Mineral balance in infantile cortical hyperostosis: effects of corticosteroids

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Abstract

The effects on mineral metabolism of therapeutic doses of corticosteroids were investigated in infantile cortical hyperostosis; in four untreated cases the calcium, phosphorus, and magnesium balances were strongly positive. In one severe case, treatment with prednisolone was associated with an alteration to negative calcium and magnesium balance, and faecal losses of calcium were particularly high. This effect persisted for at least three months after the steroids had been discontinued, and during this period there was pronounced retardation of linear growth. Six months after the treatment had been stopped mineral balance was again positive and there was rapid 'catch up' in growth.

In infancy, the negative effect of corticosteroids on calcium, phosphorus, and magnesium metabolism may contribute to inhibition of bone growth and steroid stunning.

We know of few biochemical studies in infantile cortical hyperostosis. Serum calcium and phosphorus concentrations are normal but alkaline phosphatase activity is usually increased during the active phase. Aminoaciduria has been reported in one case.

Corticosteroid treatment has been advised with reservation but recommended for all patients, particularly for severe cases. In this study of infantile cortical hyperostosis one case, whose growth was studied longitudinally, had a series of balance studies for calcium, phosphorus, and magnesium before, during, and after treatment with corticosteroids. Three other cases not receiving steroids had single balance studies.

Table 1 Calcium balances

<table>
<thead>
<tr>
<th>Case</th>
<th>Age (months)</th>
<th>Weight (kg)</th>
<th>Intake (mmol/day)</th>
<th>Output (mmol/day)</th>
<th>Gross retention (mmol/day)</th>
<th>Comment</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Faeces</td>
<td>Urine</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Case 1</td>
<td>Balance (i)</td>
<td>3.5</td>
<td>4.9</td>
<td>28.7</td>
<td>18.6 0.6</td>
<td>9.5</td>
</tr>
<tr>
<td></td>
<td>Balance (ii)</td>
<td>5</td>
<td>5.1</td>
<td>24.0</td>
<td>20.95 0.2</td>
<td>2.85</td>
</tr>
<tr>
<td></td>
<td>Balance (iii)</td>
<td>6.5</td>
<td>5.9</td>
<td>18.7</td>
<td>18.5 0.7</td>
<td>-0.5</td>
</tr>
<tr>
<td>Case 2</td>
<td>Balance (iv)</td>
<td>10</td>
<td>7.1</td>
<td>8.4</td>
<td>8.7 0.4</td>
<td>-0.7</td>
</tr>
<tr>
<td></td>
<td>Balance (v)</td>
<td>14</td>
<td>8.7</td>
<td>11.1</td>
<td>5.2 0.1</td>
<td>5.8</td>
</tr>
<tr>
<td></td>
<td>Case 2</td>
<td>5</td>
<td>5.9</td>
<td>19.0</td>
<td>9.1 0</td>
<td>9.9</td>
</tr>
<tr>
<td>Case 3</td>
<td>Case 3</td>
<td>4</td>
<td>6.8</td>
<td>21.5</td>
<td>12.2 0.9</td>
<td>8.4</td>
</tr>
<tr>
<td></td>
<td>Case 4</td>
<td>5</td>
<td>5.6</td>
<td>17.7</td>
<td>10.0 1.4</td>
<td>6.3</td>
</tr>
</tbody>
</table>

*Three day balance period, all others five day periods. ACTH, adrenocorticotrophic hormone.

Patients and methods

Clinical details of the four infants are shown in table 1. Case 1 had generalised disease and was seriously ill; a protracted course of corticosteroid was therefore necessary. Case 2 had generalised disease, and cases 3 and 4 had disease localised to the mandible that resolved without the need for steroids.

Balance studies, except where stated, all extended over five day periods. Analysis for calcium, phosphate, and magnesium was made on duplicate diets, stools, and urine. Samples were homogenised and aliquots were digested in dilute nitric acid. Calcium and magnesium were estimated by atomic absorption spectrophotometry and phosphate by a micro adaptation of the phosphomolybdate method of Fiske and Subbarow.

Results

In all four patients serum calcium, magnesium, and phosphorus were always within normal limits and there was no evidence of acid-base disturbance, hepatic, renal glomerular, or tubular dysfunction.

Changes in linear growth in case 1 are shown in fig 1. Height is shown as the difference between the observed value and the normal mean height for that age expressed as a proportion of the normal SD for that age (SD score). Balance data in the four cases are shown for calcium, phosphorus, and magnesium in tables 1, 2, and 3, respectively. Sequential balance data for case 1 are shown diagrammatically in fig 2.

When untreated (case 1, balance (i); cases 2, 3, and 4) gross retention of calcium was high for age and also in relation to intake. This excessive retention was especially pronounced in...
Changes in phosphate balance were less pronounced and consistent, but magnesium retention fell progressively and balance became negative at the time when calcium retention was lowest. Urinary magnesium seemed to rise at this stage, although in magnesium deprivation there is usually renal conservation.10

Discussion
The metabolic balance data presented here show that untreated infants with cortical hyperostosis are in strongly positive mineral balance.

Balance is usually positive in infancy with a net retention of calcium of about 5 mmol (200 mg)/day to allow for accretion into the growing skeleton.6 Anything less than this implies a net deficit of calcium, and balances that actually become negative (case 1 balances (iii) and (iv)) indicate a serious degree of mineral depletion.

The general tendency to positive balance in cases 2, 3, and 4—and in case 1 the strongly positive balance before and after treatment—suggest that corticosteroids were indeed responsible for the change to negative mineral balance that occurred in case 1.

The main reason for this negative calcium balance seemed to be an increase in faecal losses. This effect persisted for months after treatment had been stopped, suggesting some profound functional alteration in intestinal mechanisms of absorption or excretion. The tendency for faecal calcium to exceed calcium intake might indicate a state of calcium leakage into bowel ("calcium losing enteropathy"). Urinary calcium in cases 1, 3, and 4 was within the reference range,8 but in case 2 no calcium was detected in the urine over a five day period, and at other times values of 0.025 mmol and 0.15 mmol/day were obtained. In this case serum calcium tended to be near the lower limit of the reference range; as a result tubular reabsorption of calcium may have been enhanced by the effect of secondary hyperparathyroidism. The aminoaciduria reported in Caffey's disease by Campbell and Turner was renal in type, also suggesting an effect on tubular function.2

The relationship between the changes in mineral balance and the growth retardation in case 1 is uncertain, as the effects of treatment cannot be entirely separated from the effects of the disease. Negative balance coincided with maximum depression of linear growth, however, and a reversion to positive balance correlated with the period of 'catch up' growth. Prolonged courses of large doses of corticosteroids lead to stunting with impairment of bone growth and delayed skeletal maturation,11 12 but the pathogenesis of these effects remains uncertain.13 14 Among contributing factors, the adverse effects of corticosteroids on mineral metabolism and bone have been shown in children,15-17 and direct correlation shown between the degree of osteopenia and the extent of growth retardation.18 The reported effects of corticosteroids on bone include inhibition of collagen synthesis,14 decreased osteoblastic activity,19 and osteopenia.17 18 20

Calcium depletion complicating treatment with steroids in children has been
During rapid development of the skeleton in infancy, the side effects of steroids on linear growth and mineral metabolism might be especially undesirable. This study suggests that mineral depletion by corticosteroids may be severe and prolonged in infancy contributing to the inhibition of bone growth. Mineral supplements, with or without vitamin D, might help to offset these mineral losses and mitigate the stunting effects of steroids.