunoassay. Serum was obtained by allowing specimens to clot for 9 hours at 4°C. Serum insulin was determined by a modification of the double-antibody immunoassay.

<table>
<thead>
<tr>
<th>Condition</th>
<th>No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hernia or hydrocele</td>
<td>15</td>
</tr>
<tr>
<td>Hypospadias</td>
<td>11</td>
</tr>
<tr>
<td>Undescended testis</td>
<td>7</td>
</tr>
<tr>
<td>Hare-lip or cleft palate</td>
<td>7</td>
</tr>
<tr>
<td>Haemangioma</td>
<td>7</td>
</tr>
<tr>
<td>Bat ears</td>
<td>5</td>
</tr>
<tr>
<td>Syndactyly</td>
<td>5</td>
</tr>
<tr>
<td>Old polio; dislocated hip</td>
<td>10</td>
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<tr>
<td>Miscellaneous</td>
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<table>
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<th>10-25</th>
<th>25-50</th>
<th>50-75</th>
<th>75-90</th>
<th>&gt; 90</th>
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<tr>
<td>Actual number</td>
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<td>13</td>
<td>12</td>
<td>19</td>
<td>17</td>
<td>10</td>
</tr>
<tr>
<td>Expected number</td>
<td>(8·6)</td>
<td>(12·9)</td>
<td>(21·5)</td>
<td>(21·5)</td>
<td>(12·9)</td>
<td>(8·6)</td>
</tr>
</tbody>
</table>

\[ \chi^2 = 10·78; p > 0·05. \]
Blood glucose was measured on specimens collected into bottles containing sodium fluoride and potassium oxalate, using a modification of the glucose oxidase method described by Huggett and Nixon (1957).

The statistical significance of differences between mean values and the significance of regression coefficients was tested by Student's 't' test.

Results

Blood glucose. Fasting blood glucose was measured in 85 of the 86 children. The values ranged from 43 mg./100 ml. to 95 mg./100 ml., with a mean value of 73 mg./100 ml. The slope of the regression line calculated for blood glucose on age (blood glucose (mg./100 ml.) = 0·83 age (years) + 67·7) is statistically significant (p < 0·01). This rise with age is similar to the rise in total reducing substances reported by Mayer (1951).

The mean blood glucose in the 9 premature infants was 69 mg./100 ml. (49-84 mg./100 ml.), and in the 12 adults 77 mg./100 ml. (55-90 mg./100 ml.).

Serum insulin. The fasting serum insulin in the 86 children ranged from undetectable levels to 32 μu/ml., with a mean value of 8 μu/ml. The distribution of the insulin values in different age-groups and the mean values in the different groups is shown in Fig. 2. The mean values show a progressive rise with age. The regression line for insulin on age was calculated for the complete group of children. The slope of this line is statistically highly significant (insulin (μu/ml.) = 1·02 age (years) + 2·1; p < 0·001). The regression lines for boys and girls are almost identical, and the slope of the line for children under 7 does not differ significantly from the slope for children over 7.

Significant correlations between insulin and body-weight (p < 0·001), and insulin and blood glucose (p < 0·01) were also found. There was no significant correlation between the length of fast (8-14 hours) and the insulin concentration.

The fasting insulin values in the 12 adults ranged from 6 to 20 μu/ml., with a mean value of 10 μu/ml. This is significantly lower than the mean value for children over the age of 11 years (p = 0·02).

The insulin values found in 9 premature infants after a 3-hour fast ranged from 15 to 43 μu/ml., with a mean value of 29 μu/ml. This value is significantly higher than the mean value for children over the age of 11 years (p < 0·001).

Discussion

It would have been desirable to study healthy children at home, but as this was not possible the
survey was confined to hospital patients. Samples were collected within 24 hours of admission and only healthy children were studied, in an attempt to minimize any possible effects of illness, an unfamiliar diet, or restricted activity.

The data presented here suggest that the fasting serum insulin concentration rises progressively with increasing age. It is unlikely that this finding is due to changes in the precision of the assay during the period of study, as the assay conditions were kept constant and a control specimen gave consistent results. Specimens were collected from the different age-groups in a random order, and the results of repeated assays agreed closely with the initial assay results. Serum may interfere with the double-antibody assay by modifying the rate of precipitation (Soeldner and Slone, 1965). Under the conditions used, no appreciable non-specific effects were noted, and it is considered unlikely that the results are due to age-dependent serum effects on precipitation.

The interpretation of fasting insulin levels is difficult, as the factors controlling these levels are still incompletely understood. Low insulin levels have been described in adults after prolonged fasting (Yalow and Berson, 1965) and the correlation between fasting insulin level and age or body weight found in this study may merely reflect the ratio between body size and length of fast. Raised fasting insulin levels associated with an exaggerated serum insulin response to glucose have been reported in obese adults (Samaan, Brown, Fraser, and Trayner, 1965), and in patients with hypertension (Welborn, Breckenridge, Rubinstein, Dollery, and Fraser, 1966) and ischaemic heart disease (Peters and Hales, 1965), and fasting insulin levels may be related to tissue sensitivity to insulin.

The difference between the mean fasting insulin levels in the older children of 11-15 years and adults is of interest. Acromegaly (Beck, Schalch, Parker, Kipnis, and Daughaday, 1965), pregnancy (Spellacy and Goetz, 1963), and the administration of corticosteroids (Perley and Kipnis, 1966) are associated with increased serum insulin responses to glucose. The apparent rise in fasting insulin at the time of puberty may reflect the endocrine changes or growth acceleration which occur at this time. The relatively high insulin values found in the premature infants are hardly comparable with the values found in older children, as the infants were all having a

![Fig. 2.—The insulin concentrations and mean values in different age-groups. Open rectangles = ± SD, hatched rectangles = ± SE.](image)
high calorie intake and were fasted for a short period of time. No conclusions can therefore be drawn on changes in insulin levels during the first year of life.

The findings reported here differ from those reported by Slone et al. (1966), who found a low mean fasting insulin level between the ages of 3 and 9 years and higher mean levels in the younger and older children, comparable with the mean value for adults. A possible explanation for this difference is that their series may have contained a relatively high proportion of young infants and post-pubertal children.

Loeb (1966) studied the rate constants for the disappearance of glucose after intravenous injection and found a progressive rise during the first 6 months of life, followed by little change until the onset of puberty when the rate constant fell to adult levels. It seems possible that the change in insulin levels reported here may also indicate changes in carbohydrate metabolism during childhood.

Summary

Fasting serum immuno-reactive insulin and blood glucose values in 86 children, 9 premature infants, and 12 adults are given. In the 86 children fasting serum insulin appeared to rise progressively with increasing age. Blood glucose also showed a significant rise with age. Some possible implications of this are discussed.

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REFERENCES


