Nephrocalcinosis in preterm babies

A Narendra, M P White, H A Rolton, Z I Alloub, G Wilkinson, J H McColl, J Beattie

Abstract

Objectives—To determine prospectively the incidence and cause of nephrocalcinosis in preterm infants.

Study design—Inborn babies of gestation less than 32 weeks or birth weight less than 1500 g were eligible to be entered into a prospective observational study. Two renal ultrasound scans were performed, the first at 1 month postnatal age and the second at term or discharge. Data were collected on gestation, birth weight, sex, race, family history of renal calculi, oliguria on first day, respiratory support (ventilation, steroid, and oxygen dependency), and use of nephrotoxic drugs (gentamicin, vancomycin, and frusemide). Intake of fluid, calcium, and phosphate and plasma urea, creatinine, calcium, and phosphate were recorded for the first 6 weeks of life. Random urinary calcium/creatinine, oxalate/creatinine, and urate/creatinine ratios and tubular absorption of phosphate were measured once at term.

Results—A total of 101 preterm infants were studied. Twenty three (23%) had abnormal ultrasound scans. Sixteen (16%) had nephrocalcinosis. On univariate analysis, gestational age, male sex, duration of ventilation, oxygen dependency, duration and frequency of gentamicin treatment, toxic gentamicin/vancomycin levels, and postnatal dexamethasone were significantly associated with nephrocalcinosis. In addition, babies with nephrocalcinosis had a lower intake of fluid, calcium, and phosphate, longer duration of total parenteral nutrition, and higher urinary oxalate/creatine and urate/creatinine ratios than infants who did not have the condition. There was also a significant association with plasma urea and creatinine but not with plasma calcium or phosphate or urinary calcium. Multivariate analysis showed that the strongest predictors of nephrocalcinosis were duration of ventilation, toxic gentamicin/vancomycin levels, low fluid intake, and male sex.

Conclusion—16% of babies born at less than 32 weeks gestation developed nephrocalcinosis. The multifactorial origin, in particular, the association with extreme prematurity and severity of respiratory disease, is confirmed. In addition, an association with male sex, frequency and duration of gentamicin use, and high urinary oxalate and urate excretion is shown.

Keywords: nephrocalcinosis; renal dysfunction; calcium; kidney

Nephrocalcinosis in preterm babies was first described in 1982 by Hufnagel et al., who attributed the cause primarily to frusemide treatment. Since then studies published in the North American literature have shown a varying incidence of 28% to 64% in very low birthweight babies. In a previous study in the United Kingdom, the incidence of nephrocalcinosis was 21.5% and renal calcification (nephrocalcinosis and renal stones) 27%, and in a recent Scandinavian study an incidence of 20% was reported.

The role of frusemide in the development of nephrocalcinosis in preterm babies remains unclear. Jacinto et al. studied 31 babies of less than 1500 g birth weight and found a significant association between administration of frusemide and nephrocalcinosis. In a larger study of 79 babies of less than 32 weeks gestation by Short and Cooke, there was no difference in the mean total dose of frusemide given before detection of renal calcification between those with nephrocalcinosis and the unaffected group. These authors suggested that frusemide was used for infants already at risk of renal calcification and found duration of oxygen dependency to be the strongest predictor of renal calcification. In contrast, Saarela et al. found that the cumulative dose of frusemide in the first 8 weeks of life was significantly greater in the group that developed calcification.

Karlowicz et al. studied 50 babies of less than 1200 g birth weight and found white race and history of renal calculi, oliguria on first day, respiratory support, both of which have been shown to improve considerably the outcome of very low birthweight babies.

Nephrocalcinosis appears to resolve spontaneously in 40–50% of cases on follow-up, but can be associated with recurrent urinary tract infections, renal colic, and haematuria. More controversial is whether nephrocalcinosis leads to a reduction in glomerular filtration rate and tubular dysfunction or whether renal dysfunction on follow up reflects prematurity.

Our study was designed to establish prospectively the current incidence of nephrocalcinosis in a group of very preterm infants, to study the causes, and to provide a cohort for follow up.

Patients and methods

Only inborn babies of less than 32 weeks gestation or with a birth weight of less than...
1500 g from Queen Mother’s Hospital (QMH) and Southern General Hospital (SGH) Glasgow were recruited. Babies born with major congenital anomalies or those who died or were transferred before the end of the study period were excluded. The study protocol was approved by the local ethics committees, and informed written parental consent was obtained.

Data on the following were recorded: sex, gestation, birth weight, race, family history of renal calculi, oliguria on day 1, number of days ventilated, number of days in oxygen, oxygen dependency at 36 weeks postconceptional age, antenatal and postnatal use of steroids, administration of surfactant, phosphate supplementation, and treatment with frusemide, aminoglycoside (gentamicin), or vancomycin. High peak and trough levels of vancomycin and gentamicin (above normal laboratory reference) were also recorded. The actual daily intake of fluid, calcium, and phosphate was collected for the first 6 weeks of life.

Plasma concentrations of calcium, phosphate, urea, creatinine, and electrolytes were recorded daily during intensive care treatment and weekly thereafter for the first 6 weeks of life. Random urinary calcium/creatinine, urate/creatinine, and oxalate/creatinine ratios were measured once at term. Urine specimens were collected into a urine collecting bag or as clean catch specimens. Tubular reabsorption of phosphate was calculated once at term. Plasma electrolytes, including calcium and phosphate, were measured by standard dry chemistry methods with the Ortho Vitrous 750 analyser. Urinary urate, calcium, phosphate, and creatinine were measured with the Vitrous 250 analyser. Urine oxalate was measured enzymatically using the Sigma method modified for use on the Cobas Fara (Sigma UK).

Both neonatal units had a policy of providing oral phosphate supplementation. All enterally fed babies at QMH received 1.5 mmol phosphate/kg/day to maintain normal plasma phosphate. Babies from SGH received an appropriate dose of phosphate if plasma phosphate was less than 1.5 mmol/l. Total parenteral nutrition (TPN) solutions in both units provided 1 mmol phosphate/kg/day.

A renal ultrasound scan was performed at 1 month of age and once at term. The ultrasound examination was performed by one consultant paediatric radiologist at QMH (AGW) and by one consultant radiologist at SGH. When nephrocalcinosis was suspected in babies from SGH, the diagnosis was confirmed by a scan performed by AGW. Both radiologists undertaking the ultrasound examination were unaware of the past and current clinical details of the babies. The ultrasound examination was performed using either an Ultramark 4 or Acuson 128 scanner. With the Ultramark 4, a mechanical sector probe operating at either 10 MHz or 7.5 MHz was used, and, with the Acuson, either a 7.5 MHz linear array or 7.5 MHz sector probe was used. Nephrocalcinosis was diagnosed according to the criteria of Myracle et al.12

STATISTICAL METHODS

The groups of infants with and without nephrocalcinosis were compared using appropriate two sample techniques. Because of the considerable skewness shown by many of the data, the Mann-Whitney U test was performed to compare average levels of the continuous variables. The χ² test of association was used to analyse binary variables, augmented by Fisher’s Exact Test when expected frequencies were too small to justify the χ² approximation. Individual tests were taken to be significant when p < 0.05.

Variables that were significant on the univariate analysis were subsequently entered into a multivariate analysis, namely binary logistic regression, to discover whether a small number of explanatory variables may be particularly important for discrimination between babies in the two groups.

RESULTS

A total of 160 eligible babies were identified between October 1996 and May 1998. Twenty five died, and 27 were transferred to other hospitals. Five parents refused to give their consent, and two babies did not have the term ultrasound scan. This left 101 babies who completed the study.

ULTRASOUND FINDINGS

Twenty three babies had an abnormal ultrasound and 16 had nephrocalcinosis. Other abnormalities were hydronephrosis in four and bifid pelvis, ovarian cyst, necrotic papilla, and solitary kidney in one each. Two babies with hydronephrosis had antenatal pelvic dilatation. One baby with a postnatal diagnosis of unilateral hydronephrosis developed nephrocalcinosis in the contralateral kidney. The mean age of diagnosis of nephrocalcinosis was 10.4 weeks or 72.8 days (range 38–132 days). Nephrocalcinosis was detected in only four (20%) babies on the first ultrasound scan performed at 38–46 days of life, and this persisted at term. The nephrocalcinosis was bilateral in 10 and unilateral in six infants; all involved the left kidney. An approximate 95% confidence interval (CI) for the population rate of nephrocalcinosis was 8.7% to 23.0%.

BASIC DATA

Lower gestational age and male sex were significantly associated with an increased risk of nephrocalcinosis (table 1). Birth weight, growth retardation, and family history of renal calculi were not significantly associated. There were 10 babies of gestational age > 31–34 weeks and birth weight below 1500 g in the study, and none developed nephrocalcinosis. All 16 babies with renal calcification were white but this association was not significant (p = 0.234). Only seven babies in the study population were non-white. All babies with nephrocalcinosis were asymptomatic.

VENTILATION AND OXYGEN DEPENDENCY

All the babies with nephrocalcinosis were ventilated compared with 50 (59%) of those without (table 2). The median duration of ventilation was 7.5 days for babies with
Nephrocalcinosis in preterm babies

Table 1 Comparison of the basic data of the infants with or without nephrocalcinosis

<table>
<thead>
<tr>
<th></th>
<th>With (n=16)</th>
<th>Without (n=85)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gestation (weeks)</td>
<td>27 (24–31)</td>
<td>30 (24–34)</td>
<td>0.004*</td>
</tr>
<tr>
<td>Birth weight (g)</td>
<td>1170 (565–1880)</td>
<td>1270 (640–2720)</td>
<td>0.099*</td>
</tr>
<tr>
<td>Male/female ratio</td>
<td>13/3</td>
<td>41/44</td>
<td>0.015</td>
</tr>
<tr>
<td>SGA/AGA</td>
<td>2/14</td>
<td>24/61</td>
<td>0.187</td>
</tr>
<tr>
<td>F/H of renal stone (yes/no)</td>
<td>3/13</td>
<td>4/81</td>
<td>0.155</td>
</tr>
</tbody>
</table>

Values are median (range). *95% Confidence interval 49 to 390.

Table 2 Comparison of the respiratory support of the infants with or without nephrocalcinosis

<table>
<thead>
<tr>
<th></th>
<th>With (n=16)</th>
<th>Without (n=85)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ventilation</td>
<td>16 (100)</td>
<td>50 (59)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Median duration of ventilation (days)</td>
<td>7.5</td>
<td>1 &lt;0.0001</td>
<td></td>
</tr>
<tr>
<td>Reventilation, 2 or &gt;2 episodes</td>
<td>6 (37)</td>
<td>7 (8)</td>
<td>0.005</td>
</tr>
<tr>
<td>Oxygen (median days)</td>
<td>78.3</td>
<td>144</td>
<td>0.013</td>
</tr>
<tr>
<td>O, dependency at 36 wks PCA</td>
<td>11 (69)</td>
<td>25 (29)</td>
<td>0.003</td>
</tr>
</tbody>
</table>

Values in parentheses are percentages.

Table 3 Relation between medication and nephrocalcinosis (NC)

<table>
<thead>
<tr>
<th>Drug</th>
<th>NC (n=16)</th>
<th>Non NC (n=85)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Antenatal steroid</td>
<td>12 (75)</td>
<td>69 (81)</td>
<td>0.57</td>
</tr>
<tr>
<td>Dexamethasone</td>
<td>6 (37)</td>
<td>5 (6)</td>
<td>0.001</td>
</tr>
<tr>
<td>Surfactant</td>
<td>15 (94)</td>
<td>44 (52)</td>
<td>0.002</td>
</tr>
<tr>
<td>Frusemide</td>
<td>9 (56)</td>
<td>20 (23)</td>
<td>0.008</td>
</tr>
<tr>
<td>High nephrotoxic antibiotic level*</td>
<td>11 (69)</td>
<td>17 (20)</td>
<td>0.001</td>
</tr>
<tr>
<td>Phosphate</td>
<td>13 (81)</td>
<td>61 (72)</td>
<td>0.432</td>
</tr>
</tbody>
</table>

Values in parentheses are percentages. *High serum levels of gentamicin and/or vancomycin.

Nephrocalcinosis compared with one day for those without (95% CI –13 to –3). Six (37%) infants with nephrocalcinosis required reventilation twice or more compared with seven (8%) of those without. Median duration of continuous positive airway pressure was also significantly different between the two groups (95% CI 0 to 2). The median duration of oxygen dependency for babies with nephrocalcinosis was 75.5 days compared with 14 days for those without (95% CI –79.98 to –7.99). Eleven (69%) infants with nephrocalcinosis were oxygen dependent at 36 weeks postconceptional age compared with 25 (29%) of those without.

Medication

There was no association between antenatal treatment with steroids and nephrocalcinosis (table 3). Nephrocalcinosis, however, was significantly associated with the use of surfactant, postnatal dexamethasone, and frusemide. The median total dose of frusemide given to infants with nephrocalcinosis before detection of calcification on the term ultrasound scan was 16.1 mg (range 5—62), and in infants without nephrocalcinosis it was 24 mg (range 2—195.7), which is not significantly different (p = 0.75). None of the four infants in whom nephrocalcinosis was detected on the first scan received frusemide.

Eleven (69%) infants with nephrocalcinosis had high trough or peak serum levels of vancomycin or gentamicin compared with the standard laboratory reference range on at least one occasion compared with 17 (20%) of the infants without nephrocalcinosis (p = 0.001). This difference remained significant after correction for gestation by logistic regression.

Seventy three infants without nephrocalcinosis and 15 infants with nephrocalcinosis received gentamicin for a median of one and two episodes respectively (95% CI 0.0 to 1.0). The median duration of gentamicin treatment was five and 23 days respectively in the two groups (95% CI 3.0 to 20.0). On further analysis there was a significant association between the number of episodes and duration of gentamicin treatment and nephrocalcinosis (p = 0.0077 and p = 0.0002 respectively). Thirty infants without nephrocalcinosis and 11 with nephrocalcinosis received vancomycin for a median of one (range one to three) and one (range one to three) episodes respectively. The median duration of vancomycin treatment was 9.5 (range 2–70) and 22 (range 1–51) days respectively in the two groups. There was no significant difference in frequency and duration of vancomycin use between the groups (p = 0.35 and p = 0.41 respectively). The median high gentamicin trough level in the group with nephrocalcinosis was 2.6mg/l (range >2–6.0) and 2.8mg/l (>2–4.8) in the group without nephrocalcinosis (p = 0.99). No infant had high peak gentamicin level. The median high vancomycin trough was 15mg/l (range 13.7–29) in the nephrocalcinosis group and 16.5mg/l (range 12.9–22.8) in non-nephrocalcinosis group (p = 0.88). The median high peak serum level of vancomycin was 54.5mg/l (range 41.1–100) in the nephrocalcinosis group compared with 43.15mg/l (range 41.2–44) in the non-nephrocalcinosis group (p = 0.059). Eighty eight babies received gentamicin, and, of these, 41 received vancomycin, 24 concurrently and 17 consecutively.

Nо baby had vancomycin who had not received gentamicin. Receiving vancomycin in addition to gentamicin significantly increased the risk of nephrocalcinosis (p = 0.0225), but the risk was no greater when vancomycin was given concurrently with vancomycin (p = 0.486).

Nutrition

The median duration of TPN in infants with and without nephrocalcinosis was 18 and seven days respectively (p = 0.0009). The median calcium intake was lower in the group with nephrocalcinosis compared with the group without nephrocalcinosis (p = 0.0225), but the risk was no greater when vancomycin was given concurrently with vancomycin (p = 0.486).

Serum Biochemistry

Nephrocalcinosis was significantly associated with higher median levels of serum creatinine during weeks 3–5 (fig 2B) and higher median levels of serum urea in weeks 1–5 (fig 2A). There was no significant difference in median levels of plasma calcium or phosphate between babies with nephrocalcinosis and those without (fig 2C,D).

Urinary Metabolite Concentrations

The median urinary oxalate/creatinine and urate/creatinine ratios were significantly higher...
in babies with nephrocalcinosis (table 4). There was no significant difference in median urinary calcium/creatinine ratio or tubular reabsorption of phosphate between the groups. Only one infant (without nephrocalcinosis) had oliguria on day 1.

**UNIVARIATE ANALYSIS**

The most significant of all the above variables was duration of ventilation ($p = 0.0001$).

**MULTIVARIATE ANALYSIS**

This was performed by stepwise fitting of a binary logistic regression test model. The presence/absence of nephrocalcinosis was the dependent variable and the predictor variables were all those that had produced a significant result in the univariate analysis. The strongest indicators of nephrocalcinosis (in order) were: number of days ventilated ($p = 0.0001$); toxic gentamicin/vancomycin levels ($p = 0.0059$); fluid intake in week 3 ($p = 0.0059$); male sex ($p = 0.023$). These four variables correctly allocated 80.6% of the sample overall to the nephrocalcinosis or non-nephrocalcinosis group.

**Discussion**

The incidence of nephrocalcinosis (16%) in this study is lower than previously reported,\(^2\)\(^-\)\(^5\), which may reflect improvements in neonatal intensive care, in particular the antenatal use of steroids and surfactant and improved nutrition. Our policy of ensuring adequate oral phosphate supplementation may also have helped to decrease the incidence. However, we note that the wide interval estimate for this
population includes most of the values reported in previous studies and therefore our incidence may not be a true reduction.

We think it unlikely that we underdiagnosed nephrocalcinosis because ultrasound is a very sensitive diagnostic tool. Cramer et al reported a sensitivity of 96% and specificity of 85% using ultrasound compared with computed tomography and postmortem histological diagnosis in a rabbit model. We used strict criteria and a single experienced paediatric radiologist to diagnose/confirm the presence of nephrocalcinosis.

In agreement with other studies, we found nephrocalcinosis to be associated with low gestational age but not with birth weight. For the first time to our knowledge, we found male sex to be significantly associated with nephrocalcinosis but did not find an association with a family history of renal calculi or white race as reported by Karlowicz et al, although the small number of non-white infants (n = 7) in this study limited any statistical evaluation.

**Respiratory Disease**

Short and Cooke found duration of oxygen to be the strongest clinical indicator of renal calcification. We confirm that severe respiratory disease, as indicated by duration of ventilation, reintubation episodes, duration of oxygen, and oxygen dependence at 36 weeks after conception, is significantly associated. All babies in our study with nephrocalcinosis were ventilated, and duration of ventilation was the most significant variable in the univariate and multivariate analyses. Preterm infants with lung disease are reported to have decreased urinary citrate, which may predispose them to nephrocalcinosis because citrate is a known inhibitor of renal calcification in adults and children.

**Medication**

We found an association between nephrocalcinosis and postnatal dexamethasone in agreement with the findings of Saarela et al. Although corticosteroids may cause negative calcium balance, osteopenia, hypercalciuria, and nephrocalcinosis in children, few data are available on the effect of dexamethasone on urinary calcium excretion in preterm infants. Sonntag and Gaude found no increase in urinary calcium excretion in a group of preterm infants who received dexamethasone compared with a control group. Postnatal dexamethasone and surfactant may not be independent variables, but merely reflect the severity of the respiratory disease and low gestation.

**NUTRITIONAL FACTORS**

In this study, TPN was significantly associated with nephrocalcinosis, and we confirm the work of Short and Cooke showing that nephrocalcinosis is significantly associated with low intake of calcium, phosphate, and fluid and raised serum creatinine and urea. Phosphate deficiency is typically associated with hypercalciuria, whereas phosphate supplementation decreases urinary calcium excretion. Excess phosphate administration may result in renal rather than skeletal deposition of calcium. However, unlike Saarela et al, we did not find phosphate supplementation to be significantly associated with nephrocalcinosis. In our study,
although we tried to avoid phosphorus deficiency, babies with nephrocalcinosis still had a lower phosphate intake, and this was related mainly to the longer duration of TPN. The TPN solution that we used provided 1 mmol phosphate/kg/day. There are concerns about the solubility of higher concentrations of phosphate in TPN solutions, although a concentration of 1.34 mmol/kg/day has been associated with lower urinary calcium excretion.\(^{33}\)

**URINE BIOCHEMISTRY**

We found a significant association between nephrocalcinosis and urinary oxalate/creatinine and urate/creatinine ratios measured at term. None of our babies were receiving TPN when urinary oxalate and urate were being determined. The role of increased urate and oxalate in the development of renal calcification in children and adults has been well documented,\(^{32,33}\) but to our knowledge, this has not previously been shown in preterm babies in a prospective study. Calcium oxalate has been shown in a postmortem study to be the most common type of renal calcification in infants dying after intensive care.\(^{34}\) The oxalate/creatinine ratios found in our study are in broad agreement with the high levels quoted in the literature for preterm babies, who are known to have higher oxalate/creatinine ratios than those born at term.\(^{35}\) Preterm and healthy term babies have substantially higher oxalate/creatinine ratios than adults, reaching a peak at 3–4 weeks and declining thereafter.\(^{36,37}\)

We measured urinary calcium once at term, during the first month of life, and respiratory acid levels in premature babies remained increased in premature infants. Berard et al\(^ {38} \) has shown that urinary acid levels in premature babies remained significantly higher than in normal term infants during the first month of life, and respiratory failure resulted in a further increase. In our study, the urinary urate/creatinine ratios were higher oxalate/creatinine ratios than those born at term.\(^ {39}\) Preterm and healthy term babies have substantially higher oxalate/creatinine ratios at term, mainly to the longer duration of TPN. The TPN solution that we used provided 1 mmol phosphate/kg/day. There are concerns about the solubility of higher concentrations of phosphate in TPN solutions, although a concentration of 1.34 mmol/kg/day has been associated with lower urinary calcium excretion.\(^ {33}\)

We are grateful for the contributions of Dr D A R Robertson (renal ultrasound examination) and Dr R Logan (urinary biochemistry). We thank the Medical and Nursing staff of the neonatal intensive care units of The Queen Mother’s Hospital and Southern General Hospital, Glasgow.


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