—Case Reports—

Urinary Tract Malformation and Infection with Hyperkalemia and Decreased Fractional Excretion of Potassium in an Infant

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Abstract

We describe an uncircumcised male infant treated for urinary tract infection who exhibited multiple hormonal and electrolyte abnormalities consistent with a diagnosis of transient pseudohypoaldosteronism. Successful treatment of the urinary tract infection was accompanied by the resolution of all hormonal and electrolyte abnormalities, including hyperaldosteronemia, hyperreninemia, hypernatremia and hyperkalemia. Radiographic examination revealed marked left dilatation of the renal pelvis and hydrourereter but no vesicoureteral reflux. Owing to the possibilities of future renal scarring, decreased renal function, and hypertension, evaluation of urinary tract malformation and appropriate hormonal studies should be performed in infants with urinary tract infection and hyperkalemia.

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Key words: Pseudohypoaldosteronism, infant, urinary tract infection, urinary tract malformation, hypokalemia

Introduction

Numerous recent cases of renal scarring and reflux on follow-up imaging demonstrate that infant urinary tract infection is not a benign condition\textsuperscript{1}. Urinary tract infection in early life may also lead to compromised renal function and possibly to hypertension later in life\textsuperscript{2}. In newborns with obstructive uropathy, a syndrome has been described in which infant urinary tract infection is accompanied by elevated serum levels of aldosterone, hypernatremia and hyperkalemia and in which hormonal and electrolyte abnormalities are reversed after the infection is treated\textsuperscript{2}. Here, we report on a 7-month-old male infant who exhibited pseudohypoaldosteronism with hydronephrosis and infection.

Case Histories

A male infant was admitted to our hospital at 7 months of age after 7 days of persistent vomiting despite medical treatment, including oral antibiotics. Physical examination revealed an uncircumcised penis, dehydration, and a body temperature of

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38.0°C. Laboratory tests showed hyponatremia and hyperkalemia (Table 1). Urinalysis indicated urinary tract infection, but culture of urine failed to identify a pathogen. Endocrine tests revealed elevated serum aldosterone levels (2.827.6 ng/dl) and plasma renin activity (40.8 ng/ml/hour). The infant was treated intravenously with normal saline solution and antibiotics. Subsequently, serum levels of sodium and potassium returned to the normal ranges and findings of urinalysis became normal (Table 1). Radiographic examination revealed left dilatation of the renal pelvis and hydrourerter (Fig. 1), but no vesicoureteral reflex (VUR) was detected. Antibiotic prophylaxis was started at 8 months of age. At age 1 year, the infant underwent surgery for ureteral re-implantation.

**Discussion**

Urinary tract infection in infants causes renal scarring and can compromise future renal function. Infants with urinary tract infection are also reported to have elevated serum levels of aldosterone, hyponatremia, and hyperkalemia. Here, we have reported on a 7-month-old male infant who presented with pseudohypoaldosteronism. We could not isolate a specific pathogen from urine culture, and although he had taken antibiotics for several days before admission to our hospital, the urine still showed numerous bacteria. We thus believed that he had a urinary tract infection, and because he also had hydronephrosis, we considered this to be a typical case of pseudohypoaldosteronism.

Our review of 60 cases of secondary pseudohypoaldosteronism previously reported shows that urinary tract infection and urinary tract anomalies are additional factors for the development of secondary pseudohypoaldosteronism, with at least one required for its development. Temporary aldosterone resistance in infants decreases the distal secretion of H⁺ and K⁺, the distal absorption of Na⁺, and the activity of Na-K adenosine triphosphatase. This resistance to aldosterone is increased by urinary tract infection because interstitial inflammation or a bacterial toxin, either primarily or secondarily through the synthesis and release of inflammatory mediators, might cause tubular unresponsiveness to aldosterone.
On the other hand, urinary tract obstruction might increase intrarenal pressure and the intrarenal synthesis of cytokines, and the increased intrarenal pressure and cytokine production might be responsible for the renal tubular dysfunction. Also the obstruction would lead to a reduction in renal blood flow and involve the complex renin-angiotensin mechanism, resulting in elevated aldosterone levels.

In conclusion, urinary tract infection and urinary tract malformation can lead to an acute metabolic imbalance in infants with urinary tract malformation. Tract malformation should be suspected in an infant with hyperkalemia and urinary tract infection.

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References


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