Hypertension and segmental renal hypoplasia causing a syndrome of haemolysis and uraemia

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SUMMARY An 11-year-old girl presented with acute renal failure and severe hypertension. The blood film showed thrombocytopenia, numerous fragmented red blood cells, and a reticulocyte count of 10%. An intravenous pyelogram showed a small contracted left kidney, and plasma renin activity was increased in the left renal vein. Treatment with minoxidil and propranolol controlled the hypertension. After nephrectomy the hypertension resolved. Light microscopic examination of the left kidney showed a segmental renal hypoplasia. Malignant arterial hypertension can provoke a syndrome of haemolysis and uraemia in children. Aggressive lowering of blood pressure leads to an improvement in renal function.

Severe hypertension in haemolytic uremic syndrome is common, and is often a complication of the renal disease. However, in adults, haemolysis and uraemia are present in some cases of malignant hypertension.1 This case report shows that this can be so in children too.

Case report

An 11-year-old girl presented with a history of polyuria, headaches, nausea, and vomiting for one year. Blood pressure (BP) had not been measured. Gross haematuria, pallor, and purpura had developed 3 days before admission. The day she was admitted BP was 250/130 mmHg. The liver was enlarged and there was bilateral oedema on the legs. Haematocrit was 23%, haemoglobin 5 g/dl, white blood count 12.9 × 10⁹/l, platelets 10.0 × 10⁹/l. The blood film showed numerous fragmented red blood cells, and the reticulocyte count was 10%. Urine analysis showed protein (++++) and numerous red blood cells. Other values were as follows: blood urea 60 mg/100 ml (9.9 mmol/l) on admission rising to 185 mg/100 ml (31 mmol/l). Plasma creatinine level was 3 mg/100 ml (265 µmol/l), fibrinogen degradation products were between 25 and 40 mg/ml. The C3 fraction of complement was 135 mg/100 ml and the C4 was 49 mg/100 ml. Plasma renin activity, measured after the administration of 40 mg furosemide and 20 mg hydralazine, was found to be 708 ng/ml per hour. The normal value at this age, in the supine position and on a normal diet, is 3 ± 2 ng/ml per hour in our laboratory. During the next 3 weeks the child received 80 mg hydralazine (4 mg/kg) and 200 mg acetobutolone (10 mg/kg) each day, and intravenous heparin at a dose of 5 mg/kg a day, for 7 days only. After this treatment BP, blood urea, reticulocyte count, and platelet count returned to normal. However, after a few days the blood pressure again rose to 160/130 mmHg, despite continuous antihypertensive treatment. One intravenous injection of 200 mg diazoxide was given every day for 8 days. Treatment with minoxidil was started at a dose of 1 mg/kg a day with propranolol (40 mg daily) and furosemide (40 mg daily). After 3 days the BP had fallen to a normal level. The treatment was maintained at the same dose for 3 months. An intravenous pyelogram showed a small, contracted left kidney. The cystourethrogram was normal, without evidence of reflux. Left renal arteriography showed no evidence of stenosis, but
there was poor filling of vessels in the lower pole of the kidney. During treatment with hydralazine and acetobutolone, venous samples from right and left renal veins and from the inferior vena cava, below and above the renal veins, were taken. The plasma renin activity of venous blood from the inferior vena cava, below and above the renal veins was 4·8 and 7·4 ng/ml per hour. In the left renal vein plasma renin activity was 14·6 ng/ml per hour, while it was 4·6 ng/ml per hour in the right. A right kidney biopsy was performed. On light microscopical examination focal and segmental glomerulosclerosis without pronounced vessel lesions could be seen. Because of the high level of plasma renin activity in the left renal vein, a left nephrectomy was performed. A small contracted kidney weighing 20 g and measuring $6 \times 2 \times 1$ cm was removed. Light microscopical examination showed segmental areas of severe cortical and medullary abnormality. The affected segments contained a few sclerosed glomeruli, many atrophic and dilated tubules, and thickened blood vessels (Figure). After nephrectomy the hypertension resolved and the plasma renin level fell to 1 ng/ml per hour.

**Discussion**

The association between segmental renal hypoplasia (Ask-Upmark kidney) and hypertension has been known for many years. The pathogenesis of the hypertension in this disease is not perfectly understood. An abnormal renin secretion has been suggested in the course of this disease. Although the measurements were made after treatment had started in our patient, the renin level was three times the normal rate on the side of the diseased kidney. After nephrectomy the hypertension resolved and plasma renin activity fell to below the normal value. However, as is usual in the hypoplastic segment, no juxtaglomerular cells could be seen, thus raising the question of the origin of the secreted renin. Malignant arterial hypertension can provoke a syndrome of haemolysis and uraemia in adults, but this has not been reported in children. It has been suggested that an increased BP would result in accumulation of fibrinogen and in the initiation of coagulation in the wall of the arteries. The fibrinoid vascular lesion might induce mechanical damage of the red blood cells and provoke a microangiopathic haemolytic anaemia. Furthermore, reversible acute renal failure can be the consequence of malignant hypertension.

Minoxidil is a new vasodilator, and studies show that it is more effective than hydralazine. In this patient minoxidil combined with propanolol controlled hypertension. There was sodium and water retention and severe hypertrichosis. In a child with unilateral kidney disease, nephrectomy is undoubtedly the treatment of choice.

**Conclusion**

Renal segmental hypoplasia may be the cause of increased hypertension associated with haemolytic uraemic syndrome. Although treatment with minoxidil was effective for this type of hypertension, nephrectomy was performed because of side effects.

**References**


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