Effect of dystonic movements on oesophageal peristalsis in Sandifer's syndrome

SUMMARY
Children with diabetes mellitus are not significantly heavier than their peers, and those who are obese do not have poorer diabetic control.

Adults with insulin dependent diabetes are often obese, and this may adversely affect their diabetic control, resulting in complications. We are unaware of any publication on the incidence of obesity in childhood diabetes or of its effect on diabetic control. In this study we report the incidence of obesity in children attending our diabetic clinic, and relate this to their diabetic control.

Patients and methods
One hundred and six children (58 boys and 48 girls) attending the diabetic clinic at the Bristol Children's Hospital who had had diabetes for more than one year were studied over the 12 month period. Their mean age during the study period was 14·9 years (range 5–22), and their mean age at diagnosis was 8·3 years (range 1·3–15·9). The mean duration of their diabetes was 6·4 years (range 1–16·6). All the children were receiving quick and intermediate acting insulins twice daily, with the exception of three patients who used an insulin pump.

They were seen at intervals of three months, when their height and weight (measured with a Harpenden stadiometer) were recorded. Skinfold thickness (triceps and subscapular) was measured using a Harpenden skinfold caliper, and puberty was assessed clinically by the method of Tanner and Whitehouse. Each child's height, weight, and skinfold thickness was then taken as the average over visits during the year of investigation.

The children were regarded as overweight when.
Weights of 48 diabetic girls

Weights of 58 diabetic boys

Figure  Weights in diabetic girls and boys.

their absolute weight was more than 20% above the median for their age and sex (Tanner charts), or
their relative standard weight percentile (actual
weight/standard weight x 100) was greater than
120%, or their skinfold thickness, both triceps
and subscapular, was above the 70th centile (Tanner
charts).

Glycated haemoglobin (HbA1c) was measured
at each visit by electroendosmosis. The mean annual
HbA1c value was calculated and diabetic control
graded as good when the mean HbA1c value was
10.5% or less, and poor when the mean HbA1c was
over 10.5%.

Statistical analysis comparing various indices of
body fat with the quality of control (HbA1c) was by
the Mann-Whitney U test and probabilities of less
than 0.05 were considered to be significant.

Results

The scatter of children's weights (figure), skinfold
thickness, and relative standard weight percentiles
were no different from those given in the Tanner
charts. The mean annual HbA1c concentration was
11.6% (range 7.2–17.2) (boys 11.4%, girls 11.9%).

Thirty seven children (35%) had a mean annual

Table  Characteristics of good and poor diabetic control in 106 diabetic children

<table>
<thead>
<tr>
<th></th>
<th>Good control (HbA1c ≤10.5%)</th>
<th>Poor control (HbA1c &gt;10.5%)</th>
<th>95% Confidence interval</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean age at diagnosis (years)</td>
<td>8.5</td>
<td>8.3</td>
<td>7.63 to 7.34</td>
</tr>
<tr>
<td>Median absolute weight (kg)</td>
<td>53.1</td>
<td>50.0</td>
<td></td>
</tr>
<tr>
<td>Mean duration (years)</td>
<td>6.1</td>
<td>6.6</td>
<td></td>
</tr>
<tr>
<td>Median triceps skinfold (mm)</td>
<td>13.6</td>
<td>12.0</td>
<td>3.0 to 1.6</td>
</tr>
<tr>
<td>Median subscapular skinfold (mm)</td>
<td>8.0</td>
<td>8.0</td>
<td>1.0 to 2.0</td>
</tr>
<tr>
<td>Relative standard weight (%)</td>
<td>97.3</td>
<td>95.6</td>
<td>7.30 to 5.20</td>
</tr>
<tr>
<td>Insulin dosage (U/kg/day)</td>
<td>0.9</td>
<td>0.9</td>
<td></td>
</tr>
</tbody>
</table>

There were no significant differences between the groups.
HbA1c concentration of 10.5% or less and 69 children (65%) had a mean annual HbA1c concentration of more than 10.5%.

There were no significant differences in the median HbA1c among those children whose absolute weight was more than 20% above the mean (median HbA1c 11.9%), those whose relative weight was more than 120% (median HbA1c 11.1%), or those whose skinfold thickness was more than 70% (HbA1c 11.5%) when compared with their peers (median HbA1c concentrations 11.6%, 11.6%, and 11.7%, respectively). The 95% confidence interval of the difference in medians was −1.20% to 1.15% for absolute weight, −1.7% to 1.2% for relative weight, and −1.2% to 1.0% for skinfold thickness. Conversely, no significant differences were found between those with good as opposed to poor control and the various indices of obesity (table).

Discussion

Any definition of obesity is arbitrary as the distribution of weight in the general population does not segregate into distinct populations of obese and non-obese. Furthermore, methods for evaluating fatness cannot be compared directly and thus it is not easy to derive acceptable criteria for the diagnosis. Direct methods for the measurement of body fat are generally invasive, but it can be measured indirectly by estimating skinfold thickness, and it is this pain free and non-invasive method that is generally used in children.

We are unaware of any publications on the incidence of obesity in childhood diabetes, or on its effect on diabetic control. Many paediatricians believe, however, that children with diabetes are often overweight and that this may have a deleterious effect on their diabetic control. We have found the scatter of the children's weights to be no different from those of the standard national weight charts, and only 9% of children had weights in excess of 120% of relative weight compared with a mean relative weight of 117% in adults with insulin dependent diabetes.

Obesity affecting mainly the upper half of the body correlates with increased atherogenic and diabetogenic mortality and should be considered more seriously than obesity that predominantly affects the hips and lower limbs. In this study 20% of the children had excess fat affecting the upper half of the body. We found no evidence, however, of poorer diabetic control among them, and no correlation between skinfold thickness and HbA1c concentration.

Obesity leads to insulin resistance, but we have no evidence to suggest that this is the case in the obese children as their mean insulin dosage—U/kg body weight—and their mean HbA1c% were similar to those among non-obese diabetic children.

Our diabetic population is not obese and this may reflect the high fibre, low fat diet prescribed in our clinic. As work in adults has confirmed an association between obesity, poor diabetic control, and diabetic complications, it is important to emphasise the prevention of excessive weight gain.

We are grateful to Dr AO Hughes for statistical advice.

References


Correspondence to Dr DCL Savage, Bristol Royal Hospital for Sick Children, St Michael's Hill, Bristol BS2 8BJ.

Accepted 23 March 1989