Rare and Treatable Case of Hypertension in Adolescent

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Abstract
Renal segmental hypoplasia or Ask-Upmark kidney is a rare congenital kidney disorder that can be associated with hypertension.

A 12-year-old girl previously healthy was referred to our emergency department with anxiety crisis and high blood pressure. The physical examination was unremarkable except for a blood pressure extremely high (191/141 mmHg). Echocardiography revealed left ventricular hypertrophy. Renal ultrasound showed a renal asymmetry with a small left kidney. She was treated with three antihypertensive agents without blood pressure normalization. Further investigation included doppler ultrasound and computed tomography angiography which revealed atrophic left kidney and a narrow but permeable renal left artery. No radionuclide uptake was observed in captopril renogram. Patient underwent a left nephrectomy with marked improvement in blood pressure control. Macroscopic appearance of the atrophic kidney showed segmental hypoplastic area with absence of glomeruli, atrophic tubules and thick walled arteries at histological findings with associated hypoplasia of the renal artery.

Hypertension can be silent and screening children and adolescents for elevated blood pressure (BP) or hypertension has the potential to shift the management of hypertension to younger age groups and potentially reduce future cardiovascular disease risk in adults[3].

We report an unusual case of secondary hypertension, renal segmental hypoplasia, in a previously apparent healthy adolescent girl.

Case Report
A 12-year-old girl, caucasian, was referred to the pediatric department with hypertensive crisis. She had a past medical history without problems with negative prenatal diagnosis of congenital anomalies of the kidney and urinary tract. She had prior history of systolic blood pressure with pre-hypertension range in medical office which was interpreted as white coat hypertension. She never had urinary tract infections or other significant diseases. Her family history was significant for middle age onset of primary hypertension in her father.
She went to a general practice office for routine vaccination and because of anxiety, an unexpectedly high blood pressure was measured (190/141 mmHg – (> Pc 99 + 5 mmHg) - stage II hypertension) and she was admitted to the emergency department. On physical examination she maintained high blood pressure without other clinical symptoms (no associated organ damage). Neurological examination and ophthalmic fundus examination were normal. Her complete blood count, basic metabolic panel, renal function and urinalysis were unremarkable. Renal ultrasound showed a renal asymmetry with a small left kidney (4 cm) with increased echogenicity and a normal right kidney (12 cm). Because of hypertensive crisis present as hypertensive urgency she was discharged from the hospital with three antihypertensive agents (nifedipine, carvedilol and enalapril) and she was referred to hypertension consultation of a tertiary hospital.

At outpatient consultation investigations revealed she maintains high blood pressure (> Pc 95) with ambulatory blood pressure monitoring (during twenty-four hours). She repeated laboratory test at outpatient consultation with little increased plasma rennin levels (39,1 pg/ml; normal values 5,1-38,7 pg/ml). Doppler renal ultrasound (figure 1) and computed tomography angiography revealed atrophic left kidney and a narrow but permeable renal left artery (figure 2 and figure 3). Captopril renograme showed no radionuclide uptake in the left kidney (figure 4).

Echocardiography revealed left ventricular hypertrophy with good function and without structural heart disease. Investigations revealed a small, non-functioning left kidney with hypoplastic left renal artery and hypertension. A left nephrectomy was indicated.

**Macroscopic examination**

The excised left kidney was small and atrophic measuring 4,8*2,5*1,8 cm; it was attached to a hypoplastic renal artery. On sectioning the kidney showed reduction on cortical measurement, distinction between cortex and medulla was ill defined and there was a dilatation of the pelvicalyceal system.

**Microscopic examination**

The renal parenchyma appeared glomerulosclerosis and areas without glomeruli, the renal tubules are atrophic with thyroidisation and thick walled arteries (figure 5). The histological finding of renal left artery was normal. With five months of follow up she remained normotensive (119/75 mmHg; < Pc 90) with monotherapy (nifedipine).

![Figure 1: Doppler renal ultrasound revealed atrophic left kidney (5 cm) and a small renal left artery with renal left parenchyma hypoperfusion.](image1)

![Figure 2: Computed abdominal tomography angiography showed caliber reduction small of renal left artery with permeability maintained. Atrophic left kidney (4 cm) and rewarding hypertrophy right kidney (12 cm) are revealed.](image2)
Hypertension in Adolescent

Figure 3: Three dimensional reconstruction of computed tomography angiography revealed atrophic left kidney with no contrast enhancement.

Figure 4: Captopril renograme without radionuclide uptake of the left kidney.

Figure 5: Histological examination of excised left kidney described atrophic renal tubules with thyroidisation with absence or sclerosis of glomeruli.

Discussion

One of the most important components of the successful management of childhood with hypertension is determining whether or not there is an underlying cause that is amenable to treatment. Identifying children with this condition and successfully treating their hypertension should have an important impact on long-term outcomes of CVD.[6]

Screening children and adolescents for elevated blood pressure could identify hypertension at an early stage where interventions could be initiated, potentially decreasing the rate of progression of hypertension from childhood to adulthood and reducing the clinical consequences of hypertension in adulthood.[6]

The European Society of Hypertension recommend that children 3 years of age and older have their BP measured during every healthy care visit. Several steps should be followed, from screening to confirmation, to rule out secondary causes of hypertension if indicated[7]. Secondary hypertension is more common in children compared with adults. Young age, severe and uncontrolled hypertension, family history, abnormal renal function, proteinuria, high plasma rennin, signs and symptoms of syndromes associated with hypertension are suggestive of an underlying secondary cause of elevated BP. Disorders of the renal parenchyma and renal vessels account for nearly 78% of the secondary hypertension in the pediatric population[6].

Renal segmental hypoplasia or Ask-Upmark kidney has first been described in 8 patients in 1929[9] Till date, approximately 200 cases have been reported[10]. This condition is seen more frequently in females below the age of 12 years[11]. Histological, the affect kidney appears small and contains segments with decreased number of pyramids, atrophic or thyroid like tubules, no glomeruli and thick walled arteries[11,12].

Ask-Upmark kidney need to be differentiated from chronic pyelonephritis or ectopic segmentation of vascular origin. The renal artery is small and that is not secondary to vascular changes since the walls are always normal. The segments affected don’t have the fan-like distribution characteristics of chronic pyelonephritis and renal infarctions[11]. Radionuclide renal scan combined with an angiotensin-converting enzyme (ACE) inhibitor, referred to as captopril renography, may be useful in some patients, like our case. This renograme shows a decrease in the glomerular filtration rate and renal blood flow on the affected side after ACE inhibition[13].

Although the condition is unilateral like the case reported, bilaterally asymmetrical segmental hypoplasia has also been described[10]. The etiology of this renal lesion is rather controversial. Some authors propose a congenital malformation in the metanephric development, whereas others attribute the principal cause to vesicoureteric reflux (VUR) with intrarenal reflux and reflux associated pyelonephritis[13-16]. Our patient had no previous history of VUR or urinary infections but it has been argued that the absence of reflux at the time of diagnosis does not rule it out as the initial cause of the atrophic kidney[15,16].

Presenting symptoms are hypertension, haematuria, loin pain and headache in isolation or associated with hypertensive encephalopathy[16-17]. Most often Ask Upmark kidney presents as severe hypertension in pediatric or adolescent patients[17]. Nephrectomy of the affected usually normalizes the blood pressure[11]. The cause of hypertension in Ask-Upmark kidney seems related to an increased rennin levels. Normalization of BP fol-
Hypertension in Adolescent

 Following nephrectomy also suggests that rennin excess was the cause of hypertension[10-15].

 In our patient, the BP not normalized after the nephrectomy but she has the BP controlled with monotherapy and with low doses instead three antihypertensive agents as she had before. We have only five months of follow up and maybe the BP tends to normalize.

 She had prior high systolic BP which was interpreted as white coat hypertension, but she had not make an ambulatory blood pressure monitoring to confirm the diagnosis. Furthermore echocardiography showed signs of pre-existing blood pressure. This case reinforces the importance of measuring the BP in children and adolescents because she had no symptoms of hypertension and she had already target-organ damage. Owing the white coat phenomena, ambulatory blood pressure monitoring is indispensable. Hypertension-induced preclinical target-organ damage is not uncommon in children and should be evaluated in all hypertensive children[18].

 Conclusion

 This case represents a form of hypertension potentially curable which the diagnosis is crucial. In a child with elevated blood pressure and high plasma rennin levels without renal artery stenosis the possibility of an Ask-Upmark syndrome should be considered. Finally target organ damage can occur with asymptomatic hypertension. Hypertension can be silent and it’s important to measure BP in all office visits and all changes in the blood pressure profile should be valued and the ambulatory blood pressure monitoring should be obtained.

 Conflict of Interest: The authors declare no conflict of interests.

 References