Fatal perinatal nephropathy with onset in intrauterine life

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Abstract
A girl, born at 29 weeks' gestation, died of renal failure aged 16 days. Postmortem histology showed diffuse mesangial sclerosis with failure of development of the cortex. This is an unusual cause of neonatal renal failure, and it demonstrates the effect of disease arising in utero and influencing the development of the kidney.

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Renal failure in neonates is rarely due to primary renal disease. We present a case of a neonate, born at 29 weeks' gestation, in whom renal failure led to death within 16 days. The pathological findings demonstrate an effect of the disease process on the maturation of the kidney.

Case report
The mother was a 28 year old white primigravida. She was negative for syphilis antibodies and had no history of renal or autoimmune disease or of hypertension before pregnancy. The parents were unrelated; the father had sarcoidosis. Serum α-fetoprotein at week 18 was raised at 3·2×median for the gestation. Fetal ultrasound was normal.

Severe hypertension developed during the last four days before delivery, and an emergency caesarean section was performed at 29 weeks after an antepartum haemorrhage.

The infant girl weighed 1160 g and required resuscitation at birth. Her Apgar scores were 2 at 1 minute and 4 at 5 minutes; she was also ventilated.

Respiratory distress was present for the first three days and was complicated by a pneumothorax. She was treated with pancerumion, surfactant, pipercillin, and gentamicin. Her blood pressure was well maintained. Although the infant was mildly oedematous with an albumin concentration of 27 g/l, she passed urine at 5 ml/kg/hour for the first 72 hours. Urinalysis was negative and the specific gravity was below 1012. An umbilical catheter was removed at 24 hours because of left leg mottling.

At 96 hours proteinuria and haematuria were detected, but proteinuria was transient and not quantified. Her urea concentration rose to 9 mmol/l from 4·8 mmol/l at birth; creatinine was static from birth at 109 mmol/l until day 7 when it increased progressively. At that time the infant became oliguric and hypertensive and gained weight rapidly. Cardiac and renal ultrasound were normal. The baby rapidly became anuric, and peritoneal dialysis was commenced. The baby died at 16 days of respiratory failure.

Chromosome analysis was not performed and specific genetic counselling was not undertaken. A normal female sibling was born in January 1992.

POSTMORTEM EXAMINATION
A limited examination was carried out and the infant was found to be normally formed, weight 1240 g, on the 10th centile; she had normal genitalia. Internal examination showed peritoneal oedema, bile stained ascites, a congested liver, small adrenals, and two small gastric ulcers. The kidneys were enlarged, pale, and oedematous and their combined weight was 21·3 g (normal combined weight = 12·9 (±3·3–9) g for a 1250 g fetus). There were no cysts. The ureters, bladder, and urethra were normal. The bladder was empty.

The placenta (weight 215 g) was not oedematous and showed no evidence of abruption.

MICROSCOPY OF KIDNEY (FIG 1 AND 2A)
Despite the large size of the kidneys there were only 4–5 layers of glomeruli. The nephrogenic blastema was disordered and reduced in thickness. The superficial glomeruli were small and immature, with prominent primitive epithelial cells. The deepest glomeruli had developed appropriately, but they were enlarged and showed mesangial sclerosis. There was no increase in cellularity and no crescent formation. The tubules were dilated by a mass of amorphous material that stained positive with periodic acid Schiff; they were not dysplastic and there was no evidence of acute tubular necrosis. This tubular dilation appears to be the cause of the increased renal weight.

Immunostaining showed peripheral IgM and C3 involving all the glomeruli, including immature glomeruli that could barely have acquired function.

ELECTRON MICROSCOPY (FIG 2B)
In the larger glomeruli the epithelial cells were separated from the basement membrane, with some floating free in Bowman's space. Beneath these cells was a mass of amorphous material, which was not electron dense, and therefore did not represent an immune complex deposit. The basement membrane was tortuous, with
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Fatal nephropathy with onset in this unusual utero. The blastema appearance of amorphous cells (uranyl acetate lead citrate: 1800).

Discussion

The interest of this case lies both in the unusual clinical presentation and the pathological findings. In our opinion the histological appearance indicates that the renal failure in this case was a disease process commencing in utero. At 29 weeks’ gestation there are normally 9–10 layers of mature glomeruli and the nephrogenic blastema is prominent (fig 1B). In our case (fig 1A) there were 4–5 layers of glomeruli, many of which were immature, which corresponds to the stage of development at about 20–22 weeks, and the blastema was thin and disordered. The appearance on electron microscopy (fig 2B) indicates that those glomeruli that were capable of function were suffering from hyperfiltration injury, with trapping of large molecules (including the IgM and C3 seen on immunostaining) in a leaking glomerulus. It appears that a pathological process, probably a specific glomerular disease, had curtailed the formation of new glomeruli during the last seven weeks in utero.

Although there are many possible causes of renal failure in preterm neonates, in this case the insidious onset and relentless progression are unusual features. Possible acquired causes in a preterm infant are antenatal hypoxia due to pre-eclampsia or abruptio placentae, intra-partum asphyxia, polycthæmia, emboli from a catheter, or gentamicin or pancuronium toxicity. However these would all cause a more acute onset of renal failure. More importantly, no disease acquired during or after birth could have had this profound effect on development.

Many congenital causes of renal insufficiency can be excluded on morphological grounds. These include malformations of the lower urinary tract, Finnish type congenital nephrotic syndrome, polycystic disease, and renal dysplasia. The lesion in the glomeruli bears some resemblance to that seen in congenital syphilis but this can be excluded on maternal serology, and by the absence of immune deposits.

The renal lesion most closely fits into the category of idiopathic diffuse mesangial sclerosis (DMS). This is a poorly characterised condition, probably of heterogenous aetiology, and can be familial. It is rare in infancy and is associated with a less rapidly fatal course than in our case. It has not been described in appreciably preterm infants. However the glomerulosclerotic lesions are similar to ours, and the infants with DMS who present early and survive for a few months do show an excess of immature glomeruli. In one 15 month infant there was also peripheral IgM and C3 staining in the glomeruli, without electron dense deposits on electron microscopy. The precise pathogenetic mechanism remains obscure, in our case as in other cases of DMS.

We believe that this is an example of the influence on development of disease acquired during fetal life. From a practical point of view it emphasises the importance of reaching a pathological diagnosis in cases of neonatal renal failure.

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