**Unilateral Adrenal Hyperplasia: A Rare Cause of Primary Aldosteronism**

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**ABSTRACT**

A 73-year-old gentleman with a background history of long-standing hypertension and hyperlipidaemia presented with one-year history of persistent asymptomatic hypokalaemia associated with metabolic alkalosis. Within a few months, the hypertension worsened, needing increasing dose of anti-hypertensive agents. Biochemical test showed an elevated aldosterone-renin ratio but the computerized tomography (CT) scan of adrenal did not show any adrenal mass. Arterial stimulation venous sampling (ASVS) lateralized the lesion to the left side. A left retroperitoneal adrenalectomy was performed with immediate reversal of hypokalaemia and improvement of blood pressure control to only single anti-hypertensive agent.

**KEYWORDS:** primary aldosteronism, adrenal venous sampling, adrenal hyperplasia, hypertension

**INTRODUCTION**

Primary aldosteronism (PA) is characterised by aldosterone hypersecretion and represents a common cause of secondary hypertension. It accounts for 5-10% of hypertensive patients.\(^1\) The incidence is underreported perhaps due to difficulty in the screening tool, as the Aldosterone-Renin ratio (ARR) has a lot of heterogeneity in terms of result interpretation and laboratory protocol. During diagnostic evaluation, it is important to determine the aetiology of PA since the treatment of unilateral and bilateral disease differs significantly. In patients with proven unilateral overproduction of aldosterone, unilateral adrenalectomy is curative and preferred over medical therapy.

**CASE REPORT**

A 73-year-old man had a ten-years history of hypertension and hyperlipidaemia, requiring only single anti-hypertensive and lipid lowering agent with good control of blood pressure. He presented a year ago with persistent significant hypokalaemia of 2.4 mmol/L (3.5 - 5.1 mmol/L) and metabolic alkalosis. It was incidentally detected during blood screening prior to cardiac stress test for moderate risk stratification. Subsequently, he developed resistant hypertension, needing three anti-hypertensive agents. He was screened for plasma aldosterone and renin activity which showed elevated plasma aldosterone of 669.5 pg/mL with low plasma renin activity of less than 0.2 ng/mL/hr. The Aldosterone-Renin ratio (ARR) was high (> 334), suggestive of primary aldosteronism. In view of the spontaneous hypokalaemia with significant high plasma aldosterone > 200pg/mL and supressed renin, confirmatory test was not needed. Computed tomography (CT) of the adrenal failed to show any obvious lesion and this was proceeded with venous sampling. The result is shown in Table 1 which lateralised the lesion to the left adrenal gland.

Despite the CT scan not showing any nodules, in view of convincing lateralization test, a left retroperitoneal adrenalectomy was performed uneventfully.
Post-operatively, the blood pressure became better controlled with only single anti-hypertensive agent and potassium reverted to normal; 4.5 mmol/l (3.5 - 5.1 mmol/L) without any supplement or potassium sparing anti-hypertensive agent. Histopathology report showed marked expansion of zona fasciculata and reticularis, consistent with a unilateral adrenocortical hyperplasia (Figures 1 & 2). A repeat test post-operatively showed serum aldosterone of 59.45 pg/ml and plasma renin of 0.326 ng/mL/hr giving ARR of 18.2.

<table>
<thead>
<tr>
<th></th>
<th>Cortisol (Adrenal vein: peripheral)</th>
<th>Aldosterone (Adrenal vein: peripheral)</th>
<th>Aldosterone: Cortisol ratio</th>
<th>Lateralization index (Aldosterone Cortisol ratio adrenal vein: Peripheral)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Left adrenal vein</td>
<td>420</td>
<td>2511</td>
<td>1.72</td>
<td>5.97</td>
</tr>
<tr>
<td>Right adrenal vein</td>
<td>474</td>
<td>241</td>
<td>2.75</td>
<td>0.51</td>
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<tr>
<td>Left peripheral</td>
<td>244</td>
<td>306</td>
<td>-</td>
<td>1.25</td>
</tr>
<tr>
<td>Right peripheral</td>
<td>172</td>
<td>217</td>
<td>-</td>
<td>1.26</td>
</tr>
</tbody>
</table>

Table 1: Result of arterial stimulation venous sampling

DISCUSSION

Based on the latest European and American guidelines, resistant hypertension is defined as high blood pressure that remains above goal with the use of 3 or more anti-hypertensive agents. Our patient has long standing hypertension but only started to develop resistant hypertension at the age of 73, which is relatively uncommon. Thus, the evaluation of resistant hypertension requires investigation to identify and treat the secondary causes of hypertension, including primary aldosteronism, obstructive sleep apnoea and renal artery stenosis.

The approach to diagnosing primary hyperaldosteronism should be step-wise, starting with screening of at-risk populations, confirmatory testing for positively screened patients, and subtype classification in order to plan for surgical or medical management. In the case presented, it was an opportunistic screening where he was referred following routine blood test for unexplained persistent hypokalaemia needing fast correction of potassium. Only then was he screened for primary aldosteronism and ruled out for other causes of secondary hypertension. The confirmatory test was not needed in this case as it fulfilled the criteria of case confirmation which were spontaneous hypokalaemia, marked high plasma aldosterone concentration > 20ng/dL and suppressed plasma renin.

Based on current guidelines, subtype classification of primary hyperaldosteronism should be determined with both imaging and adrenal vein sampling (AVS), reserving deferment of AVS for a selective subset of patients. AVS is the gold standard diagnostic test for
surgically curable primary aldosteronism. However, it is invasive, technically challenging and difficult to interpret. Due to AVS complexity, costs and limited availability, many patients with PA are being treated based on the computed tomography (CT) findings.

A non-functioning adrenocortical adenoma is not uncommon. Hence, cases of discrepancies between CT and AVS results have been widely reported, thus demonstrating the need of arterial stimulation venous sampling (ASVS) for localising the source of aldosterone excess. Based on Wachtel et al, primary aldosteronism patients with non-localizing imaging but lateralizing AVS would definitely benefit from adrenalectomy as seen in the case presented. Regardless of imaging findings, AVS is indicated to determine whether patients may be surgically curable. Thus, a correct subtype diagnosis will determine whether the patient may be surgically curable. It is also seen as a practical treatment objective to reduce the risk associated with cardiovascular disease.

Unilateral adrenalectomy will remove the source of aldosterone and treat the hypokalaemia. Unfortunately, the hypertensive changes in the kidney may not be completely reversible but the control of hypertension might be improved. Based on a retrospective cohort study for unilateral adrenalectomy for PA from 2004 to 2015 by Shariq et al, all patients achieved biochemical normality following adrenalectomy. There was no significant difference in the rates of hypertension cure or improvement observed in comparisons across pathological subtype. From 206 patients who met the inclusion criteria, 152 (73.8%) had single adenoma, 33 (16%) had unilateral hyperplasia, and 21 (10.2%) had multiple unilateral adenomas. Those with unilateral hyperplasia were more likely to be male (81.2% vs 57.9%, P = .03) and tend to be on the left side (78.8% vs 47.4%, P < .01) as in the case presented. Unilateral adrenalectomy performed either laparoscopically or retroperitoneoscopically is generally safe, efficient and curative in primary aldosteronism with resistant hypertension.

**SUMMARY**

Unilateral adrenal hyperplasia is a rare cause of resistant hypertension. Localization of the disease by arterial stimulation venous sampling is essential before surgery. Outcome of surgical treatment is excellent following appropriate diagnosis.

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