Spontaneous Renal Artery Dissection Resulting in Renal Infarction: A Case Report and Review of the Literature

Yoorim Seo, Ji Won Min, Yong Kyun Kim, Ho Cheol Song and Myung Ah Ha

Abstract:
Spontaneous renal artery dissection is a rare disease and an uncommon cause of renal infarction. The patient was a man who presented to the emergency room with sudden-onset right flank pain. Computed tomography revealed right renal infarction; thus, anticoagulation was initiated. Renal angiography revealed luminal narrowing of the segmental artery to the superior pole of right kidney without a dissection flap or false lumen. We stopped anticoagulation due to a lack of evidence of thrombi or luminal narrowing of the dissected vessels. When patients present with acute flank pain, it is important to suspect renal infarction and to perform a correct diagnostic workup, even when the patient shows normal urinalysis results and a normal LDH value.

Key words: spontaneous renal artery dissection, renal infarction, acute flank pain


Introduction
Renal infarction (RI) is a rare disease that is most commonly caused by cardiogenic thromboembolism associated with underlying diseases such as atrial fibrillation, intracardiac thrombus, infective endocarditis, and valvular heart disease (1). Less common causes of RI include hypercoagulation disorders, hematologic disease, and spontaneous renal artery dissection (SRAD) (2). Renal artery dissection is most commonly caused by the progression of a dissecting aneurysm of the aorta (3). Most cases are accompanied by an underlying disease such as malignant hypertension, Marfan syndrome, Ehlers-Danlos syndrome, neurofibromatosis, syphilitic arteritis, tuberculosis, and polyarteritis nodosa (4). Spontaneous dissection occurring in the renal arteries without any known trauma or underlying disease is termed SRAD (5). This case report describes the diagnostic workup and treatment of an otherwise healthy man with RI due to SRAD.

Case Report
A 37-year-old man presented to our emergency room with sudden-onset right flank pain. The pain persisted for a few hours without radiation or worsening on movement. He had no complaints of symptoms such as dysuria, gross hematuria or systemic symptoms such as fever, fatigue or joint ache. A physical examination at the time of presentation revealed the following findings: blood pressure (BP), 130/90 mmHg; heart rate (HR), 78 beats per minute (bpm); and body temperature, 36.8°C. He had no history of relevant medical conditions including hypertension or recent physical trauma. On physical examination, his abdomen was soft and flat with normal bowel sounds; however, right costovertebral angle tenderness was observed. He had no features related to Marfan syndrome, Ehlers-Danlos syndrome, neurofibromatosis or Takayasu arteritis. A blood analysis revealed the following findings: white blood cell count, 12,450/mm³ (neutrophils 71.5%, lymphocytes 23.3%); hemoglobin level, 17.9 g/dL; lactate dehydrogenase (LDH), 185 IU/L; creatine phosphokinase, 124 U/L; blood urea nitrogen (BUN), 12.8 mg/dL; creatinine (Cr), 0.85 mg/dL; and C-reactive protein

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2.65 mg/L. The other results of blood and urine analyses were within the normal ranges. The patient’s urine sediment showed no evidence of proteinuria or hematuria. Electrocardiography showed rare atrial premature complexes (APCs), with an HR of 71 bpm. A plain X-ray of the abdomen showed normal findings. Abdominopelvic computed tomography (APCT) showed a wedge-shaped perfusion defect in the superior portion of the right kidney with cortical rim enhancement, which suggested acute infarction (Fig. 1). The intimal flap, a characteristic finding of vessel dissection on CT scans, was observed in an area of low mural attenuation along the superior mesenteric artery (SMA) and the superior segmental branch of the right renal artery (Fig. 2). The jejunum seemed intact without ischemic changes. Anticoagulation therapy was started with low molecular weight heparin (LMWH; 1 mg/kg per 12 hours) for 3 days together with warfarin. We performed coagulation tests to exclude coagulopathy and the results were within the normal ranges (Table). On 24-hour Holter monitoring, rare APCs were observed but no arrhythmia (e.g., atrial fibrillation or atrial flutter) were seen. Echocardiography showed no evidence of thrombi or vegetation. The patient’s vital signs were stable without fever and his flank pain subsided during hospital admission. Laboratory tests showed no deterioration of the renal function and no hematuria or pyuria; however, his CRP level was elevated from 2.65 mg/L to 9.81 mg/L. Renal artery angiography was performed on the fifth hospital day. The exam revealed luminal narrowing of the segmental artery leading to the superior pole of the right kidney without a dissection flap or false lumen. Flow limitation was not seen on angiography at this time (Fig. 3A). SMA angiography showed luminal narrowing of the proximal SMA without flow limitation and the intimal flap seen in the initial APCT was not observed on angiography at this time.
Table. Coagulation Test Results.

<table>
<thead>
<tr>
<th>Blood test</th>
<th>Test value</th>
<th>Reference range</th>
</tr>
</thead>
<tbody>
<tr>
<td>Prothrombine time(s) / INR</td>
<td>10.8/0.92</td>
<td>10.1-14.0/0.85-1.13</td>
</tr>
<tr>
<td>Activated partial thromboplastin time(s)</td>
<td>28.1</td>
<td>21.0-38.0</td>
</tr>
<tr>
<td>D-dimer (mg/L)</td>
<td>0.19</td>
<td>0-0.55</td>
</tr>
<tr>
<td>C protein activity (%)</td>
<td>116</td>
<td>70-130</td>
</tr>
<tr>
<td>S protein activity (%)</td>
<td>85</td>
<td>77-143</td>
</tr>
<tr>
<td>Lupus anticoagulant Ab</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>Anti cardiolipin Ab IgM/IgG</td>
<td>Negative/Negative</td>
<td>Negative/Negative</td>
</tr>
<tr>
<td>Anti phospholipid IgG/GPL/mL</td>
<td>1.4</td>
<td>&lt;10.0</td>
</tr>
<tr>
<td>Factor V Leiden mutation</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>Anti-thrombin III (%)</td>
<td>69</td>
<td>60-120</td>
</tr>
<tr>
<td>Fibrinogen (mg/dL)</td>
<td>283</td>
<td>180-415</td>
</tr>
<tr>
<td>Factor VIII (%)</td>
<td>85</td>
<td>60-150</td>
</tr>
<tr>
<td>Factor V (%)</td>
<td>102</td>
<td>66-126</td>
</tr>
</tbody>
</table>

Figure 3. Angiography. (A) Right renal angiography. (B) SMA angiography.

Discussion

Spontaneous renal artery dissection is a rare disease and only 200 reports have been made since the first report in 1944 (6). A quarter of these cases were discovered on autopsy (2). The condition predominantly occurs in males (4:1 ratio) in their fourth to sixth decade of life (1). The incidence of SRAD in each kidney has been considered to be similar; however, a recent case series reported that the left side is more frequently involved (2). One hypothesis suggests that physical exercise can cause renal artery dissection by stretching the arterial wall (7, 8). The right kidney artery is perhaps affected less frequently than the left side, due to its longer length, which may cause shear stress to be more evenly distributed. Bilateral involvement was reported in 10-15% of the cases and was more frequently associated with underlying arterial disease (9-11).

The clinical presentation of SRAD ranges from life-threatening hypertension to renal failure. Hypertension may in turn also act as an aggravating factor for arterial dissection by potentiating arteriosclerosis and medial degenera-
tion (4). There are three different clinical manifestations of SRAD: an indolent state with no apparent progression; a renal infarction state due to an acute occlusion; and a chronic state with renovascular hypertension (1). The formation of a true and false lumen in the renal artery due to the rupture of the intimal layer in SRAD, may result in the obstruction of all or a part of the renal artery, thereby resulting in RI. In cases of RI, such as the case of our patient, the diagnosis should be made as quickly as possible to increase the possibility of renal revascularization and spare the renal parenchyma from further ischemic damage. In previous cases of RI, the AST, ALP and LDH levels were found to be increased, with accompanying proteinuria and/or hematuria (12). However, as was observed in the present case, some patients may show normal urinalysis results and LDH levels that are within the normal range on their initial presentation to the hospital (13).

Although there have been reports of cases in which SRAD was diagnosed by intravascular ultrasound (14), ultrasound and Doppler ultrasound show poor sensitivity (4). SRAD is most commonly diagnosed by CT angiography or MRA. Invasive exploration with angiography is recommended at an early stage to demonstrate the extent and nature of the vascular lesion, and to determine whether endovascular treatment is feasible. The results may also serve as a baseline vascular state to be used for comparison in follow-up studies (9). On angiography, thromboemboli and dissection appear differently; emboli appear as a meniscus crossing the width of the artery, whereas dissection is visualized as a linear filling defect in the arterial lumen or as uniform narrowing due to non-filling of the false lumen (7).

The optimal treatment of SRAD has not been established, and treatment should be selected based on the anatomical features of the dissection, the patient’s hemodynamic status and the renal function. The importance of BP control is uncontroversial, as previous reports have shown that BP control is the only treatment to have a significant effect on patient outcomes (1, 5). Although previous studies on the treatment of RI due to SRAD with oral anticoagulation and anti-hypertensive medication showed that the renal function remained stable with no significant complications (15, 16), its usefulness remains controversial. In the case of our patient, anticoagulation was stopped after renal artery angiography because no flow limitation or thrombi were observed. A Normal renal function was maintained without subsequent recurrence. Further studies are needed to establish guidelines for the use of anticoagulants in SRAD. The failure of medical treatment for BP control may require surgical treatments or other interventions (10). Percutaneous endovascular treatment with long-term follow-up has proven to be associated with less use of antihypertensive medications, less target organ damage and lower complication rates in comparison to surgical or medical treatment alone (17). If serial angiography shows a stable lesion without aggravation of the renal function, careful follow up is reasonable. However, if repeat angiography reveals an unstable lesion or if refractory hypertension is present, some form of intervention may be required (9). Surgical management, such as aorto-renal bypass, has also been recommended for the treatment of renovascular hypertension and acute occlusion (18). Primary nephrectomy may also be considered if the kidney is already severely damaged by infarction with a poor function on isotope renography and if revascularization is considered difficult to achieve (18, 19).

In patients presenting with acute flank or abdominal pain but with normal urinalysis results and normal LDH, BUN and Cr values, it is important to still consider RI and to perform an optimal diagnostic workup. In this patient, RI occurred due to SRAD, but angiography showed resolution without further intervention such as balloonning or stenting. As in previous reports, a favorable prognosis may be expected in young patients with RI due to SRAD without any underlying disease.

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Writing - review and editing: Ha MA
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