Renal calcification in preterm infants: follow-up at 4–5 years

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Abstract

Aim—to determine the consequences of renal calcification in preterm infants. Methods—A cohort of 11 preterm babies was studied at the age of 4 to 5 years. They had had renal calcification as neonates. Seventeen matched controls were also studied. Each child had a renal ultrasound scan, a calcium load test, and a desmopressin test for renal concentrating ability (RCA). The study group also had glomerular filtration rate (GFR) estimated, using the height:creatinine ratio, and tubular phosphate reabsorption, without phosphate load, per glomerular filtration rate (Tp/GFR) calculated. Results—In the study group the median GFR was 61 ml/min/1.73m² (range 46–79 ml/min/1.73m²) and the median calculated Tp/GFR SD score was −0.94 (range −2.8–0.68). Five children out of the study group had ultrasonic evidence of renal calcification. There was no significant difference between the two groups in renal size, calcium, before or after calcium load, or RCA. Eight children (three patients, five controls) had an abnormal calcium load test. The RCA of the children in the study and control groups combined was below that of published values, with a median calculated SD score −0.71 (95% CI −1.21 to −0.23).

Conclusions—There was evidence of renal dysfunction in children who had been born preterm. Renal calcification detected in the neonatal period does not seem to be a major predisposing factor for the abnormalities of renal function subsequently observed in these infants. (Arch Dis Child 1997;76:F185–F189)

Keywords: renal calcification; glomerular filtration rate; tubular phosphate reabsorption

Renal calcification was first reported in preterm infants in 1982.1 Its presence has been linked to frusemide treatment and hypercalciuria,2,3 and to an inappropriate dietary calcium:phosphate ratio.4,5 Other associations suggested have been the use of dexamethasone6 or xanthines,7 poor urinary flow rate, alterations in urinary pH and a decrease in the urinary concentrations of inhibitors to crystal formation.8 More recently, an increase in the urinary calcium oxalate saturation observed in neonates receiving total parenteral nutrition was shown to be associated with nephrocalcinosis.9 Jacinto et al10 and Short and Cooke11 have shown that the smallest and most immature preterm infants are more likely to develop renal calcifications. In the latter study multivariate analysis showed that duration of oxygen treatment was the strongest clinical indicator of calcification. Acute complications related to renal calcification may be urinary tract infections, abdominal pain, and haematuria.

Short term complications include renal glomerular and tubular dysfunction. Downing et al showed that preterm infants, aged 1 to 2 years, who had developed renal calcifications, had decreased tubular function when compared with preterm infants who had not developed renal calcifications.12 Ezzedeen et al reviewed nine preterm infants aged 9 to 56 months, who had renal calcification in the newborn period, and found that the serum creatinine and calculated glomerular filtration rates were abnormal in four of them.13

The aim of this study was to determine the presence of medium term renal complications in children aged 4 to 5 years, who had developed renal calcification in the neonatal period.

Methods

In a study performed at Liverpool Maternity Hospital in 1989 21 preterm infants were found to have renal calcification.11 Renal calcification was identified by sonography, using strict criteria.13 Five of these infants later died. The remaining 16 children were invited to take part in the present study. The parents of three children declined to take part and two other children were untraceable. The remaining 11 children, who had ultrasonic evidence of renal calcification in the neonatal period, constituted the study group. These were matched for sex, birthweight, gestational age and chronological age with 17 controls from the same original study, who did not have renal calcification as neonates.

Both the study group and the control group had a renal ultrasound scan, a calcium load test, an intranasal desmopressin test and urinalysis. The renal ultrasound scan used an Ultra Mark 9 high definition imaging ultrasound machine (Advanced Technical Laboratories, Seattle, USA), with a 3.5 MHz curvilinear transducer and a linear 10.5 MHz transducer, to measure renal length and identify the presence of renal calcification. This was identified if there were focal areas of hyperechogenicity visible in the renal medulla. These foci did not necessarily show acoustic shadowing. The ultrasound scans were per-
Table 1  Demographic details of study group and controls

<table>
<thead>
<tr>
<th></th>
<th>Study group (n=11)</th>
<th>Controls (n=17)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Percentage male</td>
<td>36</td>
<td>35</td>
</tr>
<tr>
<td>Median (range) birthweight (g)</td>
<td>850 (580–1856)</td>
<td>982 (710–1760)</td>
</tr>
<tr>
<td>Median (range) gestational age (weeks)</td>
<td>27 (24–31)</td>
<td>28 (25–31)</td>
</tr>
<tr>
<td>Median (range) chronological age (months)</td>
<td>56 (49–61)</td>
<td>54 (52–61)</td>
</tr>
</tbody>
</table>

Aminoglycoside concentrations had been taken at the third or fourth dose either after the start of treatment or if the dose was changed. The parents were asked to complete a questionnaire. If there was a previous history of urinary tract infection the general practitioner or hospital notes were used to verify this. The study group also had the following investigations. (For ethical reasons blood was not taken from the control group.)

Plasma sodium, calcium, phosphate and creatinine were measured using a Technicon autoanalyser. Parathyroid hormone (PTH) was measured using an immunoradiometric assay recognising the intact molecule. At the same time as the plasma sample, urine was collected and analysed for creatinine, sodium, and phosphate.

Using these results, tubular phosphate reabsorption under basal conditions, without phosphate load, per glomerular filtration rate, was calculated as a measure of renal phosphate handling.18

\[
\text{Tp/GFR} = \frac{\text{Sp}}{\text{Up}} \times \frac{\text{Scr}}{\text{Ucr}}
\]

where Sp = serum phosphate in mmol/l, Up = urinary phosphate in mmol/l, Scr = serum creatinine in mmol/l, Ucr = urinary creatinine in mmol/l.

An estimate of creatinine clearance was calculated using the modified Couhan-Barratt formula.19 20

\[
\text{GFR} = \frac{40 \times \text{ht}}{\text{Pcr}}
\]

where GFR= glomerular filtration rate in ml/min/1.73m², ht=height in cm, and Pcr=plasma creatinine concentration in µmol/l.

The study was approved by the local ethics committee.

Statistical analysis was performed using the Mann-Whitney U test for continuous variables and the χ² or Fisher’s exact test using 2 × 2 tables for categorical variables.

Results

There was no significant difference between the median age, gestational age, birthweight and sex between the study group and the control group (table 1). Five out of 11 children, aged 4 to 5 years, in the study group had ultrasonic evidence of renal calcification. This was present in both kidneys in three of them. Renal calcification was not identified in any of the control group. An additional child in the study group had renal colic at the age of 3, secondary to a right sided ureteric calculus. An ultrasound scan at that time also showed calcification. A calcium load test indicated that this patient had absorptive hypercalciuria and she was then started on a low calcium diet. At the age of 5 her ultrasound scan was normal.

One child in the study group had only one kidney and another had a renal cyst in the upper pole of the left kidney. No structural renal abnormalities were found in the control group. Renal length was measured ultrasonically in each kidney and plotted against height, using a published reference range.21 Out of 55
Table 2: Results of renal investigations performed on study and control group

<table>
<thead>
<tr>
<th>Investigation</th>
<th>Study group (n=11)</th>
<th>Controls (n=17)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Renal calcification on ultrasonography</td>
<td>5</td>
<td>0</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>No of kidney &lt; 95% CI for length</td>
<td>4</td>
<td>4</td>
<td>NS</td>
</tr>
<tr>
<td>Median (range) post-load calcium/creatinine (mmol/mmol)</td>
<td>0.22 (0.03-1.15)</td>
<td>0.23 (0-1.6)</td>
<td>NS</td>
</tr>
<tr>
<td>Median (range) osmolality (mosmol/kg)</td>
<td>1022 (646-1086)</td>
<td>928 (727-1247)</td>
<td>NS</td>
</tr>
</tbody>
</table>

The median calculated SD score for Tp/GFR was −0.94. Three children in the study group had a PTH value below the normal range. The serum values of calcium and phosphate were all within the normal range. These results are summarised in Table 3.

There were no significant associations between decreased renal concentrating capacity or hypercalciuria and gestational age, birth weight, days of supplemental oxygen, retinopathy of prematurity or intraventricular haemorrhage. There was a trend towards those children who had idiopathic hypercalciuria to have received an aminoglycoside following a high trough concentration in the neonatal period, but this was not significant (P=0.07, Fisher’s exact test).

In three children (two in the study group, one control) urinary tract infection had been confirmed on a midstream urine sample since discharge from the neonatal unit. In the two children from the study group this was associated with systemic upset but did not result in renal scarring on a subsequent dimercaptosuccinic acid technetium isotope (DMSA) scan.

Discussion

Few studies have looked at the short to medium term effects of renal calcification in children who were born preterm. Ezzedeen et al looked at nine children aged between 9 and 56 months, who had been identified as having renal calcification in the neonatal period. Sonography revealed complete resolution of renal calcification in four children and improved, but persisting, calcification in five children. One child had a small kidney on one side, and the serum creatinine and calculated GFR were abnormal in four. There were no controls in this study.
In the study by Downing et al those children who had received frusemide and had developed renal calcifications had a significantly higher urinary calcium:creatinine ratio and fractional excretion of sodium compared with children who had not developed calcifications but had received frusemide. The first group also had a lower tubular reabsorption of phosphate and lower tubular acidification capacity. The average chronological age in this study was 14.1 months and some of the children would have been less than 12 months corrected age, when renal tubular and glomerular function are not yet mature. Renal calcification seems to diminish with increasing age. Downing et al suggested that as calcification improves renal function may also improve. We therefore felt that it was important to study a group of children aged 4 to 5 years old, who had been identified as having renal calcifications in the neonatal period.

In our study five out of 11 children, aged 4 to 5 years, had ultrasonic evidence of renal calcification. Ultrasound scanning is the most sensitive imaging technique for the detection of renal calcification and there is good correlation between ultrasound findings and histological evidence of nephrocalcinosis and nephrolithiasis. There was no clinical evidence of other pathological processes that might give rise to similar ultrasonic appearances. The patients classified as having renal calcification in the original study had areas of hyperechogenicity on at least two consecutive scans, performed at weekly intervals.

The age at diagnosis of renal calcification ranged from 27 days to 66 days. Hyperechogenicity associated with renal insufficiency in the preterm infant usually occurs earlier than this and tends to be transient.

Predicted GFR and Tp/GFR were reduced in some children from the study group. As blood was not taken from the control group it is uncertain whether prematurity or renal calcification was the predisposing factor for this.

Renal calcification in preterm neonates does not seem to affect subsequent renal calcium excretion or tubular concentrating ability, as there were no significant differences between the study and control groups. Twenty eight per cent of all the children (both study and control groups) had an abnormal calcium load test (four children had a fasting urinary calcium:creatinine ratio >0.7 mmol/mmol and four children had a normal fasting urinary calcium: creatinine ratio but a post load value of >1 mmol/mmol). Five children from the control group had an abnormal calcium load test, which suggests that preterm infants may have an increased risk of developing idiopathic hypercalciuria that is not associated with renal calcification. Hypercalciuria is defined in children as a urinary calcium excretion greater than 4 mg/kg/day. Other authors have demonstrated a correlation between 24 hour urinary calcium and a urinary calcium:creatinine ratio on an overnight urine sample, or a second fasting urine after liquid intake. We did not measure calcium excretion on a 24 hour urine collection because of its unreliability in this age group. We chose the calcium load test. This involves a fasting urine sample and a urine collection following a calculated dose of calcium because urinary calcium excretion is affected by dietary calcium. The upper reference limit for the second fasting urine sample was taken from a study measuring urinary calcium on a population of children from the North of England, who were eating a normal diet. The calcium load test may appear normal in children with hypercalciuria, defined by a 24 hour urinary calcium >4 mg/kg/day, if they have low salt or restricted calcium intake before the test. An abnormal calcium load test in a child with a normal 24 hour urinary calcium has been described in a child with a history of renal colic and renal stones. We are not aware of other studies that have shown an abnormal calcium load test in children with normal urinary calcium on a 24 hour collection. It is therefore unlikely that we underestimated the number of patients with raised urinary calcium excretion. One child who was felt to have absorptive hypercalciuria developed a ureteric calculus and was placed on a low calcium diet.

Urinary calcium excretion parallels sodium excretion and is affected by both dietary sodium and calcium. We did not find any difference in dietary sodium or calcium in the three days before the calcium load test between those children who had hypercalciuria and those who did not. The pathogenesis underlying idiopathic hypercalciuria is not yet fully understood. Several hypotheses have been suggested, including increased bone resorption and a disordered control of renal phosphate handling and 1,25(OH)2 vitamin D production. Two children who had a Tp/GFR <2SD also had raised urinary calcium excretion and in one of these children PTH was undetected. A further child in this group, who also had idiopathic hypercalciuria, had a SD Tp/GFR of −1.29 and a PTH value of 1. Blood was not taken from the control population, so we cannot comment on renal phosphate handling in the five controls with raised calcium excretion.

One explanation of decreased phosphate reabsorption may be aminoglycoside induced nephrotoxicity which causes tubular damage of the convoluted and straight portions of the proximal tubule. Reabsorption of phosphate predominantly occurs in the proximal convoluted tubule. Although the number of patients with idiopathic hypercalciuria was only eight, there were proportionally more patients in this group who received aminoglycosides after a high trough concentration. We are not aware of any long term studies which suggest that renal tubular function may remain altered four to five years after the administration of aminoglycosides. Our findings may be an epiphenomenon and relate to the severity of illness in the neonatal period. However, there was no difference in other indices of illness severity, including birthweight, gestational age, days in oxygen and days requiring total parenteral nutrition.
Renal calcification in preterm infants

Concentrating capacity in both the study and control group combined was below normal values. The reduction in concentrating capacity is unlikely to cause any clinical problems. There are limitations to the assessment of renal length by ultrasound scanning. We attempted to minimise these by correlating renal length to height, using one observer who was blind to the group to which the child belonged. We compared renal length to standards published by Han and Babcock for American children, and used the same scanning methods. Out of 55 kidneys, eight were on and eight were below the lower limit of the 95% confidence interval when comparing renal length to the child’s height. There are many predisposing factors early in the neonatal period which may cause renal cortical or medullary necrosis. These include renal vein thrombosis, hypoxia, hypotension and nephrotoxic drugs. They are more likely explanations than renal calcification for the reduction in renal size and GFR present in some of our children. Lerner et al reported that renal cortical and medullary necrosis at post mortem examination was associated with congenital heart disease, prematurity, respiratory distress syndrome and bleeding diathesis.

Our results indicated that renal calcification in these children decreased with increasing age. This has also been reported in the study by Downing. This may explain why the differences in renal function seen in the neonatal period and early childhood, observed in other studies, and considered to be associated with renal calcification, are not present at the age of 4 years.

In summary, the results of this study indicate that children born preterm may have small kidneys and a degree of renal dysfunction. Renal calcification detected in the neonatal period does not seem to be a major predisposing factor to the renal excretion of calcium, concentrating ability, and renal size in these children. The clinical importance and aetiology of these abnormalities of renal function in our patients are unclear.

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